

The use of *Eucalyptus* in agroforestry systems of southern Rwanda: to integrate or segregate?

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ABSTRACT

Eucalyptus tree species widely grown in the tropics and subtropics predominate in the Rwandan landscape and primarily serve to supply the firewood and charcoal, timber and other non-timber forest products that are in high demand. They also play a significant role in the conservation of the environment by protecting soils against the erosion and landslides that are common in the study area. The trees are managed in woodlots of varying sizes owned by farmers, private institutions and local governments. The scattered rural households countrywide influence the distribution of eucalypt woodlots and cropped fields, creating a mosaic distribution of the two landuse types. This results in an intimate interaction between annual crops and the woodlot trees next to the fields.

Eucalyptus trees are known to be very competitive, suppressing crops in such settings to the extent that crops fail in the field zones nearest the woodlots. On the other hand, farmers get firewood, timber and other wood products as well as income from sales of timber and firewood and prefer to retain these very competitive trees on their fields. Controversy about *Eucalyptus* in other areas led to policymakers restricting the growing *Eucalyptus*, especially near wet areas. This thesis aimed to study the interaction between *Eucalyptus* woodlot trees and crops in fields next to the woodlots in southern Rwanda.

General allometric equations were developed relating tree diameter and height to total and component aboveground tree biomass in coppice and planted stands and these were used to quantify aboveground tree biomass. A competition experiment was run in Huye and Gisagara district (southern Rwanda) to investigate competition between woodlots and crops, with maize as the example field crop. Environmental resources of soil moisture, soil nutrients and solar radiation were assessed in a spatial gradient from tree–crop interface. Variation of the resources with distance from the tree–crop interface was also assessed. This was related to maize growth and grain yield at the different distances from the tree–crop interface. Using the resulting biomass equations, woodlot biomass production was quantified and an economic evaluation was done using the net present value approach to make trade-offs of growing combinations of *Eucalyptus* woodlot for producing timber, poles and firewood and maize, considering varying area proportions and growing sole maize. Extra woodlot biomass produced as a result of the edge effect was estimated and compared to the loss in grain yield due to competition.

Eucalyptus water use was estimated using the leaf area–tree diameter allometric relationship and the rate was compared to that of *Eucalyptus* reported in other areas and that of key crops grown in the area. General equations were derived for predicting aboveground total and component tree biomass. It was found that soil moisture, nutrients and light were significantly reduced in the crop fields next to the woodlots in a zone of up to 10.5 m width. Here, grain yield was 80% lower. However, the affected zone represents 10.5% of a 1 ha cropped field and when the loss is spread over the whole crop field area, the actual grain yield loss per hectare falls to 8.4%. Trees growing within 8 m of the perimeter of the woodlots had 17.8% bigger diameter at breast height and 34.5% more aboveground biomass than trees further away from the perimeter. Combining maize cropping with eucalypt woodlots was

more profitable than solely cultivating maize, and revenue from extra wood gains due to edge effects exceeded the corresponding revenue losses in maize yield.

Crop field size and orientation were very important. The smaller the field the more the edge effects since the proportion of the total field affected becomes more significant as the field size decreases. Rectangular crop fields will suffer more when longer sides adjoin woodlots. Woodlots, especially if small, also suffered from edge effects. Farmers with bigger landholdings are likely to benefit more than those with very small landholdings and they are likely to prioritise maize production (a less profitable option) over woodlots, for food security reasons. Recorded potential tree transpiration (3 mm d^{-1}) was 10% below local annual precipitation. The observed dry month water deficit could be covered by reductions in leaf area, stomatal closure and changes in soil water storage. A sensitivity analysis showed that 50% leaf area reduction corresponded to a 32.8 mm decline of potential tree transpiration. The moderate eucalypt water use rate observed may be a function of trees' small size and low tree stocking. The observed eucalypt water use rate is smaller than the range reported for eucalypts in Africa and was also smaller than that of key annual crops in the study area.

It is concluded that *Eucalyptus* woodlots are worth maintaining in the study area to ensure the supply of firewood and charcoal, timber and non-timber forest products, and for soil erosion and landslide control, as well as for generating income for the local communities. They should, however, be managed as woodlots, since managing them as single trees in cropland may lead to severe tree–crop competition. The current practice of not planting *Eucalyptus* trees in the major catchment areas in the country should be maintained. This should be extended to riparian zones and near wetland zones, since eucalypt water use is reported to increase with water availability. Managing woodlots as short rotations and increasing initial tree spacing may contribute to resolving issues related to catchment hydrology associated with eucalypt plantations. This could be enhanced by planting mixed stands of *Eucalyptus* and other tree species, preferably N-fixing, as these would also be useful in improving soils and woodlot productivity.

It is also recommended that eucalypt species be selected to allow for undergrowth development in the woodlots that improves soil erosion control and biodiversity. Finally, a holistic, integrated planning and management approach involving different stakeholders in natural resource management is recommended, to achieve agricultural production that is more compatible with the protection of ecosystems without impairing farmers' livelihoods.

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CHAPTER 1: General introduction

General introduction

Agroforestry: its history and naissance as a scientific field

Agroforestry is a relatively young term for an old age practice, the practice probably dating back to the beginning of humanity on this earth. Cultivation of trees and crops together has been practiced throughout the world since the beginning of agriculture (Nair, 1993). Literature reports that agroforestry have been practiced in Europe since the Middle Ages (King, 1987). The practice is still ongoing in Europe today and the main combinations of most of agroforestry practices used nowadays are mainly found in two categories: silvoarable and silvopastoral areas (Rigueiro-Rodriguez et al., 2009). Cork oak agroforestry is a common practice in Spain (Campos et al., 2009). Silvoarable systems as hedgerows are seen in Brittany (bocages), Normandy, Ireland, the Knicks and Walhecken in Germany (Rigueiro-Rodriguez et al., 2009). Dispersed oak trees exist in dehesas systems in Spain or Portugal, and parkland systems in the UK, and zoned systems including the planting of olive in 5–10 m rows with cropped areas in between, and the use of shelterbelts for livestock in Northern Europe (Eichhorn et al., 2006). Riparian strips and line planting (hedgerows, shelterbelts, and windbelts) defined as trees outside forests (MCPE 2003) are classified as agroforestry (Long and Nair 1999). Silvopastoral system can be found in the Mediterranean region (Eichhorn et al. 2006). Forest farming, an agroforestry system where non woody forest products such as mushrooms, medicinal plants etc. are harvested (Lange and Schippmann 1997; Lange 1998) exist in countries like Albania, Bulgaria, Hungary and Spain (Rigueiro-Rodriguez et al., 2009).

This practice was also reported in America (Ares et al., 2006; Wilken, 1977) and Asia (Conklin, 1957), specifically in India in 1890 (Raghavan, 1960). It is said to have been “introduced” in South Africa in 1877 (Hailey, 1957) and in Nigeria in early 20th century (Forde, 1937; Ojo, 1966). Yet agroforestry is reported to be an ancient practice in sub-Saharan Africa where farmers deliberately retain and integrate trees into their farmland (Bucagu et al., 2013).

As a modern scientific study, the term ‘agroforestry’ appeared first in a review published by Bene et al. (1977). More importantly, it gave a breakthrough in recognizing the key concepts that trees and shrubs grown on farmlands played distinctive and valuable roles (Young, 1997). It was born from other agricultural sciences in 1970-80s (Torquebiau, 2000) and in the early 1980s, agroforestry research focused on conceptual issues such as the definition and classification of its different types (Young, 1997); the inventory of existing agroforestry systems (Nair, 1989) and soils research (Mongi and Huxley, 1979). It depended on information drawn from the fields from which it was born: agriculture, forestry and soil science (Young, 1997).

Agroforestry is defined as any form of land use where trees and agricultural crops and/or animals are combined, either in some form of spatial mixture or sequence in one production unit. While classifying agroforestry practice is as complex as the systems themselves, several attempts have been made to identify and name different systems. Examples include

simultaneous systems (e.g. alley cropping, savannah parklands, and multi-storey gardens) with direct interactions and sequential or rotational systems (e.g. the old shifting cultivation, improved fallows, and rotational woodlots) with indirect interactions or residual effects between trees and crops (Nair, 1993). Different classes are also recognized based on the agroforestry components, including: agrosilvicultural (crops and trees such as in alley cropping, homegardens, etc.), silvopastoral (trees and animals as in pastures) and agro-silvopastoral (crops, trees and animals as in zero grazing).

According to Radersma (2002), classification of agroforestry is a complex undertaking and its definitions range from purely physical and scientific to more “political” ones, the latter including aims of the system(s) such as ecological, social and economic gains. Such ‘aim containing’ definitions may not apply to unsuccessful agroforestry systems (Radersma, 2002), although the latter are likely to occur. Unsuccessful agroforestry systems may occur when there are wrong choices of species combinations, management practices, and lack of peoples' motivation and understanding (Nair, 1993) and when trees become invasive and hard to control.

Tree-crop interactions

Tree-cropping systems fall within a range between complete segregation and complete integration of trees in a landscape as two extremes of a spatial pattern perpendicular axis of deforestation/afforestation (van Noordwijk et al., 2012). Along this continuum, agroforestry is associated with more integrated systems, while a coarse mosaic of tree and crop monocultures form an alternative system. However, tree-cropping systems will be considered as agroforestry even when the components are segregated, as long as the components of the mosaic interact and the interaction influences the productivity of individual components, either positively in which case they are facilitative, or negatively when they are competitive.

The interaction between trees and crops when the two are grown together or in close proximity has been a subject of interest since the start of agroforestry as a scientific field. Initially, scientists thought trees in agroforestry systems could improve these systems and therefore improve crop yields directly (Kidd and Pimentel, 1992). It is however, not an easy and direct process how trees and crops on farmland interact and provide expected outputs because environmental conditions interact with plant and population traits to determine outcomes (Sheley and James, 2014). Relative plant size may be an essential factor needed to predict the outcome of plant-plant interactions. Variation in plant size creates size asymmetric interactions where larger individuals (trees) may pre-empt resources by acquiring them while they are unavailable to smaller individuals (annual crops) (Weiner and Damgaard, 2006). These interactions are among the strongest drivers of dynamics in tree-crop systems because they set the limits within which organisms can live and function (Lin et al., 2013).

Accounts on how trees and crops interact in multi-cropping systems and their farmers' benefits and shortfalls in relation to monocrops in different cropping conditions and environments exist. In their account on resource capture and crop productivity under different rainfall and management conditions in the agroforestry system in the Sahel, Coulibaly et al.

(2014) observed that more water did not in all cases lead to a better sorghum grain production. They conclude also that under the study conditions, increasing tree density does not negatively affect sorghum growth if crown pruning is applied. On the other hand, Bayala et al. (2015) also studied the processes in soil-tree-crop interactions in the parkland systems in west Africa. They conclude that, an over-all understanding of the functioning of the complex agroforestry systems is still lacking because no published studies have addressed possible interactions that take place at the same time in such systems. Garcia-Barrios & Ong (2004) and Ong et al. (2004) explain the interspecific interaction and facilitation between plants, how these interact under different environmental resource conditions and how this impose trade-offs, biophysical limitations and management requirements in tree-crop mixtures. They state that, introducing trees in croplands to promote low-input sustainable agroforestry systems is a challenging undertaking due to some reasons: (a) trees provide useful products for smallholders and strongly facilitate crops, but can also exert stronger competitive effects than previously expected; b) practices aimed at increasing trees' beneficial effects can sometimes also enhance trees' competitiveness; c) the interplay between positive and negative effects of trees change – sometimes significantly – from one environment to another. This makes it difficult to predict the consequences of extending successful agroforestry practices to new environments having different resource levels and resource balances.

In an intercrop, component crops differ in terms of patterns of resource use, which may lead to a more efficient exploitation of resource than when grown in monocultures (Willey, 1990; Francis, 1989). Environmental resources may be more completely absorbed and used to build crop biomass by intercropping (Amini et al., 2013). This is a consequence of differences in the component species capacity to compete for growth resources between them (Amini et al., 2013; Anil et al., 1998). This may imply that, the components are not competing for the same ecological niches and that interspecies competition is weaker than intra-specific competition for a given factor (Vandermeer, 1989). The same would occur if resource requirements of the component species are separated in space or in time (Cannell et al., 1996).

Agroforestry systems are designed to optimise the use of physical resources either spatially or temporally, by maximising positive interactions (facilitation) and minimising negative ones (competition) among the components (Jose et al., 2000) as well as to diversify products (Kidd and Pimentel, 1992). Successful management of agroforestry systems will essentially require the understanding of complex biophysical processes and mechanisms involved in the allocation of resources so as to get systems that are ecologically sound, economically feasible and socially acceptable (Ong et al., 1996; Rao et al., 1998). This has to be in commensurate with the central agroforestry hypothesis which states that “the benefits of growing trees with crops will occur only when the trees are able to acquire resources of water, light and nutrients that the crops would not otherwise acquire” (Cannell et al., 1996).

Ecologically as Jose et al. (2004) states, productivity of an agroforestry system is comparable to that in natural environments, and is the net result of positive and negative interactions among the components. Interactions occur as component species exploit growth resources

above- and belowground (Ong et al., 1996). The intensity of interaction, both inter- and intra-specific is size and density dependent and will increase when individuals per species increase per unit area and vice versa. Two or more species will co-exist until one or more of the resources they share spatially or temporally turn insufficient or limiting for one or both species.

The mechanism of co-existence and competition among species was illustrated by several scientists such as De Wit (1960) and Armstrong and McGhee (1980). The mechanisms are used to assess interference, niche differentiation, resource utilisation and productivity in simple mixture of species (Jolliffe, 2000). A standard replacement series involves at least two species mixed and their yield is compared per unit area and the results can be presented graphically as in Figure 1.1. To study competition in mixtures of two species, the proportion of each of the two species is varied while keeping constant the overall density. Replacement diagrams are used to interpret the results by showing relative yields of each individual in relation to the relative frequency. The relative yield total (RYT) is the index of resource competition and is calculated as:

$$RYT = \frac{Y_{AB}}{Y_{AA}} + \frac{Y_{BA}}{Y_{BB}}$$

where Y_{AB} and Y_{BA} are yields of species A and B per unit area when grown in mixtures, and Y_{AA} and Y_{BB} the yields of the two species when grown in pure stands. When $RYT=1$, it means the species compete for same resources; when $RYT>1$, it means there is resource complementarity.

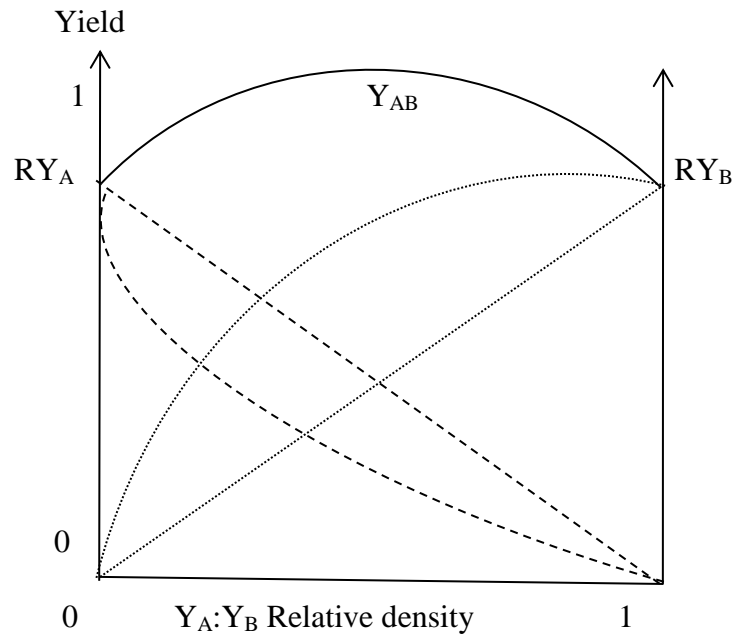


Figure 1.1. Replacement diagram in showing trends in total (Y_{A+B}) and component yields (Y_A = yield of species A and Y_B = yield of species B) (in say, dry matter, in units per unit area).

The components are mixed and a selected measure of yield of each component per unit area (e.g. biomass) is assessed while the proportions of individual components in the mixture vary from 0 to 100%. This very simplified situation is more suited to agro-ecosystems with similar component species having similar rooting architecture, expected to exhibit similar physiological needs and to respond to fulfilling the needs in a similar manner (Jose et al., 2004). The situation in agroforestry systems is far more complicated and component species differ in many various ways: species combinations, life forms and structure, resource needs in quantity, quality, spatial and temporal dimensions, etc.

Since agroforestry systems are deliberately established and managed, the idea would be to choose species combinations that will, in one way or another, ensure reduced competition and whenever possible, opt for species that will eventually lead to facilitative or complementary systems. Complementary species will be expected to use resources more effectively either by using a different niche as, for example, when the annual crop is shallow rooted while a companion tree is deep rooted, or by changing its phenology to exhibit different temporal resource use. Differences in structure and behaviour of species therefore may allow for better species combinations that may allow for better capture of environmental resources, leading to better system productivities (Ashton, 2000; Garcia-Barrios and Ong, 2004). For example, *Senna siamea* and *Leucaena leucocephala* were intercropped with maize in semiarid Machakos, Kenya. *L. leucocephala* reduced maize yield in the intercrop by 16% compared to the sole maize while *S. siamea* did not reduce the intercropped maize yield (Jama et al., 1995). Compared with the sole-crop systems, *L. leucocephala* intercropping did not affect land equivalent ratio (LER) (competitive situation), whereas LER increased by 28% with *S. siamea* intercropping (a complementary situation). The capture of growth resources by trees and crops can be grouped into three broad categories to show competitive, neutral or complementary interactions. In the neutral or trade-off category, trees and crops exploit the same pool of resources, so that increases in capture by one species result in proportional decreases in capture by the associated species (Ong and Shallow, 2003). If trees were able to tap resources unavailable to crops (Cannell et al., 1996), then the overall capture would be increased, i.e. complementary use of resources. In the third category, negative interactions between the associated species could result in serious reduction in the ability of one or both species to capture growth resources. It is important to bear in mind that tree-crop interactions may change from one category to another depending on the age, size and population of the dominant species, as well as the supply and accessibility of the limiting growth resources (Ong and Swallow, 2003).

A simplified schematic representation of complementarity in resource use in agroforestry is shown in Fig. 1.2a as illustrated by Wojtkowaki (1998) and as modified by Jose et al. (2004). The line A1-B1 shows how the total yield of species A and B as the proportion of each in sole cropping varies in a given area. The trajectory A+B represents the over-yielding compared with either of the sole crop yields of one possible mixture of species A and B. Unfortunately, this presents an ideal situation which can seldom be attained in nature. The situation is more complicated by the fact that agroforestry systems are dynamic and temporal changes occur across the rotation of one cycle. A hypothetical illustration of how situations

can change over time is presented in Fig. 1.2b (Jose et al., 2004; Wojtkowski, 1998). Changing of species needs and physical structure affect the degree of resource sharing, leading to alternating complementarity and competition. When the latter predominates, management intervention may be applied to reverse the situation so as to ensure better co-existence and productivity. The response of tree-crop mixtures to the effects of management of the trees is illustrated in Figure 1.2b.

Ong and Swallow (2003) summarise resource capture in tree-crop combination systems in three categories to show neutral, complementary or competitive interactions. In the neutral or trade-off category, trees and crops exploit the same pool of resources, so that increases in capture by one species result in proportional decreases in capture by the associated species. If trees were able to tap resources unavailable to crops, then the overall capture would be increased, i.e. complementary use of resources. This could be possible for example, through spatial differences in resource capture strategies shown by different species. Trees mainly depending on resources captured underground lower than crop roots can reach; or by increasing aboveground zone where light is captured from near the ground level by annual crops to 30 m above ground for tall trees. Light that escapes upper storey trees are then captured by shorter plants with the overall increase in the amount of light captured by the intercrop. In the third category, negative interactions between the associated species could result in serious reduction in the ability of one or both species to capture growth resources. It is important to bear in mind that tree-crop interactions may change from one category to another depending on the age, size and population of the dominant species, as well as the supply and accessibility of the limiting growth resources.

In practice, extension agents advise farmers on tree species to grow in their fields. This may not favour adoption because it may be against farmers' preferences. Wajja-Musukwe et al. (2008) suggest that farmers be allowed to select tree species they prefer, and then apply tree management practices such as root and shoot pruning to reduce tree competition. Wajja-Musukwe et al. (2008) reported very little to no competitive effects in *Grevillea robusta* and *Markhamia lutea* with crops while *Casuarina equisetifolia* and *Maesopsis eminii* affected crops significantly. This may be a result of the widely spreading canopy of *G. robusta* (Wajja-Musukwe et al., 2008) and the fact that coppice roots in *G. robusta* grow downwards and do not spread in the rooting zone (Wajja-Musukwe, 2003). Such less competitive species could lead to over-yielding since almost all tree products will add to the slightly reduced or non-affected crop yields if the two were grown together. Another example of over yielding was reported with *A. acuminata* where larger crop yields were recorded near tree lines than far from the tree. This observation was reported in south western Uganda (Okorio et al., 1994) and in northern Rwanda (Ukozehasi, 2006). Over yielding may be attributed to the downward growth of coppice roots in this species (Wajja-Musukwe, 2003) and its narrow and less compact canopy which seems to facilitate light penetration to the understorey crop. Compared with other tree species, *A. acuminata* and *G. robusta* also had small root number: trunk volume ratios (Okorio et al., 1994) which may also contribute to explain the less competition. *A. acuminata* interacted positively with maize in Thika, Kenya, and this was

attributed to its ability to fix nitrogen (Muthiri et al., 2005). Similar observations have been reported in Uganda (Peden et al., 1993) and in India (Sharma and Sharma, 1997).

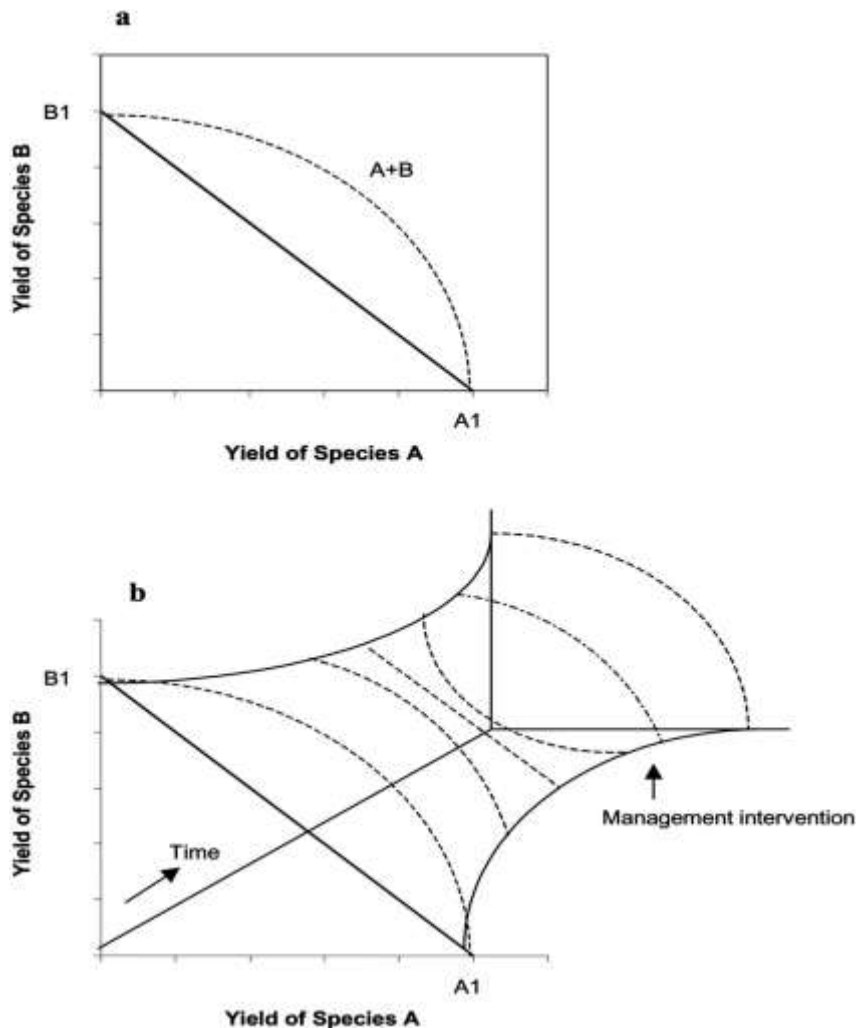


Figure 1.2. The production possibility curves for two species, A and B. (a) Points A1 and B1 represent the maximum production potential if species A and B were grown in monocultures. Line A1 to B1 represents the proportional yield of species A and B when grown in mixtures. (b) A hypothetical temporal production possibility surface for species A and B (modified from Wojtkowski 1998). As time progresses competition becomes more intense, thus resources become limiting, over-yielding gives away to under-yielding, but a timely management intervention (e.g., root pruning of trees) alleviates competitive interactions, thereby resulting in over-yielding again.

Appropriate agroforestry systems are often assumed to provide the environmental functions needed to ensure sustainability and maintain microclimatic and other favourable influences, and it is assumed that such benefits may outweigh the disadvantages of a more complicated management (Sanchez, 1995). Again, it is also assumed that agroforestry might be a practical way to mimic the structure and function of natural ecosystems, since components of the latter

result from natural selection towards sustainability and the ability to adjust to perturbations (Van Noordwijk and Ong, 1999). Natural ecosystems are known to have greater biodiversity and the latter is known to be a major determinant of community and ecosystem dynamics and functioning, and greater diversities lead to greater ecosystem stability, greater resistance to invasion by exotic species, and lower disease incidence (Tilman et al., 2014). Previous agroforestry reviews have highlighted several unexpected but substantial differences between intensive agroforestry systems and their natural counterparts that would limit their adoption for solving some of the critical land-use problems in the tropics (Ong and Leakey, 1999; van Noordwijk and Ong, 1999). Denison and McGure (2015) also observe that higher diversity may not necessarily lead to better performance of agroecosystems and that mimicry of natural ecosystems may have little value in agriculture.

Spatial arrangement of individuals in tree-crop systems differ from place to place depending on environment, farmers' preferences and traditional cultural practices. Common arrangements include: trees dispersed in the crop fields at varying tree densities; line plantings (as in alley cropping, border plantings, contour plantings and windbreaks); complex mixtures as in homegardens; or rotational or simultaneous woodlot systems. The varied arrangement structures lead to complex systems which complicate the understanding of the system performance. In west African parkland system alone, Bayala et al. (2015) identify multiple interacting components: different tree species and crop varieties with variations at field, watershed and landscape levels. There are also diverse management practices and diverse spatial and temporal scales. All these factors influence resource capture and system productivity in many different ways, and may contribute to the success or failure of the system.

This study

The current study described *Eucalyptus* woodlot-agricultural crop system using maize as an example. This system is common in areas that are characterised by high population densities with small household landholdings and where tree woodlots are an integral part of the farming systems. Rwanda is a typical example of areas where this type of tree-crop farming is dominant. Trees are grown to supply fuelwood which is the main source of domestic energy for cooking (Ndayambaje and Mohren, 2011) and wood for construction (Nduwamungu, 2011a). This field setting is such that, pure stands of eucalypt woodlots and pure maize fields are growing next to each other. The system complies with the coarse mosaics of tree and crop monocultures described by Van Noordwijk et al., (2012), hence it contains elements of segregation as well as integration. It is a typical agroforestry system because plant components in the two subsystems interact and influence each other's growth and reproduction. A system in this case is defined as – a structure consisting of interrelated elements which interact (directly or indirectly). Usually it has boundaries that are usually defined by the system observer. An agricultural system is an assemblage of components which are united by some form of interaction and interdependence and which operate within a prescribed boundary to achieve a specified agricultural objective on behalf of the beneficiaries of the system (McConnell and Dillon, 1997). Depending on the scale of the

analysis, a farming system can encompass a few dozen or many millions of households. *Eucalyptus* woodlots and maize crop make the components of our system and the interaction between them is the focus in this study.

The tree species studied, *Eucalyptus saligna*, is one of the eucalypt species commonly planted in Rwanda, especially in farming systems in the southern Province where it serves to supply firewood to the local community (Nduwamungu et al., 2008). While pointing out that there is information gap on fuelwood consumption rate data in Rwanda, Ndayambaje and Mohren (2011) report that 92.2% of the population in rural areas and 93.5% in urban areas rely on firewood and charcoal as the sources of domestic energy. An estimated 72% of this comes from eucalypts (MINERENA, 2013). In addition, the soils of the study region are known to be very poor, and together with hilly topography drive farmers to undertake some soil erosion control practices. Among these, tree planting is very important and farmers in Nyamagabe area, southern Rwanda, use eucalypt leaves and branches as mulch to reduce soil loss by reducing kinetic energy of rain drops and nutrient losses through litter decomposition (Nzeyimana et al., 2013).

The urgency of tree planting coupled with small landholdings per household, make the conditions unavoidable to grow crops near woodlots, irrespective the fact that farmers are aware that these trees compete with their crops. The subject of this study was to evaluate the tree-crop interface in this setting as described later in the next sections of this thesis, by evaluating tree growth, tree-crop interaction and their trade-offs; and tree water use since these aspects are interrelated and contribute substantially to the farmers' livelihoods in the area.

The study site

The study was carried out in the southern part of the central plateau agro-ecological zone in southern Rwanda (Delepierre, 1975), corresponding to the southern central zone according to agro-ecological classification by Clay and Dejaegher (1987) in Huye and Gisagara districts (Fig. 1.3). In this zone, *Eucalyptus saligna* is one of the most dominant plantation tree species. According to the classifications, the zone is characterised by a bimodal rainfall (March-May and September-December) and the mean annual rainfall is about 1200 mm. Monthly means of daily temperature maxima range from 28.5 °C (April) to 32 °C (September) and corresponding minima from 10 °C (April) to 9 °C (September). The soils are derived from granitic rocks and are classified as oxisols or ultisols on the hills (Birasa et al., 1990; Champelle, 1985).

Problem description

About 1,369,576 ha, equivalent to 52% of the total Rwandan surfaced area is arable land. According to the countrywide forest inventory done in 2007, *Eucalyptus* plantations covered 63,600 ha and coppice woodlots on 39,200 ha (MINIRENA/CGIS-NUR, 2008). In 2010, new eucalypt plantings plus the coppice woodlots covered 62,700 ha, making total eucalypt cover to be 126,300 ha (Nduwamungu, 2011a). This inventory however, excluded <0.05 ha

woodlots and any forest whose cover was $<10\%$ and trees with DBH <7 cm. According to FAO (2010), small woodlots and trees outside forests (including minor quantities of non-*Eucalyptus* species) covered about 6.6% of the country area, equivalent to 173,830 ha. *Eucalyptus* contributes 90% of the woodlots and trees outside forests (Ndayambaje et al., 2014), leading to 156,447 ha, making a total eucalypt coverage of 282,747 ha, equivalent to about 21% of the total arable land. In the past, tree planting on peoples' fields were facilitated by such factors as: 1. Community works practice locally called *Umuganda*; 2. Readily availability of free planting material and 3. A tree tenure system which allowed growers to own trees they planted (GTZ, 2008).



Figure 1.3. Rwanda map showing the study sites (marked with a red star)

Household eagerness to be self-sufficiency in tree resources to meet demand for firewood and for other wood products led to growing trees on every land available, sometimes including the best available land for food production. Land fragmentation as a function of population growth led to progressive decline in land size per household, the latter being reported to have decreased from 3 ha in 1960's to <1 ha today (Verdoodt and Van Ranst, 2006). Mpyisi et al. (2003) report average landholdings per household to be 0.5 ha in 2002. To meet household food and wood requirements necessitated intercropping food crops with trees. This resulted in intensive agroforestry practices where trees are either directly mixed with, or grown in very close proximity to crops. For the same number of trees, a decrease in farmland size following fragmentation means increased tree density and therefore a more intense tree-crop competition. An inverse relationship between farm size and the amounts of woody biomass

was reported in Rwanda, where more average wood biomass per farm was observed in the lower average farm size, high altitude region than in the higher farm size, low altitude region (Ndayambaje et al., 2014). Similar relationships between farm size and on-farm tree biomass production were observed in other areas as well (e.g. Chand et al., 2011; Thapa, 2007). Under such circumstances, farmers' possible strategy becomes to shift from woodlots to growing fewer trees along lines or single trees scattered on crop field.

Intercropping trees with crops is known to reduce crop yield severely, especially in water scarce areas (Ong and Leakey, 1999; Rao et al., 1998; Cooper et al., 1996). An expected tendency would be the use of non- or less competitive tree species in such agroforestry systems (Schroth, 1995). Farmers sometimes integrate under certain conditions, surprisingly fast growing and presumably very competitive tree species in their agroforestry systems (Schaller et al., 2003). Examples include the use of extremely fast growing *Paraserianthes falcataria* (L) I.C. Nielsen and *Eucalyptus deglupta* Blume in coffee plantations in Indonesia and Costa Rica respectively (Schaller et al., 2003), and *Eucalyptus* species in Rwanda. When trees provide higher economic advantages for example, farmers usually ignore tree competitiveness and integrate them with crops. This has been observed in a range of study conditions across countries and continents. Examples include in Africa: Rwanda (Ndayambaje et al., 2014), Ethiopia (Zerihun and Kaba 2011) and Ghana (Isaac et al. 2009); in Asia: China (Tang et al. 2012) and India (Banyal et al., 2011; Palsaniya et al., 2010), and Europe: France (Mary et al. 1998). *Eucalyptus urophylla* was rated as very good by farmers in Kageyo area (northern Rwanda) owing to its wood characteristics but very poor in terms of compatibility with crops, yet 50% of all farmers grew the species (Bucagu et al., 2013). *Eucalyptus* trees are reported to affect the environment negatively (Lemenih et al., 2004), to suppress crops in the farming systems (Forrester et al., 2010) and to impart allelopathic effects to crops growing in their vicinity (Hartemink, 2003).

Rwandan policy makers, based on allegations on eucalypts raised elsewhere that they use much water and dry sites, suggested that the spp. should be eliminated from marshlands and near other water bodies, and restricted their use in reforestation in the country (Gahigana, 2006). They are concerned with water availability for the irrigation of rice fields and for hydropower generation.

Despite the alleged negative effects of eucalypt on water discharge and biodiversity, some studies appraise eucalypts and report that growing them improve soil structure, organic matter content and soil nutrients (Parrotta et al., 1997). The species are further reported to improve native biodiversity restoration on degraded lands (Senbeta and Tekatay, 2001; Yirdaw, 2002).

With such contradictory reports on the effects of eucalypts, there is a need to conduct studies to bridge the knowledge gaps for policy makers to make sound decisions to solve locally specific problems rather than basing on exotic information which may not be appropriately suited. Scanty information on the effects of eucalypts on the environment in Africa complicates further the situation. The lack of information becomes even poorer in eastern African region as revealed by Dye (1996). In the whole of Africa, only few studies have been

done in South Africa (Dye, 1996), in Ethiopia (Dessie and Erkossa, 2011) and in the Congo (Bernhard-Reversat, 2001). No such studies have been done in Rwanda and in the east African region. The aim of the present study is to contribute to the knowledge on eucalypt growth, their effects on crop yields and thus on food security and their water relations as well as the contribution to the livelihoods of rural farmers as income generators.

Research objectives and hypotheses

The main objective of this research is to contribute to the knowledge on the growth of eucalypts under local conditions, and its effects on crop productivity and the environment. The availability of this information and its dissemination will enable different stakeholders to improve land use management practices for the sound environment. To this end, the following series of specific objectives and hypotheses were put forward:

Objectives

- To develop a general equation for the estimation of aboveground tree biomass applicable to both planted and coppice stands of *Eucalyptus saligna* in the study area
- To quantify the extent to which the competition for environmental resources manifest itself in maize grain yield and how this varies with distance from *Eucalyptus* woodlots along a line perpendicular to the tree-crop interface under the conditions of the study site
- To determine the area of maize field/eucalypt woodlot affected by the edge and the extent of maize grain loss or gain in aboveground biomass, and the trade-offs between the two components in an agroforestry system
- To estimate the water use rate of eucalypt woodlots and compare this to the local precipitation to learn if eucalypts transpire much water than the rain supplies

Hypotheses

The central agroforestry hypothesis by Cannell et al. (1996) is the overall hypothesis of our study. It states that “*benefits of growing trees with crops will occur only when the trees are able to acquire resources of water, light and nutrients that the crops would not otherwise acquire*”. Under this, we further hypothesize that:

- *Coppice shoots and stems in coppice and planted stands of eucalypts differ significantly in size yet total aboveground tree biomass estimation in both stand types can be done by using a common general allometric equation*
- *Eucalypt woodlots negatively affect crops grown in adjacent fields and both crops and biophysical resources spatially differ with distance from the woodlots*
- *Financial losses due to crop yield reduction in farmlands next to eucalypt woodlots can be compensated for by extra biomass production in woodlot periphery trees to maintain an overall system productivity beneficial to farmers*
- *Trees in short rotation eucalypt woodlots use much water as to affect catchment water balance negatively*

Methodological Approach

The aim of this thesis was to evaluate this tree-crop combination biophysically to see if the system benefits smallholder farmers practicing it. The field setting of the experiment involved two apparently monoculture components, which would fall under the category of fields + forests, as defined by Van Noordwijk et al. (2012). Due to an intense competition between the eucalypt woodlots and maize crop in these mosaics, we consider the mosaic to constitute an agroforestry system of our interest. What makes the two components one system is the interaction between them, which was our main subject of investigation. The approach is explained further in the next section, the thesis outline and more precisely under respective chapters.

Thesis outline

This thesis is made up by six chapters outlined below. After Chapter one which introduces the whole study, chapter two deals with the growth of eucalypt trees and develops a general equation for use in predicting above ground tree biomass in both the coppice and the planted stands of *Eucalyptus saligna*, a species most dominant in the study area among the eucalypts. Separate equations have also been developed for the different tree components of stem, branch leaves. Component models are useful in determining component biomass since in practice; different biomass components represent different market commodities. Tree growth reflects its capacity to capture and use resources, therefore its competitive power with other plants. Chapter three evaluates the interaction between the maize as a representative annual crop and the eucalypt woodlots. An analysis of tree-crop competition is made by examining woodlot effects on soil physical (particle distribution) and chemical (nutrients), soil moisture and solar radiation and how these change with distance from the tree-crop interface, and also how soil pH, soil nutrients and soil moisture change with soil depth from the surface to 80 cm below ground. These were consequently related to maize plant growth and grain yield.

Chapter four is also about tree growth and tree effects on maize crop. A comparison is made on interspecific competition in the woodlot trees looking into the growth of individual trees at the woodlot centre and those at its edge. The intraspecific competition (on the woodlot trees) is combined with the interspecific competition observed in the previous chapter (the effects of woodlot trees on maize crop) to evaluate the edge effects. A trade-off analysis is provided on the eucalypt woodlot-maize agroforestry system by comparing in monetary terms, the gains or extra biomass of the edge trees in the woodlots and the losses in maize yield when the two components comprise a cropping system, and show whether the losses are compensated for by the gains in this type of agroforestry system.

In Chapter five, eucalypt woodlot water relations is investigated at catchment scale. The tree or woodlot capacity to capture water is an index of its competitive capacity. Tree water use was estimated based on tree leaf area index relationship with tree diameter. A monthly water balance was developed for two representative catchments in the study area whose main land use is mixed crop production and eucalypts woodlot management. A comparison of the water

use by eucalypts was made with that of key crops in the area and also with that of eucalypts in other areas worldwide.

The last chapter provides the general discussion and synthesises the main findings of this study. Basing on the findings, conclusions and recommendations related to eucalypt growing in relation to food crop yield and environmental protection are given.

**CHAPTER 2: Above ground biomass prediction in coppice and planted stands of
*Eucalyptus saligna***

CP Mugunga KE Giller & GMJ Mohren

Above ground biomass prediction in coppice and planted stands of *Eucalyptus saligna*

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Abstract

We derived general equations to predict total tree and tree component aboveground biomass for planted and coppice stands of *Eucalyptus saligna* in Southern Rwanda. The equations were developed using data collected from 14 coppice and planted stands across the region. Diameter alone or diameter and height were regressed against biomass and the best fitting general allometric equations were selected. Variance was observed to increase with tree diameter, necessitating the use of non-linear regression and Cunia's regression weight of D^4 . Biomass did not differ significantly between coppice and planted stands. Site differences observed were attributed to cattle grazing practised in the past. Existing equations were not applicable because they combined several eucalypt species, developed from a small sample from a small area or were developed for other sites characteristically different from those of the study region. While using both height and diameter only slightly improved the equation precision, using both parameters in biomass equation development is recommended since this usually allows for equation application to other regions. It is easy to measure height in this species stands and errors associated with height measurement in this species may be minimal due to bole straightness and light crowns.

Key words: Allometric equations, model performance, total tree above ground biomass, tree component biomass, short rotation coppice systems

Introduction

Eucalyptus species are grown in Rwanda mainly as a source of firewood but also for the supply of timber and other wood products (Nduwamungu et al., 2008). The trees are either managed as short rotation (5-7 years) coppice or long rotation (≥ 20 years) stands for firewood and timber production respectively. The eucalypt plantations typically belong to a variety of owners from farmers through different private groups to the government (Nduwamungu, 2011b). The ownership pattern, among other factors affect planting and harvesting regimes, leading to changes in stocks over time. Standing volume and aboveground biomass (W_{AGB}) are two main measures of forest stocking that are typically considered within the framework of sustainable forest management and for carbon accounting purposes (Brandeis et al., 2006). Accurate estimations of tree volume and forest biomass is crucial for assessing expected yields from commercial and subsistence harvesting (Mwakalukwa et al., 2014). It is also important for carbon storage assessments in relation to global climate change mitigation measures (Husch et al., 2003; Freer-Smith et al., 2007). For this purpose, allometric estimates of forest biomass can be applied to estimate carbon stocks and carbon fluxes when measured repeatedly, thus providing means for estimating the amount of carbon dioxide released into or removed from the atmosphere.

The basic principle of allometry can be used to predict a tree variable (typically its biomass) from another dimension (e.g. its diameter), and an allometric equation quantitatively

formalizes this relationship (Picard et al., 2012). For instance, the proportions between height and diameter, between crown height and diameter, between biomass and diameter, etc. follow rules that are the same for all trees, big or small, as long as they are growing under the same conditions (Bohlman & O'Brien, 2006; Dietze et al., 2008).

Furthermore, guidelines have been published in forest inventory for setting up permanent plots, counting trees correctly (Sheil, 1995; Condit, 1998), and for estimating above ground biomass stocks and changes, from these data (Clark et al., 2001; Philips et al., 2002; Chave et al., 2004). However, one of the largest source of uncertainty in all estimates of carbon stocks in planted and natural tropical forests is the lack of standard models to convert tree measurements to biomass estimates (Chave et al., 2005).

A general practice has been to estimate volume and biomass from tree dendrometric characteristics such as diameter and height, using established, general, or site-specific allometric equations (De Gier, 2003; Navar, 2010). The selection of an appropriate allometric equation is a key element in the accurate estimation of forest yield and stand productivity as well as carbon stocks and changes in stocks from basic variables such as diameter and height (De Ridder et al., 2010). Combining tree allometry and stand based inventory is probably the most accurate technique to detect the biomass change in short to medium term and are often used to validate model outputs and remotely sensed spatial predictions of biomass (Burrows et al., 2000).

Allometric models have been used in estimating tree biomass in *Eucalyptus* for different areas (Saint-Andre et al., 2005; Montagu et al., 2005; Williams et al., 2005). The equations can be species specific when applied to one species, or general when developed for more species as in forests or woodlands of mixed species (Williams et al., 2005). Very few allometric equations exist for sub-Saharan Africa (Henry et al., 2011). The equations have only been very rarely used in agroforestry systems where biomass from trees grown in crop fields is estimated (Lott et al., 2003). Even in forestry, such equations are usually used on trees managed under planted systems and are less applied to trees from coppice stands (Antonio et al., 2007).

Unfortunately, allometric equations often produce biased results when applied outside the forest area or region where they were developed (Litton, 2008; Mwakalukwa et al., 2014). Direct measurement of plant biomass is by destructive sampling and is a tedious and time-consuming process. If high accuracy is required, it is recommended to develop local biomass and volume equations or at least to harvest and measure a few trees, representing the range of tree sizes typically found in the forest, and use these to check the validity of the applied equation under local conditions (Brown, 2002; Henry et al., 2011). Once an allometric model is developed, biomass estimation will only need a set of few sample trees for validation so as to apply it in a given area.

Although eucalypts are very important in agroforestry and plantation forestry in Rwanda, no readily available models exist locally that can be used to estimate biomass. Only a limited number of species specific volume equations were developed for different eucalypt species

but these were based on data collected from a few 0.25ha plots from one small, experimental location, Ruhande Arboretum (Burren, 1995). This paper presents allometric models for estimating whole tree and tree component biomass from both coppice and planted stands of *E. saligna* in southern Rwanda, where this species dominates. Specific objectives were: a) to develop allometric model(s) for use in predicting total tree and tree component biomass applicable to both coppice and planted stands of *Eucalyptus saligna*; b) to explore if the variance increases with tree diameter; c) to study if there are any differences between coppice and planted stands or among sites; and d) to explore ways to improve the model(s).

Materials and methods

Study site

The study was carried out in the southern part of the central plateau agro-ecological zone in southern Rwanda (Delepierre, 1975), where *Eucalyptus saligna* is one of the dominant plantation tree species. According to the classification, the zone is characterised by a bimodal rainfall (March-May and September-December) and the mean annual rainfall is about 1200 mm. Monthly means of daily temperature maxima range from 28.5 °C (April) to 32 °C (September) and corresponding minima from 10 °C (April) to 9 °C (September).

Data sources and biomass estimation

Tree measurements and biomass harvests were taken from woodlots owned by farmers, private organisations and local governments. Due to diverse ownership, it was not possible to get exact information on individual stand background as regards to the establishment (and therefore, the age) and management practices carried out on most of the study woodlots. However, different size classes were ensured during sample selection which assumed to cover a range of age classes available in the area; for the woodlots whose age was known, it ranged from 6-22 years.

Descriptive data for the stands used in this study are summarised in Table 2.1. Destructive sampling was done for all sample stems, with diameter at breast height (D) and total tree height (H) measured prior to felling. Tree cutting was done as close to the ground as possible, but in coppice stands, this was sometimes as high as 20 cm above the ground, depending on the heights of the previous cuttings. All sample trees were separated into stem, branch, bark and leaf components based on the procedure by Montagu et al. (2005), and immediately measured for fresh weight using a spring balance. For each sample tree, sub-samples of each component were taken for dry matter determination in the laboratory at the Faculty of Agriculture of the University of Rwanda, Huye Campus, where they were oven-dried at 100°C until the mass remained constant. To estimate the mean ratios estimator R per site and biomass for a given sample were calculated as follows:

$$R_{cj} = \frac{\sum_{i=1}^{n_j} W_{dry_{ci}}}{\sum_{i=1}^{n_j} W_{fr_{ci}}} \quad 1$$

where $W_{dry_{ci}}$ = the dry weight of the i^{th} sample within site j of component c

$W_{fr_{ci}}$ = the fresh weight of the i^{th} sample within site j of component c

n_j = the number of samples per component within site j

With this ratio, the tree component dry weights were estimated as:

$$\hat{W}_{dry_{cij}} = \hat{R}_{cj} \cdot W_{fr_{cij}}$$

where $\hat{W}_{dry_{cij}}$ = the dry weight of component c of the i^{th} tree within site j

$W_{fr_{cij}}$ = the fresh weight of component c of the i^{th} tree within site j 2

\hat{R}_{cj} = the mean Ratio for component c between dry weight and
fresh weight within site j (estimated with Formula 1)

For simplicity, the dry weight for whole tree will here-in-after denoted as W_{AGB} and that of different tree components of stem, branch, bark and leaf as W_{ST} , W_{BR} , W_{BA} and W_{LF} respectively.

Table 2.1. Tree size ranges expressed as diameter at breast height (D), total height (H) and the above ground biomass (W_{AGB}) recorded in *Eucalyptus saligna* woodlots in southern Rwanda

Site	M ^a	Sample size (n/plot)	Breast height diameter (D) (cm)			Total tree height (H) (m)			Total tree biomass (W_{AGB}) (kg/tree)		
			min	max	mean	min	max	Mean	min	max	Mean
Cyarwa	1	18	12.1	27.8	16.1	15.0	27.2	18.6	116	516	203
Gishamvu II	1	16	8.7	15.6	12.5	13.4	18.3	16.4	61	202	132
Gishamvu III	1	11	8.8	12.8	10.7	12.2	19.4	15.6	63	146	99
Mbazi I	1	14	25.7	30.1	27.8	26.8	28.6	27.6	368	469	413
Mbazi II	1	19	25.7	30.1	27.8	28.1	32.5	29.8	354	576	476
Rango	1	49	9.5	29.2	16.1	12.2	30.9	18.2	41	543	187
Ruhashya I	1	43	16.0	29.6	24.5	19.3	30.5	26.2	214	661	401
Ruhashya II	1	14	9.0	15.9	12.0	11.9	18.7	15.9	69	182	125
Ruhashya III	1	12	11.9	17.2	14.8	13.7	19.1	16.9	70	215	168
Save I	1	37	5.8	26.5	16.6	9.6	28.2	19.8	24	452	235
Sovu	1	50	19.0	29.6	25.3	21.2	31.0	27.1	238	633	376
Gisororo	0	12	17.9	27.4	21.6	19.5	28.0	24.3	247	472	334
Mbazi III	0	15	11.6	19.5	15.2	14.4	22.8	17.6	58	201	120
Save II	0	13	19.0	28.0	22.5	21.1	28.2	24.9	274	498	364

^aManagement practice 1 = coppice and 0 = planted stands.

Allometric equation development

Diameter at breast height (D) and total tree height (H) and their combinations have traditionally been used in the estimation of tree volume and biomass using allometric relationships. This is basically due to two reasons: one, the parameters, especially D , can be easily measured in the field and two; usually conical shaped tree stems constitute a very

significant portion of the whole tree biomass. Basal area alone, or in combination with height therefore, can explain most of the variation in total above ground volume or biomass of a tree. Basically, the models tested in this study were based on those developed many years back by Berkhout (1920); Schumacher and Hall (1933); Hohenadl (1936). These are in line with those used in other places for the same purpose, example, those used by Zianis et al. (2005). The models are of the form:

$$V = \begin{cases} a \cdot D^b & \text{(Berkhout, 1920)} \\ a \cdot D^b \cdot H^c & \text{(Schumacher and Hall, 1933)} \\ a + b \cdot D + c \cdot D^2 & \text{(Hohenadl, 1936)} \end{cases} \quad 3$$

conforming also to the model types used by Whitesell et al. (1988) in Hawaii and Faunweban and Haullier (1997) in Cameroon, and by Burren (1995) and Nduwamungu et al. (2008) in Rwanda, and by Fantu et al. (2010) in Ethiopia.

The following research questions were posed: which model performs best for *E. saligna* W_{AGB} estimation?; does variation increase with increasing tree diameter in which case a regression weight is needed?, and is there any difference(s) between planted and coppice stands or between sites?

Data analysis

Data were analysed using IBM SPSS 22 software. The data set comprised of 323 stems and coppice shoots from 11 and 3 coppice and planted stands respectively. The following five models were fitted to the data.

$$\text{model 1 } W_{AGB} = b_1 \cdot D^{b_2}$$

$$\text{model 2 } W_{AGB} = b_1 \cdot D^{b_2} \cdot H^{c_1}$$

$$\text{model 3 } W_{AGB} = a + b_1 \cdot D + b_2 \cdot D^2$$

$$\text{model 4 } W_{AGB} = a + b_1 \cdot D^{b_2}$$

$$\text{model 5 } W_{AGB} = a + b_1 \cdot D^{b_2} + c_1 \cdot H^{c_2}$$

where W_{AGB} = above ground biomass for the whole tree (kg/tree), D = tree breast height diameter (cm) and H = total tree height (m). The models were used for both the total aboveground biomass (W_{AGB}) and for the different tree component biomass of stem (W_{ST}), branches (W_{BR}), bark (W_{BA}) and leaf (W_{LF}). About 20% of the leaf component was made of flowers, fruits and twigs which were considered as part of the leaf component. When applied to the components, W_{AGB} in the models was then replaced by a specific tree biomass component being tested.

Model testing

All models were judged basing on the coefficient of determination (R^2), but because not all models had the same number of parameters the R^2_{adj} was used. For the analyses, the IBM SPSS 22 software was used. The parameters were estimated in this package by finding parameters under the condition that the sum of the residues is 0 and the sum of the squares of

the residues is minimal. But these conditions give no guarantee that there is no bias within the data range of D and H . This means that problems may arise when extrapolating the function for trees with different dimensions. So an additional criterion is needed and the following model was used to control the diameter related bias.

$$residual_i = W_i - \hat{W}_i = a + b \cdot D_i$$

where W_i = the biomass observation the i^{th} tree

4

\hat{W}_i = the model prediction for the biomass for the i^{th} tree

For a good model, a and b do not significantly differ from 0. Also, residuals were plotted against diameter to examine the goodness of fit of the models.

To calculate the model parameters appropriately assumptions had to be made whether equal or unequal variance is expected. Cunia (1964) stated that all biomass equations are heteroscedastic and he advised to use a regression weight, $rgw = D^4$. Thus,

$$W_i = \text{model}_i + \text{error}_i$$

where $\text{error}_i = D_i^2 \cdot \sigma \cdot \text{standard error}_i$

5

the standard error has expectation = 0 and variance = 1

the variance of $W = D^4 \cdot \sigma^2 = rgw \cdot \sigma^2$

The analysis of the models 1 to 5 was done in the following five steps:

Step 1. First the models were fitted with the data set without any regression weight. To check if there is heteroscedacity, model residuals were plotted against D . By doing this for model 1 in a preliminary analysis a regression weight (rgw) = $D^{3.3}$ was found, the estimated value of the exponent was 3.3 but Cunia's value 4 was within the confidence interval. For the other models, comparable results were found. Since it is not recommended to calculate a regression weight in the same study data set as it will be applied to, Cunia's $rgw = D^4$ was adopted.

Step 2. The models were again fitted with a rgw of D^4 .

Step 3. The relationship between the residuals of models 1 and 2 was examined as a linear function in D .

Step 4. Site factor was introduced to the predicted value of model 2. In addition, quadratic mean diameter (d_q), the mean height (h_m) and the coppice factor, were incorporated in the analysis for coppice stands.

As will be seen in the results section, some sites were more similar than others and in general, two site groups were observed. By splitting the data in the two groups, parameters for the models per group could be calculated. But in that case, it was difficult to judge if the parameters per group really differ. With the help of the auxiliary variable *group*, it was possible to estimate the parameters in one analysis per model. In Formula 6 this is demonstrated for model 1. If the parameters b_{11} and/or b_{21} are significant, the groups have different parameter sets.

1. Model 1 (new): $W = b_1 \cdot D^{b_2}$

where $b_1 = b_{10} + b_{11} \cdot \text{group}$

$b_2 = b_{20} + b_{21} \cdot \text{group}$

6

$\text{group} = \begin{cases} 1 & \text{for sites of Mbazi I, Mbazi II, Mbazi III and Sovu} \\ 0 & \text{for other sites} \end{cases}$

In Formula 7, the same procedure for model 2 was done:

$$\begin{aligned}
 &2. \text{ Model 2 (new): } W = b_1 \cdot D^{b_2} \cdot H^{c_1} \\
 &\quad \text{where } b_1 = b_{10} + b_{11} \cdot \text{group} \\
 &\quad \quad b_2 = b_{20} + b_{21} \cdot \text{group} \\
 &\quad \quad c_1 = c_{10} + c_{11} \cdot \text{group} \\
 &\quad \text{for group see Formula 6}
 \end{aligned}
 \tag{7}$$

For the models 3 to 5 comparable adaptations could be made.

Step 5. Bias correction was done by using the following non-linear model:

$$\begin{aligned}
 &W = cor \cdot \hat{W} \\
 &\quad \text{where } \hat{W} = \text{the predicted value for } W \text{ in a certain group in Formula 6 and 7}
 \end{aligned}
 \tag{8}$$

Analysis of tree biomass components

Tree biomass components of stem, branch, bark and leaf were analysed in the same manner and all the steps above were followed. However, the sum of unbiased estimates of the four components will be unequal to the unbiased estimate of the total dry biomass (as calculated before). Therefore, a much simpler procedure was followed by estimating the fraction of each component with a simple linear fit in D :

$$\begin{aligned}
 &F = W_c / W = a + b \cdot D \\
 &\quad \text{where } F = \text{fraction of the biomass component} \\
 &\quad \quad W_c = \text{dry weight of component } c \\
 &\quad \quad W = \text{the combined total dry weight of the four biomass components.}
 \end{aligned}
 \tag{9}$$

As for the total tree biomass, the site effect was analysed for the different biomass components.

Results

Tree variation and relationship between W and predictor variables

The variation in tree size in the study area was very high and differences were significant between sites but not between management systems. Size ranges for D , H and W_{AGB} were presented in Table 2.1 and tree D ranged from 5.8 to 30.1 cm, H from 9.6 to 32.5 m and W from 24.0 to 633.0 kg/tree.

Model performance comparisons

Plots of total tree biomass as related to tree D and the predicted and observed total tree biomass are presented in Fig. 2.1 and 2.2 respectively. In step 1 of model analysis, it is clear that there is heteroscedacity in the data as observed in the plot of residuals against D for model 1 (Fig. 2.3).

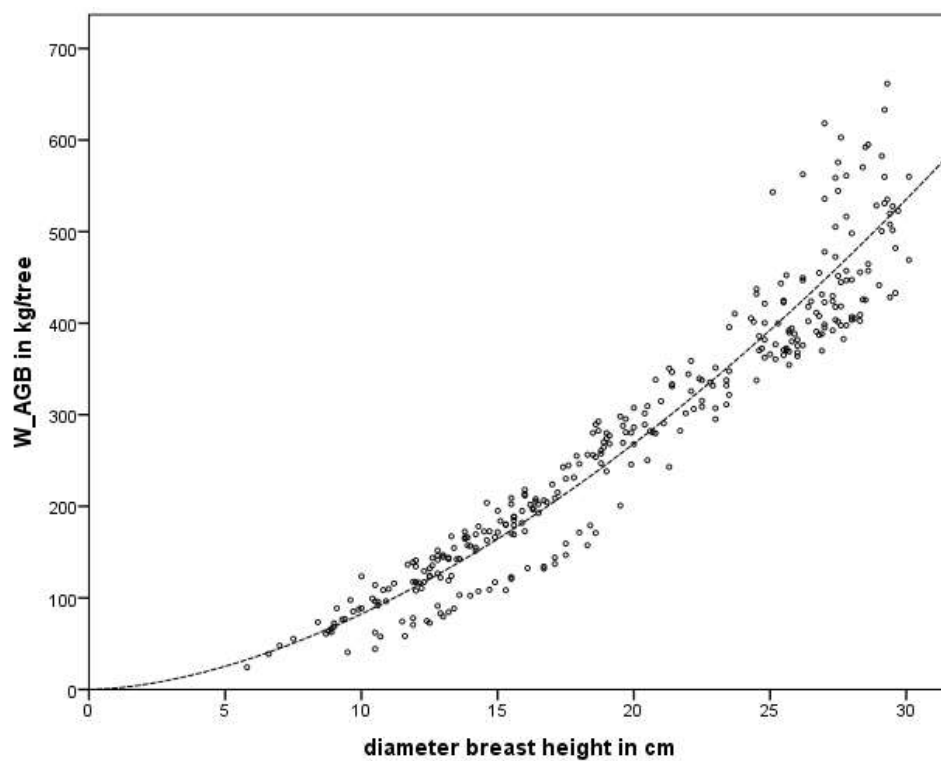


Figure 2.1. Observed total tree biomass as related to tree diameter at breast height in *Eucalyptus saligna* woodlots in southern Rwanda

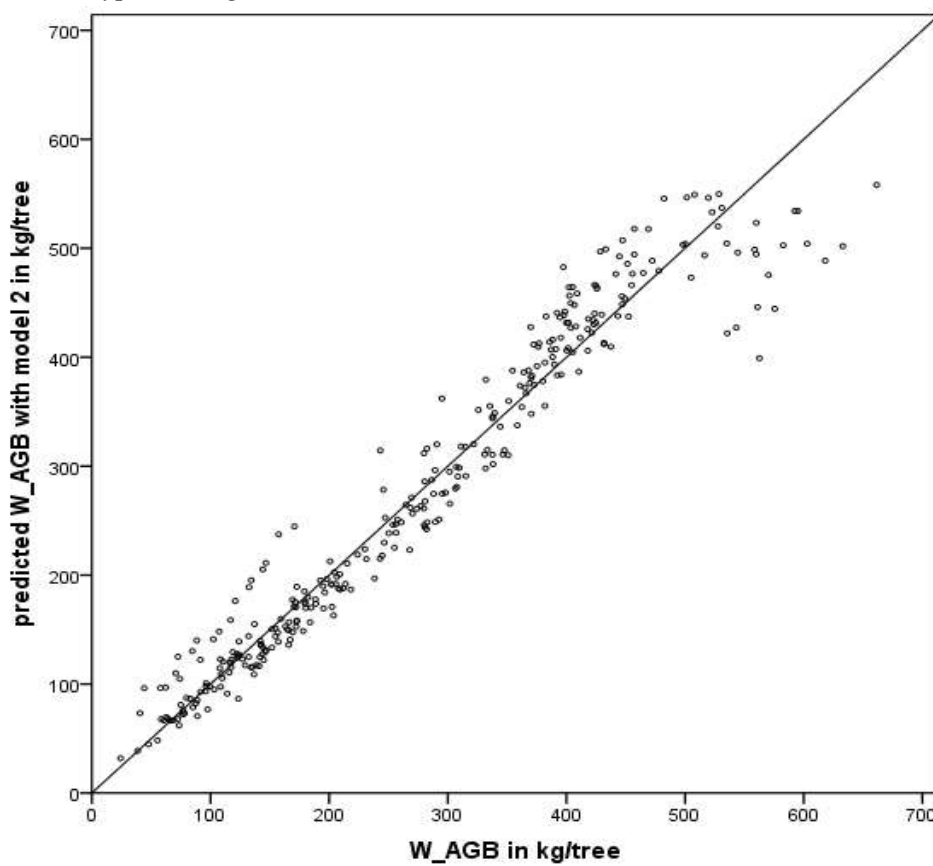


Figure 2.2. Predicted total tree biomass plotted against the observed total tree biomass in *Eucalyptus saligna* woodlots in southern Rwanda

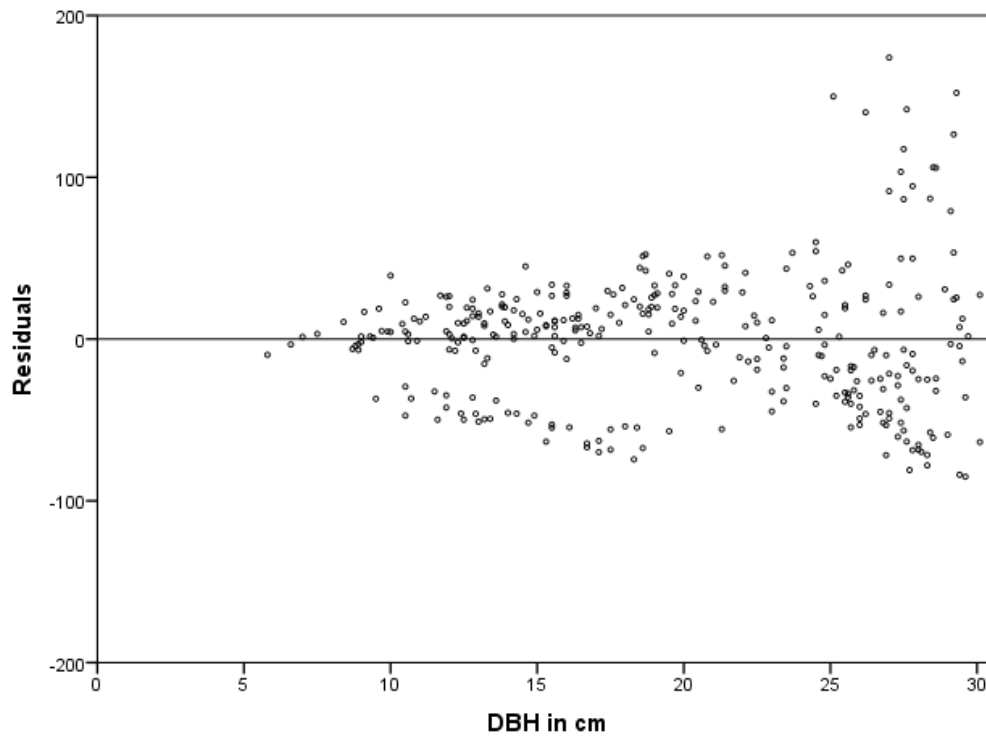


Figure 2.3. Residuals of model 1 in relation to tree diameter clearly show heteroscedacity.

For the other models, comparable results were found. The results of model 2 are presented in Table 2.2. In both steps 1 and 2, the intercepts a of models 4 and 5 were not significant. Also in model 5 the parameter c_1 was not significant and the parameter c_2 was very high (3.259). Model 3 had a rather irrelevant a parameter, because for $D = 0$ and $H = 1.3$ the parameter combination generates a negative weight, but since for tree D above 2.9 cm it generates positive weights, the combination is accepted. This means that model 4 is equivalent to model 1 and therefore the latter was skipped from further analyses.

Table 2.2 Results for the weighted linear and non-linear models for total tree biomass in *E. saligna* woodlots in southern Rwanda

Model	A	b_1	b_2	c_1	c_2	Adj R^2 of y	Adj R^2 of W_{AGB}
1		1.6367	1.7024			0.303	0.921
2		1.0746	1.4085	0.4179		0.328	0.925
3	-21.616	6.4269	0.4028			0.307	0.921
4	-5.997 ^{NS}	2.0270	1.6395			0.306	0.921
5	-12.125 ^{NS}	2.8822	1.4754	0.0016 ^{NS}	3.2589	0.333	0.925

^{NS} denote non-significant values.

Model 5 had no relevant combination of parameters and therefore it was also skipped. On the other hand, if free parameters b_2 and c_2 in Table 2.2 were used with fixed values (for example $b_2 = 2$ and $c_2 = 1.5$), the other parameters would become significant and the model would

therefore be suitable for this study. In this case however, the adjusted R^2 ($= 0.921$) is lower than that obtained in model 2 ($R^2 = 0.925$). The R^2 for all models was rather low since in most biomass studies the R^2 is 0.95 or higher. Model 2 performed best for models that combine D and H but model 1 was the best of the tested models when only D was used. In step 3 the bias as a function of tree diameter (the relationship between the residuals of model 1 and 2) is presented in Fig. 2.4 and was not significantly different from the x-axis.

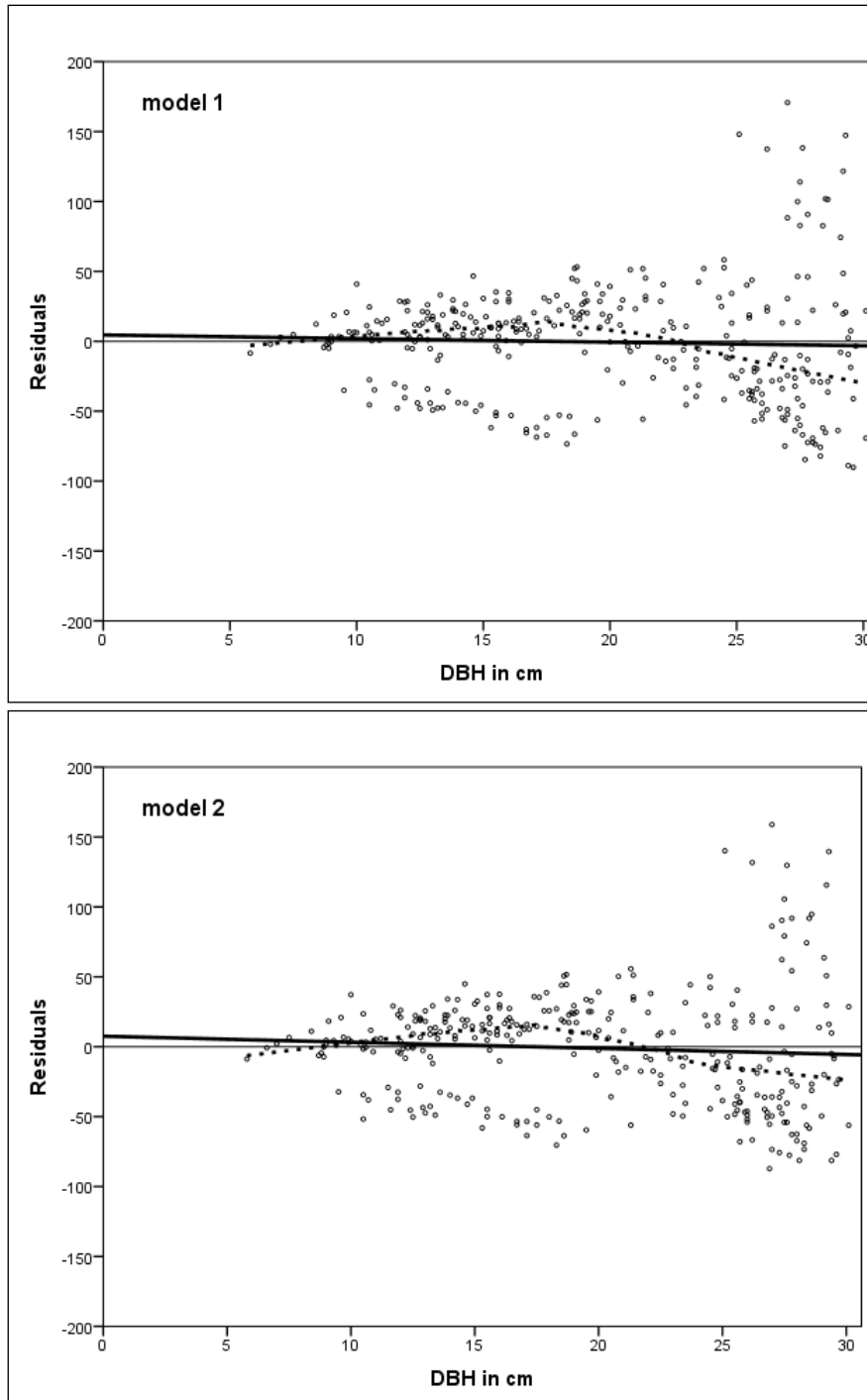


Figure 2.4. Residuals for model 2 (left) and model 3 (right) as a function of the diameter. The bold straight line is the linear fit, the hatched line is a Loes smoothing curve with 50% of the data and an Epanechnikov kernel Smoother in SPSS Graph.

Likewise, a smoothing curve drawn to the data showed the lack of fitness as there was no difference between model 1 and 2. But the curving below and above the x-axis can be related to other factors, such as site effects. The incorporation of site effect in model 2 gave very significant results and the adjusted R^2 was raised to about 0.944. The incorporation of the quadratic mean diameter (d_g) and the mean height (h_m) for coppice stands gave significant outcomes but the adjusted R^2 was only slightly improved (from 0.925 to 0.929) (Table 2.3).

Table 2.3. ANOVA of Test Between-Subjects Effects when model 2 includes site, quadratic mean D, mean H and coppice effects in *Eucalyptus saligna* woodlots in southern Rwanda.

Source	Type III Sum of Squares	Df	Mean Square	F	Sig.
Model	31,607,322	5	6,321,464	4,029.39	0.000
Coppice	498	2	249	0.16	0.853
$W_{AGB_2}^b$	2,312,457	1	2,312,457	1,473.99	0.000
d_g	13,552	1	13,552	8.64	0.004
h_m	6,565	1	6,565	4.18	0.042
Error	498,891	318	1,569		
Total	32106214	323			

^bPredicted tree biomass using model 2

However, this improvement is not good enough to necessitate the use of complicated models with d_g and h_m . The coppice factor did not give significant results (Table 2.4). Biomass prediction basing on site effect gave more than one group in all the sites, and generally, two groups were observed namely: group I. Mbazi and Sovu sites and group II. the other sites (Fig. 2.5 & 2.6). This procedure raised the adjusted R^2 of model 1 from 0.921 to 0.938 when D alone was used in the model and from 0.925 to 0.944 when both D and H were applied. With this procedure, separate values for the parameters of both models for both groups were calculated and they are given for the parameters b_2 and c_1 (Table 2.4). The values for b_1 (corrected to avoid bias through weighted non-linear regression) are also presented in Table 2.4.

Table 2.4. Total tree biomass calculated using models 1 and 2 with and without site grouping by location in *Eucalyptus saligna* woodlots in southern Rwanda

model	Group	b_1	b_2	c_1	Adjusted R^2 of y	Adjusted R^2 of W_{AGB}	σ_y	σ_w
1	Mbazi & Sovu	0.3438	2.1584		0.467	0.938	0.1017	$0.1017 \cdot D^2$
	Other sites	1.4750	1.7527					
	All sites	1.6367	1.7024					
2	Mbazi & Sovu	0.2526	1.8551	0.3917	0.488	0.944	0.0997	$0.0997 \cdot D^2$
	Other sites	0.9703	1.4851	0.3917				
	All sites	1.0746	1.4085	0.4179				

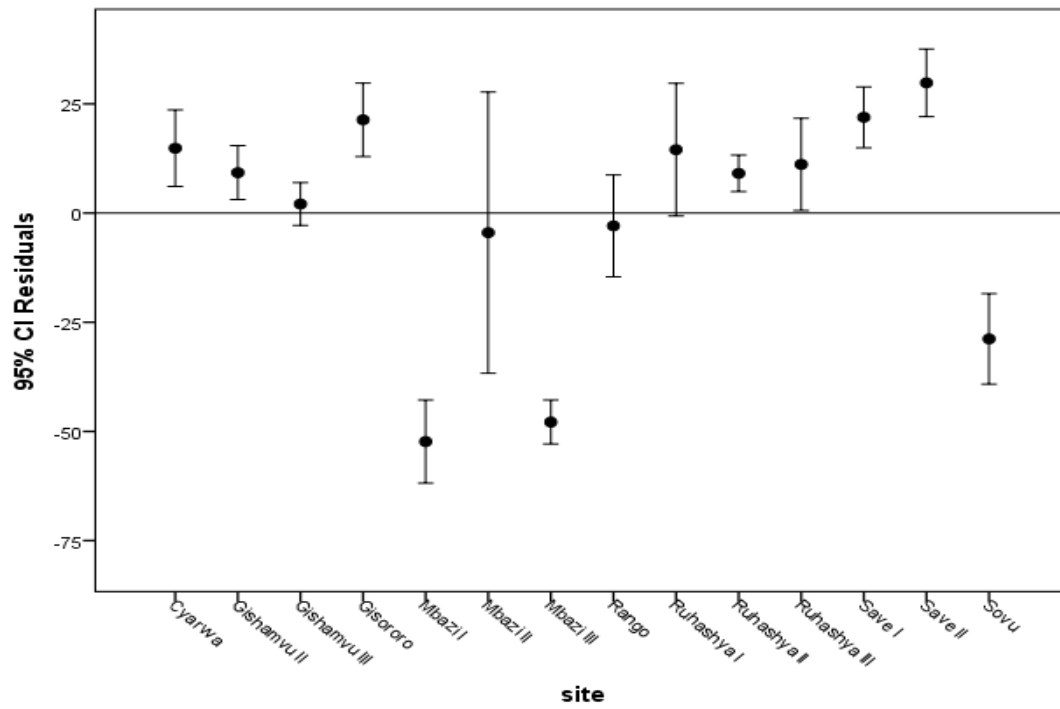


Figure 2.5. Error bar with 95 % confidence interval for residuals with model 2 by site.

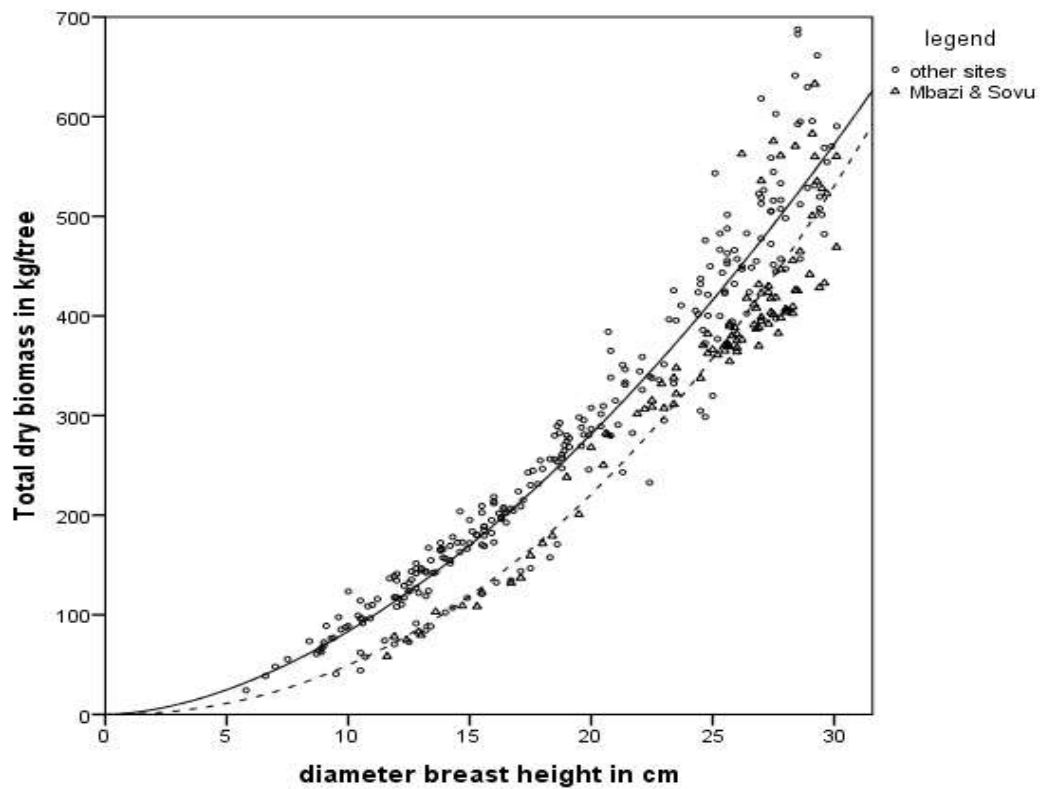
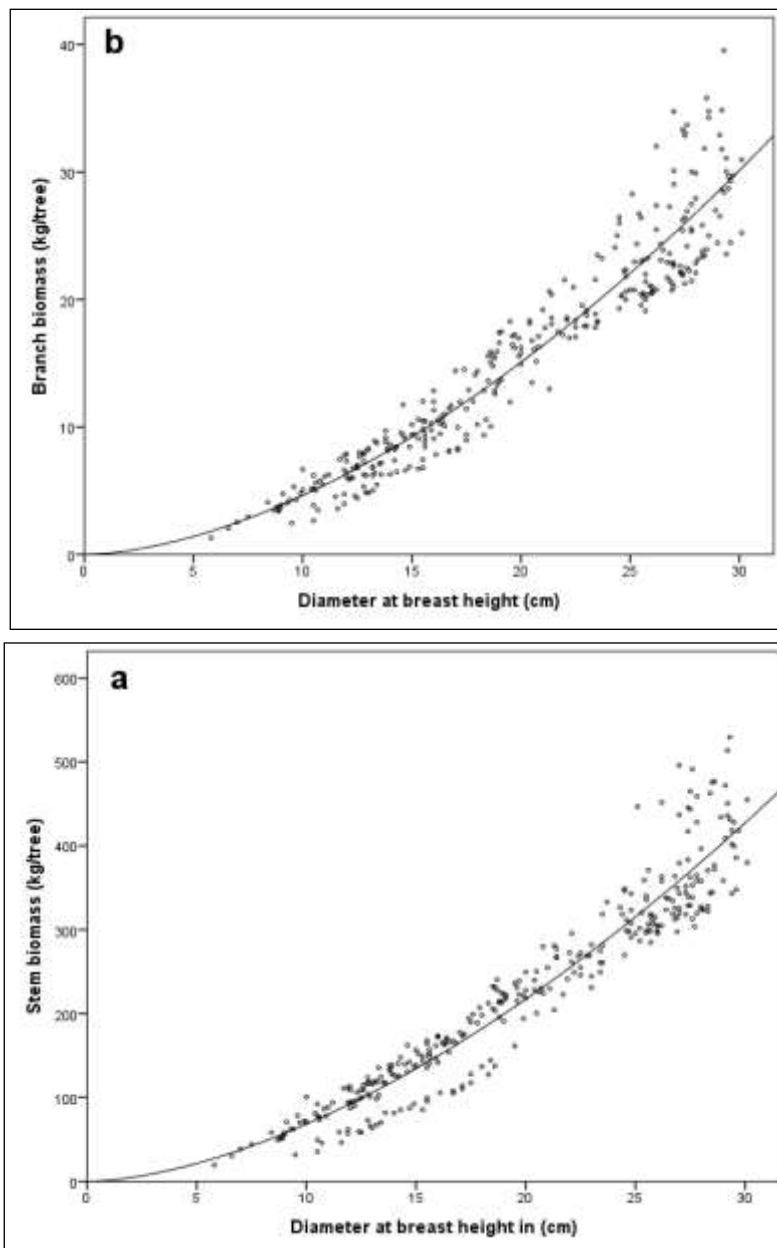


Figure 2.6. Total tree dry biomass as function of the diameter at breast height with the curves of model 1 (the hatched line is for Mbazi & Sovu).

Analysis for the weights of the tree components

The relationship of the different tree biomass components and tree diameter at breast height obtained by applying step 5 above are presented in Figures 2.7a-d while the plots of the observed against predicted values for the same components are presented in Figures 2.8a-d below. In figure 2.9, the observation and the regression line are drawn for the bark component. Also the group effect tried out in an earlier version of equation 9 for total tree biomass was not significant. For this reason, only the means were presented as percentage for the four tree component biomass (Fig. 2.10) and there is hardly any difference when either equation 9 or the means are used to determine total tree biomass. Stem contributed significantly to the total tree above ground biomass production, accounting for about 80% with the remaining proportion being shared among the other tree components.



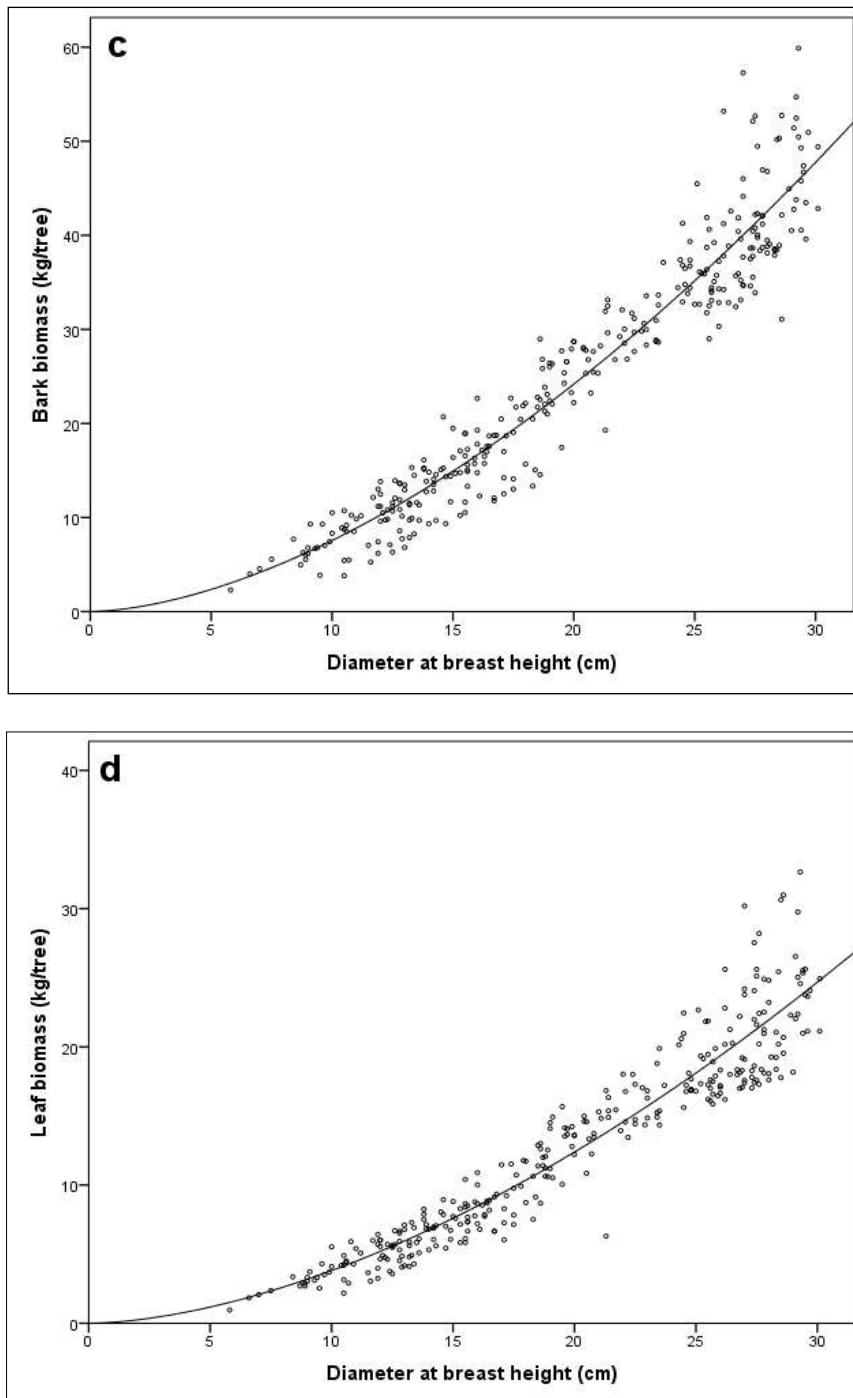


Figure 2.7. Relationship between dry biomass for the stem (a), branch (b), bark (c) and leaf components and tree diameter at breast height in *E. saligna* woodlots in southern Rwanda

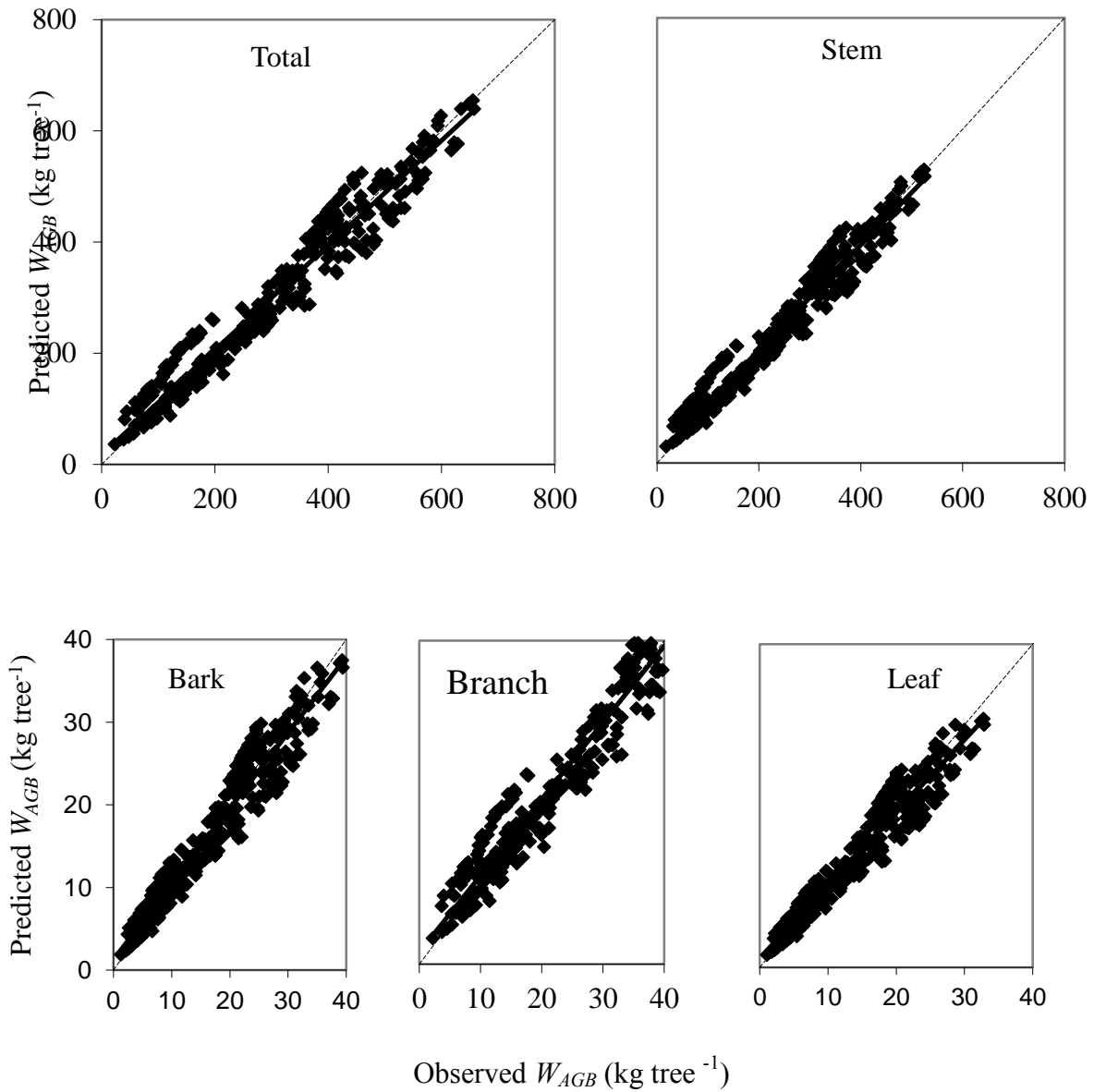


Figure 2.8. Predicted versus observed dry biomass for total tree, stem, branch, bark and leaf components in *E. saligna* woodlots in southern Rwanda

Accuracy of the models

Predictions of total tree biomass (W_{AGB}) and related statistics for the weighted non-linear regression analysis are presented in Table 5 for an average tree ($D = 19.7$ cm, $H = 19.1$ m). So for example such a tree in Mbazi calculated with model 1 and the parameters for site group “Mbazi & Sovu” applied, the predicted total dry weight of $0.3458 \cdot 19.1^{2.1584} = 214$ kg with a 95% confidence interval $\{214 - 1.96 \cdot 0.1017 \cdot 19.7^2; 214 + 1.96 \cdot 0.1017 \cdot 19.7^2\} = \{137; 291\}$, so 214 ± 77 . This may seem to be inaccurate with an error percentage of $\pm 35\%$ for model 2 and 36% for model 1. However, for use in a forest inventory with a minimum of 52 trees, the overall confidence interval (error) will decrease to 5% (Table 2.5).

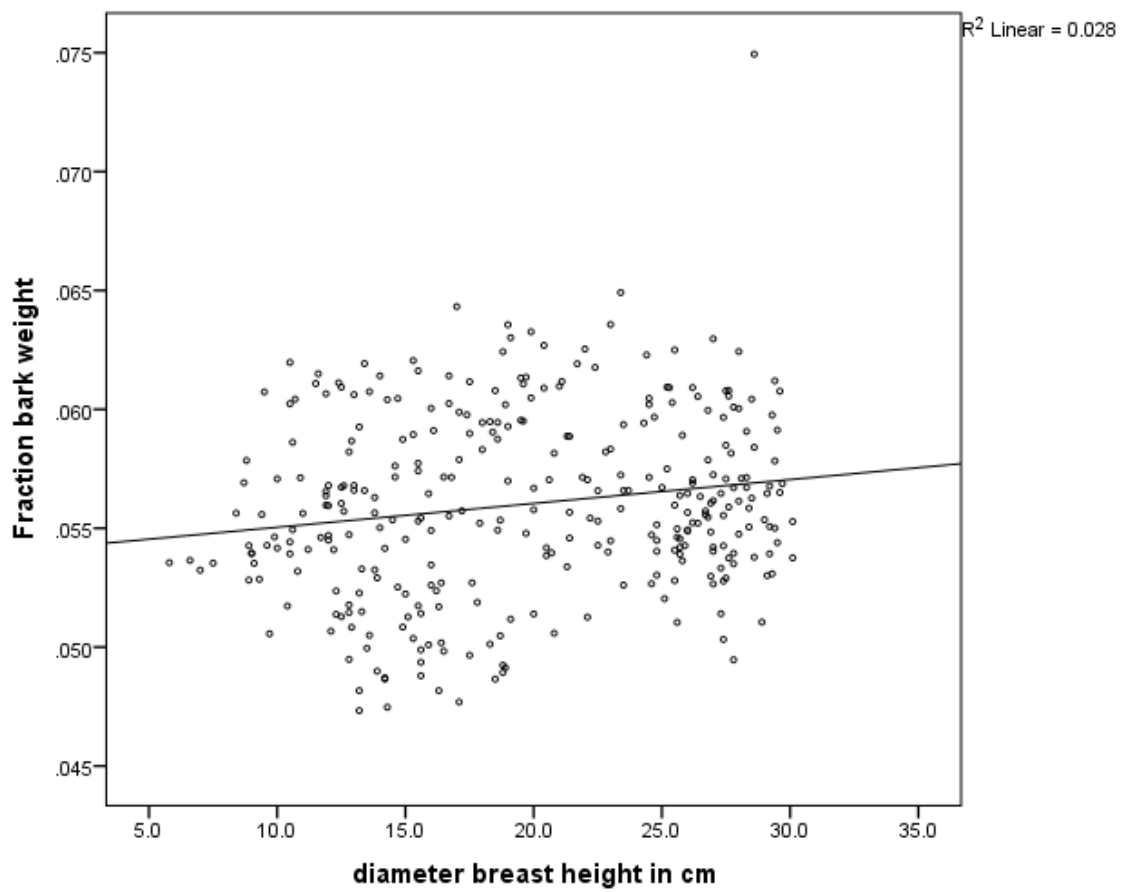


Figure 2.9. Relationship between bark fraction biomass and tree diameter at breast height in *E. saligna* short rotation woodlots in southern Rwanda.

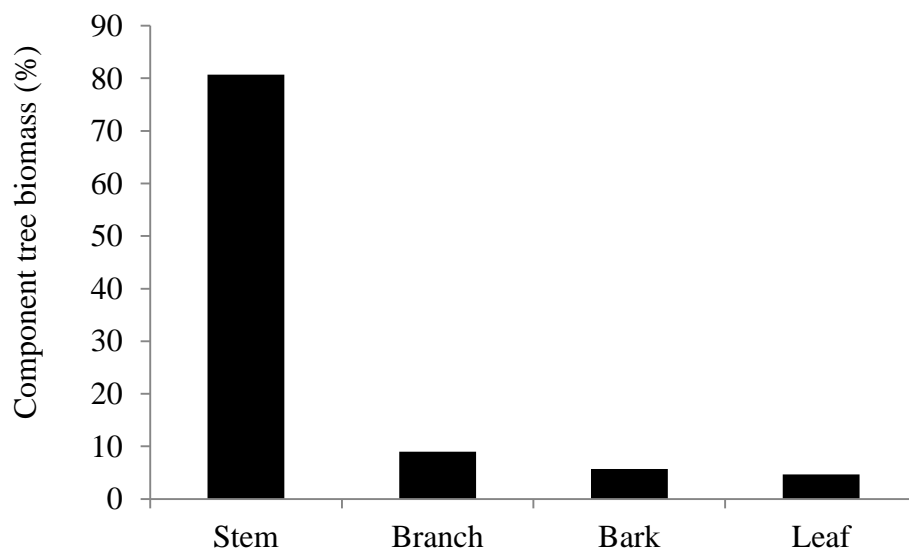


Figure 2.10. Percentage partitioning of total tree biomass by components

Discussion

The present study was done on a variety of trees from both planted and coppiced stands. The observed range in tree size is expected, since the age of trees and coppice shoots also varied. Although no site quality and its effect on tree growth was evaluated, variation is expected to be high due to such differences in topography, position of stands on the hillside, soil types and quality, etc. which varied in different stands in the study area. This variation in environmental factors is reflected in growth and biomass production of *Eucalyptus* (Eldridge et al., 1994; Bouvet et al. 2005).

Table 2.5. Predicted biomass and related statistic values for an average tree with D = 19.7 cm and H = 22.1 m in *Eucalyptus saligna* woodlots in southern Rwanda.

Model	Group	W	σ_w	95% confidence interval	Error %	n (5% error)
1	Mbazi & Sovu	214	39	{137,291}	36%	52
	Other sites	274	39	{197,351}	28%	32
	All sites	262	45	{137,350}	34%	46
2	Mbazi & Sovu	214	39	{138,290}	35%	50
	Other sites	273	39	{197,349}	28%	31
	All sites	261	44	{174,348}	33%	45

The observed variation in tree dimensions in the study woodlots where some coppice stands had shoots bigger than some trees from planted stands may reflect, in addition to tree age and site differences, the varied and changing management objectives. Some managers may establish eucalypts primarily for fuelwood production. Later in the process, they may decide to change management objectives to timber production depending on the market situation or household wealth status or financial needs. This leads to increasing coppice rotation longer than normal and thus, larger coppice shoots. This fact may have contributed to the observed lack of significant differences in biomass production between coppice shoots and planted stand stems. Harrington and Fownes (1993) found that the allometry of woody biomass and leaf area did not differ between planted and coppice stands all studied tree species namely: *Acacia auriculiformis*, *Eucalyptus camaldulensis*, *Gliricidia sepium* and *Leucaena diversifolia*.

Nduwamungu et al. (2008) report that about 75-80% of all eucalypt woodlots in the country are managed as short rotation stands. Precise allometric models have been developed using fewer trees than those used from planted stands in this study (Dudley and Fownes, 1992). In agroforestry systems, some authors report the successful application of a total of as few as 12 trees in developing allometric models for biomass estimation (Dossa et al., 2008). A sample size of ≥ 50 trees in plantation forestry is considered large enough to provide accurate parameter estimates by Coomes and Allen (2009) while Saint-Andre et al. (2005) considers as low as 12 trees to be enough for this purpose. While the data from planted stands for model development used in this study is less than that from coppice stands, it is about four times the acceptable minimum number reported by Saint-Andre et al. (2005) and Dossa et al. (2008).

The main objective of this study was to develop a general allometric model that would be applied to both coppice and planted stands of *E. saligna* in southern Rwanda. The observed site effects and consequent subgroups is not unexpected since species and site specific models are known to be more accurate and efficient predictors of tree biomass than general models combining more than one species or sites. Why the data separates into two subgroups is unclear. All woodlots in Group I (Sovu and Mbazi group) showed consistently lower W_{AGB} than trees in the other group at the same D . Group I sites are geographically located closer to each other compared to the remaining sites and therefore, site conditions may be similar. Again, the woodlots in these sites belonged to Benedictine and Franciscan nuns and were commonly used for free cattle grazing in the past, which may have contributed to the observed similarity in biomass production. This needs further investigation.

High variability in site conditions, stand management, age and tree density negatively influence the precision of a model to predict biomass (Lin et al., 2001). To reduce errors in our study, we opted to apply non-linear regression because it is better suited to data which is likely to have heteroscedacity (Xiao et al., 2011). Heteroscedacity is known to be high for individual trees with large diameter in a population (Litton, 2008). The obtained models in the current study are reasonably accurate irrespective the high variability in site conditions, stand management, age and cutting cycle for short rotation coppice stands. This, according to Medhurst et al. (1999), may imply fitness and the power of the predictive capacity of a model in question.

Using quadratic mean diameter (d_g) and quadratic mean height (h_m) only improved the model efficiency very slightly and thus does not necessitate the use of such complicated procedure to estimate tree W_{AGB} . In our study therefore, only D or the combination of D and H remained the choice parameters to be used. Combining the two parameters however, gave better results than when D alone was used. The two traits are routinely used to predict above ground tree biomass in forestry (De Gier, 2003; Navar, 2010).

Several studies have concluded that the use of D alone for estimating tree W_{AGB} is reasonable since D is the easiest parameter to measure and accounts for most of the variation (Williams et al., 2005). In addition, the use of D alone avoids errors associated with height measurements, which may be up to 10-15% (Landsberg and Waring, 1997; Verwijst & Telenius, 1999; Brown, 2002). Sometimes the use of height as a predictor variable to develop a general allometric model results in an underestimation compared to when D alone is used (Montagu et al., 2005). On the other hand, other authors have found a significant improvement of model fitting when H is also used as a predictor variable (Reed & Tome, 1998; Monserud & Marshall, 1999). Ketterings et al. (2001) suggest the use of height as a predictor variable irrespective the fact that it is tedious to measure and may not explain more of the variance at the sites where the data has originated. The authors justify the incorporation of H saying that it has the advantage of increasing the equation's potential applicability to other sites, as height is usually correlated with site quality. This is supported by the fact that the residuals are usually reduced when both D and H are combined compared to when D alone is used.

Prediction of biomass in the different tree components of stem, branch, bark and leaf were reasonably and accurately predicted by using the same equations as the ones used to estimate total tree biomass. The use of one model for the different tree components makes biomass prediction much easier. Tree component biomass obtained in this study is within the range reported in other places for *E. saligna* (Fantu et al., 2010). This proportionality in biomass production by different tree components seems to be similar in different *Eucalyptus* species as reported by others (Zewdie et al., 2008; Akindele et al., 2010; Razakamanarivo *et al.*, 2012). Similar findings have been reported in terms of sequestered carbon (Montagnini and Nair, 2004; Ryan, 2010; Campoe et al., 2012) assuming carbon sequestration to be 50% of total tree dry biomass (Nair, 2012).

The combination of D and H has been reported to be more important for W_{AGB} estimations in stem-wood than in other tree biomass components. Tree height was reported to improve the accuracy of the stem models but not of the crown models (Lambert et al., 2005; Bi *et al.*, 2004). Pitt and Bell (2004) also found large increases in model predictive capacity for stem biomass estimates but the addition of H was not necessary for estimating needle and branch biomass. It has been argued that the H variable can take into account the effects of stand age, density, and site quality (e.g., Ter-Mikaelian and Parker, 2000). Antonio (2007) reports a similar observation and the same may apply in the present study.

Several models developed in this study generally behaved in a similar manner through different evaluation and validation tests of fitness. The accuracy observed in the prediction of W_{AGB} in total tree and stem component was similar to that reported in other studies (Fantu et al., 2010). The observed under- or over-estimation of W_{AGB} in some models is not surprising as this is also reported in other studies (Akindele et al., 2010) and is known to vary with stand age (Fantu *et al.*, 2010). Precision of carbon stock estimates were observed to decline with plantation age reflecting the variation of tree growth across an estate that occur in its early stages than at older ages (Specht and West, 2003). This may be more expected when dealing with coppice stands, especially when the number of cutting cycles increases (Zewdie et al., 2008).

The application of models developed by others in the study area or elsewhere was not successful. Reasons for this may be that 1) the model by Burren (1995) was developed using data collected from a few trees harvested from a very limited area and was for application in planted stands only without considering coppice shoots; 2) the model by Nduwamungu et al., (2008) was developed from a comparatively larger number of trees specifically for use in planted stands of several eucalypt species and 3) the model by Fantu et al. (2010) was specifically for *E. saligna* planted stands but in an environment (Ethiopia) different from the current study area. Equations from harvests of small numbers of trees may produce good estimates for a given site, but these equations may not extrapolate satisfactorily to other sites (Dudley and Fownes, 1992).

Conclusion

The biomass estimation of *Eucalyptus saligna* in southern Rwanda from both coppice and planted stands using a general allometric model was possible with an adjusted R^2 for the best model of 0.944. The incorporation of quadratic mean diameter and mean height for coppice stands gave significant outcomes but the adjusted R^2 was only slightly improved (from 0.925 to 0.929) and did not justify the use of a more complex model. The variance increased with increasing diameter and this was addressed by applying non-linear regression approach for data analysis and using Cunia's regression weight (rgw) of D^4 . Site differences were observed and stratifying the study site into two subgroups lead to an increase in the adjusted R^2 from 0.921 to 0.938 when D alone was used and from 0.925 to 0.944 when both D and H were applied. With this procedure, separate values for the parameters of both models and for both groups were calculated. The prediction of W_{AGB} of the different tree components of stem, branch, bark and leaf was also possible across the region using the same general models for W_{AGB} in coppice and planted stands. This tool may be very useful in the valuation of tree and tree products since different tree components may present independent market commodities at different periods. While the use of D and H only slightly improved the model precision in biomass predictions, it is advised that, both parameters be used in W_{AGB} estimation for two reasons: 1) due to trees' straight boles and light crowns in *E. saligna* pure stands, tree height measurements is easy and the errors associated with the measurement may be minimal. 2) The use of D and H is reported to be more important when the model is applied to a range of sites including those from outside the area where the model was developed. This may increase the possibility of applying the models developed in this study to parts of the country other than the study area.

CHAPTER 3: Tree-crop interactions in maize-eucalypt woodlot agroforestry systems

CP Mugunga, KE Giller and GMJ Mohren

Tree-crop interactions in maize-eucalypt woodlot agroforestry systems

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Abstract

We studied the interaction between *Eucalyptus saligna* woodlots and maize crop in southern Rwanda. Three sites were selected and in each, a eucalypt woodlot with mature trees and a suitable adjoining crop field of 12.75 m × 30 m was selected. This was split into two plots of 6 m × 12 m and further subdivided into nine sub-plots running parallel to the tree-crop interface. Maize was grown in both 6 m × 12 m plots and one of these received fertiliser. Soil moisture, nutrients and solar radiation were significantly reduced near the woodlot, diminishing grain yield by 80% in the 10.5 m crop-field strip next to the woodlot. This reduction however affects only 10.5% of the maize crop field, leaving 89.5% unaffected. Spreading the loss to a hectare crop field, leads to an actual yield loss of 0.21 t ha⁻¹, equivalent to 8.4%.. Expressing yield loss in tree-crop systems usually presented as a percentage of yield recorded near the trees to that obtained in open areas may be misleading. Actual yields should be reported with corresponding crop field areas affected. Coincidentally, soil moisture, soil N and K increased from the woodlot-maize interface up to 10.5 m and thereafter, remained similar to the values in open areas. Harvest index in unfertilised maize exceeded that in the fertilised treatment reflecting the crop's strategy to allocate resources to grain production under unfavourable conditions. Solar radiation continued to increase with distance up to 18 m from the woodlot-maize interface. Fertilisation increased maize yield from 1.3-2.6 t ha⁻¹ but the trend in the woodlot effects on maize remained unaltered.

Key words *Eucalyptus* woodlots, environmental resources, agroforestry system, tree-crop interface

Introduction

Steady population growth in the tropics has led to an increased demand for agricultural products and for timber and fuelwood. In the eastern and central African highlands, this wood demand has bypassed the capacity of forest to supply wood, and has resulted into the degradation of many natural forests (Burnett, 1985; Thorhaug and Miller, 1986). A possible means to reduce pressure on natural forests and meet the wood and food demand is to include trees in the farming systems (Chamshama, 2011). To provide fuelwood and timber, establishment of eucalypt plantations was promoted since the colonial era (Munyarugerero, 1985). *Eucalyptus* was widely adopted, and in 2008, 64% of the total plantation area (102,800 ha) in Rwanda was covered by eucalypts (Nduwamungu et al., 2008). Most of the plantations are owned by local governments (65% of the area), followed by smallholder farmers (26%) and private institutions (9%). The plantations mainly consist of small woodlots and occur in all ecological zones, with more plantations in the southern and western Provinces (Nduwamungu et al., 2008).

Food production is limited by the availability of land for annual crops (Hauser, 2006). A continuous growing population led in Rwanda to land fragmentation and small average land holdings per household of around 0.5 ha (Bucagu et al., 2014; Mpyisi et al., 2003). The urge to be self-sufficient in wood products drives farmers to grow trees on their farms. As a result, small crop fields and eucalypt woodlots are intermixed in a mosaic in the landscape. Because of the proximity of crops and trees, and the tree-crop interactions, this can be classified as an agroforestry system, containing some features of short-rotation woodlots.

In agroforestry systems, trees and crops are commonly grown in different spatial arrangements where trees provide various benefits to the crops. Windbreaks for example are beneficial to crops, especially in dry areas where they are useful for the control of wind erosion, soil conservation, and the amelioration of the microclimate (König, 1992; Mayus et al., 1999). Trees grown in hedgerow intercropping systems may improve soil conditions through nitrogen fixation and increase crop yield (Akyeampong et al., 1995). However, trees also compete with crops for environmental resources and their negative effects may outweigh the potential benefits (Rowe et al., 2005).

Eucalyptus trees may be such a mixed blessing; they are fast growing even on marginal sites, produce a wide range of valuable products in short periods, are easy to manage and can be coppiced several times giving several yields at one planting (Casson, 1997). They are however, known to be especially competitive, and to significantly reduce productivity of companion plants. They may severely compete with crops for soil moisture (Kidanu, 2005) and soil nutrients (Harrison et al., 2000), and are thought to have allelopathic effects on crops growing in their vicinity (Lisanework and Michele, 1993).

Low crop yields are common on the highly weathered and nutrient poor soils of the study area (Rushemuka et al., 2014), constraining food production and food security. Although food security has been improving significantly in Rwanda, 21% of households were still food insecure in 2012 (WFP, 2012). The dominance of eucalypt woodlots in the agricultural landscape calls for a need to better understand the effects of the trees on crop production and food insecurity. The woodlots are expected to influence crop production in the agroforestry systems. Relevant studies have been done in tropical countries (Kidanu et al., 2005; Sudmeyer et al., 2015). However, these studies evaluated the effects of the spatial arrangement of tree lines rather than the effects of the woodlots. The latter is the most common in Rwandan farming systems.

The aim of this study is to analyse the eucalypt woodlot-maize system, and to quantify how resource availability (water, nutrients and light) and crop performance (height growth, stover production and grain yield) vary with the distance from the eucalypt woodlots. Fertilization was added to analyse whether eucalypts and maize mainly compete for nutrients, and whether fertilization can mitigate for the negative woodlot effects.

Methods

Site description

Three sites were selected in southern Rwanda, with two (Cyarwa and Mukura) located in the periphery of Butare town, Huye district, and the other (Save) in the neighbouring district of Gisagara. More specifically, Cyarwa is located at 2.61591S and 29.76660E at 1655 m asl elevation; Mukura at 2.64804S and 29.73008E with the altitude of 1667 m asl and Save at 2.57384S and 29.77603E and 1694 m elevation. According to the agro-ecological classification defined by Delepierre (1975), the sites are located in the central plateau and hills zone of Rwanda. The area has a mean annual rainfall of 1200 mm y^{-1} ; mean annual temperature of 21°C and the soils are derived from granitic rocks and are classified as ferralsols in the hills (Birasa *et al.*, 1990).

The choice of sites and study woodlots was based on the presence of a woodlot with mature trees of *Eucalyptus saligna*, with an adjacent crop field. One woodlot was selected in each of the three sites. To ensure representative coverage of the study zone, the three sites used were located at least five km apart.

Maize Katumani variety was grown in the crop fields adjacent to *Eucalyptus* woodlots. The woodlot was located on the eastern side of the crop fields in two sites (Cyarwa and Mukura) and on the northern side of the crop field in the Save site.

Experimental design and field procedures

The experiment was conducted in three consecutive seasons, that is, from January-May and September-December 2007 and January-May 2008. Characteristics of the woodlots used are provided in Table 3.1. In each of the three experimental sites, three crop field replications adjoining one *E. saligna* woodlot were selected and crop plots of 12.75 m \times 30 m were pegged. Each replication was split into two smaller plots of 6 m \times 30 m separated by 0.75 cm, a space equal to the interline spacing; and further subdivided into subplots with the following dimensions running parallel to the tree-crop interface into the farmland: 0-2, 2-4, 4-6, 6-9, 9-12, 12-16, 16-20, 20-25 and 25-30 m (Fig. 3.1). One of the two 6 m \times 30 m plots received NPK composite fertiliser at a rate of 300 kg ha^{-1} , equivalent to 51 kg of N, 22.2 kg of P and 42.3 kg of K per hectare. Sites were tilled manually and sown with maize on 25 February 2007 in the first season, on 30 September 2007 in the second season and on 26 January 2008 in the third season.

Abnormally little and erratic rains in the first season led to poor establishment and the experiments were replanted at the end of March 2007. The spacing was 0.5 m between plants and 0.75 m between lines. Three maize seeds were sown per hole and these were later thinned to leave two plants per hole. The plots that received fertiliser at sowing were top dressed with an N based urea (46%) at the rate of 100 kg ha^{-1} (MINAGRI, 2000). To avoid edge effects, two guard (outer perimeter) rows were not considered for assessment, leaving the six innermost maize rows in each subplot for evaluation. It was assumed that the effect of trees did not extend as far as 25 m from the tree-crop interface; the farthest sub-plot located at 25-30 m from eucalypt woodlot was therefore taken as a control.

Table 3.1. Tree sizes in terms of diameter at breast height, total tree height and the above ground biomass recorded in the three *Eucalyptus saligna* woodlots in eucalypt-maize agroforestry systems in southern Rwanda

Site	Density (Stems ha ⁻¹)	Breast diameter (cm)			Total tree height (m)			Total biomass (kg tree ⁻¹)		
		min	max	mean	Min	max	mean	min	Max	Mean
Cyarwa	970	15.1	24.8	19.8	15.0	23.2	18.6	116.5	376.4	203.0
Mukura II	773	8.7	15.6	12.5	13.4	18.3	16.4	61.0	202.0	132.0
Save I	1005	5.8	26.5	16.6	9.6	28.2	19.8	24.0	452.0	235.0

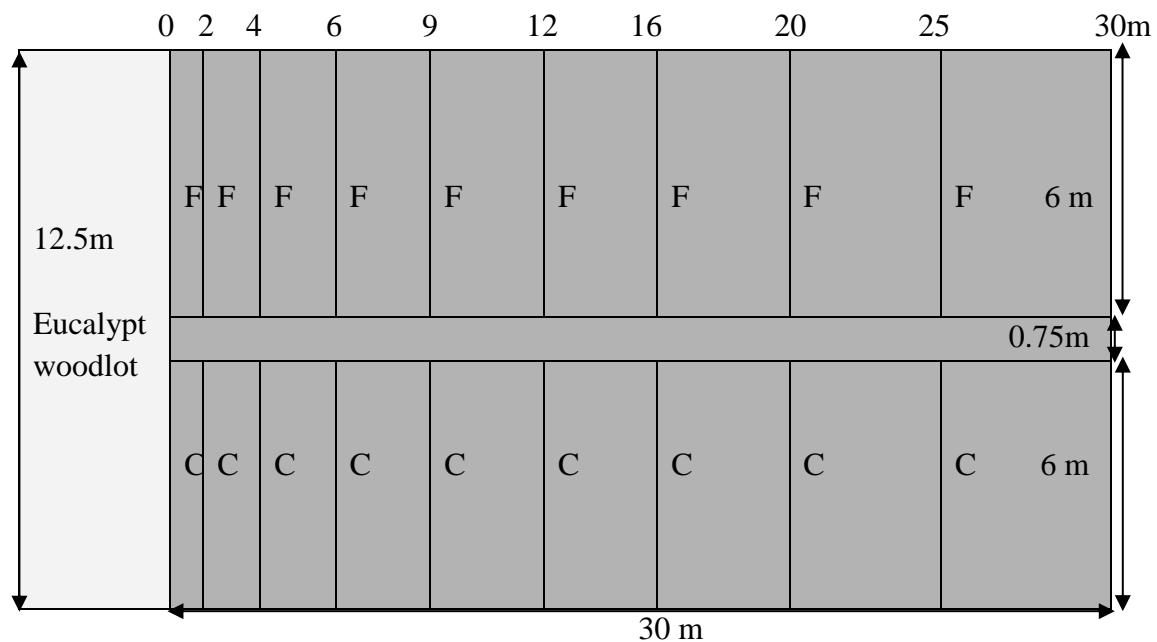


Figure 3.1. Field layout (not drawn to scale) of the eucalypt-maize experiment in Butare, southern Rwanda. F and C denote fertilised and non-fertilised treatments respectively; numbers 0- 30 refer to subplot distances (m) from eucalypt-maize interface

Data collection and analyses

Maize traits studied included growth (height and biomass) and grain yield. Field procedures for each item are described below but in principle, all measurements were done in the nine subplots explained above. For solar radiation, an extra location was added at 5 m inside the woodlot from the tree-crop interface to capture the light intensity under the woodlot canopy.

Soil properties and soil nutrients

Soil samples were collected only once at the beginning of the growing season. According to Schroth and Sinclair (2003), soil properties change most rapidly closer to trees and therefore, it is advisable to collect samples at smaller distance intervals here and increase the intervals with increasing tree distance. For this reason, samples were collected from the crop fields using a graduated auger at the midpoints of each of the nine sampling points corresponding to

1, 3, 5, 7.5, 10.5, 14, 18, 22.5 and 27.5 m from tree-crop line respectively. Simultaneously, soil samples were taken in each sampling location at four soil depths of 10, 30, 50, and 70 cm representing 0-20, 20-40, 40-60 and 60-80 cm respectively below the soil surface. The soils were later analysed for macronutrients (N, P and K), particle size distribution and pH using standard procedures (Okalebo et al., 2002).

Soil moisture

Soil moisture was assessed three times per season: at the beginning, at the mid-season and at the end and was repeated for three consecutive seasons. Soil moisture was determined at the nine sampling points by gravimetric method. Samples for soil moisture assessment were taken at the same four soil depths explained above. Soil cores were taken at every assessment occasion using steel cylinders of uniform size. The cylinders were forced into an intact soil profile at the point of sampling, completely filled with soil and immediately removed and wrapped in an aluminium foil to conserve moisture. The samples were labelled and moisture contents determined in the laboratory after drying samples to constant mass at 105 °C.

Light measurements were assessed by using LI-COR quantum sensors (Li-COR 1000). At every measurement occasion, one sensor was placed in each of the nine subplots. Data were recorded every 30 minutes from 6.30 am to 6.30 pm in terms of photosynthetically photon flux density (PPFD) ($\mu\text{mol m}^{-2} \text{s}^{-1}$). At the beginning of the season when maize seeds were not germinated or when maize plants were still very young and very short, sensors were placed on the ground. Later after the maize plants grew taller, sensors were placed on wooden sticks above the maize plants to avoid any shading other than from eucalypt trees.

Light measurements

Maize growth

Maize plants were measured for total height once in every season at approximately 50% tasselling time which corresponded to the end of vegetative growth. For this purpose, three sample plants selected randomly from the innermost rows in each subplot were measured. Height was measured from root collar to the tip of the sample plant by using a graduated measuring staff.

Maize stover

Above ground biomass was estimated from the same maize plants which were sampled for the assessment of total height since the sampling. Above ground parts for the sample plants were carefully detached from below ground parts in the field at root collar by using a sharp machete. Each sample plant was chopped carefully and immediately wrapped in an aluminium foil for oven dry weight determination in the laboratory where sample plants were oven dried at 65°C to constant weight.

Grain yield

Maize grain yield were obtained only during the second and third season since season one was severely affected by the halt of rains experienced during the period. Maize were harvested from all plots after they were fully mature and had started drying. Cobs were collected, separated from all leaves and air dried. Grain was separated from cobs using a threshing machine and was weighed using an electronic balance at the Faculty of Agriculture laboratory, the University of Rwanda, Huye Campus. The yield was recorded for each subplot. Harvesting index was then computed as the ratio of the grain dry weight to the dry weight of grain yield plus stover. Air dried grain yield was converted to dry matter by multiplying by a factor of 0.85.

Data analysis

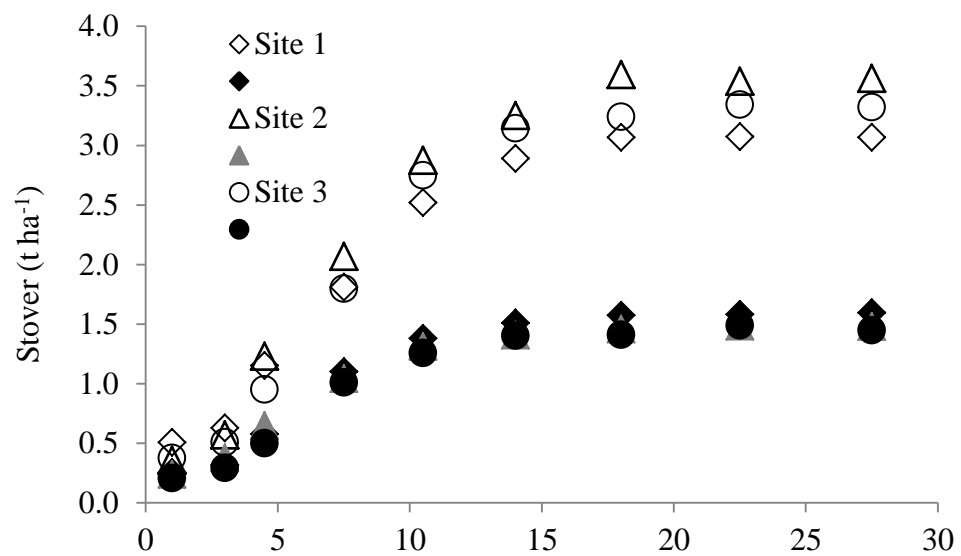
Data were analysed using IBM SPSS 22 software where one way ANOVA was used to study the differences between treatments and LSD test was used to separate statistically different means. Pairwise comparisons using t-tests were used to study the effect of fertilisation because only two levels of fertiliser were used in this experiment. Similarly, grain yield data were available for seasons two and three only, as the poor rainfall in the first season prevented any harvest.

Results

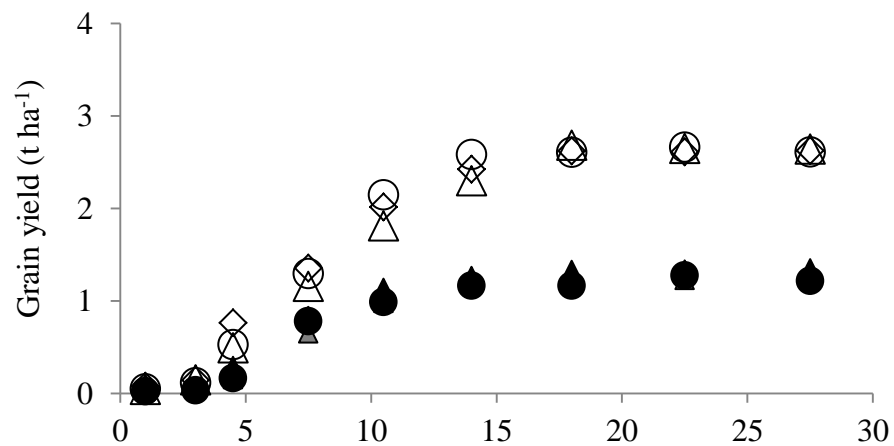
Maize height, stover and grain yield varied significantly ($P < 0.001$) with distance from eucalypt woodlots, fertilization, and often with season, but they did not vary significantly ($P > 0.05$) with site or crop field (Table 3.2; Fig. 3.2a-c). Maize harvest index varied only significantly with distance from the woodlots and not between sites or seasons. Maize height growth, stover, grain yield and the harvesting index in the 9-12 m subplot (10.5 m) from the woodlot increased by 65.7, 74.1, 97.2 and 81.5 % respectively compared to the values at 1 m (in the 0-2 m). On average, grain yield in this zone ($10.5 \times 12 \text{ m}^2$ next to the woodlot) was 0.0063 t while that in the remaining crop field ($19.5 \times 12 \text{ m}^2$) was 0.0585 t. These yields are equivalent to 0.5 and 2.5 t ha⁻¹ for the crop zones affected and that not affected by eucalypt woodlot respectively, making the edge effect prone zone to produce only 20% of the yield obtained in the open areas (27.5 m away). Beyond 10.5 m, the effect of the woodlot ceased to be evident, and maize performance did not differ significantly from the values in open areas anymore. Harvest index for the unfertilised maize was higher than that obtained in fertilised treatments in all three sites (Fig. 3.2c). Maize height was strongly and positively correlated with grain yield and this was observed to be similar in all the three sites (Fig. 3.3). The addition of fertiliser doubled maize growth and the grain yield (Fig. 3.2) but did not alter the trend in the eucalypt woodlot effects on maize performance.

Table 3.2. Probability values of ANOVA procedure for the test of significance for maize growth in a maize-eucalypt woodlot cropping system in three sites in southern Rwanda. Distance in the Table below refers to distance from eucalypt woodlots.

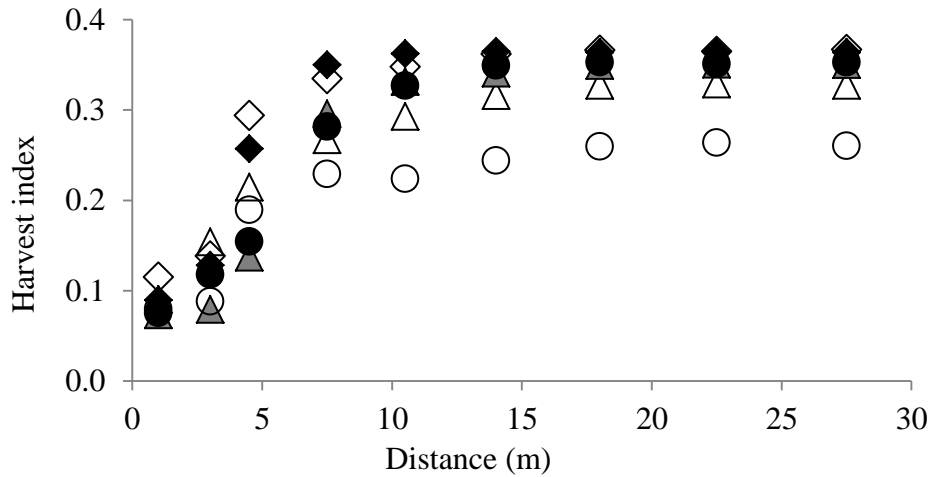
Source of variation	Maize performance			
	Height (m)	Stover (m)	Grain yield (Mg ha ⁻¹)	HI
Site	ns	Ns	Ns	Ns
Season	*	***	ns	Ns
Crop field	ns	Ns	Ns	Ns
Fertiliser dose	***	***	***	Ns
Distance (m)	***	***	***	***



(a)



(b)



(c)

Figure 3.2. Three seasons average variation of Katumani maize variety in stover (a), grain yield (b) and harvest index (c) with distance from *Eucalyptus saligna* woodlots a eucalypt-maize cropping system in three sites in southern Rwanda. Empty and filled marks denote fertilized and unfertilized treatments respectively.

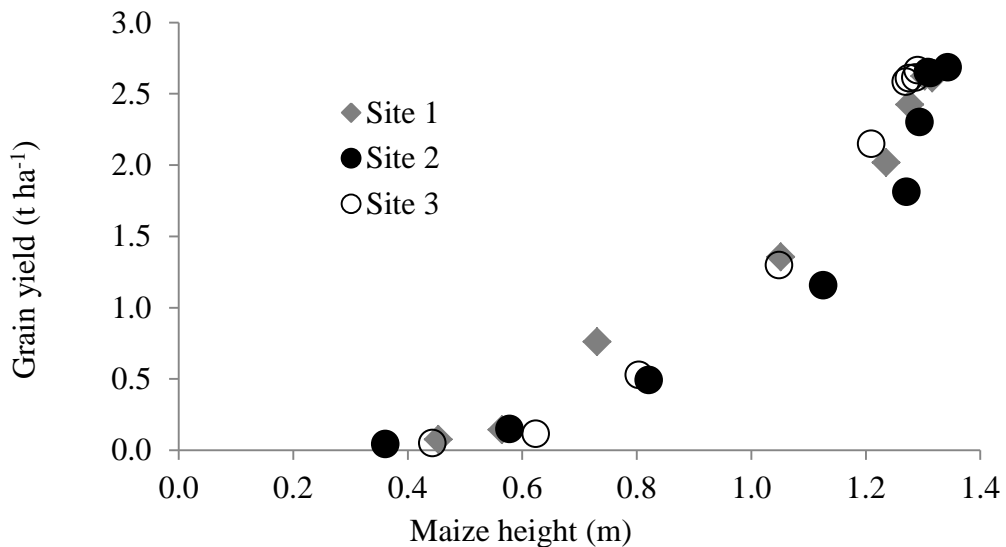


Figure 3.3. Relationship between maize grain yield and maize height observed in a eucalypt woodlot-maize cropping system in three sites in southern Rwanda.

Both solar radiation and soil moisture varied significantly with distance from eucalypt woodlots as well as between sites (Figs. 3.4 & 3.5), and often with seasons and dates of measurements (Table 3.3). Moisture content and solar radiation were 18 and 68% respectively lower near the woodlots compared to the values in open areas. Soil nutrients varied significantly ($P < 0.001$) and increased with distance from eucalypt woodlot (Fig. 3.6). Concentrations near the woodlot were smaller than those recorded in the open areas by 38% for N, 27% for P and 38% for K. Significant effects of the woodlot were detected up to 7.5 m for soil P, 10.5 m for soil N and K and soil water; and 18 m for solar radiation.

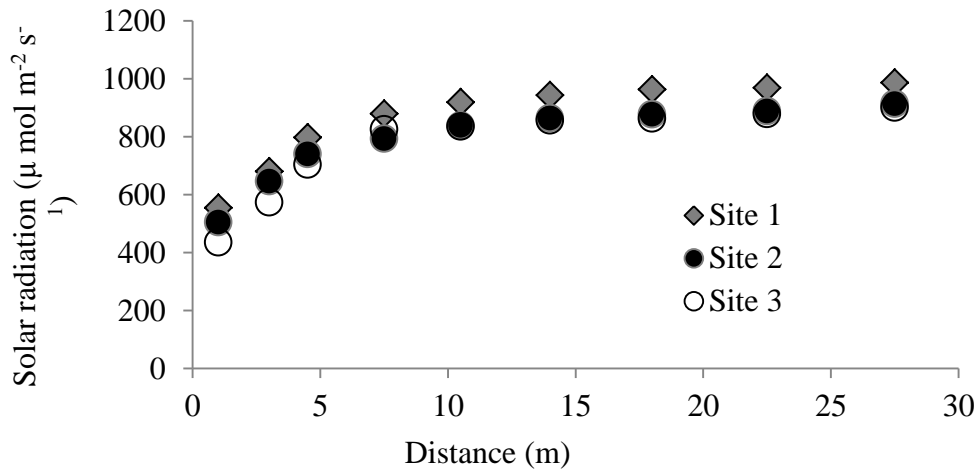


Figure 3.4. Variation of solar radiation (photosynthetic photon flux density) with distance from *Eucalyptus saligna* woodlots in a eucalypt-maize cropping system in three sites in southern Rwanda.

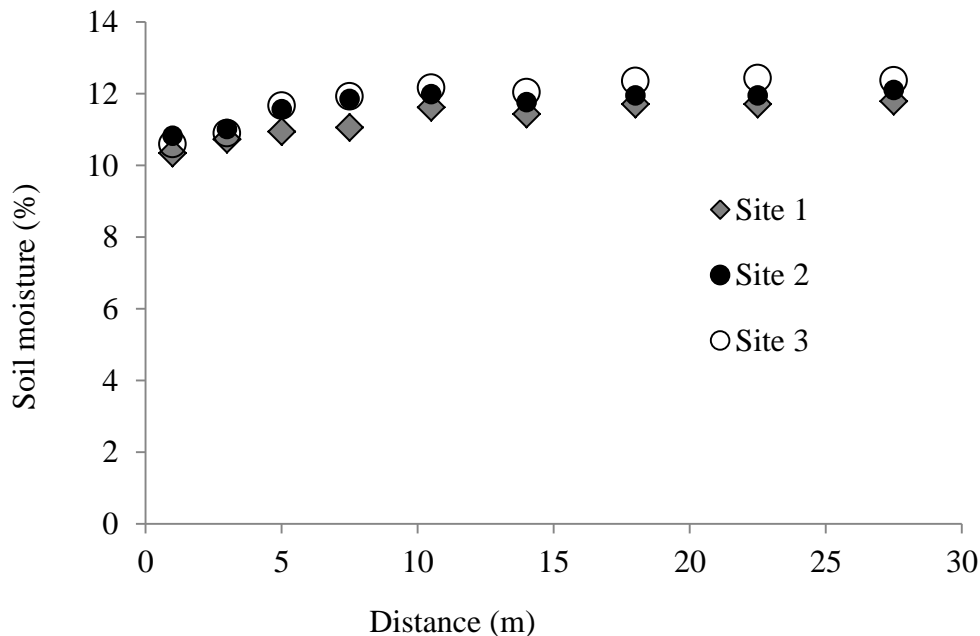


Figure 3.5. Variation of soil moisture in maize crop fields with distance from *Eucalyptus saligna* woodlots in a eucalypt-maize cropping system in their sites in southern Rwanda.

Soil pH was 4.67 at the woodlot edge and increased significantly with distance from eucalypt woodlot, reaching 4.88 at 27.5 m away from the woodlot. However, the difference was only 4% (Fig. 3.7). Clay concentration increased with distance from eucalypt woodlot as did the sand proportion, although only slightly, while silt concentration decreased with distance from the woodlot (Fig. 3.8). Clay and sand proportions near the woodlots were 30 and 4% respectively smaller near the woodlot than in the open areas, while silt was 9% higher near the woodlot than in the open areas. However, the variations found in soil particle distribution may not have been influenced by the woodlot but rather the site characteristics. Soil nutrients,

soil particle size distribution and soil pH varied significantly ($P<0.001$) between sites and soil moisture, soil nutrients and soil pH did so with soil depth (Table 3.4). Soils near the woodlots were therefore slightly more acidic and nutrient-poor, and contained less water than soils far away. Probability values of ANOVA procedure for the test of significance for the different parameters and the interaction between them are presented in Table 3.5. Mostly, there were no interactions between different parameters.

Table 3.3. Variation of soil moisture and solar radiation between seasons and dates of measurements in a eucalypt-maize cropping system in three sites in southern Rwanda.

	Season			Date			
	Jan-May 2007	Sep-Dec 2007	Jan-May 2008	Beginning of season	Mid- season	End of season	
Soil moisture (%)	10.8	13.5	13.8	12.6	12.8	12.7	
Significance level		***			**		
Solar radiation ($\text{umol m}^{-2} \text{s}^{-1}$)	756.1	794.5	699.8	789.7	711.4	749.2	
Significance level		***			***		

** and *** denote significance differences between means at $P<0.001$ and $P <0.0001$ respectively

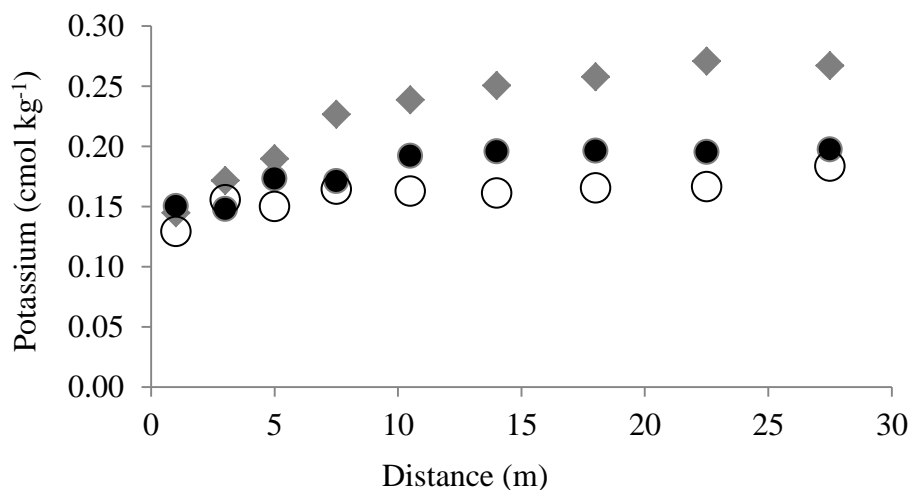
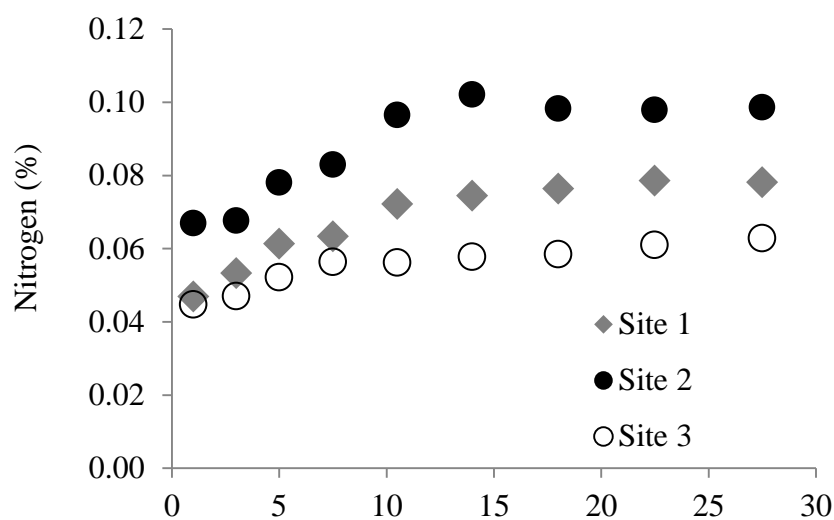


Figure 3.6. Variation of soil nutrients with distance from *Eucalyptus saligna* woodlot in a eucalypt-maize cropping system in three sites in southern Rwanda.

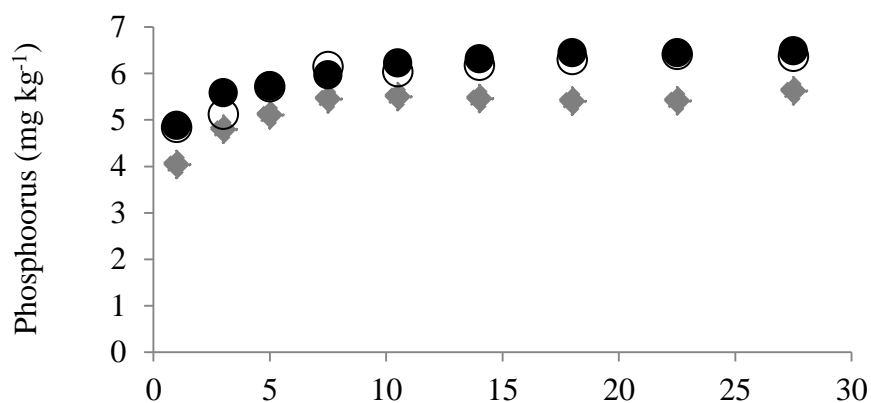
Table 3.4. Site variation in soil nutrients, particle size distribution and soil pH; and variation of soil nutrients, soil moisture and soil pH with soil depth in a eucalypt-maize agroforestry system in three sites in southern Rwanda.

Soil Nutrients				Particles (%)			pH
(a)Site	N (%)	P (mg kg ⁻¹)	K (cmol kg ⁻¹)	Clay	Silt	Sand	
Save	0.06	5.05	0.22	23.0	65.4	11.6	4.7
Mukura	0.08	5.90	0.18	27.1	61.7	11.2	4.9
Cyarwa	0.05	5.79	0.16	20.3	67.7	12.0	4.6
Significance level	*	***	***	***	***	ns	***

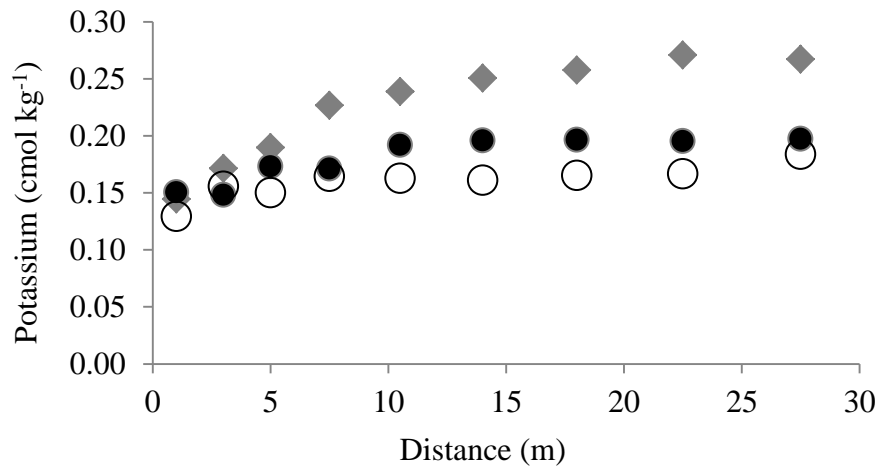
Soil moisture				pH		
Soil depth(cm)	(% by volume)					
0-20	0.09	7.23	0.24	11.6	4.9	
20-40	0.08	6.12	0.22	12.3	4.8	
40-60	0.07	5.01	0.16	13.1	4.7	
60-80	0.05	3.95	0.12	13.7	4.6	
Significance level	***	***	***	***	***	



(a)



(b)



(c)

Figure 3.6. Variation of soil nutrients: (a) Nitrogen, (b) Phosphorus and (c) Potassium with distance from *Eucalyptus saligna* woodlot in a eucalypt-maize cropping system in three sites in southern Rwanda.

Table 3.5. Probability values of ANOVA procedure for the test of significance for soil characteristics and solar radiation in a maize-eucalypt woodlot cropping system in three sites in southern Rwanda. Distance in the Appendix below refers to the location of cropping zones of 0-2, 2-4, 4-6, 6-9, 9-12, 12-16, 16-20, 20-25 and 25-30 m from *Eucalyptus saligna* woodlots. Four soil depths were used: 0-20, 20-40, 40-60 and 60-80 cm below the ground.

Source of variation	Particle size	pH	Moisture	Nutrients			Solar radiation ($\mu\text{mol m}^{-2} \text{s}^{-1}$)
				N (%)	P (ppm)	K (mg kg^{-1})	
Site	***	***	*	***	*	***	***
Season	-	-	***	-	-	-	***
Crop field	ns	*	ns	ns	ns	ns	-
Distance (m)	***	***	***	***	***	***	***
Soil depth (cm)	-	***	***	***	***	***	-
Site * Farm	ns	ns	ns	ns	ns	ns	Ns
Site * Distance	ns	ns	ns	ns	ns	ns	Ns
Farm * Distance	ns	ns	ns	ns	ns	ns	Ns
Site * Crop-field*	ns	ns	ns	ns	ns	ns	Ns
Distance							

ns denotes non-significant while * significant differences at $P < 0.05$ level and *** $P < 0.001$

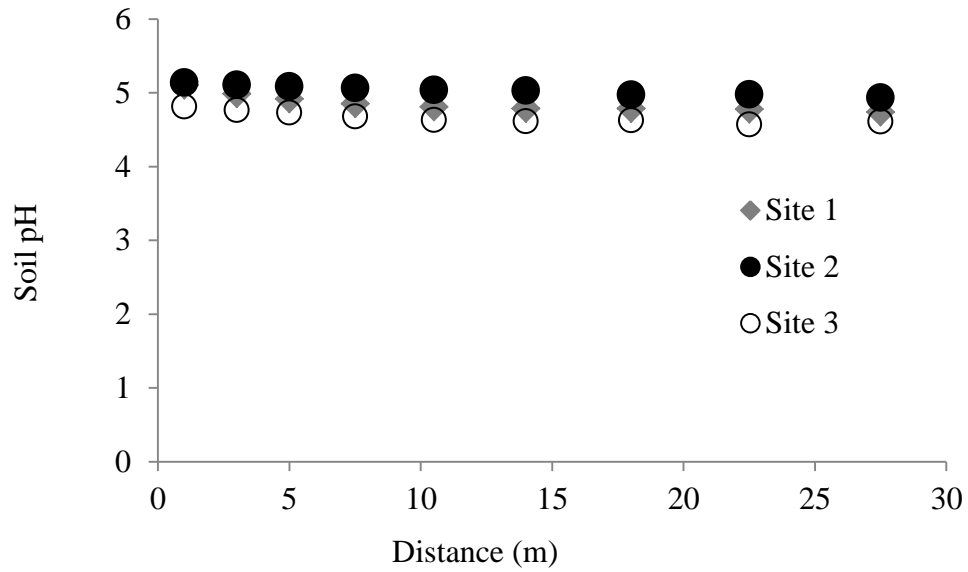
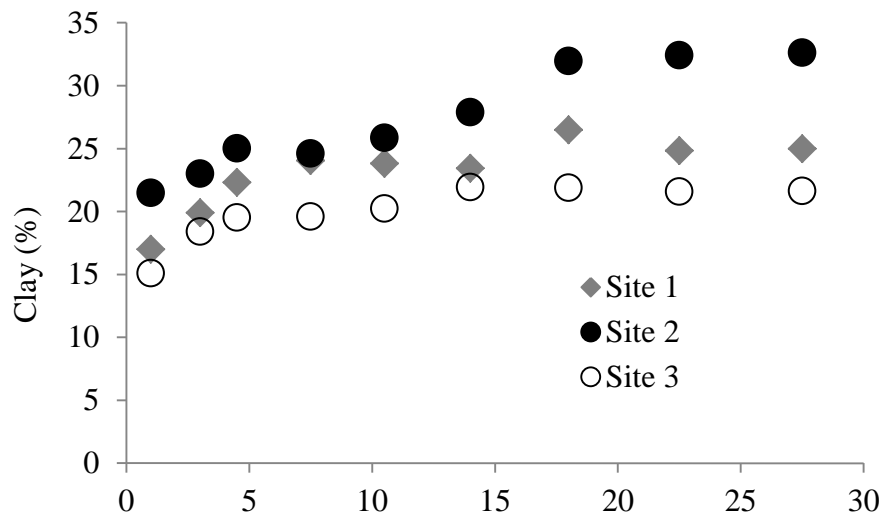


Figure 3.7. Variation of soil pH with distance from *Eucalyptus saligna* woodlot in a eucalypt-maize cropping system in three sites in southern Rwanda.



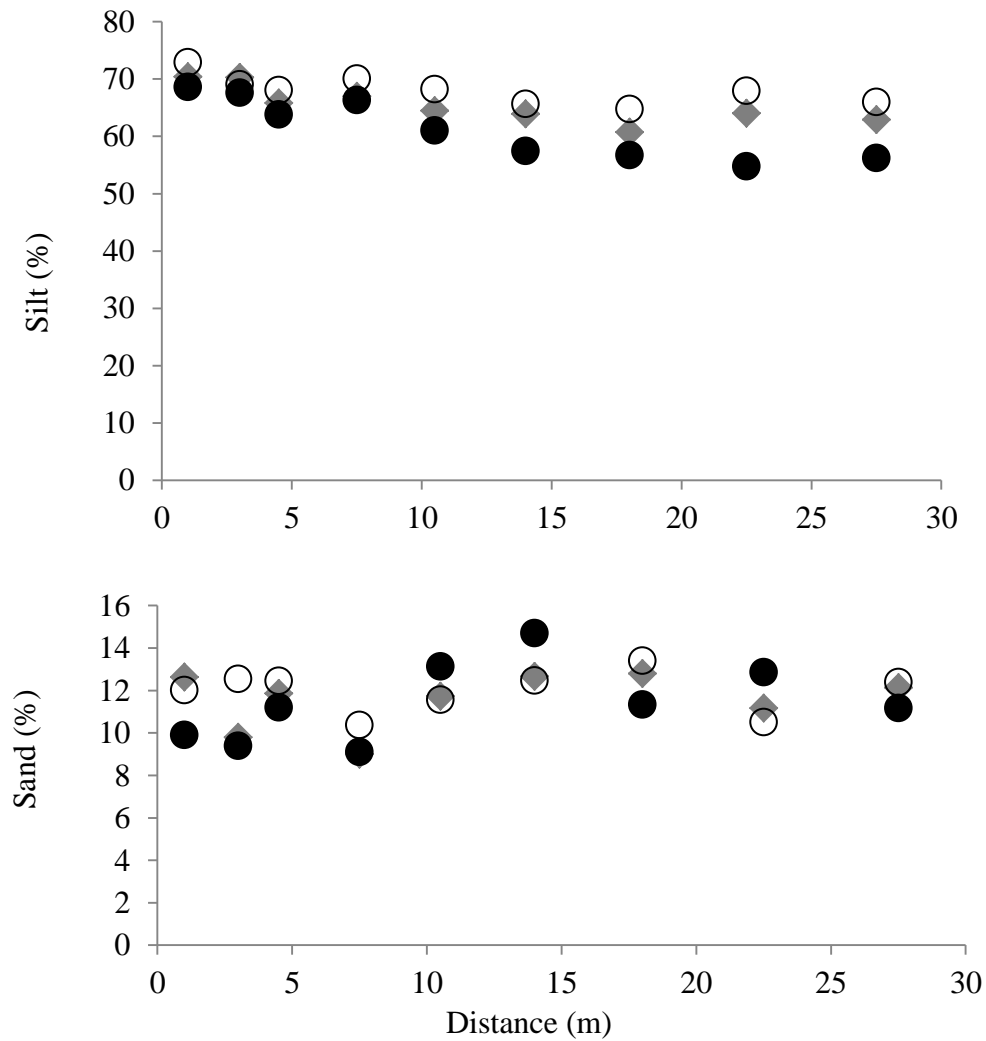


Figure 3.8. Variation of soil particle size distribution with distance from *Eucalyptus saligna* woodlot in a eucalypt-maize cropping system in three sites in southern Rwanda.

Fertilisation improved maize performance significantly and stover yield increased from 1.5 to 3.3 t ha⁻¹ in fertilised and unfertilised treatments respectively. Similarly, grain yield increased from 1.3 t ha⁻¹ for unfertilised and to 2.6 t ha⁻¹ for fertilized treatments, a two fold increase.

Discussion

The aim of this study was to analyse the eucalypt woodlot-maize system, and quantify how resource availability (water, nutrients and light) and crop performance (height growth, stover production and grain yield) vary with the distance from the eucalypt woodlots. with the aim to analyse different geometrical configurations of woodlot vs. crop field. Fertilization was added to analyse whether eucalypts and maize mainly compete for nutrients, and whether fertilization can mitigate for the negative woodlot effects.

Eucalypt and maize competition

The crop field zone affected by the woodlot was $10.5 \times 12 \text{ m}^2$ which makes 35% of the total crop zone ($30 \times 12 \text{ m}^2$), leaves out $19.5 \times 12 \text{ m}^2$ crop field unaffected by the woodlots. The affected zone produced grain yield at the rate of 0.5 t ha^{-1} which is only 20% of the yield recorded in the open areas. By comparing these two yields, we would conclude that the woodlot competition reduced maize grain yield by 80%. However, this competition with the trees affects only a small part (35%) of the crop field. By spreading the maize grain loss over the $30 \times 12 \text{ m}^2$ plot, the loss reduces from 80 to only 28%. The proportion of the crop field that will be affected will depend on the size of both the crop field and the woodlot; and the length of the tree-crop interface. If we consider a square hectare plot crop field which has one side adjoining a eucalypt woodlot, only a strip of 10.5 m wide and 100 m long next to the woodlot would be affected by the latter, the remaining $89.5 \times 100 \text{ m}^2$ being free from woodlot effects. In this case, the affected crop field zone would only be 10.5% of the whole crop field.

This means that, the 10.5% of the whole crop field area will give grain yield at the rate of 0.5 t ha^{-1} , and 89.5% at 2.5 t ha^{-1} ; equivalent to 0.0525 and 2.2375 t ha^{-1} respectively. This adds to 2.29 t ha^{-1} , which is 0.21 t (8.4%) less compared to the yield of unaffected hectare crop field.

Many reports on crop yields as affected by trees when these are grown next to each other are usually presented as percentages or yields ha^{-1} relating values nearest to trees and those in open areas, far away from the trees (Sudmeyer and Hall, 2015; Tadele and Teketay, 2014; Chanie et al., 2013). This is very informative since it does not show actual proportional sizes of crops fields affected and those not affected, and the percentages are usually high, giving a wrong impressions to the readers of the magnitude of tree effects. It would be better if the rates of yield reduction are given together with the sizes of crop fields affected and those unaffected to give a clear picture of the magnitude of the overall effect.

Reductions in crop growth and corresponding yield losses are known to occur when eucalypts and crops are grown in close proximity. *Eucalyptus* are known to have high water uptake which is primarily a strategy enabling them to grow even under harsh environmental conditions (Akhter et al., 2005). This high water uptake capacity enables them to outcompete other plants occurring in their vicinity. The suppressive effects by eucalypts in agroforestry systems have been reported in other studies (Sudmeyer et al., 2015; Kidanu et al., 2005; Malik and Sharma, 1990). The species rooting pattern consists of deep and far reaching lateral roots so as to cover a bigger soil volume to capture more resources. Depending on the environment, rooting depth for *Eucalyptus* spp. may reach 20-30 m (Akinnifesi et al., 2004) while lateral roots were observed to spread as far as 20 m in the adjacent crop fields (Kidanu et al., 2005).

The general low yield observed in our study is not unexpected since low crop yields are common on highly weathered and nutrient depleted soils in east Africa, mainly the acrisols, ferralsols, nitisols and alfisols (Woomer and Muchena, 1996). Maize biomass produced in this study was comparable to that reported by Tittonell et al. (2007) for the outfield farmer

managed sites in Kenya. It was also comparable to that reported in the study area (Muberantwari et al., 2009) and was 1.6 fold higher than the average maize grain yield reported for entire Africa (Baligar et al., 2001).

Soil moisture, soil nutrients and solar radiation decreased significantly near eucalypt woodlots in this study (Figs. 3-5). However, it is generally assumed that competition for soil moisture and soil N are more important in eucalypt-crop systems (Sudmeyer and Hall, 2015; Ong et al., 1999; Sudmeyer, et al., 2002). Water and nitrogen (N) are reported to be the most limiting factors for maize grain yield in the tropics (Moser et al., 2005). Radiation is reported to be rarely the most limiting growth resource in the tropics (Ong and Black, 1994), and belowground competition between perennial and annual crops is known to be severer than the above-ground (Nissen et al., 1999).

Fertilization had influenced plant growth and biomass production but did not change the trend in tree-crop competition and the effect on crop biomass production and grain yield (Fig. 3.2). The increase in yield in the fertilized treatment may be due to the added nutrients N, P and K on the soils in the experiment, that are known to be very unproductive. These results are consistent with those revealed by Lisuma et al. (2006) in Tanzania who concluded that in Sub-Saharan Africa, application of fertilizer can increase maize yields up to three times. The higher grain production in relation to the total biomass (harvest index) observed in unfertilized treatments over the fertilized one (Fig. 3.1c) could be an indication of more allocation of assimilates in favour of developing grains. Some crops have the capacity to preferentially allocate assimilates to grains at the expense of the biomass under unfavourable conditions. This was reported in some variety of sorghum especially under low soil moisture conditions (Wenzel et al., 2000). The precedence of the harvest index of unfertilized treatment over the fertilized one being persistent in open areas may be attributed to the general poor soil state of the study sites as earlier stated.

Eucalypt woodlots reduce water availability

Lower soil moisture values observed nearest to the woodlots than far away may suggest the increased moisture uptake by the tree roots. Crop suppression by *Eucalyptus* trees is reported even in young stands. Ceccon (2007) reported a crop decrease in rice and beans intercropped with young *E. urophylla* in Minas Gerais state of Brazil. Lateral roots in *E. saligna* were observed to be concentrated in the top 61 cm soil layer (Skolemann, 1974). This may be a common phenomenon in eucalypts since similar results were found in other eucalypt species (Kidanu, 2005). A high density of fine and medium-sized roots were reported in *E. urophylla* in the first 20 cm of the upper soil, and a marked decrease of the root density in the lower soil layers (Cohen et al., 1997), which may explain the reduced crop yield due to competition for soil water. It should be noted here that clay particle distribution in the soil increased with distance from eucalypt woodlots. Less clay near the woodlots may have contributed to the observed less soil water since the coarser texture may lead to increased water drainage.

Eucalypt woodlots reduce soil nutrient levels

Poor and variable soil fertility was reported in Rwandan soils by Verdoodt and Van Ranst (2006). These soils are known to originate from granites mixed with mica schist and quartzite (Steiner, 1998). The soils derived from these parent materials are highly acidic, deficient in basic nutrients, have a high level of exchangeable aluminium and therefore are not productive (Bourleigh and Yamoah, 1997; Sanchez et al., 2003). The observed small concentrations of soil nutrients close to the trees is commensurate with literature and may be a function of the increased capacity of the eucalypt trees to take up nutrients in their vicinity (Kidanu, 2005; Sudmeyer, 2002). *Eucalyptus* species are reported to be efficient miners of soil nutrients, thereby suppressing other vegetation that grows underneath or near them (Harrison et al., 2000). Sang et al. (2013) reported that *E. urophylla* plantations were associated with lower soil C levels and observe that these results indicate that unfertilised *E. urophylla* plantations in short rotations are both ineffectual soil rehabilitators and potentially unsustainable in the longer term. This corroborates previous conclusions (Behera and Sahani 2003). The small nutrient concentrations are at odds with the fact that eucalypts produce high amounts of litter expected to enrich soils through nutrient recycling. This may however be a consequence of local management practice where leaf litter from the woodlots is collected together with branches and used as fuel (Ndayambaje and Mohren, 2011) or as mulch for soil conservation (Nzeyimana et al., 2013) due to critical needs of firewood and soil erosion control respectively.

Soil nutrient concentrations decreased with soil depth (Table 4). Soil organic matter contents which are usually higher at the topsoil are responsible for the retention and release of nutrients, and nutrients added to soil by litter fall and fertilization usually accumulate at the topsoil (Lehmann, 2003). Subsoil nitrogen was observed to decrease under natural and *Sesbania* fallows compared to the sole maize and uncultivated bare soil in Kenya (Hartemink et al., 2000). Phosphorus and potassium were observed to concentrate near the soil surface compared to the subsoil (Jobbágy and Jackson, 2001).

Below ground competition may be explained by the hypothesis that roots grow only as deep as needed to fulfil tree resource requirements (Schenk and Jackson, 2002). Trees prefer to root shallowly since (a) energy costs for construction, maintenance and resource uptake are lower for shallow roots (Adiku et al., 2000); (b) oxygen is usually likely to be less deficient in shallow soil layers (Hillel, 1998); and (c) the concentration of nutrients is often greater in the upper soil layers (Jobbágy and Jackson 2001). The soils of the study site were generally acidic. Soil pH was only slightly higher near the woodlot compared to open areas, and thus, in agreement with other studies pH (Jobbágy and Jackson, 2003).

Eucalypt woodlots reduce light availability

The three season average PPFD recorded at the woodlot flow ($279.7 \mu\text{mol m}^{-2} \text{s}^{-1}$) and in the crop field in this experiment ($499.9\text{-}933.8 \mu\text{mol m}^{-2} \text{s}^{-1}$), even at points furthest from the woodlots, is lower than the normal range for full light which ranges from $1200\text{-}2000 \mu\text{mol m}^{-2} \text{s}^{-1}$ (Berlin and Cho, 2000). The normal range is usually reduced to lower values under the forest canopy to values around $800 \mu\text{mol m}^{-2} \text{s}^{-1}$ (Berlin and Cho, 2000). The lower values recorded here are therefore not surprising since the amount of light received was measured in

the crop fields which are variably shaded by eucalypt woodlots. The amount of light received by vegetation has several influencing factors including altitude, canopy structure, clouds, latitude, time of the year, time of the day and topography (Jones, 1992). Morphological factors such as leaf orientation, shape, pubescence, thickness, etc. also affect the amount of light received by plants (Berlin and Cho, 2000). The amount of solar radiation received in the crop fields were higher than the usual values received at the forest canopy because the amount of shade cast by *E. saligna* woodlots is not very intense. This may partly be attributed to the species leaf area index which is low and partly to the dropping nature of the species' leaves. Leaf area index of *E. saligna* was reported to be among the smallest in several exotic and indigenous tree species investigated in the study area (Nsabimana et al., 2008).

Conclusions

Eucalyptus woodlots significantly reduced environmental resources in the crop field strip adjoining the woodlots. Significant effects of the woodlot were detected up to 7.5 m distance from the woodlots for soil P, 10.5 m for soil N and K and soil water; and 18 m for solar radiation. Soil moisture and solar radiation were reduced near the woodlots by 18 and 68% compared to the values in open areas. Soil nutrient levels recorded near the woodlots were lower than those in the open areas by 37.5% for N, 27% for P and 38% for K. In this study, 10.5% of the crop field nearest to the woodlot (10.5 m wide) produced grain yield of 0.5 t ha⁻¹ while the remaining field (89.5%) produced 2.5 t ha⁻¹. The grain loss in the 10.5 m strip therefore was 80% less than the yield in open areas, which is equivalent to an average yield loss of 0.21 t ha⁻¹. Since tree competition affects only 10.5% of a 1 ha crop field, expressing this as the loss over the whole field becomes a yield reduction of only 8.4%. Reporting grain yield losses need to be quantified by actual yield values and corresponding crop field area affected rather than using percentages alone since the latter may be misleading. The magnitude of grain yield loss will depend on the size of the crop field and *Eucalyptus* woodlot and the length of tree-crop interface since these will determine the crop field proportions that will be affected/not affected by the edge. Maize harvest index was relatively higher in the unfertilised than in the fertilised treatment, reflecting the crop's capacity to preferentially allocate resources to grain production when growing under harsh environments. While fertilisation increased grain yield from 1.3 to 2.6 t ha⁻¹, it did not change the trend in the effect of eucalypt woodlot on maize growth and performance.

CHAPTER 4: Edge affects both maize and *Eucalyptus* trees in maize-eucalypt woodlot agroforestry systems

CP Mugunga, GMJ Mohren and KE Giller

Edge affects both maize and *Eucalyptus* trees in maize-eucalypt woodlot agroforestry systems

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Abstract

Agriculture is the hub of livelihoods in the east African highlands and in Rwanda. Firewood shortage and the need of wood for construction drove people to grow trees; consequently eucalypt woodlots are found interspersed with annual crops on the hilly landscape across the country. This study aimed at characterising the tree-crop interface between eucalypt woodlots and maize as a representative field crop. The specific objectives were: i) to quantify the crop field area affected by woodlot trees and yield losses of maize crops; ii) to quantify the differential growth and biomass production in eucalypt woodlot edge trees vis-à-vis those at the woodlot centre; iii) to quantify the area of the woodlot subject to edge effects and effects on tree biomass; and iv) to evaluate the trade-offs of crop yield losses in relation to the gains of wood production. Maize was grown adjacent to three eucalypt woodlots and three additional woodlots were sampled, the tree edge effect was evaluated for all six. Maize grain yield within 10.5 m of the woodlots was 0.5 t ha^{-1} against 2.5 t ha^{-1} in open areas. Grain yield loss increased with tree-crop interface length. Field orientation was very important in none square crop fields and smaller fields suffer more from edge-effects when longer sides adjoined woodlots. Trees in the outer eight meters of the woodlots had 17.8% bigger diameter at breast height and 34.5% more aboveground biomass than interior trees. Woodlots were affected by the edge and small woodlots were more affected. Combined maize-wood products were more profitable than sole systems and revenue from extra wood gains due to edge effects exceeded corresponding revenue losses in maize yield. However, land shortage may limit farmers to grow *Eucalyptus* woodlots.

Key words: Short rotation eucalypt woodlots, intra-specific competition, inter-specific competition, trade-off analysis

Introduction

Agriculture forms the mainstay of the economy in the East African highlands. In Rwanda, 90% of the population lives in rural areas with agriculture practiced on 52% of the land overall (MINITERE, 2004). Rapid population growth has led to intense pressure on land for agriculture and forestry. In 2012, Rwanda had 10,515,973 people with a growth rate of 2.6% (NISR, 2014). The average population density of about $407 \text{ inhabitants km}^{-1}$ (Bucagu et al., 2014) makes Rwanda the most highly populated country in Africa (Warnest et al., 2011). Natural forests declined by 65% in area between 1960 and 2007 (MFM, 2010). This renders Rwanda strongly dependent on growing trees on-farm for the supply of fuelwood, timber and poles. In 2007, forests contributed 80% of total national domestic energy as firewood (57%) and charcoal (23%) (MFM, 2010).

Tree planting is seen as a solution to the problem of firewood and shortage of other wood products (Ndayambaje and Mohren, 2011). In Rwanda and other countries of East Africa, there has been widespread planting of fast-growing eucalypt species. In 2008, 64% of the total plantation area in the country was covered by eucalypts with ownership of 65%, 26% and 9% for the local governments, smallholder farmers and private institutions respectively (Nduwamungu et al., 2008). On-farm tree planting is mainly in the form of woodlots ranging from very small to large stands which are scattered countrywide, though most are found in the southern and the western Provinces (Nduwamungu et al., 2008).

Given that very small fields are devoted to crop production and are intermixed with woodlots of fast-growing eucalypt tree species, competition between woodlot trees and field crops at the tree-crop interface assumes major importance. Competition between trees and annual crops is key to understanding the benefits of agroforestry. Trees compete with crops for light, water and nutrients reducing crop yields significantly (Rowe et al., 2005). *Eucalyptus* spp. are known to be particularly competitive trees that significantly reduce crop growth due to severe competition for soil moisture (Kidanu, 2005), nutrients (Okorio et al., 1994) and light (Bertomeu, 2012). Some trees may exert allelopathic effects on crops growing in their vicinity (Lisanework and Michele, 1993).

Apart from the competition between food crops and woodlot trees, trees in mono-specific plantations compete among themselves thereby reducing tree growth within a stand as illustrated by the larger size of trees at the edge of plantations compared with those at the centre. Differential growth between trees at stand edges and in the inner zones has been observed in agroforestry systems (van Noordwijk, 1999), in forest plantations (McJannet and Vertessy, 2000) and in natural forests (McDonald and Urban, 2004). The growth of dominant trees in a forest plantation results from superior resource use-efficiency that leads to greater tree growth (Harris, 2007). However, the additional tree growth exhibited by edge trees in plantations results from the extra resources gained from crop fields adjoining them (Weathers et al., 1997). This effect may be due to differences in microclimate (Voicu and Comeau, 2006) and light (Dignan and Bren, 2003), which influence nutrient exchange (Sanou et al., 2012) thereby enhancing growth of trees on the edge compared with those in the interior (Wright et al., 2011).

Exploitation of interactions between woody and non-woody components is the key to the success of agroforestry systems (Rao et al., 1998). This necessitates understanding of the role of inter- and intra-species competition at the tree crop interface. We studied interactions between eucalypt woodlots and maize as a representative field crop. The specific objectives were: i) to quantify the differential growth and biomass production in *Eucalyptus saligna* edge trees in relation to those at the woodlot centre; ii) to quantify the crop field area affected by woodlot trees and yield losses of maize crops; iii) to quantify the area of the woodlot subject to edge effects and effects on tree biomass; and iv) to evaluate the trade-offs of crop yield losses in relation to the gains of wood production.

Materials and methods

Site description

Site description was fully presented in chapter 3, and is only briefly summarized here. The maize experiment was conducted in three sites where there were *E. saligna* woodlots and adjacent crop fields. Two sites, Cyarwa and Gishamvu were located in Huye district and one, Save, in Gisagara district, southern Rwanda. Maize (Katumani variety) was grown in crop fields adjacent to *Eucalyptus* woodlots. The woodlot and cropland together comprise an agroforestry system with woodlot and crop fields as subsystems. The edge effects on both components as expressed in tree sizes and maize biomass production and yield, and the profitability of the products were the main focus of this study.

Experimental design and field procedures

The field procedure and experimental design are described in detail in chapter 3. The experiment commenced during the Jan-May 2007 season and ran for three consecutive cropping seasons. The maize experiment was done on three sites and in each of these, three crop fields belonging to farmers were used. To ensure representative coverage of the study zone, the three sites used were located at least five km apart. Fields incorporating woodlots with trees of *E. saligna* were selected and experimental plots of 12.5 m × 30 m marked by pegging. In all sites, the tree-crop interface was along the 12.75 m side of the crop field length and the sides of the woodlots were longer. Woodlot sizes were 60 x 70 m for Save site, 50 x 50 for Cyarwa and 60 x 65 m for Mukura. Each crop field was divided into subplots running parallel to the tree-crop line and away into the farmland with the following dimensions: 0-2, 2-4, 4-6, 6-9, 9-12, 12-16, 16-20, 20-25 and 25-30 m. Site characterisation for soil pH and soil particle size distribution were presented in chapter 3. Soil pH ranged from 4.67-4.88, clay from 20.1-26.3, clay from 62.2-68.9 and sand from 10.9-11.5. Katumani maize variety was sown in the crop plots adjacent to the woodlots after site preparation by manual tillage. The planting was done at a spacing recommended at the time of experiment by Rwanda Agriculture Board (RAB), of 0.5 m between plants and 0.75 m between lines, with three seeds planted per hole, later thinned to leave two plants per hole. Land preparation was done by hand hoe and plots were hand weeded three times during the season to avoid weed competition.

Data collection and analyses

For biomass and grain yield determination, the inner 2 × 2 m area was demarcated in each of the nine plots after the maize were fully mature and had started drying. Six plants were harvested from this area and the above ground parts of each were cut at the root collar by using a sharp machete. For each plant, the parts were labelled, oven dried at 65°C to constant weight and weighed. All cobs from this inner plot were dried under full sunshine for 20 consecutive days. Grain was separated from cobs by using a threshing machine and weighed using an electronic balance in the faculty of agriculture laboratory at the University of

Rwanda. The air dry grain weight was reduced to dry weight by multiplying with a factor of 0.85.

Six woodlots were used to quantify tree growth in terms of diameter at breast height (DBH) and biomass (W) and the reduction of these traits in relation to distance from the woodlot edge. These included the three woodlots Cyarwa, Save and Mukura that were used to investigate the interaction between woodlot trees and maize crop, and an additional three (AR^I , AR^{II} and AR^{III}) woodlots sampled from the Ruhande Arboretum (Butare, southern Rwanda). Woodlots AR^{II} and Save were planted, while the rest were coppice stands. For tree biomass estimation, DBH was measured and recorded by tree rows. From each woodlot, twelve consecutive tree rows from the edge running parallel to the tree-crop interface towards the woodlot centre were selected. From each of the 12 tree rows in the three sites, thirteen trees were selected and measured for DBH. Tree spacing was 2×2 m, and the measurements extended from the woodlot edge to 22 m inside the woodlot. The DBH of the surviving trees were recorded and total biomass per tree (W , kg tree⁻¹) of each tree measured for DBH was estimated using an allometric equation developed for the species in the study area (Mugunga et al., submitted). Differences in DBH and W between edge trees (from the four outer lines of the woodlots) and the average for four innermost trees inside the woodlots (assumed to be free from the edge effect) were taken as the DBH and W gains due to edge effect. Tree weight was converted to a hectare basis based on tree density. Average annual biomass increment per ha was then deduced from total biomass by dividing by stand age.

To analyse the trade-off between maize yield (including stover biomass) and wood production, maize stover and grain yield obtained in one season were extrapolated to annual values by multiplying by two as there are two cropping seasons in the study area, with the remaining months consisting of the dry season. Basing on our observation on maize growth and grain yield, the crop field was divided into two zones: zone A, the one affected by the woodlot edge and zone B, the one not affected (Fig. 4.1). Losses in maize stover and grain yield in zone A were computed as differences between stover and grain yield in this zone and corresponding values in zone B. For the woodlots, a similar zonation was made such that the edge affected or the outer part of the woodlot closest to the woodlot-crop interface was denoted as zone C while the one not affected was zone D (Fig. 4.1).

Data were statistically analysed by using IBM SPSS 22 software. One way analysis of variance was used to determine significant differences between treatments and means were separated by using least significant difference method ($P < 0.05$).

In addition to the effects on biomass increment, the long term economic feasibility of maize and *E. saligna* woodlots was addressed using a net present value (*NPV*) approach based on the work of Whittock et al. (2004). The *NPV* shows the total amount of surplus (profit) or loss that a given project or enterprise is expected to generate over its lifetime and a positive *NPV* indicates a profit since the expected net cash inflows over the total project lifetime are higher than the cost of financing the project (van Eijck et al., 2012). Costs of the different activities and commodity prices for revenue estimations were based on those provided by the Rwandan Agriculture Board (RAB), Ruhande Arboretum station located in the study area.

Conversion of biomass estimated in this study to harvestable products of timber, pole and firewood were based on the growth estimates by Burren (1995). All costs and incomes were discounted to the time of plantation establishment (year 0), and the present value was calculated using the standard formula:

$$NPV = \sum_{t=0}^T (R_t - C_t) (1+r)^{-t}$$

where the index t represents time measured in years; R_t and C_t represent revenues and costs in year t respectively. A discount rate r of 10% was adapted and a 21 year project life equal to one timber tree rotation (three short rotations for firewood and pole production) was used in this study.

NPV analysis allows the projection of economic costs and benefits attributable to different cropping systems over multiple cropping seasons. A comparison of the *NPV* of maize and alternative *E. saligna* production systems of firewood and poles (short rotation) and timber (long rotation) were tested. The analysis of changes in costs and returns over time used by Vosti et al. (2000) were applied to identify which alternative cropping system could be more profitable for the farmers in the area. Input production costs for the different alternative systems are shown in Table 4.1. Fixed cost of land was ignored for two practical reasons: firstly, it appears in all investment options at the same cost and secondly, it belongs to the farmers and the latter do not need to pay for land every year. Costs of producing maize were kept constant each year and those of producing firewood and poles were kept constant in each of the three rotations. Rotation cycles for both firewood and poles was fixed at seven years, leading to three rotation cycles for these commodities for one timber rotation cycle of 21 years. Revenue was assumed to be constant in all three firewood and pole harvesting cycles and also in all 21 years in the case of maize grain.

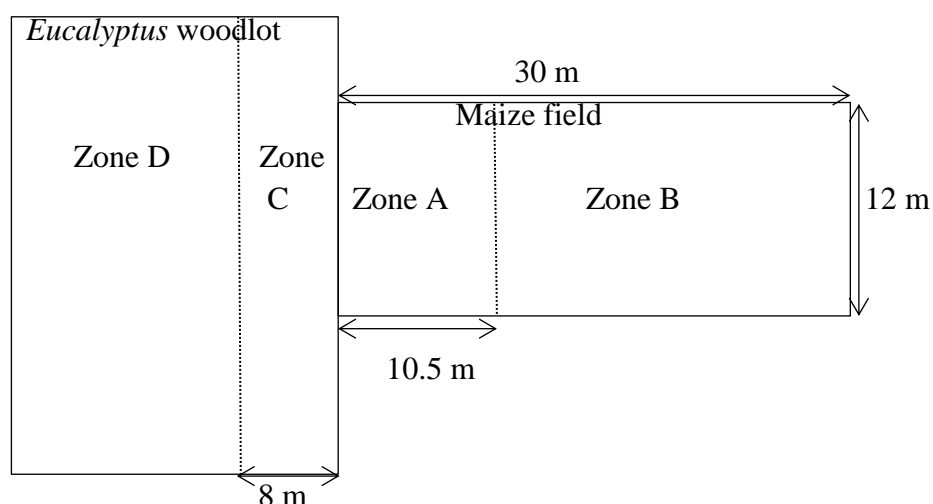


Figure 4.1. Schematic presentation the experiment showing zones affected by the edge: A for the maize field and C for the woodlot; and the zones not affected: B for the maize field and D for the woodlot.

Table 1 Input cost of maize and *E. saligna* products in eucalypt woodlot-maize agro-ecosystem in southern Rwanda – calculations based on 1 ha smallholder land area

A. Maize					B. <i>E. saligna</i> woodlot				
Item/activity	Unit	Quantity	Cost (US \$/year)		Item/activity	Unit	Quantity	Cost (US \$)**	
			Unit	Total				Unit	Total
1. Seed purchase	kg	30	0.83	25	1. Seedling production	Number	2,500	0.08	208
2. Land preparation	Person days	150	1.67	250	2. Site preparation	Person day	80	1.67	133
3. Sowing		75	1.67	125	3. Field planting	Person day	55	1.67	92
4. Fertiliser purchase					4. Weeding	Person day	45	1.67	75
4.1 NPK (17%)	kg	300			5. Thinning - timber production*	Person day	1,215		
			0.67	200				1.67	2,025
4.2 Urea (45%)	kg	100	0.70	70	5. Harvesting	Person day	2,945	1.67	4,908
5. Fertiliser application	Person days	66	1.67	110	5.1 Timber	Person day	300	1.67	500
6. Weeding	Person days	105	1.67	175	5.2 Firewood	Person day	350	0.08	208
7. Harvesting	Person days	90	1.67	150	5.3 Poles				
8. Drying & processing	Person days	60	1.67	100					
Total				1,205					7,942

*combines cost of three thinnings, **1 US \$ = 600 Rwandan Francs (FRW)

An analysis was done to study whether revenues from additional tree growth due to edge effects compensate for the crop loss due to edge effects. For the case of maize grain, the loss in revenue was taken as the difference between the revenue obtained from maize grown in open areas (zone B) and the average obtained from maize in the area close to the woodlot (zone A). Likewise, the average gains in the different wood production alternatives were simply the difference in average revenue between the products obtained from trees in zone C and those in zone from zone D. A trade off analysis was made by comparing revenues obtained from both maize and woodlot sole cropping and from the combined maize-woodlot production alternatives (of timber, pole and firewood). A comparison was also made on revenues obtained from edge tree gains (i.e., revenue obtained from extra biomass in zone C) and revenue losses incurred in maize yield (i.e., revenue losses obtained due to yield decline in zone A).

Results

The results on maize growth and grain yield, and the crop field area affected by the woodlot referred to here are fully reported in a different paper (Mugunga et al., submitted). The proportion of the crop field affected by the woodlot, zone A, was observed to be a crop field strip whose width perpendicular to the tree-crop interface is 10.5 m. In the whole maize field of 12×30 m therefore, the edge affected area is equivalent to 10.5×12 m (35% of the total maize field) and the remaining portion, zone B, of 19.5×12 m was free from woodlot effect, and represents the open area as illustrated in Fig. 4.1 above.

Maize stover and grain yield in zone A declined significantly ($P < 0.001$) nearest to the eucalypt woodlots as a result of inter-specific competition (Fig. 4.2). The average stover and grain yield recorded in this zone was 1 t ha^{-1} for stover and 0.5 t ha^{-1} for grain yield while respective values recorded in open areas were 3.3 and 2.6 t ha^{-1} . The yield in zone A therefore was only about one third for stover and one fifth for grain yield of the values in open areas.

Characteristics of woodlots used in this study are presented in Table (2). The average heights of trees in the woodlots were: 17 m for site AR^I, 22.7 m for AR^{II}, 18.5 m for AR^{III}, 18.6 m for Cyarwa, 16.4 m for Mukura and 24.9 m for Save sites. Tree survival rate ranged from 56% for Mukura site to 88.5% for AR^I site. Intra-specific competition was also evident within the eucalypt woodlots where the edge trees were much larger than those at the centre of the woodlot (Fig. 3a & b). Tree sizes differed significantly ($P < 0.001$) between woodlots and more so with distance from woodlot edge. Older woodlots where trees had thicker trunks seemed to be more affected than the younger ones with smaller DBH. Generally, the edge effect was observed in the outer six to eight m band of the woodlots closest to the tree-crop interface (zone C) irrespective the size of the woodlot.

Table 4.2. Characteristics and growth of 12 outer tree lines in six *Eucalyptus saligna* woodlots used to study the edge effects in southern Rwanda

Woodlot	Stems ha ⁻¹	DBH (cm)	Height (m)	Biomass (kg tree ⁻¹)	Biomass (t ha ⁻¹)
*AR ^I	1209	13.4	13.8	141.7	171.3
AR ^{II}	560	23.8	22.6	360.3	201.8
*AR ^{III}	1152	13.8	14.9	147.7	170.2
*SAVE	1005	16.6	19.8	235.0	236.2
*CRWA	1205	16.1	15.0	203.0	244.6
*MUKU	1376	12.5	16.4	132.0	181.6

*Coppice stands; AR^I, AR^{II} and AR^{III} denote eucalypt woodlots from Ruhande Arboretum while SAVE, CRWA = Cyarwa and MUK = Mukura woodlots belonged to the farmers together with the maize fields.

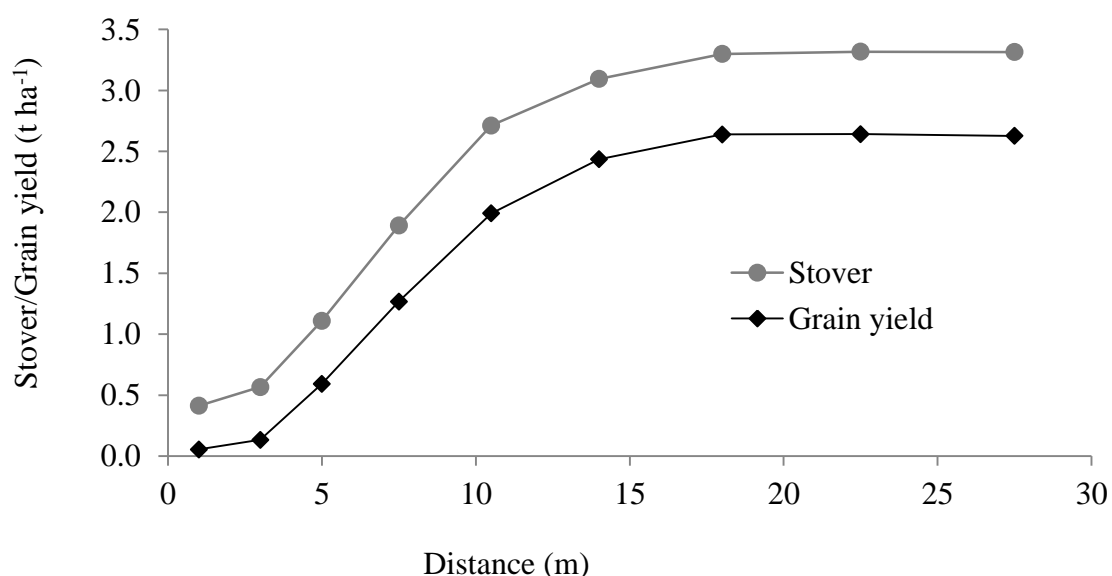
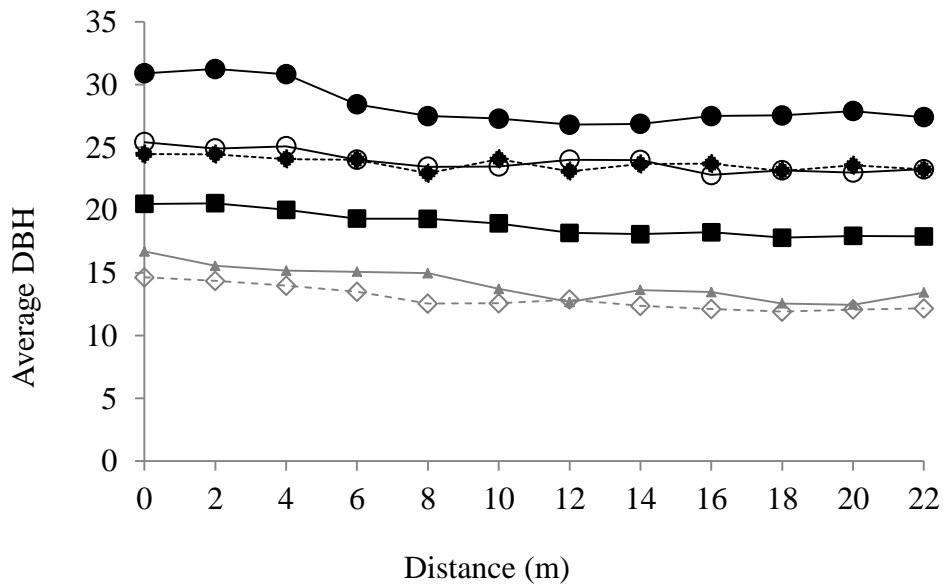
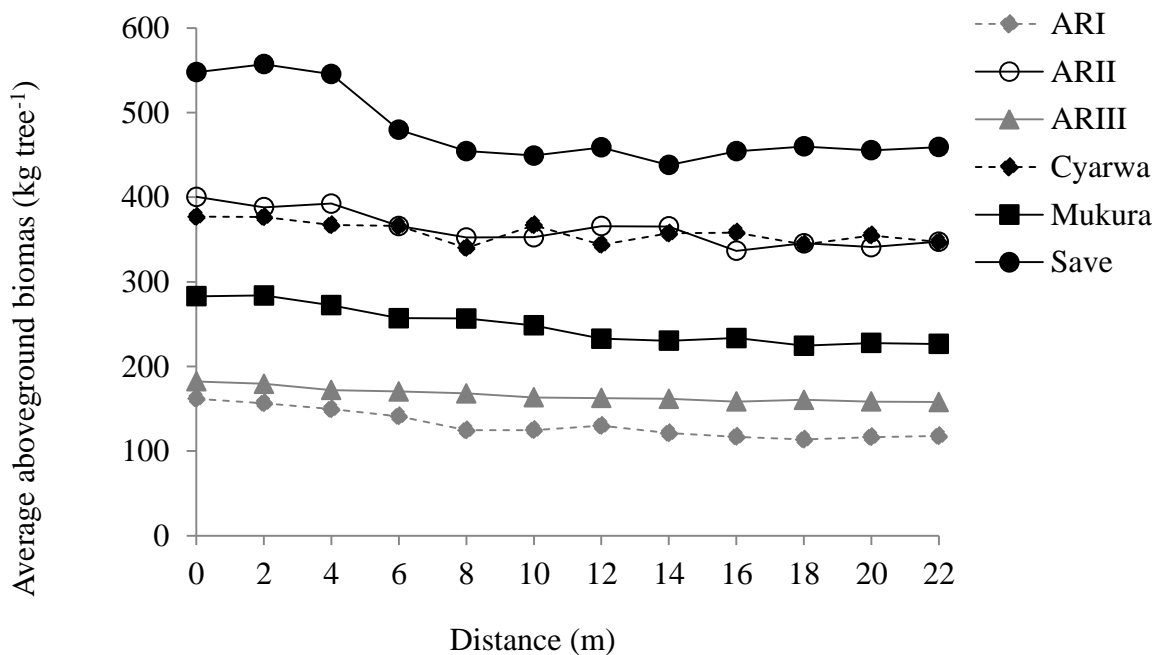


Figure 4.2. Variation in maize stover and grain yield with distance from *Eucalyptus saligna* woodlots and maize interface at three sites (Save, Cyarwa and Mukura sites in Table 4.1) in southern Rwanda). Horizontal axis indicates the position of maize sampling for yield determination as distance from tree-crop interface. The standard error of the mean (SE) ranged from 0.02-0.8, n = 9. Figure adapted from chapter 3.

On average, tree DBH and above ground biomass in zone C were respectively 17.8% and 34.5% larger than at the woodlot centre (zone D). For a woodlot of 50 × 50 m, zones C and D would have respective sizes of 8 × 50 and 43 × 50 m. Zone C therefore is equivalent to 18.6% of the whole woodlot. This estimation however considers only the woodlot side located close to the crop field for the purpose of this study, but the effect could actually be up to four times if the whole woodlot perimeter was not adjoined by other tree stands.



(a)



(b)

Figure 4.3. Variation in tree diameter at breast height (a) and above ground tree biomass (b) with distance from the woodlot edge in six *Eucalyptus saligna* stands in southern Rwanda.

The area of the crop and the woodlot impacted by the edge effect (zones A and C respectively) depend on the size of the two sub-components of the maize-woodlot agro-ecosystem. In general, the two increase with the area occupied by the two components since the length of the tree-crop interface increased as the two sub-systems increase in size. Logically, a farmer would locate a woodlot at the corner of her/his plot and not at the centre, amidst the crop so as to minimise the tree-crop interface and the associated crop yield. To

illustrate the effects, assume a plot of 1 ha was used to produce both maize and eucalypts in a woodlot such that maize is grown in one corner of the 1 ha plot, in a series of sizes varying, say from 15×15 m (225 m^2) to 50×50 m ($2,500 \text{ m}^2$) (Fig. 4.4).

The area of maize field affected (zone A) will range from 0.04-0.12 ha for the 225-2,500 m^2 plots respectively and at the same time, the woodlot area affected (zone C) will range from 0.02-0.07 ha (Fig. 4.5). Maize yield in a eucalypt woodlot-maize cropping system decreases as the crop area declines since, for a fixed total area, say one hectare plot, increasing woodlot size means leaving less area proportion made available for maize production but the proportion of the crop field affected by the tree-crop interface increase (Fig. 4.6). The economic analysis of net present value (*NPV*) of sole maize and that of all alternative investment options that combined maize and eucalypts were profitable since their *NPV* exceeded zero. Combinations of maize-firewood and maize-pole enterprises provided about the same revenue but that of timber was higher by about 25% than that of firewood and poles (Fig. 4.7a and b).

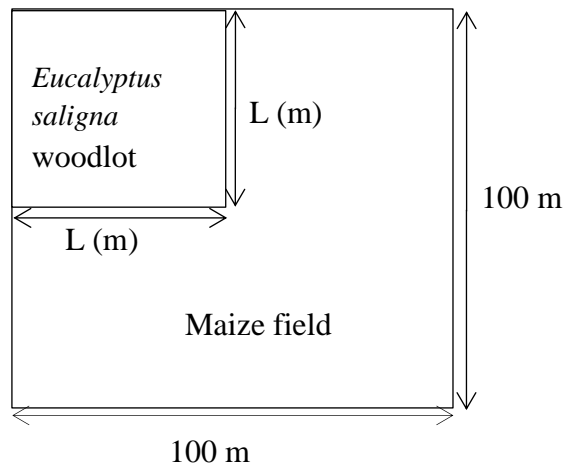


Figure 4.4. Schematic presentation of one hectare plot used for growing both maize and a square *Eucalyptus* woodlot, showing locations of each component. Woodlot side length is varied to explore the edge effects on both components of the agroforestry system.

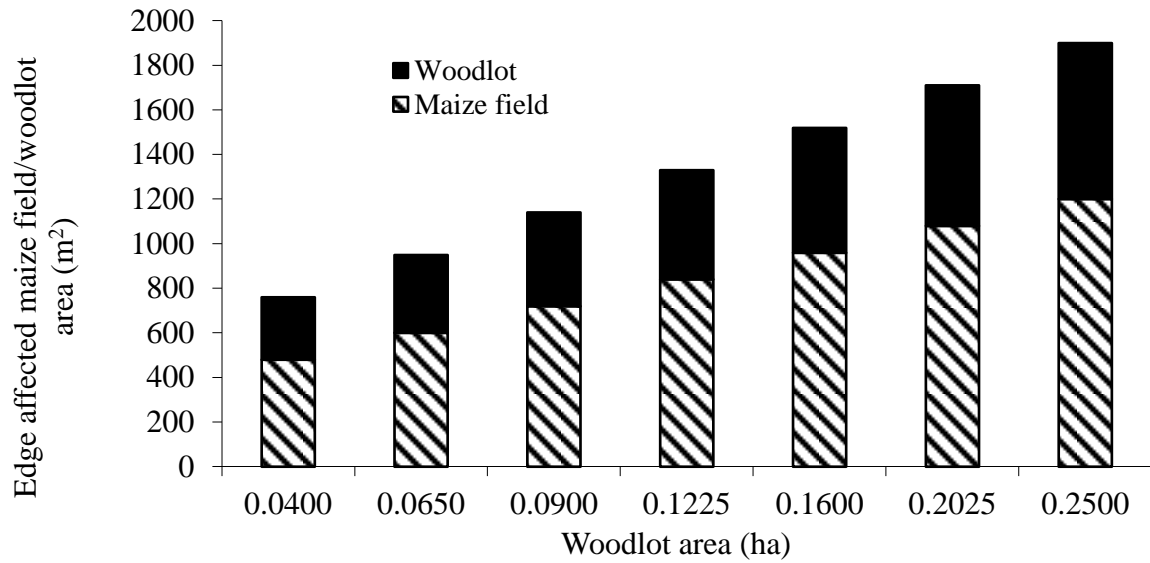


Figure 4.5. Variation of the edge affected area of maize fields and *Eucalyptus* woodlots with increasing woodlot size in one 1ha plot devoted to growing *Eucalyptus* trees and maize in southern Rwanda.

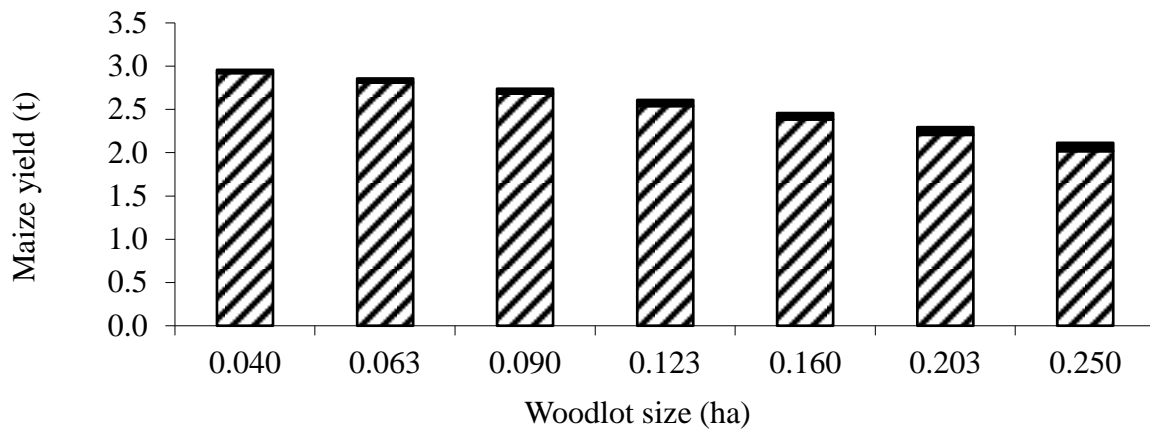
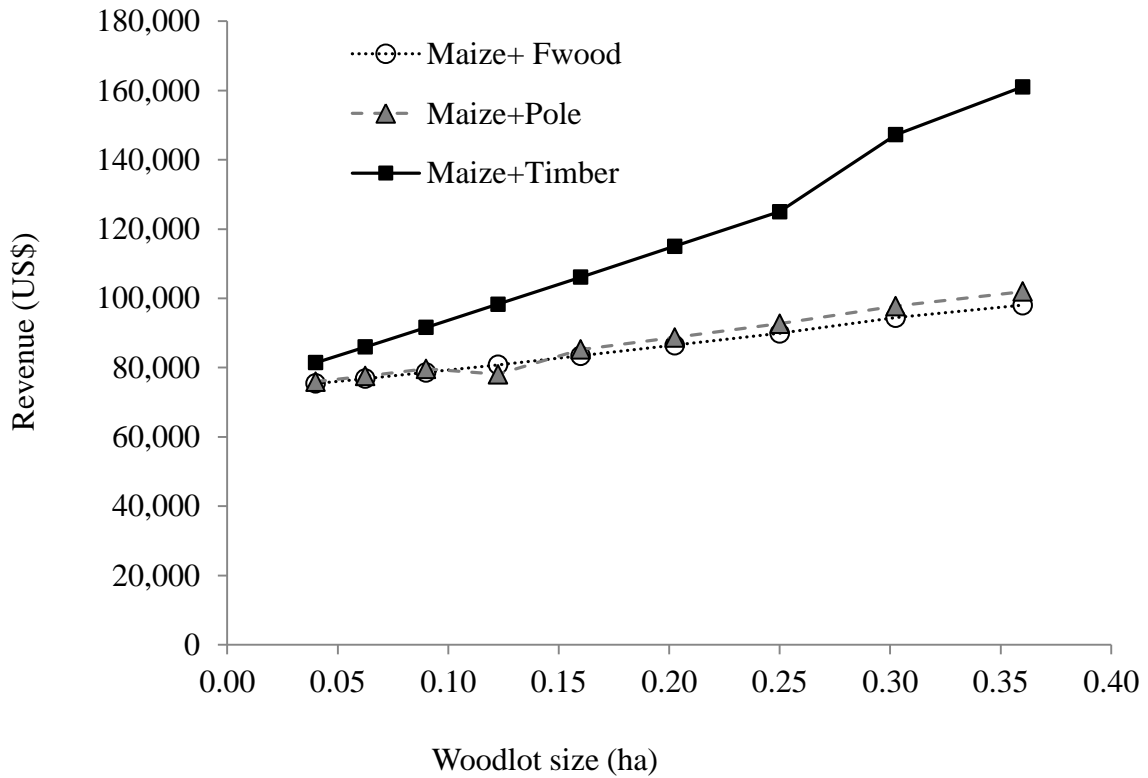
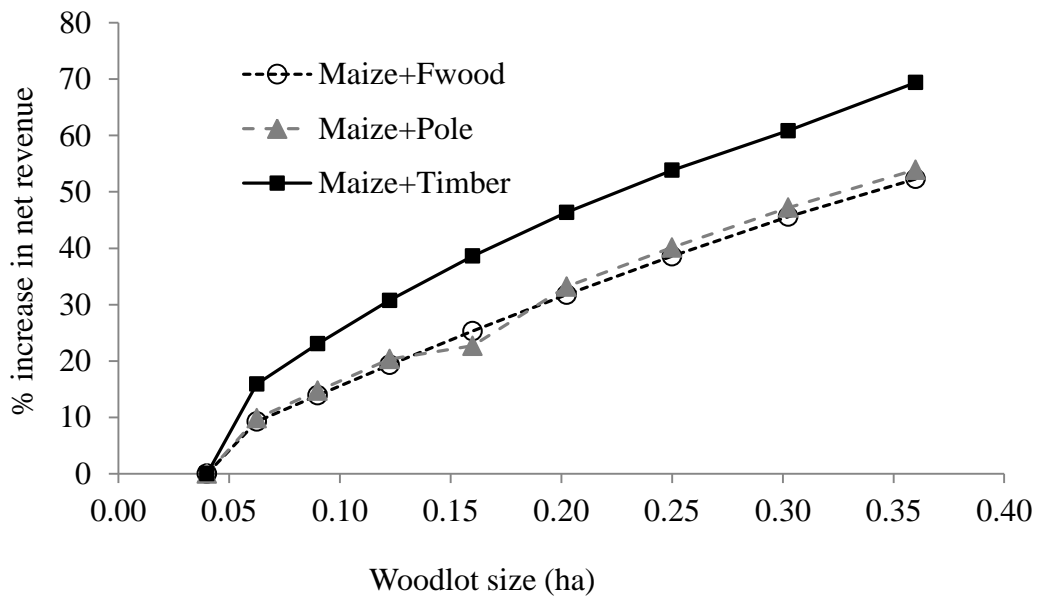


Figure 4.6. Total maize grain yield obtained from 1ha plot with increasing land area devoted to *Eucalyptus* woodlot in southern Rwanda. Dark bars denote maize yield from crop fields affected by the woodlot while striped bars represent grain yield in open areas unaffected by the woodlot



(a)



(b)

Figure 4.7. Revenue from three alternative production options of maize-firewood, maize-pole and maize-timber in a woodlot-maize system in southern Rwanda spread over a 20 year timber rotation period. (a) net revenue (US\$) (b) percent increase in revenue over sole maize of three different wood end use production options. The grey line denotes maize-timber; dark, broken line maize-pole and the grey, dotted line maize-firewood options.

For all production options, revenue from wood components increased with the woodlot size and this lead to a decrease in revenue contribution from maize as a function of the decrease in the field allocated to maize production. At a given point, revenue from both components in a given maize-wood product combination were equal. This point occurred at woodlot sizes of 0.16, 0.275 and 0.33 ha for timber-, pole- and firewood-maize combinations respectively (Fig. 4.8). In woodlot sizes below these critical sizes, maize revenue exceeded that of wood product and vice versa.

The monetary value of the mean biomass gain obtained from the edge trees in eucalypt woodlots was about five times higher for firewood and poles and 10 times higher for timber than the financial losses in maize grain yield encountered in the crop fields adjacent to the woodlots (Fig. 4.9).

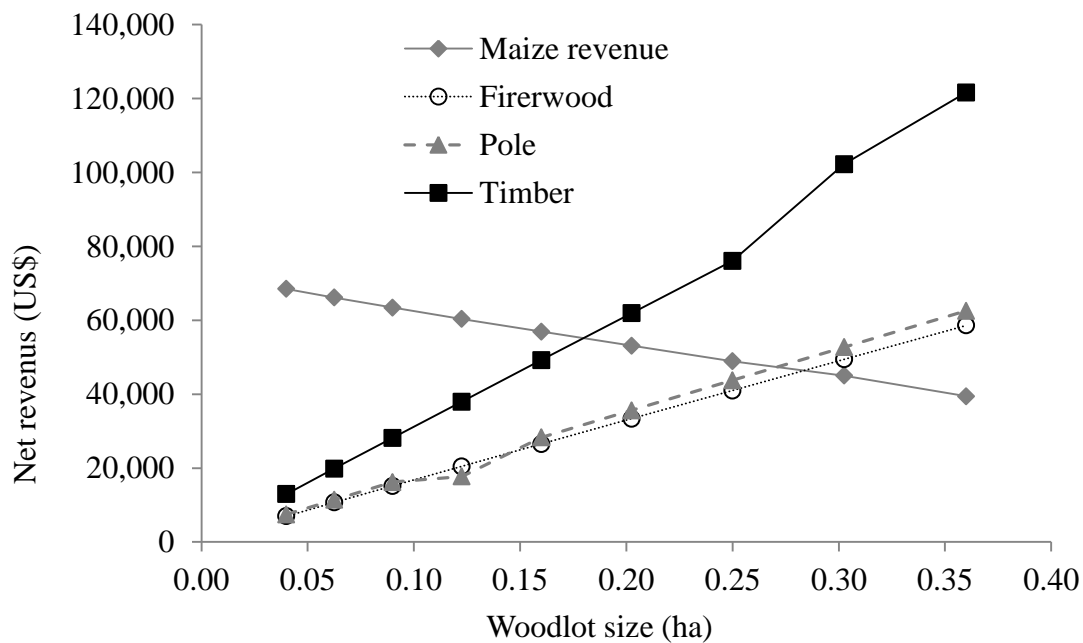


Figure 4.8. Net revenue for the different production option combinations of maize and firewood, pole and timber in a maize-eucalypt woodlot system in southern Rwanda spread over a 20 year timber rotation period.

Discussion

The suppression of maize plants by eucalypt woodlots is not surprising since trees are known to compete with other plants for resources necessary for plant growth (Sudmeyer and Hall, 2015, also chapter 3). Traditional agroforestry trees species such as *Grevillea robusta* and *Leucaena leucocephala* were observed to cause little maize grain reductions (Muchiri et al., 2002a; Immo and Timmer, 1999) and competitive effects on grasses differed among tree species (Samra et al., 1999). Under a drier environment in semi-arid Kenya, *Gliricidia sepium* and *Grevillea robusta* caused more crop yield decline up to 50 and 40% respectively relative to crop yield in control plots lacking trees and reductions of crop yield were greatest close to

trees (Odhiambo et al., 2001). Forest trees (*Cupressus lusitanica* and *Pinus patula*) in a taungya system in Kenya reduced yield in adjacent crops up to 41 and 48% respectively when trees were two years old in a deep soil and from 16-26% respectively in a shallow site (Imo, 2008).

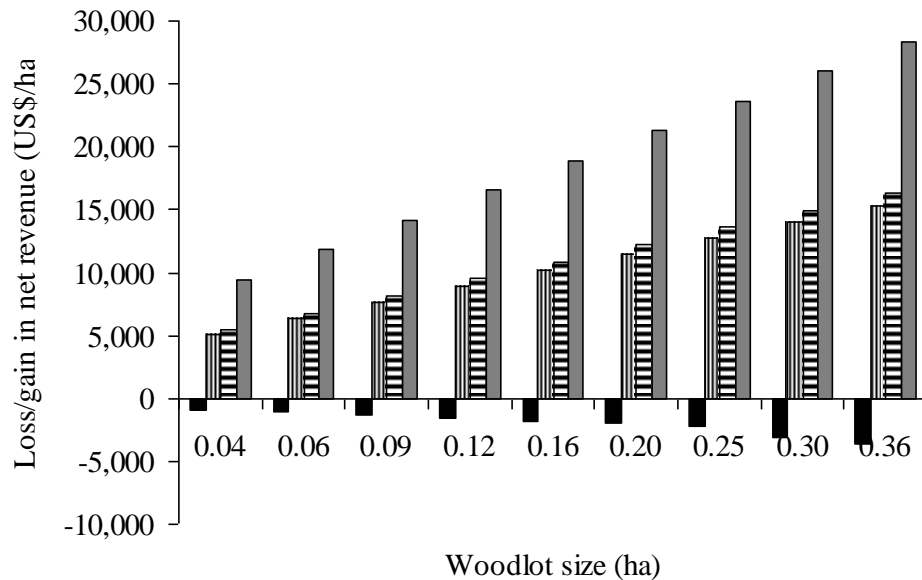


Figure 4.9. Losses in net revenue of maize (dark bars) and respective gains in timber (dark grey bars), pole (black and white with horizontal line bars) and fuel wood (black and white with vertical line bars) production alternative options as a result of edge effects in the crop field and eucalypt woodlots in southern Rwanda.

Eucalyptus trees being very fast growing would be expected to be more competitive than many other tree species. Their suppressive effects on maize was observed (chapter 3), to be primarily due to severe competition for soil moisture, soil nutrients. Similar results have been reported in other studies elsewhere (Bertomeu, 2012; Kidanu et al., 2005; Okorio et al., 1994). Root concentration of eucalypts is reported to be highest in the upper soil layers and this would be expected to explain the observed species' severe competition for soil moisture (Grant et al., 2012). However, Radersma and Ong (2004) indicate that there is a lack of a clear relation between root length density and water extraction near *Grevillea robusta* tree lines and conclude that the competition can be explained by a decrease in water-potential gradient between root and soil at increasing distance from the tree base. Smaller grain yield observed closest to the trees not only resulted from trees' competitive capacity but also from the crop's tendency to use available energy and water for survival rather than for seed production at the tree-crop competition front (Oliver et al., 2005).

The proportion of the crop field affected by trees in woodlot-cropping systems may depend on the species, size and age of the woodlot, field orientation and the local climate among other factors (Kidanu et al., 2005). To illustrate the effect of plot orientation, with the field layout in Fig. 1a, zone A, the maize field affected by the edge is 10.5×12 m (which is equivalent to 35% of the total maize field) while the remaining 65% (19.5×12 m) was edge effect free. By

changing the maize field orientation such that the longer maize crop side (30 m) adjoins the woodlot, zone A would increase by 2.5 times to 10.5×30 m, equal to 87.5%. This means the total loss in maize grain yield per plot would be 2.5 times higher. On the woodlot side, extra wood will be produced in the outer perimeter of 8 m width surrounding the woodlot. Assuming that all four sides of a woodlot are affected, that is, the woodlot is not surrounded by other woodlots or tree stands, small woodlots up to 16×16 m² will be completely prone to the edge effects. Square woodlots of 17 m length will have only one square metre unaffected but this will have no tree since the spacing is commonly 2×2 m. Woodlots of 20×20 m² will have 16 m² unaffected, which is only 4% by area. The woodlot area affected by the edge will increase linearly with the increasing size and will reach 50% for a woodlot of 54×54 m² (about 0.3 ha) (Fig. 4.10). Timber increase due to edge effect will therefore increase with the woodlot size and will be that of unaffected woodlot area plus that of woodlot affected, the latter being 34.5% higher than the former for an equal size proportion.

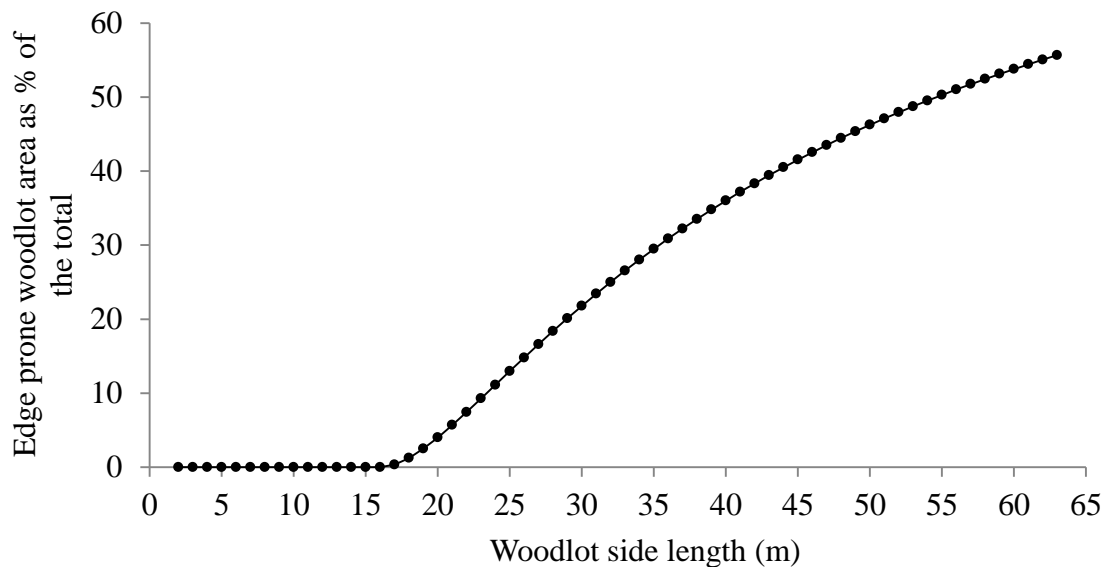


Figure 4.10. Woodlot area affected by the edge area in relation to the woodlot size. Square woodlots of lengths of 16 m will be 100% totally be prone to the edge effect and thereafter, edge affected area will increase linearly until it is 50% by area when a woodlot has $54 \text{ m} \times 54 \text{ m}^2$.

Some authors report tree effects on crops in fields adjacent to trees to extend to a distance equal to/or several times higher than the tree heights from the tree-crop interface (Pinto et al., 2005; Chanie et al., 2013). In this study, the effect of woodlots on maize yield was limited to a distance about half the average woodlot tree height. This could probably (Carberry et al., 2007; Hou et al., 2003) and for eucalypts, the common range was observed be attributed to the distance to which the shade extends to. The edge effect distance from tree-crop interface differ between species and sites to be 12 m in Ethiopian highlands (Kidanu et al., 2005) and up to 44 m in Australia (Sudmeyer et al., 2004). The crop field area prone to woodlot tree suppression observed in this study (zone A) is slightly lower than this range. The differential growth rate observed in this study between trees at the woodlot edge and those at the woodlot centre are the result of intraspecific competition. Competition in plant communities occurs when individuals use resources that would otherwise have been used by their neighbour had

they not been present (Donald, 1963; Cannell et al., 1996). The onset of competition is usually hastened by greater initial plant population densities and, in the case of light competition, by greater stand growth rates due to better site quality (Weiner, 1986), leading to earlier crown closure. The development of structural variation within even-aged plant populations is considered to be due to a hierarchy of exploitation following the onset of intra-specific competition (Harper, 1967; Weiner, 1986), whereby larger plants are able to capture a relatively greater ratio of resources than smaller plants, resulting in greater relative growth rates for larger plants (Ford 1975; Weiner, 1986).

Intraspecific competition has been recognised as a key factor in variation of growth rates, leading to variation in tree sizes within stands (Weiner et al., 2001). Trees in zone D experience severer competition than those in zone C and resources gained from crop fields adjoining the woodlots (Weathers et al., 1997; Pinto et al., 2005). In agroforestry systems where N_2 -fixing tree species are usually grown, the extra growth of border trees may not simply be seen as a positive effect of N-fixation (Van Noordwijk, 1999) since this could be a result of additional resources obtained from adjacent fields, which could otherwise be used by the crops if trees were not there (Cannell et al., 1996). This has been observed to be the case in fallow systems under which below ground resource capture was observed to extend up to 10 m in the crop field from the fallow edge or even beyond in older trees (Van Noordwijk et al., 1996). Roots of some plantation species may extend even further in the crop fields compared to traditional agroforestry fallow species. For example, the lateral extent of tree roots was observed to range from 10 m for *Eucalyptus kochii* to 44 m for *Pinus pinaster* (Sudmeyer et al., 2004).

Woodlot size influences its proportion that is prone to the edge effect. Although the area of the woodlot affected by the edge increases with increasing woodlot size, smaller woodlots are more susceptible than the big ones. A bigger proportion of stems in woodlots with areas of ≤ 0.05 ha for example will be highly affected by the edge, and in fact, woodlot sizes of 14.1×14.1 m (approximately ≤ 0.02 ha) will show 100% edge effects. Most woodlot sizes commonly owned by farmers in the study area fall under this category. Estimation of woodlot biomass in these woodlots without considering edge trees would not be possible. This is not uncommon in small scale agroforestry systems and Van Noordwijk (1999) observes that biomass production of a fallow plot will depend on the fraction of border trees in the plot as a whole. Economic and financial analyses of agroforestry systems in most cases have revealed greater benefits when trees are incorporated in the cropping systems compared to the sole crops. This was observed in eastern Africa with *Eucalyptus* trees where household income were increased by far over sole crops in Ethiopia (Jagger and Pender, 2000; Kidanu et al., 2005; Kebebew and Ayele, 2010) and in Kenya (Peralta and Swinton, 2009). Kebebew and Ayele for example reported that eucalypts contributed 50% of household income relative to major agricultural crops and by allocating 12% of available land to eucalypts, returns were increased from the same unit of land by 90%. They further report a reduction in the returns from the land of 127 and 34% for teff and barley respectively when eucalypt contribution was not considered. Similar results were also reported in the region when eucalypts and another tree species (jatropha) gave superior income compared to crops (cassava) (van Eijck et al., 2012). In

financial terms, the tree component in these systems were observed to adequately compensate for crop yield reductions and even generate additional income (Kidanu et al., 2005). In addition to generating much income, *Eucalyptus* is less sensitive to changes in wages and has yields more than the other crops and can be highly profitable for smallholders (van Eijck et al., 2012). Contradicting observations were made in the Philippines, where maize monocropping was more profitable over timber-maize intercropping (Bertomeu, 2006) while the reverse was true in a study by Magcale-Macandog et al. (2006) in the same country. Bertomeu (2006) however realises that this is not expected and attribute this to a widespread planting of a few fast growing tree species which has led to oversupply and drastic decline in the price of home-grown timber.

Long rotation cycles is another issue in making investment choices by farmers in tree-crop enterprises in high potential areas where landholdings are very small. Annual crops provide yield and income only within three to four months after planting and thereafter two to three times every year while early returns from trees may take at least five years. Farmers in the central highlands of Kenya for example decided to cultivate almost all their farmland with maize and not *Grevillea robusta* alone or grevillea-maize intercrops because of this problem, irrespective their great need for firewood and construction material (Muchiri et al., 2002b). The observation by Muchiri et al. (2002b) which holds true in our study area also is that, although pure tree or tree-crop combinations may be economically more viable than sole crops, farmers are likely to opt for the sole crop alternative because this option provides insurance as far as food security is concerned.

Conclusions

In the crop close to the woodlot (zone A), yield was reduced to 0.5 t ha^{-1} due to inter-specific competition between *Eucalyptus saligna* woodlots and maize compared to the yield of 2.6 t ha^{-1} observed in zone B. Beyond 10.5 m away, the maize yield was not significantly affected by the woodlots. The grain yield increased with the tree-crop interface and therefore with increasing size of woodlot and maize field. The orientation of a crop field is very important and in rectangular plots, the proportion affected by the edge effect may be very significant, especially in very small fields. Intra-specific competition was observed to be high in eucalypt woodlots as evidenced by bigger individual trees in zone C compared to those in zone D. Tree DBH and biomass in zone C were 17.8% and 34.5% higher respectively compared to the values of trees in zone D. In small woodlots of 0.05 ha, 40% will be affected by the edge while those of 0.02 ha will be 100% affected. The effect seemed to be more apparent in older woodlots with bigger individual stems than those with younger, smaller stems. Ignoring edge effects in forest inventories as it is traditionally done in standard forest inventory practice may lead to significant under-estimation in small woodlots. The area of the crop field (and that of the woodlot) subject to the edge effect increased with the size of the woodlot as a result of the increased length of tree-crop interface.

Trade-off analysis showed that, while investing in growing sole maize is profitable ($NPV > 0$), integrating maize and woodlots is far more profitable with the profit increasing with the woodlot area. Average revenue gained from positive tree biomass increments resulting from

edge effects compensated for the corresponding revenue losses due to grain yield decline resulting from the edge effects in these systems. Basing solely on this criterion, integration of maize and eucalypt woodlots is justifiable. However, the seemingly viable option of income generation through combined eucalypt woodlot-annual crop combination may be hampered by the constraint of land availability which is a serious issue in the study area. Farmers who own larger plots are more likely to adopt the crop-eucalypt woodlot combination and therefore more likely to obtain the benefits associated with the practice and vice versa.

CHAPTER 5: Water use by short rotation *Eucalyptus* woodlots in southern Rwanda

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Water use by short rotation *Eucalyptus* woodlots in southern Rwanda

CP Mugunga, D Kool, MT Van Wijk, GMJ Mohren, KE Giller

Abstract

Eucalyptus is abundant in Rwanda, mainly planted in short rotation woodlots, scattered in small clusters over the hilly landscape. A study was done in Butare and Busoro catchments, southern Rwanda from May-November 2007 to estimate water use of eucalypts in representative catchments in Rwanda, establishing a monthly water balance. We compared eucalypt water use to water use of other key crops in the study area and to that of eucalypts elsewhere. The woodlots had small coppice shoots ranging from 2-36 cm breast height diameter and potential tree transpiration recorded was 3 mm d⁻¹. The annual potential tree transpiration was 10% below annual precipitation. Dry month water deficit observed could be covered by reductions in leaf area, stomatal closure and changes in soil water storage. A sensitivity analysis showed that 50% leaf area reduction corresponded to potential tree transpiration decline of 32.8 mm. The deficit may not impact tree growth negatively since dry seasons are usually not active for tree growth. The moderate eucalypt water use rate observed in this study may be a function of trees' small size and low tree stocking since such woodlots had less potential transpiration. The observed eucalypt water use rate is smaller than the range reported for eucalypts in Africa and was also smaller than that of key annual crops in the study area. Managing woodlots as short rotations and increasing initial tree spacing may contribute to resolving issues related to catchment hydrology associated with eucalypt plantations.

Key words: Leaf area index; potential tree transpiration; catchment water balance; Busoro catchment; Butare catchment

1. Introduction

The genus *Eucalyptus* (Myrtaceae) comprises more than 700 species and an unknown number of hybrids and varieties (Boland et al., 2006). Only a few of these have potential in industrial plantations (FAO, 2000). *Eucalyptus* species occur naturally in Australia and in the Philippines, Papua New Guinea, Indonesia and Timor. They grow in diverse ecological conditions with some hardy species growing in semi-arid areas, while others are able to grow on marshy and swampy sites. Owing to their wide ecological adaptation, *Eucalyptus* species are among the most widely cultivated forest trees in the world. In the late 1990s, *Eucalyptus* plantations were estimated to cover at least 12 million ha throughout the tropical zone, 90% of which had been established since 1955 (Turnbull, 1999). There are now >20 million hectares of *Eucalyptus* plantations around the world and interestingly, more than 50% of these plantations occur only in three countries: Brazil (4.2 M ha), India (3.9 M ha) and China (2.6 M ha) (Iglesias-Trabado and Wilstermann, 2008).

In Africa, there is little quantitative information on the planting of *Eucalyptus*. Several *Eucalyptus* species were introduced to the continent in the second half of the nineteenth

century. South Africa has the largest area under *Eucalyptus* plantations of about half a million hectares (Turnbull, 1999). Their first planting included introductions to botanical gardens and private arboreta and once in cultivation, their potential was recognized and they were taken to many parts of the world and planted for ornamental value, windbreaks, land reclamation, and oil production (Turnbull, 1999). In South Africa, commercial plantations were intensified from 1930 onwards, to meet the demand for wood destined for underground mining.

Eucalyptus species were introduced to the eastern Africa region at about the same time as in South Africa (Dessie and Erkossa, 2011). More specifically, plantation forests were established in the early 1900s in Rwanda (Nduwamungu, 2011a), in Kenya (Mathu, 2011) and in Tanzania (Ngaga, 2011); in 1919 in Burundi (Nduwamungu, 2011b) and most recently in Uganda, since the earliest report was in Kigezi District in 1940 (Kaboggoza, 2011). Large scale establishment of industrial plantations in the region began during the period 1911-1960, motivated by the realisation that the slow growing and uneasily propagated indigenous forests would not meet future wood and non-wood forest products (Chamshama, 2011). The introduction of the *Eucalyptus* to East Africa was aimed to meet this demand (Nduwamungu et al., 2008).

As the first step to prepare for plantation forestry expansion in Rwanda, a 200 ha Arboretum was established in Ruhande, Butare (southern province) in which exotic tree species were tested, among which 63 eucalypt species were introduced (Burren, 1995). Plantations expanded over time (Oballa et al., 2005), and by the early 1970s the area of *Eucalyptus* in Ethiopia, Rwanda, Uganda, Kenya and Sudan had reached 95,684 ha (FAO, 1979). The largest plantations at that time were in Ethiopia and Rwanda, at 42,300 ha and 23,000 ha, respectively (Dessie and Erkossa, 2011).

Plantation trees and *Eucalyptus* spp. in particular have provoked controversy regarding their effects on the environment (Vanclay, 2009). Some authors support *Eucalyptus* planting for their economic, social and environmental benefits (Ferraz et al., 2013). On the other hand, *Eucalyptus* are said to mine large amounts of nutrients, rendering the soils poor and unfit for other use (Lemenih, 2004). Plantation trees are reported to take up much water, drying up sites or interfering with water supply for other land uses (Jobbágy et al., 2012). *Eucalyptus* trees are reported to use water in excess of supply from rainfall (Dye and Versfeld, 2007), thereby lowering water tables due to water extraction (Dye, 2013). They also reduce water availability for irrigation due to soil hydrophobicity (water repellence) of their litter and their deep and dense root network (Lane et al., 2004). Replacement of natural vegetation by the exotic eucalypts reduces biodiversity (Brokerhoff et al., 2013) and is said to prevent understorey vegetation growth by exhausting soil water and nutrient resources, which also may affect biodiversity negatively (Bouvet, 1998). These factors, especially the impacts on water relations, raise considerable concern regarding the long-term sustainability of eucalypt plantations.

In Rwanda, as in many high population density areas in the highlands of eastern Africa, most forest plantations are privately-owned small woodlots. Principal owners include individual farmers and businessmen, and institutions like churches, schools, cooperatives and tea

plantation companies (Nduwamungu, 2011a). This ownership pattern renders much of the landscape of Rwanda to be covered predominantly by trees in small groups, rows or single trees scattered on farms. Forest plantations are dominated by *Eucalyptus* species which comprise 78% of all plantations, mostly managed as short rotation coppice (Nduwamungu, 2011a). Woodlots cover about 51% of plantation forests and are also dominated by *Eucalyptus* spp. followed by *Grevillea robusta* and *Pinus* spp. (Nduwamungu et al., 2008; Nduwamungu, 2011a). According to Nduwamungu et al. (2011a), preliminary results of surveys carried out by FAO in 2010 to determine the extent of tree resources outside forests (including woodlots below 0.5 ha) show that these resources cover about 6.6% of total country land area.

Recent policy briefs in Rwanda recommend that the species should be uprooted from wetlands and wherever they grow near water bodies, and their use prohibited in further reforestation (Gahigana, 2006). Unfortunately, information on water use by *Eucalyptus* in Africa is very scanty (Hailu et al., 2003) and particularly lacking in eastern Africa (Bayabil et al., 2010). In Africa, water use by *Eucalyptus* has only been studied extensively in South Africa, in the Congo, and to a lesser extent in Ethiopia, regions with a clearly different agro-ecological environment from the East African highlands.

To quantify water use by *Eucalyptus* trees in the East African highlands we estimated the transpiration of the species in the locally dominant management practice, i.e., small woodlots scattered over the landscape by using a leaf area index (LAI) – diameter at breast height (DBH) relationship. On the basis of the derived relationship, we quantified the water use by short rotation *Eucalyptus* woodlots in representative catchments in southern Rwanda. A monthly water balance of such woodlots was estimated and the overall water use by *Eucalyptus* trees was compared with other key crops in the study area and to that of *Eucalyptus* growing in other regions of the world.

Materials and methods

Study site

The water use by *Eucalyptus* woodlots was estimated in two catchments in southern Rwanda by employing a LAI –DBH relationship. The woodlots in Busoro catchment were planted by farmers for fuelwood production in the first place but also for soil protection against erosion. Those in Butare catchment were planted by the local government primarily for protection purposes. The catchments were chosen because of the availability of clear Google Earth images, the general representativeness of land use, and the clear demarcation of the boundaries of both catchments. One of the catchments borders the town of Butare, and is referred to as the “Butare catchment”. It is located at 2.61° S, 29.72° E and covers an area of 472 ha. The other catchment is situated to the south near the village of Busoro, and is referred to as the “Busoro catchment” (Fig. 5.1). This catchment covers an area of 618 ha and is located at 2.67° S, 29.70° E.

The landscape of the study area is hilly with the villages mostly located on top of the ridges. The predominant land use in both catchments is small-scale agriculture, with short rotation *Eucalyptus* stands for firewood and timber production as well as prevention of soil erosion. The woodlots were primarily planted by farmers. Based on the agro-ecological classification of Delepierre (1982) and the elevational classification of Ndayambaje et al. (2012), the catchments are located in the central plateau and hills zone of Rwanda. The climate is sub-humid with moderately high rainfall of 1200 mm y⁻¹ and mean annual temperature of 21°C. The elevation is around 1700 m above sea level. The soils are derived from granitic rocks and are classified as oxisols or ultisols (Delepierre, 1982).

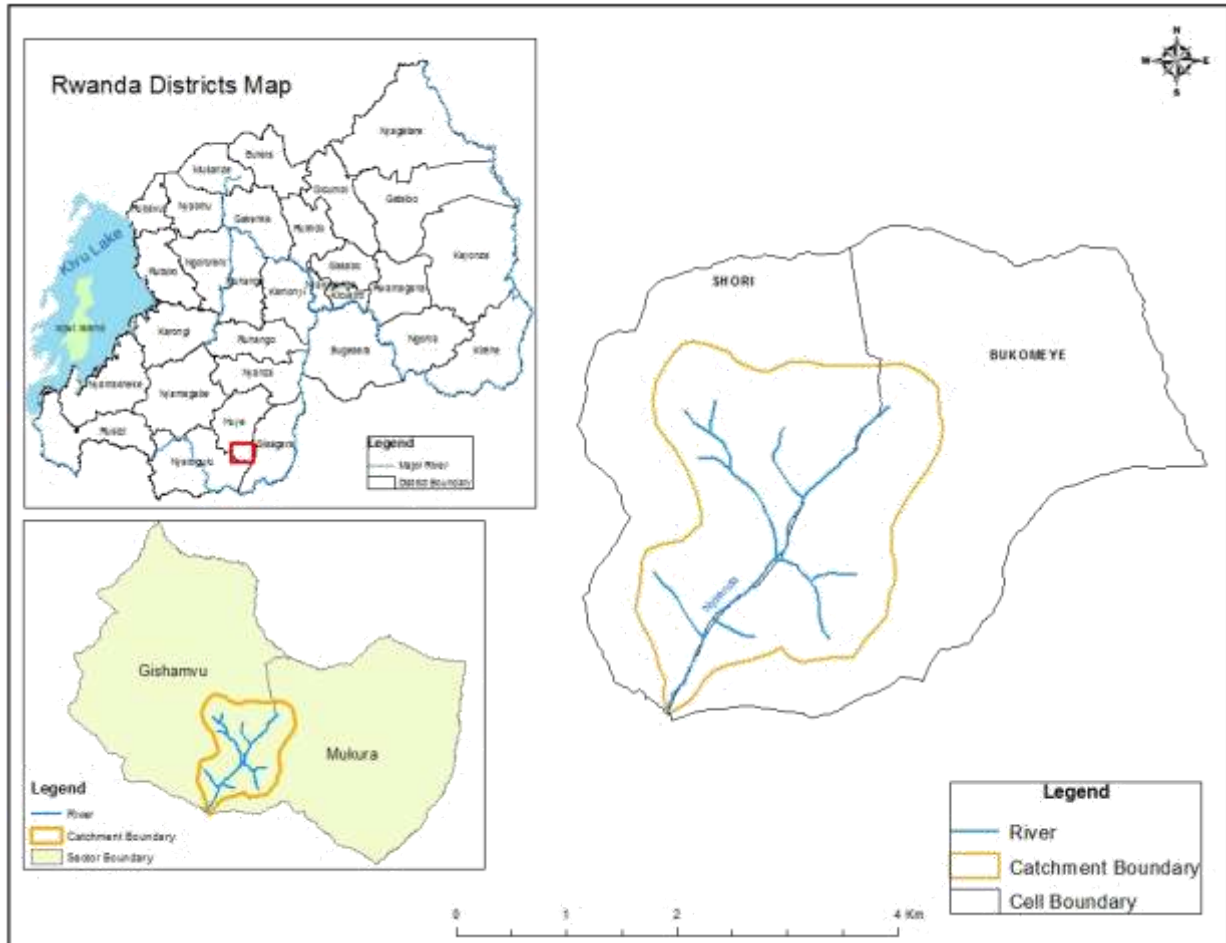


Figure 5.1. The map of Rwanda showing Busoro catchment using ArcGIS. The catchment is located in Huye (former Butare) district and is bordered by Shori cell of Gishamvu sector on the west and Bukomeye cell of Mukura sector on the east.

This study was carried out in one year, 2007, and the LAI-DBH relationship was established during the period May-August which is a transition between a wet and dry season, and the woodlot sampling for LAI determination was done in October-November of the same year. This sampling period is during a short rainy season. The difference in the months of field work may not be important since DBH is not expected to change over such a short period. This period starts at the end of the cropping season and at its beginning, most annual crops were being harvested. Most major crops in the upland fields were perennial, including banana, cassava and some sweet potato. In the relatively flat valleys rice was grown. Most households had an avocado tree and a small *Eucalyptus* plot for domestic wood supply. *E. saligna* was the

predominant species, with a few small stands of *E. tereticornis*, *E. maculata*, *E. maidenii*, *E. microcorys*, and *E. citriodora*. Some woodlots were comprised of mixed species of *Eucalyptus*. Besides *Eucalyptus*, there were a few small stands of other species. *Pinus patula* and *Grevillea robusta* were frequently found scattered on farms, *Euphorbia tirucallii* was used to mark farm boundaries and *Jacaranda mimosifolia* and *Ficus thonningii* were grown in home gardens. *Eucalyptus* woodlots were mostly coppice stands and were located on the steepest parts of the landscape on infertile soils.

Estimation of the water use by *Eucalyptus* trees

We estimated the water use of *Eucalyptus* woodlots in the catchments by quantifying the most important components of the water balance. The water balance can be determined by calculating the amounts of water entering, stored and leaving a system. The inputs and outputs of the water balance of the woodlots taken into account are based on Whitehead and Beadle (2004):

$$P = ET + \Delta S + D + R \quad 1$$

where P is precipitation, ET is total evapotranspiration, ΔS is the change in total soil water storage, D is drainage and R is surface run-off, all in mm d^{-1} .

Subsequently, ET was divided into tree transpiration (E_t), evaporation of leaf interception (E_i), and evaporation from understory and the soil surface (E_u , Whitehead and Beadle, 2004). Precipitation (mm d^{-1}) was derived from daily climate data from the weather stations of Butare and Nyakibanda. These were used to quantify a monthly water balance, thereby capturing the variations in the water balance throughout the year. Generally, E_u is estimated at 10% or less of E_t (Whitehead and Beadle, 2004), which is the value we used. E_u is known to be directly related to the LAI (Callister and Adams, 2006).

E_t and E_i are considered the major components of evapotranspiration (Whitehead and Beadle, 2004). E_i can be estimated as a percentage of annual rainfall, and is normally affected by the leaf area and the shape of the leaves of the vegetation, as well as by the rainfall distribution. Interception losses in *Eucalyptus* woodlots are generally less than those from woodlots of other tree species, which can be explained by the relatively small LAI of *Eucalyptus* trees compared to other tree species (Calder, 1986). LAI for *E. saligna*, one of the most dominant eucalypt species in the study area, was found in a different study in the same area to range from $2.5\text{--}3.5 \text{ m}^2/\text{m}^2$ (Nsabimana *et al.*, 2009). *Eucalyptus* interception losses were reported to range from 4% in South Africa to 34% of the annual rainfall in Australia, India and Israel (Hall *et al.* 1992; Calder, 1986). Most studies report E_i to range from 10–11% of annual rainfall (Whitehead and Beadle, 2004). In South Africa interception losses were higher for large trees than for small ones. As the woodlots in our study areas are mostly coppice stands of several generations, dominated by very small coppice shoots of small DBH ranging from 2–36 cm, a value on the lower side of the range for interception losses seemed most appropriate. We therefore assumed interception losses to be 10% of annual rainfall.

E_t was estimated for individual trees using the model of Radersma et al. (2006). The model describes individual tree transpiration on the basis of an estimation of the leaf area. The model assumes well-watered conditions, and therefore describes the potential tree transpiration (E_{tp} , Radersma et al., 2006; Whitehead and Beadle, 2004). Radersma et al. (2006) used sap-flow measurements to quantify the relationship between leaf area and E_{tp} for *Eucalyptus* trees in sub-humid Kenya, an area with similar agro-ecological conditions to our study site in Rwanda. The variance in measured transpiration accounted for by leaf area, was compared with the variance accounted for by leaf area together with saturation deficit, radiation, and the direct effect of soil water content. Leaf area was the most important determining factor for transpiration across a large range of leaf. The equation that was established for E_{tp} in relation to leaf area is:

$$E_{tp} = c \times LA^{0.63} \quad 2$$

where E_{tp} is in g d^{-1} , c is coefficient $1058 (\text{g d}^{-1} \text{m}^{-2})$, and LA is leaf area (m^2). The explained variance for this equation was 0.62. This equation does not take into account that during the dry season *Eucalyptus* trees normally shed their leaves and thus the relationship between DBH and leaf area changes across seasons (Roberts et al., 1992). During prolonged dry periods, shedding of leaves is common in *Eucalyptus* (Ladiges, 1974) and this can result in large seasonal changes in LAI, depending on the timing and magnitude of leaf production and shedding (Whitehead and Beadle, 2004). As a result, transpiration of *Eucalyptus* during the dry season could be overestimated. Although this equation was established specifically for *E. grandis*, water-use per unit LA is known not to vary greatly among *Eucalyptus* species (Radersma et al., 2006; Whitehead and Beadle, 2004). Therefore the same equation was applied to all *Eucalyptus* species. *E. saligna* dominated in the study area and is morphologically similar to *E. grandis* (Burren, 1995).

The leaf area of individual trees was derived from the leaf biomass, based on the relationship between leaf biomass and DBH. Specific leaf area (SLA) or the ratio of leaf area to leaf mass ($\text{m}^2 \text{kg}^{-1}$) or its inverse, leaf mass per area (LMA) are important Eco physiological parameters widely used to derive canopy leaf area from leaf biomass (Diao et al., 2010). The relationship between leaf dry matter and DBH was established for *E. saligna* by Mugunga et al., (in prep.) as:

$$W_L = \alpha + \beta D^2 \quad 3$$

where W_L is leaf biomass (kg), α is intercept ($0.881461 \text{ kg cm}^{-2}$), β is the coefficient ($0.026311 \text{ kg cm}^{-1}$) and D is the DBH in cm. The explained variance of observed W_L using this equation was 0.88. After calculating the W_L for each tree, LA was obtained by multiplying the leaf biomass by the SLA ($\text{m}^2 \text{kg}^{-1}$). To obtain the values of the SLA , a sample of 80 leaves was taken for the test species. Each leaf was measured for projected leaf area and fresh weight, and the corresponding dry weight was calculated by multiplying with a dry weight:green weight conversion factor of 0.3 derived from the study by Mugunga et al. (in prep.). The steps taken to estimate the E_{tp} of individual trees are shown in Figure 5.2.

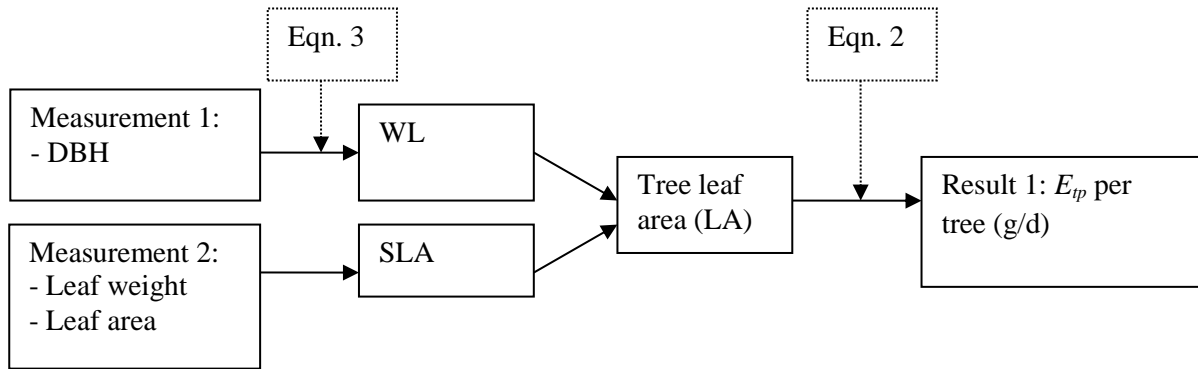


Figure 5.2. Determining the leaf area and estimating the potential transpiration (E_{tp}) for individual trees. DBH is tree diameter at breast height, W_L leaf biomass and SLA is specific leaf area.

After determining the E_{tp} for the individual trees, the results were aggregated to establish the E_{tp} of *Eucalyptus* woodlots across the two catchments. Google Earth images of the catchments showed a very patchy landscape, with small fields of grain crops interspersed with banana fields and *Eucalyptus* woodlots. Some of the *Eucalyptus* woodlots were dense, although bare soil around the trees was visible in most of them. Busoro catchment was divided into two parts: the northern and the southern sub-catchments. Sampling was done by taking one or more 0.02 ha (8 m radius) plots per woodlot depending on woodlot size (Fig. 5.3), while some small woodlots with widths smaller than 16 m were sampled entirely.

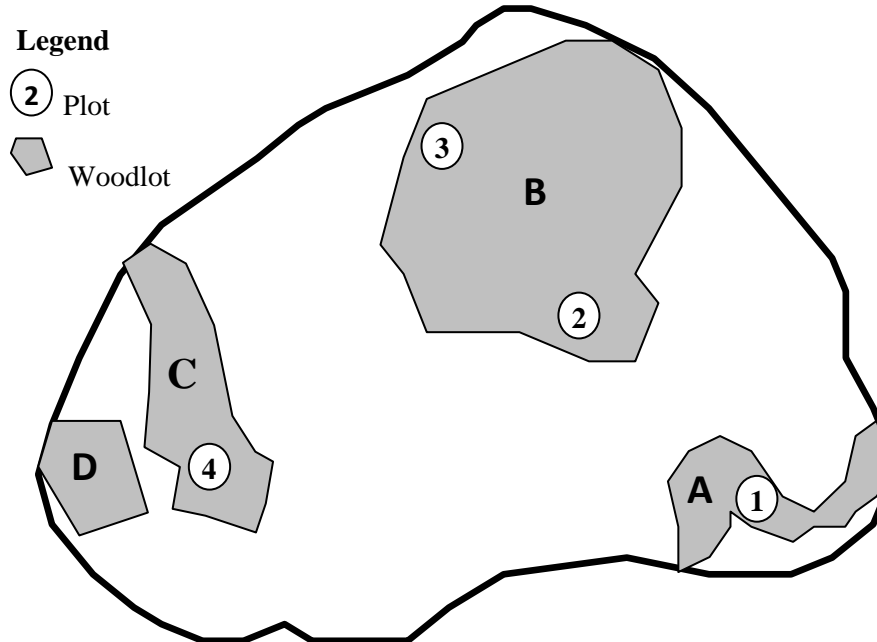


Figure 5.3. Sampling strategy: The grey areas represent a number of woodlots in a catchment. In this example, 4 plots were measured. The first plot was taken in woodlot A. The calculated total tree potential transpiration (E_{tp}) for this plot was extrapolated to the entire area of A. In woodlot B two plots were taken, as this was a larger woodlot with more variation. The average E_{tp} of the two plots was used to compute the total E_{tp} of B. Woodlot C and D are very similar. Only one plot was taken and E_{tp} extrapolated to represent both areas.

In each plot, all tree species were identified and measured for DBH. A total of 59 samples were taken in the Busoro catchment, representing 15 woodlot areas in the southern sub-catchment and 36 woodlot areas in the northern sub-catchment. Seven samples representing 7 woodlot areas were taken from the Butare catchment (Fig. 5.4), giving a total of 66 plots representing 58 woodlot areas.

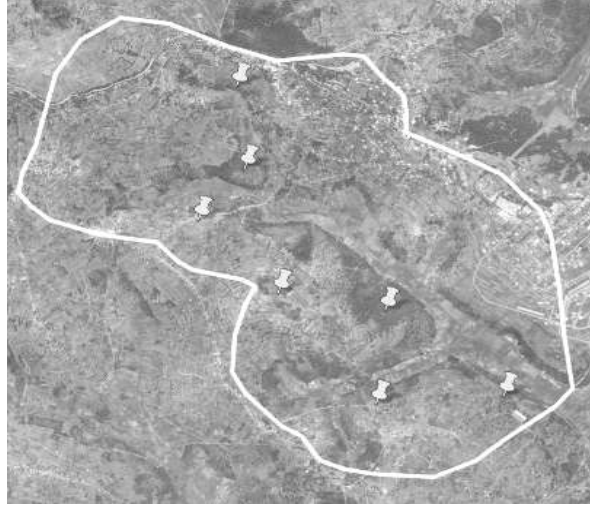


Figure 5.4. Butare catchment. In this catchment 7 plots were measured

The E_{tp} of each individual tree was calculated using the equation of Radersma *et al.* (2006). Subsequently the sum of the E_{tp} of all individual trees was divided by the plot area to determine the E_{tp} per m^2 for each plot. Areas of the woodlots were determined using the Google Earth images. The total E_{tp} of *Eucalyptus* woodlots at catchment level was calculated by aggregating the E_{tp} of all the individual woodlots and converting them to mm d^{-1} (Fig. 5.5). Catchment water balance was developed for the Busoro catchment by considering P as the sole input and the E_{tp} , E_i and E_u the outputs.

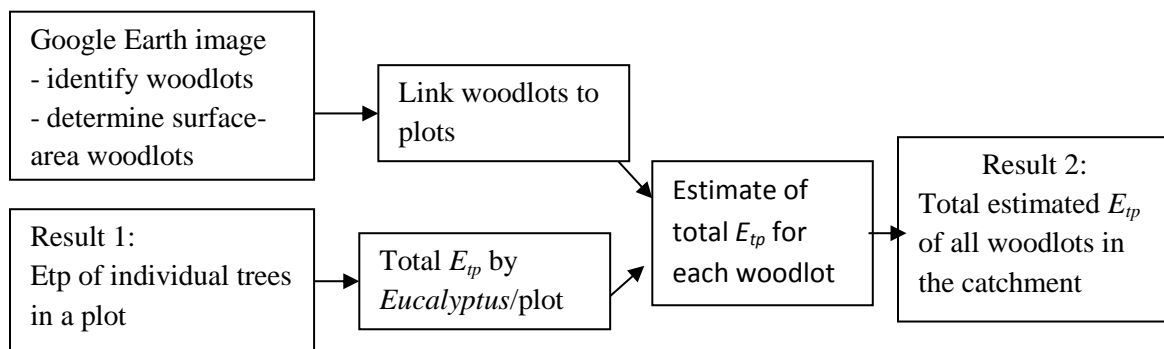


Figure 5.5. Procedure for determining potential tree transpiration (E_{tp}) on catchment level.

For data analysis, linear regression was used to explore the correlation between tree DBH and total tree water use. Mean DBH of each plot was plotted against the total water use of each plot to determine if the total water use per unit area was affected by the size of the trees. It is well known that LAI during the mid- to late-dry seasons is usually 40-50% smaller than the LAI during the wet- and early-dry seasons in *Eucalyptus* forests (Nouvellon *et al.*, 2010).

Reductions in the LAI were reported to be up to 50% even in moist tropical forests in Congo-Guinean forest (de Wasseige et al., 2003), in Mexico (Mass et al., 1995) and in Thailand (Tanaka et al., 2003), while leaf shedding during a severe drought in Australia ranged from 50-97% (Pook et al., 1985).

or this reason, we wanted to adjust the E_{tp} following LAI reductions in the driest months. Unfortunately, the exact LAI variation in our study area is unknown and we did not find reports on LAI variations for the East-African region. To capture seasonal variations in LAI and the resulting changes in E_{tp} , a sensitivity analysis was done by assuming reductions in the amounts of LAI in the driest months of June, July, August and September. To calculate the reductions in E_{tp} , equation 2 above was applied with an assumption that dry season reductions in LA corresponded to reductions in LAI since the trees covered the same area. We used LA reductions of 15, 30 and 50% and calculated the related reductions in E_{tp} here-in-after referred to as adjusted E_{tp} ($Adj.E_{tp}$). The new (adjusted) E_{tp} values were computed as:

$$Adj.E_{tp} = ((c*LA)^{0.63}*((1-r)^{0.63})) = E_{tp.orig}*((1-r)^{0.63}) \quad 4$$

where c is a constant described in equation 2, r is % reduction in LA expressed as a fraction (50% reduction is entered in the equation as 0.5) and $E_{tp.orig}$ = the original E_{tp} (mm month^{-1}). The available water was estimated by assuming that the water accumulated in the root zone during the wetter months was available for evaporation in the drier months. The soil water storage (ΔS) and physiological regulation on water loss by closing stomata were assumed to further reduce the water deficit during dry months, thereby contributing to the closing of the catchment water balance. Annual difference between potential ET and P were attributed to runoff and drainage.

Results

Results on the occurrence of tree species, number of stems ha^{-1} , tree DBH and leaf area are summarized in Tables 5.1, 5.2 and 5.3 for southern Busoro sub-catchment, Butare catchment and the northern Busoro sub-catchment respectively. Woodlots in the study area were dominated by *Eucalyptus* species and about 43 and 35% of all the standing stems per hectare (N) identified were *E. tereticornis* and *E. saligna* respectively. The number of stems ha^{-1} ranged from 50 to 11,500, DBH from 2-36 cm and leaf area from 8-211 m^2 . The SLA's for *E. saligna* and *E. tereticornis* were 12.2 and 16 ($\text{m}^2 \text{kg}^{-1}$) respectively. The total forested area, E_{tp} and relative cover by eucalypt woodlots for Butare catchment, and northern and southern Busoro sub-catchments are presented in Tables 5.4, 5.5 and 5.6 respectively.

In northern Busoro, the sampled woodlots were relatively small with areas ranging from 130 m^2 to 8,000 m^2 , and four larger woodlots with areas of 14,300 - 41,700 m^2 (Table 5.4). There was a wide range in E_{tp} , with 4.58 mm d^{-1} for Plot 38 and only 0.41 mm d^{-1} for Plot 17. This can be explained by the size and number of trees in the respective plots. Plot 17 was a thinly spread coppice stand with a mean DBH of 4-6 cm while Plot 38 represented an area with full grown trees with a mean DBH of 10-36 cm. In the southern Busoro sub-catchment the

samples represented areas ranging from 3,000-62,000 m², and one area of 154,000 m² (Table 5.5).

Table 5.1. No. of sample plots/woodlot, tree species, number of stems, tree DBH and leaf area of woodlot trees in Busoro southern sub-catchment , southern Rwanda

Woodlot	Plot no.	Species	No. of stems		Tree DBH (cm)				Tree leaf area (m ²)		
			(plot ⁻¹)	(ha ⁻¹)	Min.	Max	Av. Std	±	Min	Max	Av. ± Std
*SB-1	35+37	<i>Eucalyptus saligna</i>	47	1,150	1	8	4 ± 2		6	49	18 ± 10
		<i>E. tereticornis</i>	68	1,700	1	8	3 ± 2		5	34	11 ± 7
		<i>E. microcorys</i>	18	450	2	5	3 ± 1		7	24	13 ± 5
		Other	1	<50	20	20	20		176	176	176
SB-2	47	<i>E. saligna</i>	34	1,700	3	20	12 ± 4		4	12	96 ± 43
		<i>E. microcorys</i>	1	50	11	11	11		72	72	72
SB-3	48	<i>E. saligna</i>	56	2,800	1	26	9 ± 6		6	306	71 ± 65
		<i>E.tereticornis</i>	6	300	2	15	10 ± 5		7	99	56 ± 36
SB-4	49	<i>E. saligna</i>	19	950	1	20	5 ± 6		6	212	38 ± 56
		<i>E. tereticornis</i>	2	100	4	5	5 ± 1		16	21	18 ± 3
		<i>E. microcorys</i>	100	4,950	1	23	6 ± 5		6	213	39 ± 43
		Other	1	50	14	14	14		103	103	103
SB-5	50	<i>E. tereticornis</i>	106	5,250	1	13	4 ± 3		5	77	18 ± 15
		<i>E. microcorys</i>	5	250	6	9	8 ± 1		27	51	46 ± 10
		<i>E. saligna</i>	5	250	2	9	4 ± 3		8	54	20 ± 19
		<i>E. tereticornis</i>	13	650	1	4	2 ± 1		5	16	8 ± 4
SB-6	51	<i>E. microcorys</i>	74	3,700	1	20	6 ± 4		6	174	33 ± 35
SB-7	52	Other	1	50	14	14	14		100	100	100
SB-8	53	<i>E. saligna</i>	23	1,150	1	16	8 ± 5		6	146	58 ± 43
		<i>E. tereticornis</i>	45	2,250	2	17	6 ± 4		6	119	29 ± 24
SB-9	54	<i>E. saligna</i>	20	1,000	2	19	10 ± 5		11	192	76 ± 54
		<i>E. Microcorys</i>	71	3,550	1	20	8 ± 5		6	178	52 ± 47
SB-10	55	<i>E. saligna</i>	5	250	6	17	11 ± 4		30	161	87 ± 47
		<i>E. tereticornis</i>	83	4,150	1	13	5 ± 3		5	80	21 ± 17
SB-11	56	<i>E. saligna</i>	68	3,400	2	24	7 ± 6		8	275	54 ± 65
SB-12	57	<i>E. microcorys</i>	15	750	5	19	11 ± 4		19	156	73 ± 37
		<i>E. tereticornis</i>	221	11,000	1	8	2 ± 1		5	34	9 ± 4
		<i>E. saligna</i>	84	4,200	1	21	8 ± 5		6	229	62 ± 54
		<i>E. microcorys</i>	6	300	2	13	8 ± 4		7	84	45 ± 31
		Other	11	550	1	27	8 ± 9		6	287	64 ± 95
SB-13	58	<i>E. saligna</i>	141	7,000	2	8	3 ± 1		10	45	16 ± 8
SB-14	59	<i>E. tereticornis</i>	90	4,500	2	6	3 ± 2		6	26	12 ± 7
SB-15	58-59	<i>E. saligna</i>	141	3,500	2	8	3 ± 1		10	45	16 ± 8
		<i>E. tereticornis</i>	90	2,250	2	6	3 ± 2		6	26	12 ± 7

*Southern Busoro sub-catchment

Table 5.2. No. of sample plots/woodlot, tree species, number of trees, tree DBH and leaf area of woodlot trees in Butare catchment, southern Rwanda

Woodlot	Plot no.	Species	No. of stems		Tree DBH (cm)				Tree leaf area (m ²)		
			(plot ⁻¹)	(ha ⁻¹)	Min.	Max	Av. Std	±	Min	Max	Av. ± Std
*Bt-1	1	<i>Eucalyptus saligna</i>	145	7,200	1	9	2±1		5	47	8±5
Bt-2	2	<i>E. saligna</i>	14	700	1	5	2±1		8	28	12±5
		<i>E. tereticornis</i>	110	5,450	1	7	3±1		5	27	10±4
		<i>E. microcorys</i>	10	500	2	6	4±1		1	3	1±1
		Other	1	50	1	2	1±0		1	1	1±0
Bt-3	3	<i>E. saligna</i>	22	1,100	1	14	3±3		6	119	16±23
		<i>E. tereticornis</i>	33	1,650	1	8	3±2		5	37	10±7
		<i>E. microcorys</i>	1	50	11	11	11±0		7	7	7±0
Bt-4	4	<i>E. saligna</i>	20	1,000	3	28	8±8		12	353	66±95
		<i>E. tereticornis</i>	3	150	3	10	10±4		9	50	27±21
		<i>E. maculata</i>	37	1,850	2	23	8±7		1	23	5±6
		Other	3	150	3	18	9±8		1	15	6±7
Bt-5	5	<i>E. saligna</i>	19	950	1	4	2±1		6	16	10±3
		<i>E. tereticornis</i>	6	300	1	1	1±1		5	5	5±0
		<i>E. citriodora</i>	80	4,000	1	4	3±5		6	308	18±54
Bt-6	6	<i>E. saligna</i>	4	200	2	3	2±0		9	13	11±2
		<i>E. tereticornis</i>	17	850	1	5	2±1		5	19	8±4
		<i>E. microcorys</i>	5	250	2	7	5±3		7	33	22±14
		<i>E. citriodora</i>	61	3,050	1	14	5±4		1	101	25±25
		<i>E. maidenii</i>	18	900	1	14	4±3		6	97	18±21
		Other	4	200	1	2	1±0		6	8	7±1
Bt-7	7	<i>E. saligna</i>	10	500	2	6	4±2		10	34	21±9
		<i>E. tereticornis</i>	62	3,100	1	25	7±7		5	215	45±5.3

*Butare catchment

The highest E_{tp} was found in Plot 57, with a value of 6.44 mm d⁻¹. This plot was located in a very dense *E. saligna* and *E. microcorys* stand, with an average DBH of 8 cm. *E. microcorys* stands were in general more leafy, denser and darker than woodlots with other species. This value may be over-estimated since the water use: leaf area relation of Radersma et al. (2006) is based on the assumption that there is little or no light competition between trees. Linear regression of tree DBH and total tree water use showed no correlation between these two variables, indicating that an increase in mean DBH was compensated for by a decrease in the number of trees per plot (data not shown). For Butare catchment (Table 5.6) the areas covered by a plot range from 50,000-172,000 m². There was less variation than in the Butare catchment because the plots here represented large areas. Actually the woodlots in this catchment belong to the local government and not to the smallholder farmers.

Table 5.3. Number of sample plots/woodlot, tree species, number of stems, tree DBH and leaf area of woodlot trees in Busoro northern sub-catchment (NB), southern Rwanda

Woodlot No	Plot no	Species	No. of stems		Tree DBH (cm)			Tree leaf area (m ²)		
			(plot ⁻¹)	(ha ⁻¹)	Min	Max	Av ± Std	Min	Max	Av ± Std
*NB-1	1	<i>E. saligna</i>	83	450	5	36	17 ± 6	22	520	166 ± 99
NB-2	2	<i>E. tereticornis</i>	280	800	2	28	12 ± 6	8	267	80 ± 56
NB-3	3	<i>E. saligna</i>	8	250	2	6	3 ± 2	10	36	17 ± 9
		<i>E. tereticornis</i>	24	800	2	10	5 ± 2	7	52	19 ± 10
		<i>E. maidenii</i>	10	350	2	11	6 ± 4	8	69	32 ± 24
NB-4	4	<i>E. tereticornis</i>	17	850	2	13	6 ± 3	7	74	28 ± 20
NB-5	5	<i>E. saligna</i>	62	3,100	2	10	5 ± 2	10	69	26 ± 12
		<i>E. microcorys</i>	7	350	5	6	5 ± 1	19	28	22 ± 3
NB-6	6	<i>E. saligna</i>	29	1,450	2	17	6 ± 3	8	164	33 ± 30
		<i>E. tereticornis</i>	10	500	4	12	7 ± 2	15	66	32 ± 15
NB-7	7	<i>E. tereticornis</i>	28	1,400	3	5	3 ± 1	9	19	12 ± 2
NB-8	8	<i>E. saligna</i>	26	150	8	8	8 ± 0	49	49	49 ± 0
		<i>E. microcorys</i>	3	<50	5	5	5 ± 0	22	22	22 ± 0
		Other	27	200	4	30	18 ± 7	16	346	157 ± 81
NB-9	9	<i>E. saligna</i>	13	650	2	9	5 ± 2	10	57	26 ± 15
		<i>E. microcorys</i>	1	50	3	3	3	12	12	12
		Other	7	350	12	23	18 ± 4	75	226	155 ± 51
NB-10	10	Other	14	700	8	25	19 ± 5	42	248	165 ± 63
NB-11	11	<i>E. saligna</i>	33	1,000	2	10	5 ± 2	9	73	25 ± 15
NB-12	12	<i>E. tereticornis</i>	38	1,150	3	12	6 ± 3	9	65	28 ± 17
NB-13	13-14	<i>E. saligna</i>	39	950	2	11	5 ± 2	10	80	26 ± 17
		<i>E. tereticornis</i>	55	1,350	2	8	4 ± 2	6	36	12 ± 7
		<i>E. citriodora</i>	5	100	2	9	6 ± 3	8	46	30 ± 20
		<i>E. maidenii</i>	1	<50	7	7	7	35	35	35
		Other	13	300	2	7	3 ± 2	8	36	15 ± 8
NB-14	15	<i>E. tereticornis</i>	49	1,000	2	8	4 ± 2	6	38	14 ± 8
NB-15	16	<i>E. tereticornis</i>	210	3,500	2	19	5 ± 4	6	142	24 ± 27
NB-16	17	<i>E. tereticornis</i>	31	600	3	6	4 ± 1	9	24	13 ± 4
		<i>E. maidenii</i>	4	100	6	6	6 ± 0	25	25	25 ± 0
		Other	1	<50	9	9	9	54	54	54
NB-17	18	<i>E. tereticornis</i>	37	2,850	2	6	2 ± 1	6	24	9 ± 3
NB-18	19-20	<i>E. tereticornis</i>	134	3,350	1	8	2 ± 1	5	37	10 ± 5
		<i>E. maidenii</i>	2	50	3	10	6 ± 5	12	56	34 ± 31
NB-19	21-22	<i>E. saligna</i>	3	50	8	12	10 ± 2	52	88	66 ± 19
		<i>E. tereticornis</i>	35	850	1	11	7 ± 3	5	64	33 ± 17
		Other	2	50	3	4	4 ± 0	14	15	14 ± 1
NB-20	23-24	<i>E. saligna</i>	75	1,850	2	15	7 ± 3	8	125	41 ± 26
NB-21	25	<i>E. tereticornis</i>	59	2,950	2	5	2 ± 1	6	18	9 ± 3
NB-22	26-27	<i>E. tereticornis</i>	29	700	2	28	2 ± 1	7	310	161 ± 69

		<i>E. camaldulencis</i>	35	850	4	36	17 ± 7	15	454	147 ± 90
NB-23	28-29	<i>E. saligna</i>	64	1,600	2	13	5 ± 3	8	101	27 ± 21
		<i>E. teriticornis</i>	33	800	2	11	5 ± 3	6	60	20 ± 16
NB-24	30	<i>E. saligna</i>	11	550	2	4	2 ± 1	10	20	12 ± 3
		<i>E. teriticornis</i>	65	3,250	1	5	2 ± 1	5	20	8 ± 3
		<i>E. microcorys</i>	2	100	2	2	2 ± 1	7	10	8 ± 2
NB-25	31-36	<i>E. saligna</i>	103	850	1	10	4 ± 2	6	69	19 ± 14
		<i>E. teriticornis</i>	281	2,350	1	19	3 ± 2	5	142	15 ± 16
		<i>E. microcorys</i>	2	<50	2	2	2 ± 1	7	10	8 ± 2
		Other	9	50	5	14	8 ± 4	22	100	46 ± 37
NB-26	35+37	<i>E. saligna</i>	47	1,150	1	8	4 ± 2	6	49	18 ± 10
		<i>E. teriticornis</i>	68	1,700	1	8	3 ± 2	5	34	11 ± 7
		<i>E. microcorys</i>	18	450	2	5	3 ± 1	7	24	13 ± 5
		Other	1	<50	20	20	20	176	176	176
NB-27	38	<i>E. saligna</i>	31	1,550	1	17	7 ± 4	6	158	46 ± 36
		<i>E. teriticornis</i>	63	3,150	1	12	5 ± 3	5	64	21 ± 14
		<i>E. maidenii</i>	2	100	7	17	12 ± 7	32	130	81 ± 69
		Other	25	1,250	1	4	2 ± 1	6	19	10 ± 4
NB-28	39	<i>E. teriticornis</i>	75	3,750	2	9	4 ± 3	6	41	20 ± 15
NB-29	40	<i>E. teriticornis</i>	79	3,950	1	7	3 ± 1	5	27	10 ± 5
		Other	10	500	3	15	8 ± 3	12	107	50 ± 28
NB-30	41	<i>E. saligna</i>	101	5,000	1	7	3 ± 1	6	36	14 ± 7
		<i>E. teriticornis</i>	48	2,400	1	10	3 ± 2	5	48	13 ± 8
		Other	3	150	2	3	3 ± 1	10	13	11 ± 2
NB-31	42	<i>E. saligna</i>	3	150	1	2	2 ± 1	6	11	9 ± 2
		<i>E. teriticornis</i>	89	4,450	1	6	2 ± 1	5	24	8 ± 4
NB-32	43	<i>E. teriticornis</i>	153	7,600	1	5	2 ± 1	5	20	8 ± 3
NB-33	42-43	<i>E. saligna</i>	3	50	1	2	2 ± 1	6	11	9 ± 2
		<i>E. teriticornis</i>	242	6,000	1	6	2 ± 1	5	24	8 ± 3
NB-34	44	<i>E. teriticornis</i>	169	8,400	1	6	2 ± 1	5	22	9 ± 4
NB-35	45	<i>E. saligna</i>	63	3,150	1	16	3 ± 2	6	136	18 ± 19
		<i>E. teriticornis</i>	26	1,300	1	7	3 ± 2	5	33	13 ± 8
		<i>E. maidenii</i>	3	150	5	7	6 ± 1	24	34	29 ± 5
		Other	7	350	2	15	6 ± 4	9	106	35 ± 33
NB-36	46	Other	90	300	20	25	22 ± 2	176	253	211 ± 38

*Northern Busoro sub-catchment

Interestingly the total E_{tp} of the Butare catchment was similar to that of the Busoro catchment, that likely management differences between the different types of owners in the two study areas did not have a major impact on the total water use. The E_{tp} here ranges from 1.34-3.44 mm d⁻¹. The percentage of the area covered by trees in the three sub-catchments ranged from 10.9-17.4% of the total catchment. Table 5.7 gives a summary of the weighted average estimates of E_{tp} for *Eucalyptus* woodlots in the whole study area.

Table 5.4. No. of sample plots/woodlot, woodlot area, estimates of potential tree transpiration (E_{tp}) and extent of cover by *Eucalyptus* woodlots in northern Busoro sub-catchment

Woodlot No	Plot No	Woodlot Area (m ²)	Potential tree transpiration (E_{tp})			Cover (%)
			(g d ⁻¹)	(g d ⁻¹ m ²)	(mm d ⁻¹)	
*NB-1	1	1,800	2.11E+06	1173	1.17	100
NB-2	2	3,400	4.39E+06	1291	1.29	100
NB-3	3	540	5.32E+05	984	0.98	100
NB-4	4	2,100	1.45E+06	692	0.69	98
NB-5	5	530	1.46E+06	2761	2.76	100
NB-6	6	710	1.24E+06	1745	1.74	98
NB-7	7	440	3.10E+05	704	0.7	100
NB-8	8	1,500	1.01E+06	671	0.67	34
NB-9	9	1,600	2.26E+06	1411	1.41	38
NB-10	10	760	1.37E+06	1801	1.8	0
NB-11	11	330	2.55E+05	774	0.77	100
NB-12	12	3,680	3.49E+06	949	0.95	100
NB-13	13-40	2,700	4.69E+06	1738	1.74	92
NB-14	15	500	2.65E+05	530	0.53	100
NB-15	16	600	1.46E+06	2437	2.44	100
NB-16	17	500	2.07E+05	415	0.41	94
NB-17	18	130	1.52E+05	1166	1.17	100
NB-18	19-20	4,000	5.95E+06	1488	1.49	100
NB-19	21-22	670	6.38E+05	953	0.95	100
NB-20	23-24	2,000	3.86E+06	1932	1.93	100
NB-21	25	1,800	2.15E+06	1195	1.19	100
NB-22	26-27	14,300	5.54E+07	3874	3.87	100
NB-23	28-29	3,700	6.66E+06	1799	1.8	100
NB-24	30	7,800	1.23E+07	1580	1.58	100
NB-25	31-36	31,400	5.51E+07	1754	1.75	97
NB-26	35+37	20,000	3.69E+07	1847	1.85	97
NB-27	38	6,800	3.11E+07	4575	4.58	88
NB-28	39	4,100	9.84E+06	2399	2.4	100
NB-29	40	5,600	1.28E+07	2279	2.28	74
NB-30	41	4,200	1.65E+07	3939	3.94	98
NB-31	42	1,800	3.17E+06	1760	1.76	100
NB-32	43	900	2.69E+06	2989	2.99	100
NB-33	42-43	41,700	9.90E+07	2375	2.37	100
NB-34	44	7,400	2.56E+07	3464	3.46	100
NB-35	45	8,000	2.45E+07	3064	3.06	89
NB-36	46	6,000	5.52E+06	920	0.92	0
Total		193,990	4.36E+08	2,250	2.25	95
Total area catchment (m ²)			1,7850,000			
% coverage by trees			10.9			

*Northern Busoro sub-catchment

Table 5.5. Woodlot area, estimates of potential tree transpiration (E_{tp}) and extent of cover by *Eucalyptus* woodlots in southern Busoro sub-catchment

Plot No.	Woodlot area (m ²)	Tree potential transpiration (E_{tp})			Cover (%)
		(g d ⁻¹)	(g d ⁻¹ m ²)	(mm d ⁻¹)	
35+37	28,500	5.26E+07	1847	1.85	100
47	19,000	6.02E+07	3168	3.17	100
48	3,700	1.59E+07	4284	4.28	100
49	38,000	2.16E+08	5682	5.68	98
50	154,000	5.44E+08	3534	3.53	100
51	30,000	1.10E+08	3676	3.68	97
52	74,000	2.44E+08	3299	3.3	100
53	3,000	1.71E+07	5684	5.68	100
54	82,900	2.63E+08	3177	3.18	100
55	51,000	2.56E+08	5015	5.02	100
56	62,000	2.80E+08	4509	4.51	100
57	3,000	1.93E+07	6439	6.44	90
58	3,500	1.43E+07	4100	4.1	100
59	6,500	1.45E+07	2224	2.22	100
58+59	22,000	6.96E+07	3162	3.16	100
Total	581,100	2.18E+09	3746	3.75	100
Total area catchment (m ²)		4,392,000			
% coverage by trees		13.2			

Table 5.6. Woodlot area, estimates of potential tree transpiration (E_{tp}) and extent of cover by *Eucalyptus* woodlots in Butare catchment, southern Rwanda

Plot no.	Woodlot area (m ²)	Potential tree transpiration (E_{tp})			Cover (%)
		(g d ⁻¹)	(g d ⁻¹ m ⁻²)	(mm d ⁻¹)	
1	101,000	280,116,110	2,773	2.77	100
2	120,000	349,608,055	2,913	2.91	100
3	50,000	66,989,693	1,340	1.34	100
4	131,000	247,621,809	1,890	1.89	98
5	155,000	394,592,172	2,546	2.55	100
6	172,000	591,586,594	3,439	3.44	98
7	90,000	298,106,349	3,312	3.31	100
Total	819,000	2,228,620,782	2,721	2.72	99
Total catchment area: 4,717,000 (m ²)					
% coverage by trees: 17.4%					

Precipitation of the study site recorded at Butare and Nyakibanda meteorological (southern Rwanda) stations from 1971-1992 gave a monthly averages ranging from 8 mm in July to 219 mm in April (Figure 5.6). There was no rainfall data recorded beyond 1992 to date following the 1994 genocide. Rainfall is bi-modal with a long dry period in June-September and a short one in February, with December and January also being relatively dry. The rainfall observed

here is similar to that recorded previously for the central plateau agro-ecological zone (Delepierre, 1982). Applying the unadjusted models of Whitehead and Beadle (2004) and Lane et al. (2004), the annual difference between P and E_{tp} was estimated to be -84.0 mm y^{-1} .

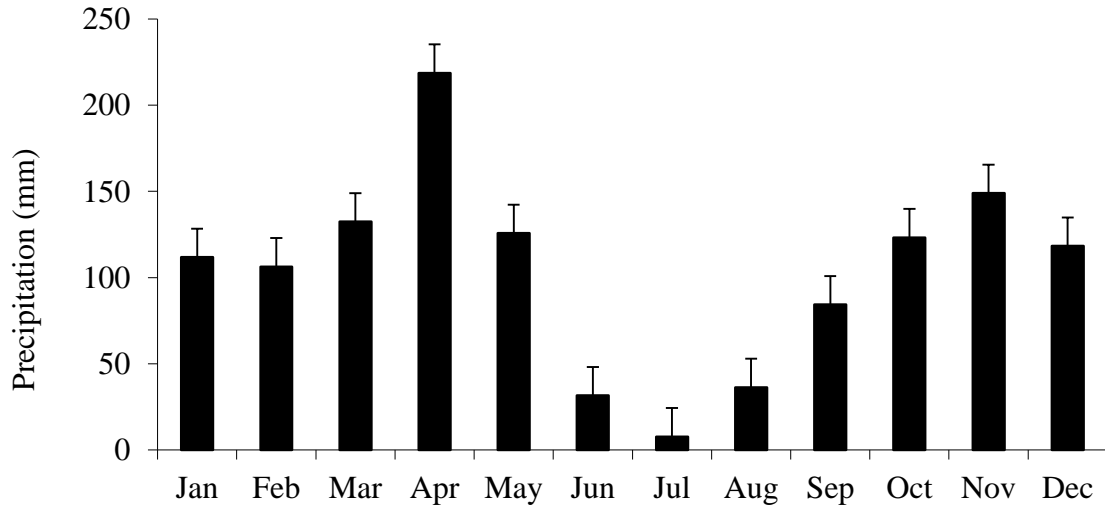


Figure 5.6: Monthly mean precipitation of Butare area, southern Rwanda based on precipitation rates collected in Butare in 1971-1993.

Table 5.7. Weighted average estimates of potential tree transpiration (E_{tp}) for *Eucalyptus* woodlots in Busoro and Butare catchments in southern Rwanda

Catchment	Catchment area	Total area under trees	% area covered by trees	Tree transpiration			Eucalypt cover /woodlot
	(m ²)	(m ²)	(%)	(g d ⁻¹)	(g m ²)	d ⁻¹ (Mm d ⁻¹)	(%)
Busoro							
Northern sub-catchment	1,785,000	193,000	10.8	4.36E+08	2,250	2.25	95
Southern sub-catchment	4,392,000	603,200	13.7	2.18E+09	3,612	3.61	100
Total	6,177,000	796,200	12.9	2.61E+09	3,371	3.37	99
Butare							
	4,717,000	819,000	17.4	2.26E+09	2,721	2.72	99
Weighted average			14.6		3,037	3.04	99

This was obtained as follows: Rainfall = 1245 mm; average tree water consumption (E_{tp}) = $3 \text{ mm d}^{-1} \times 365 \text{ days y}^{-1} = 1095 \text{ mm}$; E_i (10% of P) = 124.5 mm; E_u (10% of E_{tp}) = 109.5 mm. Subtracting the amount of water that potentially left the system from water that entered the system gave: $P - E_{tp} - E_i - E_u = 1245 - 1095 - 124.5 - 109.5 = -84.0 \text{ (mm/y)}$ (-7.0 mm/month or -2 mm d^{-1}). Since water outflow cannot exceed water inflow, this indicates that annual actual E_t did not equal annual E_{tp} , likely due to drought stress in the drier months. Rainfall in the region may seem not to be enough to support a year-round actual tree transpiration of 3 mm d^{-1} . Estimated E_{tp} was much larger than P in the months of June, July, August and September and

to a lesser extent in January (Table 5.8). To obtain a more accurate estimation of the actual E_t , both E_{tp} and E_u were adjusted according to the assumed 15, 30 and 50% reductions in the LAI during June-September. A reduction in LAI of 15% cut the yearly water balance by half from -84.0 to -39.3. Decreasing LAI further resulted to a neutral to positive yearly water surpluses of 2 and 62.8 mm y^{-1} for LAI reductions of 30 and 50% respectively (Table 5.9). Transpiration deficits during the drier months were assumed to be buffered by both the soil storage accumulated in wet months and stomatal closure as trees' physiological response to water deficits. Even in drier years, the differences in soil water storage are not likely to affect the water balance on an annual basis (Everson, 2001).

Table 5.8. E_{tp} in a monthly water balance (WB) estimated for *Eucalyptus* woodlots in Busoro catchment, Butare, southern Rwanda based on Whitehead and Beadle (2004). P = precipitation, E_{tp} = potential tree transpiration, E_i = interception losses, E_u = evaporation from the undergrowth and soil surface

Month	* P (mm)	E_{tp} (mm)	E_i (mm)	E_u (mm)	$D+R$ (mm)
January	112	93.0	11.2	9.3	-1.5
February	106	84.0	10.6	8.4	3.0
March	132	93.0	13.2	9.3	16.5
April	219	90.0	21.9	9.0	98.1
May	126	93.0	12.6	9.3	11.1
June	32	90.0	3.2	9.0	-70.2
July	8	93.0	0.8	9.3	-95.1
August	36	93.0	3.6	9.3	-69.9
September	84	90.0	8.4	9.0	-23.4
October	123	93.0	12.3	9.3	8.4
November	149	90.0	14.9	9.0	35.1
December	118	93.0	11.8	9.3	3.9
Total	1,245.0	1,095.0	124.5	109.5	-84.0

* $P = ET + \Delta S + D + R$, where P is precipitation, ET is total evapotranspiration, ΔS is the change in total soil water storage, D is drainage and R is surface run-off, all in mm d^{-1}

Discussion

Eucalyptus woodlots in the study area mainly comprised of coppice stands of varying rotations of small young shoots with small DBH. Coppice shoot sizes are known to decline with the number of harvesting cycles in short rotation *Eucalyptus* plantations. Stand above ground biomass was observed to decrease as the number of cutting cycles increased (Zewdie, 2008). Specific leaf area (SLA) obtained for *E. saligna* in this study (16 m² kg⁻¹) is only slightly higher than that obtained for *E. grandis* in Nyabeda, western Kenya (Radersma et al., 2006). This is not surprising since the two species are known to be morphologically similar (Burren, 1995).

The estimated water use of *Eucalyptus* during the rainy season was generally less than that of key crops in the study area (Table 5.9). The average potential evapotranspiration (ET_c) of *Eucalyptus* was reported to be 3.6 mm d^{-1} (1,329 mm y^{-1}) (Gislain, 2008). In comparison, the

reference ET , ET_o^1 is between 4 and 5 mm d⁻¹ (i.e. 1,460 and 1,825 mm y⁻¹) (Table 5.10). Crops such as banana, cassava, beans and maize generally have a potential evapotranspiration (ET_c) slightly greater than 5 mm d⁻¹ in the mid-season (Gislain, 2008). There are few studies of *Eucalyptus* water use in Africa (Hailu et al., 2003). The water use observed in this study (3 mm d⁻¹) is on the lower side of the range observed for *Eucalyptus* and for forests of other tree species.

Table 5.9. The adjusted potential tree transpiration through LA reductions of 15, 30 and 50% in dry seasons in a monthly water balance in Butare, southern Rwanda. P = precipitation, $Adj E_{tp}$ = adjusted potential tree transpiration, E_i = interception losses, $Adj.E_u$ = adjusted evaporation from under storey and soil surface, and R and D which add up to give water surplus.

Month	* P (mm)	$Adj.E_{tp}$ (mm)			E_i (mm)	$Adj.E_u$ (mm)			Surplus ($R+D$)		
		$LA_{85\%}$	$LA_{70\%}$	$LA_{50\%}$		$LA_{85\%}$	$LA_{70\%}$	$LA_{50\%}$	$LA_{85\%}$	$LA_{70\%}$	$LA_{50\%}$
January	112.0	91.6			11.2	9.2			0.0		
February	106.0	84.0	84.0	84.0	10.6	8.4	8.4	8.4	3.0	3.0	3.0
March	132.0	93.0	93.0	93.0	13.2	9.3	9.3	9.3	16.5	16.5	16.5
April	219.0	90.0	90.0	90.0	21.9	9.0	9.0	9.0	98.1	98.1	98.1
May	126.0	93.0	93.0	93.0	12.6	9.3	9.3	9.3	11.1	11.1	11.1
June	32.0	81.2	71.9	58.2	3.2	8.1	7.2	5.8	-60.6	-50.3	-35.2
July	8.0	82.1	72.7	58.8	0.8	8.2	7.3	5.9	-83.2	-72.8	-57.5
August	36.0	82.1	72.7	58.8	3.6	8.2	7.3	5.9	-58.0	-47.6	-32.3
September	84.0	81.2	71.9	58.2	8.4	8.1	7.2	5.8	-13.8	-3.5	11.6
October	123.0	93.0	93.0	93.0	12.3	9.3	9.3	9.3	8.4	8.4	8.4
November	149.0	90.0	90.0	90.0	14.9	9.0	9.0	9.0	35.1	35.1	35.1
December	118.0	93.0	93.0	93.0	11.8	9.3	9.3	9.3	3.9	3.9	3.9
Total	1245.0	1054.4	925.1	869.9	124.5	105.4	92.3	87.0	-39.3	2.0	62.8

* $P = ET + \Delta S + D + R$, where P is precipitation, ET is total evapotranspiration, ΔS is the change in total soil water storage, D is drainage and R is surface run-off, all in mm d⁻¹

Water use ranged from 0.4 to 9.9 mm d⁻¹ in Africa (Fritzsche et al., 2006; Radersma et al., 2006) and from 0.4 to 49.0 mm d⁻¹ around the world (Morris et al., 2004). The small rate observed in the study area is attributed to the small tree size and LAI of the frequently coppiced *Eucalyptus*. Tree water use is strongly related to heartwood area which in turn is directly dependent on tree DBH (Vertessy et al., 1995). *E. saligna* was observed to have a smaller LAI than other *Eucalyptus* and other tree species in Rwanda (Nsabimana et al., 2009). *Eucalyptus* woodlots generally have smaller LAI than other tree species (Nsabimana et al.,

¹ ET_o is the reference evapotranspiration widely used in the field of irrigation as a reference surface for potential crop evapotranspiration. "The reference surface is a hypothetical grass reference crop with an assumed crop height of 0.12 m, a fixed surface resistance of 70 s m⁻¹ and an albedo of 0.23. The reference surface closely resembles an extensive surface of green, well-watered grass of a uniform height, actively growing and completely shading the ground. The fixed surface resistance of 70 s m⁻¹ implies a moderately dry soil surface resulting from about a weekly irrigation frequency." ET_o can be computed from meteorological data or estimated from pan evaporation. (FAO guideline, Allen et al., 1998)

2009; Calder, 1986) which may explain their lower water use rates. The observed rates in this study are within the range of values given by Rutter (1968) who report a relatively good agreement in studies that compared extrapolated results of stand transpiration with those obtained by the Bowen ratio (Dunin and Greenwood, 1986) and eddy correlation methods (Berbigier et al., 1996).

Table 5.10. Comparison of the potential evapotranspiration of *Eucalyptus* with ET_0 and the potential evapotranspiration (ET_c) of key crops in Rwanda. The ET_c of the reference crops for the first and second year of growth are derived from a research conducted by the former Institut des Sciences Agronomiques du Rwanda, now Rwanda Agriculture Board (RAB) (Gislain, 2008)

Month	1ET_0 (mm)	Banana ET_c (mm)		Cassava ET_c (mm)		Maize ET_c (mm)		Beans ET_c (mm)		<i>Eucalyptus</i> ET_c (mm)
		yr1	yr2	yr 1	yr 2	yr 1	yr 2	yr 1	yr 2	Pot.
January	149	78	155	124	171	0	0	47	0	114
February	148	84	140	112	154	56	0	0	0	103
March	155	130	171	109	140	124	0	0	0	116
April	141	150	165	60	66	135	0	0	0	121
May	130	155	171	0	47	149	53	0	0	115
June	123	150	165	0	45	105	105	0	60	102
July	130	155	171	0	47	0	140	0	155	103
August	143	155	171	0	47	0	155	0	155	106
September	147	150	165	0	45	0	60	0	0	107
October	161	78	155	47	109	0	0	62	0	115
November	141	75	150	90	165	0	0	171	0	114
December	140	78	155	124	171	0	0	155	0	114
Total	1,708	1,437	1,932	665	1,204	569	512	435	370	1,329

In addition to the observed low water use rates, *Eucalyptus* species are efficient users of water. *Eucalyptus* water use range from 0.3 to 0.8 g of water g⁻¹ of dry matter produced; water use efficiency by other trees is up to about 0.9 g while that of some crops is 1.1 to 3.6 fold higher (Davidson, 1993). Biophysical water use efficiency however may need to be supported by an economic water use efficiency, which considers the financial returns of an investment rather than relying on biomass produced, since not all biomass may be converted to monetary terms. Water use efficiency is broadly defined by Hood (2002) as maximising returns and minimising environmental impacts of every mega litre (ML) of water used by plants. This is seen as an important concept for informing different aspects of land and water resource management including reducing overall water use and producing high and/or better quality yields (Wise et al., 2001). Hood (2002) developed an economic water use efficiency applied on irrigated crops by measuring the net economic return per ML of water, estimated by subtracting the costs of all non-water inputs from the total per hectare returns to production and dividing by the average annual quantity of water evapotranspired per hectare. This concept was successfully applied by Wise et al. (2011) in comparing water use by indigenous and introduced forests.

Transpiration is strongly dependent on LAI (Benyon and Nambiar, 2006) which declines during the dry seasons (Shi et al., 2012; Morris et al., 2004) as it depends on available water (White et al., 2010) and temperature (Gholz, 1982). LAI of fast growing eucalypt forests is also known to be dynamic both seasonally and between years (Le Maire et al., 2011), especially in evergreen forests prone to seasonal droughts (Turner et al., 2008). Large structural changes of the canopy that occur over the rotation affect LAI, crown cover, leaf angle and specific leaf area (*SLA*) (Laclau et al., 2010). There is a strong linear relationship between tree LAI and mean daily water use for a wide range of *Eucalyptus* species grown under similar climatic conditions (Hatton et al., 1995). It was observed that, for each additional 100 mm y^{-1} in rainfall, LAI increased 0.3 units and productivity increased 2.3 Mg $ha^{-1}y^{-1}$, resulting in an additional increment of 7.6 Mg $ha^{-1}y^{-1}$ per added unit of LAI (Ryan et al., 2010). *E. saligna* (Ares and Fownes, 2000) and *E. grandis* (Kallarackal and Somen, 1997) are known to exhibit stomatal closure in response to stronger atmospheric vapour pressure deficits to reduce the E_{tp} .

LAI during the mid- to late-dry seasons is reported to decline following water shortages in the soil, and is usually reported to be 40-50% smaller than its value during the wet- and early-dry seasons in *Eucalyptus* forests (Nouvellon et al., 2010). Reductions in the LAI were reported to be up to 50% even in moist tropical forests in Congo-Guinean forest (de Wasseige et al. 2003), in Mexico (Mass et al., 1995) and in Thailand (Tanaka et al., 2003). In this study, an assumed dry season reduction in LAI (E_{tp}) of 15% was observed to lead to a deficit in catchment water balance but a further increase in the reduction (30%) lead to a neutral catchment water balance where $P-ET$ gave a sum of $R+D$ of about 2 mm. A higher rate LAI (E_{tp}) reduction which may be expected to reach a maximum of 50% would lead to a more positive surplus. To a lesser extent, stomatal closure during the dry season as the soil water declines may reduce tree transpiration further, thereby reducing water loss by trees (Wullschlegel et al., 1998) and thus, leading to a closed catchment water balance. The effect of this on tree growth was assumed to be minimal or none since dry seasons are usually dormant periods for tree growth.

According to climate data from 1982-1993 collected from Butare and Nyakibanda meteorological stations in the study area, the area experiences annual dry seasons of up to 4 months during which mean monthly temperatures range 17.1-20.6°C. This agrees with the general temperature for the central plateau agro-ecological zone reported by Delepierre (1982) that the mean annual temperature is 19.5°C, with mean annual maxima and minima of 25.7 and 13.3°C respectively. The temperatures are most likely to have increased following global climate change in the region, as affected by land cover change. In Africa, the role of cover changes, both natural and human induced, in modifying regional climates is perhaps most marked (Xue, 1997). A study on African climate concludes that additional stresses on water resources are expected regardless of the rainfall alterations which are not clearly defined currently (Hulme et al., 2001).

Considering plantation water use at catchment level, there is a general agreement in most of the findings that tree species produce less surface run off, ground water recharge and stream

flow than shallow rooted vegetation such as crops, pastures and grasslands (Benyon et al., 2007). Crops have a transpiring leaf area for only part of the year (Robinson et al., 2006) and consequently, recharge under agro-ecosystems may be one to two orders of magnitude greater than under natural vegetation (Tennant and Hall, 2001). This may be a result of the trees' capacity to improve soil physical properties due to the addition of large quantities of litter fall and root biomass (Scroth et al., 1995), root activity of trees, root induced soil biological activity and tree roots leaving macropores following their decomposition (Van Noordwijk et al., 1991). Litter fall of 6-9 Mg ha⁻¹ y⁻¹ was reported in *Eucalyptus* plantations at canopy closure in a large range of tropical environments (du Toit, 2008). Likewise, large quantities of fine roots were reported in *Eucalyptus* plantations in the Congo with wide and deep soil volume exploration, which were thought to prevent water and nutrient losses by deep drainage and to contribute to the maximisation of resource uses (Laclau et al., 2003).

If well situated within a catchment, trees may improve instead of negatively affect catchment hydrology. Trees located close to rivers may affect stream flows more severely owing to their large size and more abundant leaves with greater leaf area (Dvorak, 2012). Dvorak et al. (2012) further observed that the severity of problems associated with water extraction are greater where plantations are large in size and cover most of the catchment area or in places with seasonally low rainfall. In this way, well planned agroforestry practice may provide an option of locating trees on sites where they can provide net hydrological benefits, such as controlling run off rates where this is a problem (Keenan et al., 2004). In our study, most woodlots were growing on the hillside, reasonably far from the River Nyiranda. Negligible impacts are expected in catchments in which plantations are established on sites which are hydro-geologically isolated from streams (O'Loughlin and Nambiar, 2001).

The potential impact of land use change on catchment hydrology may depend on the type of native vegetation replaced by exotics and on the extent of the land area covered by the new species (Albaugh et al., 2013). Unfortunately, there are no records of the vegetation types that were replaced by *Eucalyptus* woodlots in the study area and the respective water use. Generally, indigenous vegetation was reported to have less annual evapotranspiration than exotics since they usually show shallow roots and are seasonally dormant (Albaugh et al., 2013).

The rate of water use by *Eucalyptus* trees depends on the amount of rainfall and soil water available (Dye et al., 2001) and more water is transpired when water is plenty and vice versa (White et al., 2010). Similarly, wetter catchments were observed to have greater water flow reductions in South Africa (Scott et al., 2000). With this fact, more water flow reductions may be expected in the northern and western Provinces of Rwanda where *Eucalyptus* populations are larger (Nduwamungu et al., 2008) and the two regions have higher elevation and higher average annual rainfall compared to the eastern and the southern Provinces (GoR, 2011). However, Gush et al. (2002) cautioned that the results from one catchment cannot be extrapolated to other areas with confidence because of differences in local climates.

Conclusion

Short rotation eucalypt stands in the study area had small shoot sizes with DBH ranging from 2-36 cm due to frequent coppicing. Potential tree transpiration (E_{tp}) for eucalypt woodlots ranged from 0.41-6.44 mm d⁻¹ (mean 2.93 mm d⁻¹) for the Busoro catchment and from 1.34-3.44 mm d⁻¹ (mean 2.72 mm d⁻¹) for the Butare catchment. The overall mean E_{tp} for eucalypt woodlots in the study area was about 3 mm d⁻¹, equivalent to 1,095 (mm y⁻¹). It should be noted that this is a conservative estimate, based on the assumption that runoff and drainage only occurred when available water exceeded annual E_{tp} . Long term precipitation of the study site was 1,245 mm y⁻¹ on average, which is 150 mm y⁻¹ greater than the observed potential tree transpiration. The observed surplus annual catchment water balance suggests that short rotation eucalypt woodlots may not negatively affect the hydrology. Small deficits observed during the driest months of June, July, August and September may be eliminated by physiological responses of trees through reducing LAI and to a lesser extent, stomatal closure. A sensitivity analysis on the variation of the LA during the dry seasons resulted into E_{tp} decreases. A possible maximum 50% LA reduction for example lead to a corresponding decrease of 32.8 mm in E_{tp} . Extra shortages if any may be evened out by soil storage, which can be recharged in the wetter months. The observed deficit in soil water during dry seasons is not expected to impact tree growth negatively since trees usually do not grow during the dry seasons. The moderate eucalypt water use rate observed in this study may be a function of the small size of trees and low stand density since less dense woodlots with smaller trees had much less potential transpiration. Managing the woodlots at short rotations (to produce small stems), plus increasing initial tree spacing, may contribute to the reduced catchment hydrology problems associated with eucalypt plantations. The observed water use rate of eucalypt woodlots was less than that of many annual crops in the study area and eucalypt trees in Africa and elsewhere in the world. In addition to biophysical water use efficiency, economic water use efficiency may be a useful option for water use comparisons among different production options.

CHAPTER 6: SYNTHESIS

SYNTHESIS

Forestry and land use dynamics

Land use in Rwanda has been influenced by a number of factors, the main ones being climate, socioeconomic (culture and population dynamics) and government policies (REMA, 2009). During the period 1990-2002, cultivated land increased from 782,500 to 899,133 ha in absolute terms between 1984 and 2002 (Fig. 6.1) (Mpyisi et al., 2003). This increase occurred at the expense of land under pasture, fallow and woodlots. The shifting from one land use to another has stabilised over the near past but land is being farmed intensively without fallow at all (REMA, 2009). Example is the extent of eucalypt woodlots of areas <0.5 ha which were estimated to occupy 7% of the total area in 2002 (Mpyisi et al., 2003) (Fig. 6.1) and 6.6% by FAO in 2010 (FAO, 2010). The forest policy reports this to be 8% of the total country area (MFM, 2010).

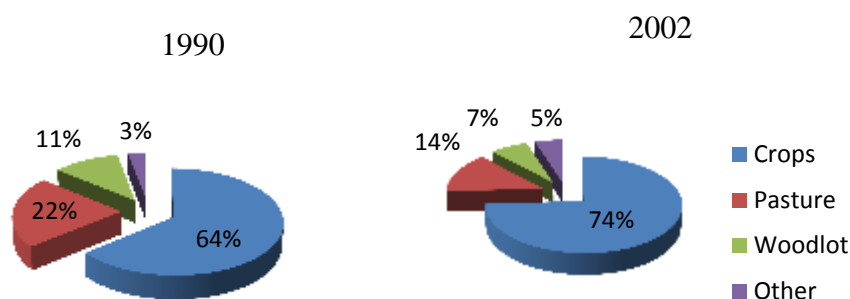


Figure 6.1. Land use dynamics in Rwanda during 1990-2002 – after Mpyisi et al. (2003). The two pies from left to right show the situation in 1990 and 2002 respectively.

Land pressure remains in an interplay where different land uses compete for the limited land available. Land scarcity limits the scope for further expansion of farming into uncultivated lands except in the marshlands which are also limited in size. To improve crop productivity the focus should be on optimizing the use of inputs and natural resources for sustainable food production. For this purpose, Rwandan government has adopted a simplified land use consolidation model in which case farmers in a given area grow the priority food crops in a synchronised way while keeping their land rights unchanged (Kathiresan, 2012). This author reports that crop intensification under this program has increased by 18 fold between 2008 and 2011. As a result, an impressive improvement in productivity has been observed in the past decade which has resulted in significant crop yields as illustrated in Figure 6.2 (WFP, 2012). Forestry on the other hand needs to increase in order to meet a target cover of 30% by 2020 (MFM, 2010).

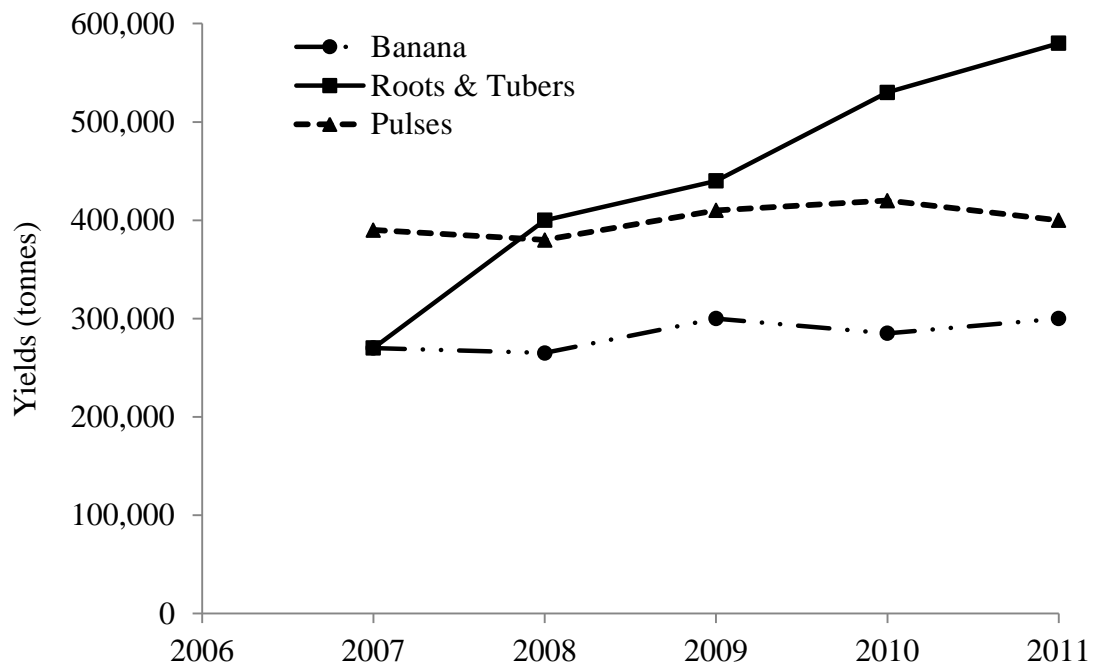


Figure 6.2. Production of main staple crops (2007-2011). (Adapted from WFP, 2012).

Maize is among the top priority crops in the current Rwandan crop intensification program agriculture due to increased importance in food security assurance and the crop has seen an increasing area under cultivation (Fig. 6.3) and grain yield. Average maize grain yield increased from 600 to 2,600 kg ha⁻¹ (WFP, 2012).

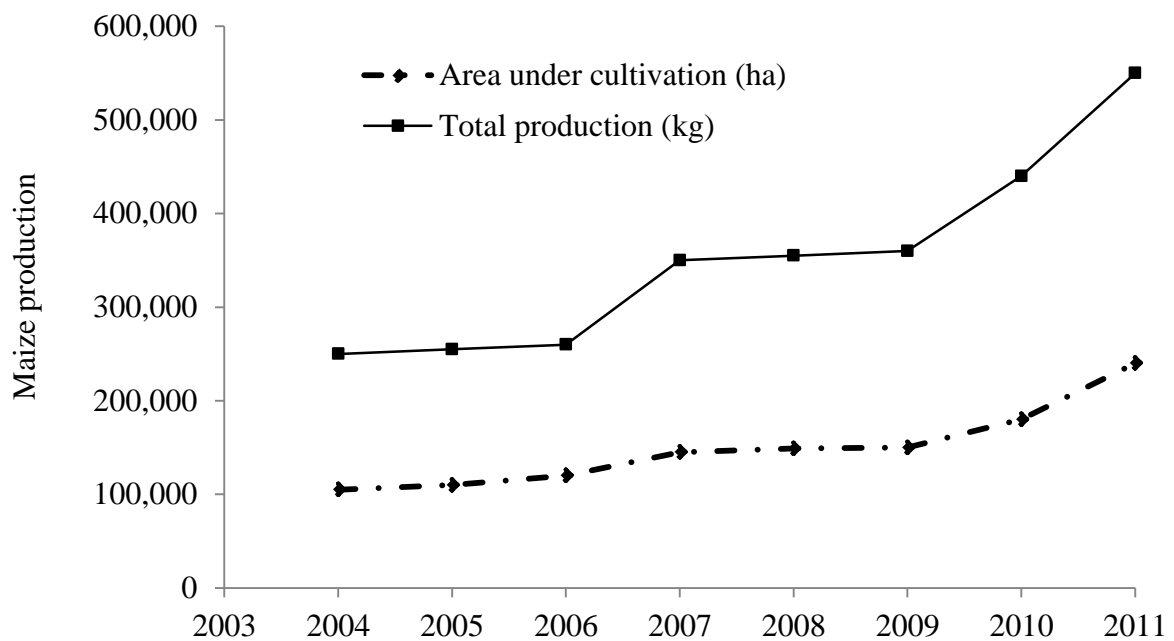


Figure 6.3. Trends in maize farming area and grain yield (Adapted from Kathiresan, 2012).

Forests only cover a total area of 330,576 ha, of which 215,739 ha are natural forests and 114,837 ha represent forest plantations (MFM, 2010). *Eucalyptus* cover the largest plantation area compared to other exotic timber species and in 2008, it was reported to be 64% of the total plantation area in the country, with 65% of it owned by the government; 26% by smallholder farmers and 9% by private institutions (Nduwamungu et al., 2008). On-farm tree planting is mainly in the form of woodlots ranging from very small to reasonably large stands which are scattered on the landscape countrywide, though most are found in the southern and the western Provinces (Nduwamungu et al., 2008). Small woodlots with areas of <0.5 ha and trees on farm occupy 222,520 ha equivalent to conventional forests (FAO, 2010). This raises the total area of national forest cover to 553,096 ha, equivalent to 21% of total country area such that, in total, natural forests occur on 8% of total country area while plantations on 13% (MFM, 2010).

Plant biomass is the main source of domestic energy, and is used by 96% of Rwandans (MFM, 2010). While pointing out that there is information gap on fuelwood consumption rate data in Rwanda, Ndayambaje and Mohren (2011) report that about 93% of Rwandan population rely on firewood and charcoal as the sources of domestic energy. An estimated 72% of this comes from eucalypts (MINERENA, 2013). Data on this is however scanty and none consistent. An example is depicted in Figure 6.4 below, where firewood as a source of domestic energy is reported to be 57 and 80% by two different sources. This difference is unlikely a result of different dates but rather, errors in the estimations. Tree planting has been practiced to meet wood demand and to conserve soils against erosion by the government on public land and by farmers and private organisations on their respective land holdings. In most plantings, *Eucalyptus* species have been favoured. The proportion of *Eucalyptus* in the total exotic plantation species used in reforestation and afforestation was cited to be 65% by Ndayambaje and Mohren (2011) and $\geq 80\%$ by MFM (2010).

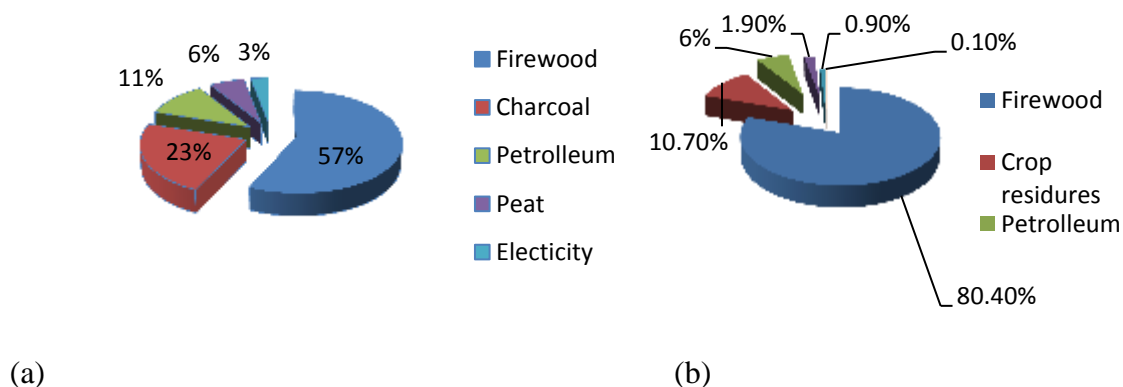


Figure 6.4. Primary energy balance for Rwanda (a) Adapted from: MFM (2010), (b) from MINEFI-DGTPE (2005).

Small size of the woodlots is borne from the fact that each household grow trees in aspiration to become self-sufficient not only in food production but also in firewood and other wood product requirements. As a result, woodlots are scattered on the landscape in the same fashion the households are distributed. This makes an interspaced woodlot-crop field mosaics on Rwandan landscape in which trees and crops adjoin. Interspecific competition between very

eucalypt woodlots and crops becomes a very important attribute of this farming system. Competition between trees and annual crops is key to understanding the benefits of agroforestry. Trees compete with crops for light, water and nutrients reducing crop yields significantly (Rowe et al., 2005). Severe competition is recorded in areas with low water areas (Ong and Leakey, 1999; Rao et al., 1998). *Eucalyptus* spp. are particularly competitive that they significantly reduce crop growth due to severe competition for soil moisture (Forrester et al., 2010), for nutrients (Harrison et al., 2000) and for light (Bertomeu, 2012) depending on local conditions where they grow.

On-farm single tree and woodlot planting in agroforestry systems seen as a viable option to contribute to the alleviation of the acute fuelwood crisis in the country was observed to be only possible on farms of at least 0.76 ha to allow also for food production (Ndayambaje and Mohren, 2011). The authors further suggest that for this purpose, the choice of tree species is necessary, as well as consideration of multipurpose tree species having fuelwood attributes, high biomass production rates and increased positive effects on crop yields. Whilst most of these criteria apply to eucalypts, the latter being very competitive excludes them. This also was the basis for the Rwandan policy makers' decision to restrict eucalypt planting to areas far from marshlands (Gahigana, 2006). However, farmers keep silent on the no use of eucalypts and continue to plant them, simply because they get multiple products from the trees at a comparatively shorter period in relation to other timber species. It has been observed that, when trees provide higher economic advantages, farmers usually ignore their competitiveness and integrate them with crops (Ndayambaje et al., 2014; Bucagu et al., 2013; Tang et al. 2012). *Eucalyptus* ensure readily available income to farmers which can be used at any time of need. An example is the seasonal selling of timber to cover for school fees when school year or term starts. The same applies during peak periods of food deficit to enable households buy food from local markets.

Increased competition for land resource will continue to grow with increased pressures from intensive and varied land use in the east African highlands and in Rwanda in particular. Food production needs expansion to ensure food security for a growing population. Cash crops need to be intensified to increase the country's economy through export. Forestry needs to expand afforestation and reforestation programmes in Rwanda to meet the national goal of attaining 30% forest cover (MFM, 2010) and elsewhere in the region to cope with increasing wood demand. It therefore becomes imperative to undertake a systematic evaluation to analyse trade-offs among different land uses so as to wisely meet different community production goals with less or no conflicts among the different land uses. Resource assessment and quantification of the extent of competition between different plants in an agroforestry system and the analysis of the benefits of different production alternatives in relation to plant growth and yields becomes necessary as tools to aid in proper planning.

This thesis focused on two levels: First, plot/field/stand level to study tree allometry in biomass estimation, the edge effects as interspecific (tree-crop) and intraspecific (tree-tree) interaction; and the trade-offs of maize and eucalypt woodlot monocultures versus combined production alternatives and second, regional/catchment level for making use of stand level

allometric equation to developing a regional allometric equation to estimate tree above ground biomass in both coppice and planted stands of *Eucalypt saligna* common in the study region, and to estimate eucalypt woodlot water use.

The use of *Eucalyptus*

***Eucalyptus* in plantations**

Eucalyptus species have been in use as plantation trees for many years in the tropics. Native in Australia and in the Philippines, Papua New Guinea, Indonesia and Timor, they grow in diverse ecological conditions ranging from semi-arid to marshy and swampy sites. They are adapted to almost all types of soils (KEFRI, 2010), making them to be most widely grown plantation trees in the world. In the late 1990s, *Eucalyptus* plantations covered at least 12 M ha throughout the tropics, 90% had been established since 1955 (Turnbull, 1999). In 2008, there were >20 million hectares of *Eucalyptus* plantations around the world, $\geq 50\%$ of these occurring only in Brazil (4.2 M ha), India (3.9 M ha) and China (2.6 M ha) (Iglesias-Trabado and Wilstermann, 2008).

Eucalyptus species were introduced to the eastern Africa region in the early 1900s (Nduwamungu, 2011a). Large scale establishment of industrial plantations in the region began during the period 1911-1960, motivated by the fact that the slow growing and uneasily propagated indigenous forests would not meet future wood and non-wood forest products (Chamshama, 2011). They were to meet this demand (Nduwamungu et al., 2008; Thorhaug and Miller, 1986). Plantations expanded over time (Oballa et al., 2005), and by the early 1970s the area of *Eucalyptus* in Ethiopia, Rwanda, Uganda, Kenya and Sudan had reached 95,684 ha (FAO, 1979). The largest plantations at that time were in Ethiopia and Rwanda, at 42,300 ha and 23,000 ha, respectively (Dessie and Erkossa, 2011). Figure 6.5 shows the extent of forest and eucalypt plantations in the region.

The eucalypts are mainly managed for timber production but also as short rotation woodlots for fuelwood supply as well as for the supply of fibre for pulp and paper industries (Cuoto et al., 2001). Recently, eucalypts have gained impetus as short rotations for bioenergy (Couto et al. 2003). Renewable energy contributes about 19% of global energy matrix after fossil fuel (78%) (Renewables, 2015). Rwandans depend on wood biomass as their primary source of energy, and in 2007, forests contributed 80% of total national domestic energy as firewood (57%) and charcoal (23%) (MFM, 2010). With the increasing eucalypt planting, there is a need to develop a management tool for estimating biomass. Since a trend shows an increasing state of managing the plantations both as coppice and short rotation stands, allometric equations for both stand types are required. It was seen necessary to develop a general allometric equation for both coppice and planted stands for southern Rwanda. This is detailed in Chapter 2, where general allometric equations were developed using easily measured DBH and height either singly or in combination for estimating above ground biomass for *E. saligna* tree species for total tree and for the components of stem, branch, leaf and bark. The general equations developed at stand level were more precise with R^2 values of 0.97-0.98 and this became comparatively less precise when applied to the regional scale, where R^2 reduced to

0.93-0.95. These are however reasonably good and can be used successfully. With expanding eucalypt planting in the region, the equations may be used across the region as a tool in forest resource management and assessment programmes. Equations developed for this species can be applied to the same species in a broader extent in the east African region, especially using equations combining DBH and height, either directly or with parameterisation. Using equations that combine both height and DBH has been recommended especially when applying allometric equations in regions other than where the equations were developed (Ketterings *et al.*, 2001).

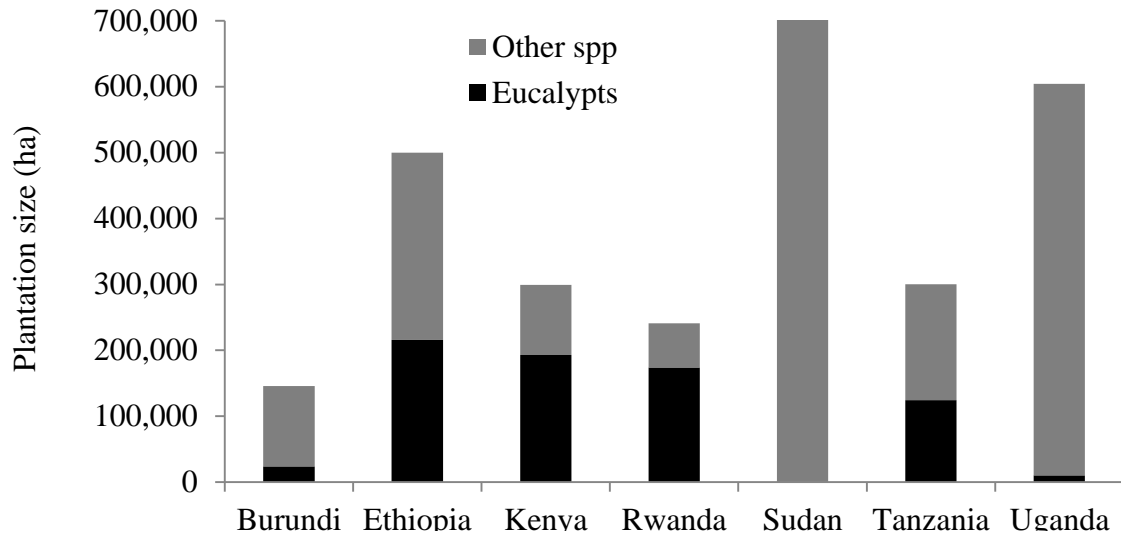


Figure 6.5. Areas of *Eucalyptus* and total plantation forests in the east African countries. *Figures for Sudan include planted gum arabic plantations and managed gum gardens.

Sources: Chamshama (2011); FAO (2009); Kabogoza (2011); Mathu (2011); Nduwamungu (2011a, b); Ngaga (2011).

***Eucalyptus* in agroforestry systems**

Eucalyptus are competitive tree species and therefore not preferred in agroforestry systems. However, in some situations, especially of land and wood shortages resulting from population growth, these species find their way to the croplands. This is a typical case in the east African region, especially in Rwanda and Ethiopia (Dessie and Erkossa, 2011; Kidanu et al., 2005).

Tree-crop interaction and competition mechanism

In agroforestry systems, valuable trees or shrubs are combined with annual crops on the same plot of land, and this may be a viable alternative for some smallholder landowners to produce food and obtain wood products for domestic use (Hagan et al., 2009; Jose and Gordon, 2008). In these systems however, component plants differ in terms of patterns of resource use, which may lead to a more efficient exploitation of resource than when grown in monocultures (Willey, 1990; Francis, 1989). Environmental resources may be more completely absorbed

and used to build crop biomass by intercropping (Amini et al., 2013; Anil et al., 1998). This may imply that, the component plants are not competing for the same ecological niches and that interspecies competition is weaker than intra-specific competition for a given factor (Vandermeer, 1989). The same would occur if resource requirements of the component species are separated in space or in time (Cannel et al., 1996). Trees being larger usually outcompete the annual crops, leading to yield reductions. Trees and crops compete more for soil moisture and nutrients below ground and for light above ground (Jose et al., 2000). Below ground interactions, particularly the competition for water and nutrients are more likely to occur when the root systems of the associated plants grow into the same area of the soil profile (Wanvestraut et al., 2004; Gillespie et al., 2000). Light becomes more limiting on plant growth only when there is sufficient water and nitrogen available to the crop (Zamora et al., 2008; Monteith et al., 1991).

Eucalyptus have developed mechanisms which enable them to adapt to various soil and environmental conditions. At the same time, these adaptation mechanisms enable the species to be very efficient competitors against other plants they happen to grow together or in close proximity. Such mechanisms include a fast root growth which spreads out (lateral growth) and deep (vertical growth) to explore a considerable volume of soil so as to increase accessibility to soil moisture and soil nutrients. A one year eucalypt plantation in the Congo showed a quickly developing root system which extended to a depth beyond 3 m deep and up to the middle of the inter-row (Bouillet et al., 2002). Similar findings were reported in Kenya, where eucalypt root system reached 4 m depth in 11-months-old trees (Jama et al., 1998). A 2-year-old eucalypt plantation indicated had roots down to 6.2 m in Brazil (Pacheco and Louzada, 1991). In *E. saligna* roots were observed to be concentrated in the top 61 cm soil layer (Skolemann, 1974). Cohen, 1997 report a high density of fine roots in the top 20 cm soil for *E. urophylla*, while the same was reported in different eucalypt species in different place (Radersma and Ong, 2002; Kidanu, 2005; Nyadzi, 2003). Stone and Kalisz (1991) identified more than 30 tree species that develop roots over long distances, and to a depth of up to 61 m below the ground. Lateral extent of tree roots was observed up to 10 m in *Eucalyptus hockii* (Sudmeyer et al., 2004). They also observed that root density declined with distance from the trees, and that rate of decrease was less for *E. globulus* and least for *P. pinaster*. This means that *E. globulus* had more roots extending far away from the trees than *P. panister* did.

Spatial arrangements

Owing to their competitive nature, *Eucalyptus* species are not usually spatially mixed with annual crops and whenever it is done, it is when the trees are still very young. Few such cases have been reported, examples include intercropping rice and beans with young *E. camaldulensis* (Ceccon, 2005) and the same crops with *E. urophylla* (Ceccon, 2007) both in Minas Gerais state of Brazil. Intercrops of *E. camaldulensis* with maize and beans were also reported in Morogoro, Tanzania (Chingaiepe, 1985), that of cabbage and *E. torelliana* in the uplands of Mindanao, the Philippines (Nissen et al., 1999). Spatially zoned system is the most commonly practiced system where crop-eucalypt combinations are managed, especially tree lines or linear tree planting and to a lesser extent, the woodlots. Presumably, farmers expect

that this setting will minimise tree-crop competition and therefore yield decline and the associated losses.

Eucalyptus tree lines can be intended for wood production like in Ethiopia where they are used to produce poles and firewood (Kidanu et al., 2005) or established as windbreaks like in Australia to protect crops against wind damage (Sudmeyer et al., 2005; 2002). Woodlots and adjacent annual crops is another spatially zoned system which is predominant in Kivu region (Rwanda, eastern DRC and east western Uganda) (Majaliwa et al., 2015). At times, monoculture plantations of eucalypts have crop fields adjacent to them, where a tree-crop interface exist with interaction between trees and crops, making two the components of an agroforestry system. An example is the sugarcane-eucalypt systems in Sao Paulo as described by Pinto et al. (2005). Sequential eucalypt-crop systems also are known in which case crops are grown in areas previously occupied by eucalypt plantations.

Why farmers grow Eucalyptus trees?

Eucalypts are believed to perform better than indigenous species and most other exotics in height and diameter growth, giving a variety of products (Casson, 1997) in 5-7 year rotations (Better et al., 2001). The trees can be easily cultivated and managed, they are not palatable to animals and therefore easy to protect, they produce superior short fibre for paper making, make high quality charcoal, provide durable furniture timber, and are useful for shelterbelts, soil erosion control, land reclamation and drainage. They can be managed as short rotation coppice stands and coppice cycles of up to seven cutting times are possible (Zewdi et al., 2009). In addition, eucalypts provide valuable non-wood forest products such as honey, tannins, essential oils, medicines and others. It is also promoted as a source of livelihood for underprivileged in many developing countries. Farmers are aware that eucalypts are least capital intensive since it needs minimum maintenance with assured returns.

Farmers usually intend to be self-sufficient and integrate trees and crops in the same land unit so as to get crop yield for food (and sometimes for cash) to ensure food security and wood and non-timber products from trees. Firewood requirements is one of the reasons that drive farmers to grow trees on their fields. To mix trees and crops on the same piece of land, the expected tendency would be for the farmers to grow trees which are compatible with crops. Surprisingly however, they sometimes grow very competitive species (Schaler et al., 2003). When trees provide higher economic advantages for example, farmers usually ignore tree competitiveness and integrate them with crops. This has been observed in a range of study conditions across countries and continents. Examples include in Africa: Rwanda (Ndayambaje et al., 2014), Ethiopia (Zerihun and Kaba, 2011) and Ghana (Isaac et al., 2009); in Asia: China (Tang et al., 2012) and India (Banyal et al., 2011; Palsaniya et al., 2010), in Europe: France (Mary et al., 1998) and in America: Costa Rica (Schaller et al 2003). *Eucalyptus urophylla* was rated as very good by farmers in Kageyo area (northern Rwanda) owing to its wood characteristics but very poor in terms of compatibility with crops, yet 50% of all farmers grew the species (Bucagu et al., 2013).

Why *Eucalyptus* are said to be bad?

The eucalypt controversy

The eucalypts have been receiving criticisms and support from different people. The controversy on eucalypts started in India in more than two decades ago, when issues on social forestry as a strategy to provide rural poor with fuel rose (Casson, 1997). The critics allege that eucalypts affect the environment negatively, some claiming that their benefits are outweighed by their negative impacts. The main problem cited is the ecosystem hydro-ecological imbalances. Other queries such as the loss of soil fertility, allelopathy and replacement of conventional forests are sometimes raised (Casson, 1997).

Eucalyptus water use

Following controversy on water use by *Eucalyptus*, many studies have been directed toward this aspect at the individual tree and stand levels with fewer studies at the landscape (catchment or watershed) level (Stanturf et al., 2013). Very high water use rates have been reported in some studies. Examples include tree daily water rates of 90 litres (7.35 mm) under high water supply situations, dropping to 40-50 litres (3.3-4.4 mm) under water scarce conditions (Mukundi and Palanisami, 2011). In the Argentine pampas, Engel (2005) studied the impact of a 40 year *E. camaldulensis* plantation and reported a transpiration rate of 2-3.7 mm d⁻¹. They further observed lowering of the ground water levels of 0.5 m with respect to the surrounding grassland and a resultant hydraulic gradient which induced flow from the grassland areas into the plantation that lead to the rising of the plantation water table at night. They conclude that *E. camaldulensis* used both groundwater and vadose zone moisture sources, depending on soil water availability, with an estimated 67% of total annual tree water use contributed from groundwater sources. Tree use of ground water was reported for the same species in Karnataka, India (Calder et al., 1997). Some studies, example in South Africa and South America have shown faster water use by *Eucalyptus* than by the native grasslands and in some cases the agricultural croplands they replace (Le Maitre et al., 2002; Nosoetto et al., 2005; Dye, 2013). Many studies in south Africa have revealed that eucalypts use much water and affect stream flow as a result of using ground water (Gush, 2006; Scott et al., 2004; Scott et al., 2000).

On the other hand, eucalypts have been reported in other areas to use less or comparable water quantities as other tree species or other vegetation types. A study was carried out in Ethiopia to compare water use by exotic tree species (*Eucalyptus globulus* and *Cupressus lusitanica*) and the indigenous trees (*Croton macrostachys* and *Podocarpus falcatus*). Depending on the phenology, *C. macrostachys* showed higher water use rates than *Eucalyptus globulus* when conditions were favourable (Fetene and Beck, 2004). *Eucalyptus camaldulensis* and *E. tereticornis* had comparable water use rates with *Tectona grandis*, an indigenous tree species in India, only that the latter shed its leaves during pre-monsoon periods showing no transpiration during this period (Kallarackal, 2010). Myers et al. (1996) observed that water use by *Eucalyptus* does not exceed that of *Pinus* when soil water is not limiting. *Eucalyptus camaldulensis* and *Casuarina cunninghamiana* used equal amounts of water per tree in

Girgarre, Victoria, Australia (Morris and Collopy, 1999). Some studies in China also indicated that eucalypt water use did not exceed the local rainfall (Yan, 2009; Lane et al., 2004). In Karnataka, India, water use of young *Eucalyptus* plantation on a medium depth soil was not greater than that of indigenous, semi-arid dry deciduous forest and both eucalypt and the indigenous forest water use did not exceed annual rainfall (Calder, 1992). This was also observed by (Salemi et al., 2013; Noretto et al., 2005), who attribute this to be in part due to lower interception of precipitation in *Eucalyptus* canopies offsetting their higher transpiration rates. One assessment concluded that total water loss from *Eucalyptus* stands in the tropics is often no greater than from native hardwoods but is greater than from (non-irrigated) agricultural crops (Cannell, 1999).

Eucalyptus water use has been observed to depend on water availability, that is, the amount of rainfall and the soil moisture (Ferraz et al., 2013; Dye et al. 2001) and more water is transpired when water is plenty and vice versa (White et al. 2010). Similarly, wetter catchments were observed to have greater water flow reductions in South Africa (Scott et al. 2000). Myers et al. (1996) and White et al. (2014) report annual water use in two sites of 975 and 1394 mm in rain fed treatments versus 1102 and 1779 mm in irrigated treatments, accounting for approximately 67% and 58% of annual precipitation and irrigation inputs respectively. They associated the increased water use in the irrigated stands with higher sapwood area, leaf area index and transpiration per unit leaf area but there was no difference in the response of canopy conductance with air saturation deficit between treatments. In the same line, the effects on stream flow was reported to be most apparent in dry regions and years than in wetter ones (Scott, 2005) and on sites with coarse-textured soils (Noretto et al., 2005; Busch, 2009).

Considering plantation water use at catchment level, there is a general agreement in most of the findings that tree species produce less surface run off, ground water recharge and stream flow than shallow rooted vegetation such as crops, pastures and grasslands (Benyon et al., 2007). Crops have a transpiring leaf area for only part of the year (Robinson et al., 2006) and consequently, recharge under agro-ecosystems may be one to two orders of magnitude greater than under natural vegetation (Tennant and Hall, 2001). Forests are deep rooted compared to most other vegetation types (e.g. grasses). In addition to their high leaf area index, forests tend to consume more water (and intercept more rainfall, which is lost to the atmosphere through evaporation) than other vegetation types. This is illustrated in a review by Gilmour (2014) using data adapted from Zhang et al. (2001), relating water use (expressed as annual evapotranspiration) to the prevailing rainfall as summarised in Figure 6.6a. Under drier conditions, the amount of water use does not differ between vegetation types (Figure 6a, up to 500 mm per annum), showing the dependence of water use rate on the amount of water available (Ferraz et al., 2013). This high water use generally leads to lower total water yield from forested catchments than with grass catchments as illustrated by Zhang (2001) (Fig. 6.6b).

Studies in other areas agree with this observation. An annual water loss from a 100% forest catchment was observed to use much more water than a grassland under the same conditions

in the UK (Marc and Robinson, 2007). Forest plantations and short rotation woodlot systems established on grasslands, arable lands, and native forests reduced streamflow and lowered the water table in some situations due to a combination of higher transpiration rates and, compared to grassland and cropland, higher interception and evaporation of precipitation (Rodriguez-Suarez et al., 2011; Updegraff et al., 2004). Rising water table following clearance of natural forests for agriculture purposes in Australia (MDBC, 1992) and replanting using eucalypt species to lower the ground water (Morris and Collopy, 1999) clearly show the potential of trees to take much water from the soil. Eucalyptus high transpiration rates associated with the rapid growth is a primary factor affecting water use (Cannell, 1999; Vance et al., 2014). The increasing rate of water use with water availability may mean that forests in humid temperate and tropical regions lead to lower total yield compared with pasture and mixed vegetation catchments (Gilmour, 2014).

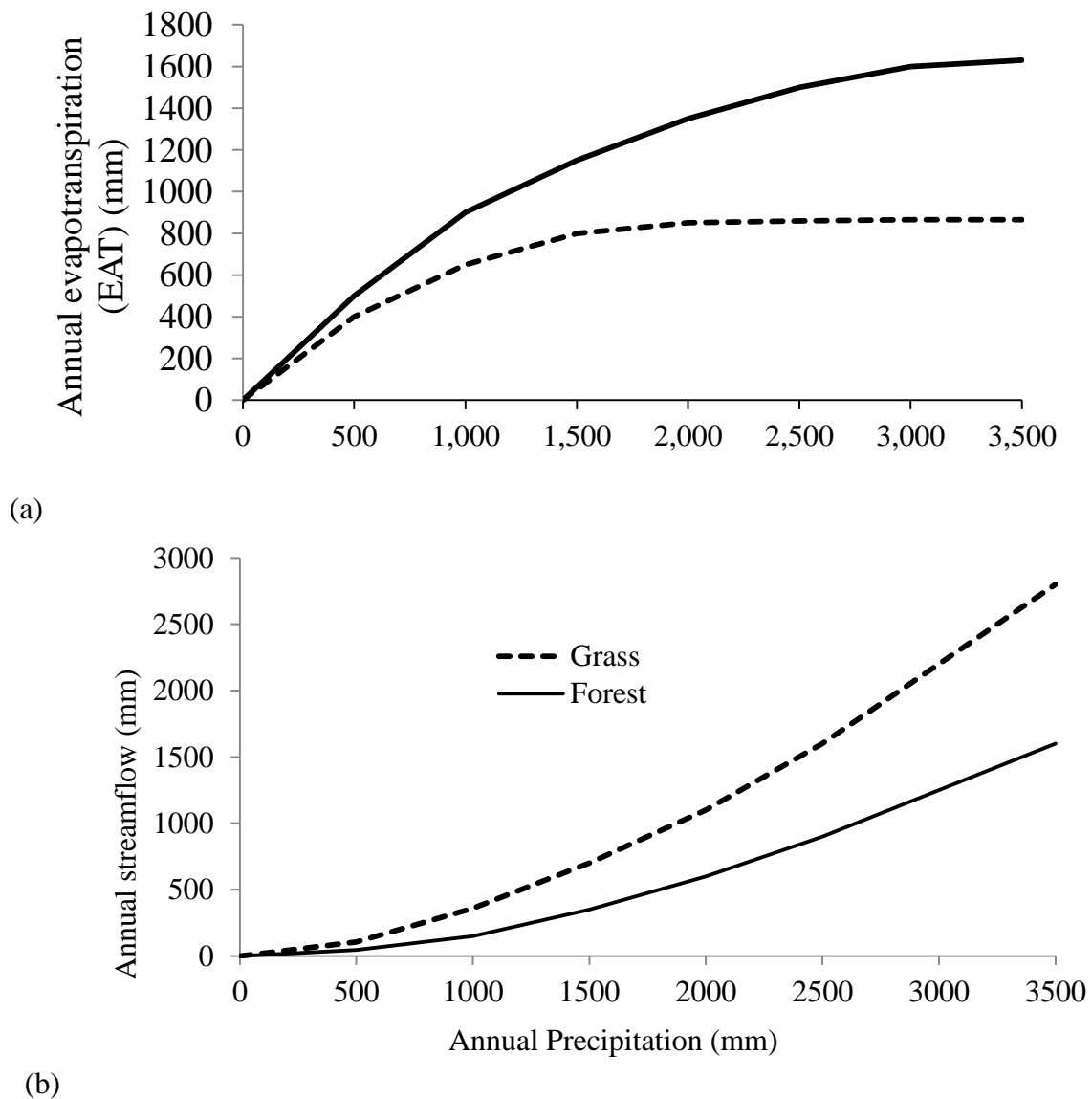


Figure 6.6. (a) Relationship between annual evapotranspiration and rainfall, by vegetation type and (b) Mean annual streamflow (calculated from the same data set used in figure 6a). A in Fig. 6a represents the difference in evapotranspiration by the grass and forest vegetation types (adapted from Gilmour, 2014 with some modifications).

Effects of combinations of plant traits have been found to influence the impacts of exotic plants on water resources relative to natural vegetation (Le Maitre et al., 2015; Le Maitre 2004; Calder, 2005). Strong impacts are related to plant size and physiology, and will occur where there are significant trait differences (e.g. evergreen versus deciduous, deep versus shallow roots). The more contrasting these are, the greater the difference is likely to be. In South Africa the most marked contrasts were fairly shallow-rooted, winter-deciduous grasslands when replaced by tall, deep-rooted, evergreen trees (Everson et al., 2011). In contrast, invasions by tamarisks in North America have had little impact because they are similar to poplars in their growth form, rooting depth and leaf seasonality (Doody et al., 2011; Hultine and Bush, 2011).

Eucalyptus water use has been reported to be comparable to, or less than that of some annual crops, especially those irrigated ones. Yan (2009) report a eucalypt water-use rate that was not more than that of sugar cane crop. Most crops such as rice, citrus, alfalfa, sugar cane, coffee and cotton are particularly water demanding. Furthermore, comparing water use rates of annual crops and trees may need to refer to the unit area since comparing water use per plant may be inappropriate. While a daily water use rate may seem to be smaller for some crops, a collective amount used per hectare will be very huge due to the number of plants per unit area. Cotton for example, is reported to use 11.4-16.9 litres (0.9-1.4 mm) of water per plant per day (Silvertooth, 1999). Considering a eucalypt stand with a full stocking of 1,680 stems ha^{-1} (although it is usually less especially in mature stands) and a water use rate of 3 mm d^{-1} per tree, total daily water use ha^{-1} will be 5,040 mm. A cotton field with an average plant density of 50,000 ha^{-1} and a daily water use rate per plant of 0.9-1.4 mm will be 40,000-70,000 mm d^{-1} ha^{-1} ; which is about 8-14 times that of the trees. Even if the annual crop was occupying the field for 30% of the year (cotton is not since it is perennial), the annual water use would still be large compared to that of the eucalypts.

Furthermore, when efficiency of water use is considered, eucalypts are more efficient users of water than crops when total biomass produced is expressed as a function of water used (Davidson, 1993). The author gives the following information (see Table 6.1) taken from results of crop water use in Australia, which clearly shows that eucalypts are generally more efficient water users than a number of commonly grown crops based on total biomass. For example, 2,500 to 5,000 litres of water are needed to produce 1 kg of rice to counteract losses from evapotranspiration, percolation and seepage (Bouman, 2009). Across a *Eucalyptus* productivity gradient in Brazil, Stape et al. (2004) found that although more productive stands used more water at the stand level than less productive stands, they also had higher water and nutrient use efficiencies and could produce the same quantity of wood with about half the land area and water than required for less productive stands. Differences in the amount of water available does not affect the water use efficiency and results from a study by Myers et al. (1996) and from other studies as well, suggest that improved resource availability does not negatively impact water use efficiency but increased productivity of these plantations is associated with higher water use resulting from increased individual tree dimensions. To this effect, water use efficiency was not influenced by irrigation and was similar to the rain fed treatment (White et al., 2014; Myers et al., 1996).

Plant	Water use per total biomass (litres kg ⁻¹) (mm kg ⁻¹)
Cotton/Coffee/Bananas	3200 (264)
Sunflower	2400 (198)
Field Pea	2000 (165)
Paddy Rice	2000 (165)
Horse Bean	1714 (141)
Cow Pea	1667 (138)
Soy Beans	1430 (118)
Potato	1000 (83)
Sorghum	1000 (83)
Eucalypts	785 (65)
Finger Millet	592 (49)

Table 6.1. Water use by plants through evapotranspiration measured in total biomass and expressed in litres/kg (adapted from Davidson, 1993).

***Eucalyptus* and soil fertility**

Soil nutrient availability is known to be poor under eucalypt plantations. This is not only attributed to the high nutrient uptake capacity of eucalypts (O'Connell and Grove, 1999; Laclau et al., 2005) but also a result of the large amounts of non-recyclable nutrients exported at the end of each rotation in harvested biomass (Laclau et al., 2010; Gonçalves et al., 2008). The amount of nutrient exported will depend on the frequency of harvesting and thus, short rotation cycles are expected to affect soils more (Poore & Fries 1988; Du Toit 2003) compared to the longer rotations. Nandi et al. (1991) and Bernahard-Reversat (1993) report a decline in soil properties under eucalypt plantations including soil pH, organic carbon and exchangeable cations.

Continued export of nutrients cause other negative effects on the soils such as increased acidity levels under eucalypt forest flow which may affect site productivity negatively. Nsabimana et al. (2008) reported high acidity under eucalypt stands in Ruhande Arboretum than under other forest tree species within the study region. This is attributed to the degree of neutralisation of the soil exchange complex (Rhoades and Binkley, 1996). Biological soil acidification under forest ecosystems has been previously reported in the tropics: in Argentina (Jobbagy and Jackson, 2003), in India (Misra et al., 2003) and in Brazil (Lilienfein et al., 2000).

Comparing a natural forest and five common plantation species including a eucalypt and a pine, Michelsen et al. (1996) revealed a higher nutrient content under natural forest and attribute this to the loss of organic matter during conversion of natural forest to plantations, increased leaching in young plantations, and low nutrient demand by natural forest trees as compared with fast growing exotics. Forrester et al. (2006) and El-khawas and Shehata (2005)

reported the accumulation of phytotoxins in soils under the plantations resulting into soil degradation and loss of productivity. Chanie et al. (2013) reported that eucalypt plantations exhausted sites which were productive before being planted with eucalypts. Gindaba (2006) proposed to discourage farmers from planting eucalypts due to the fear of long term site degradation in Ethiopian highlands.

On the other hand, eucalypt plantations have been reported to improve soil characteristics especially in degraded sites in which case abandoned lands were brought back to agriculture. An example includes in Ethiopia, where after five successive annual crop cycles, farmlands reclaimed from eucalypt stands increased total biomass and grain yield of finger millet by 31.8 and 25.4% respectively; and that of barley by 32.8 and 37% respectively (Tadele and Teketay, 2014). The authors observe that the soil conditions did not deteriorate in farmlands reclaimed from eucalypt plantations. Positive outcomes of eucalypt growing on site quality were also reported by other studies in the area. Lemma et al. (2006) reported a total soil organic carbon increase to nearly pre-deforestation levels in *Eucalyptus grandis* plantations during 20 years of establishment, after 20 years of cultivation and 35 years of pasture. Hailu (2002) also reported improvement in total soil N following land use changes from cropland to eucalypt plantations in the highlands of Ethiopia. Alem et al. (2010) found improved soil nutrients and total carbon in *E. grandis* plantation compared with adjacent sub montane rainforest in southwestern Ethiopia. It was observed in Cameroon by (Tchienkoua and Zech 2004) that planting croplands with eucalypts can improve soil productivity through translocation of nutrients from deeper horizons to the soil surface. Guo et al. (2006) reported import of up to 24% total N uptake to the soil surface via litter fall in short rotation eucalypt forests. Gindaba (2003) recorded lower soil N and P concentrations 10 m away from eucalypt edge of the canopy as compared to beneath the canopy where litter fall accumulated. Very little influence of *Eucalyptus* plantations on the chemistry of surface waters were reported in the savanna of Guinea (Laclau et al., 2007) and in Brazil (Lima and Zakia, 2006).

Realising constraints of site degradation following plantations, attempts to manage soil nutrient levels under eucalypt plantations has traditionally been to apply fertilisers. The approach has limitations due to their being expensive (Bouillet et al., 2013) as well as causing negative environmental effects in terms of pollution (Epron et al., 2013). Alternative practice was suggested as to mix eucalypts with other plantation tree species. The latter are thought to have three broad advantages over monocultures: greater productivity (Montagnini et al., 1995; Kelty, 2006), environmental services due to increased diversity (e.g. soil and water protection) (Erskine et al., 2006; Kanowski et al., 2005) and risk aversion (resistance to pests and pathogens, climate change, species failure, and market fluctuations) (Bosu et al., 2006; Nichols et al., 2001). Although managing mixed species is more difficult compared to managing simple monocultures, advantages the former gives may necessitate opting for mixed stands. Nitrogen fixing species such as acacias have been tried and shown to improve biomass, soil nutrient levels and soil properties (Bouillet et al., 2013; Epron et al., 2013).

Again, small size woodlots as traditionally grown in the study region offer a chance to alternate stands of different tree species. Mixed stands are already managed in some locations

which may also contribute to the improved biomass production per unit area as well as in managing soil nutrients. Small landholders have been observed to be the main innovators in establishing and demonstrating mixed species plantations in Australia (Nichols et al., 2006). An issue difficult to solve in the study region is the recycling of plant remains of the harvests because any plant material available including leaf litter is used as firewood due to a great deficit in the source of domestic energy for cooking (Ndayambaje and Mohren, 2011).

***Eucalyptus* allelopathy and effects on biodiversity**

Interaction between plants in agro-ecosystems is a complex aspect and among strategies for competition is allelopathy. Allelopathy is a result of chemical release by one plant species and may affect companion plants, usually negatively (Abugre et al., 2011). Generally, allelochemicals are released into the environment through leaching, root exudates, residual decomposition and volatilisation and occurs both in natural and cropping systems (Fang et al., 2009; Zhang and Fu, 2009). Different chemicals affect different plant species, meaning that allelochemicals are species specific.

Many studies have shown that *Eucalyptus* species cause losses of biodiversity of the understorey vegetation in plantations (He et al., 2014; Wang et al., 2010; Gareca et al., 2007). This has been a crucial issue for long term sustainability of native ecosystems and allelopathy is considered a factor for the losses (Zhang and Fu, 2009; Ahmed et al., 2008). Understorey plants are very important in ensuring the overall species diversity in plantations where in most cases indigenous, multispecies vegetation are replaced by a single, in most cases, exotic species. In such plantations, many species are restricted to the understorey layer and others must pass through it during their seedling stage (Ramovs and Roberts, 2003).

In experiments with rice seedlings and their reaction to leachates from the litter of *Eucalyptus* hybrids Bernhard-Reversat (1999) noticed a sharp decrease of root length and an increase of root weight/root length ratio with a decrease in the number of lateral roots of the seedlings. However, the effect may substantially vary with the quality of litter as well as with the quality of the soil. In the above mentioned experiment leachates from litter older than five days did not exhibit any effect. Also the soil obtained under a *Eucalyptus* plantation used in a greenhouse experiment to grow beans did not show any difference in comparison to trials on other soil (Couto & Betters, 1995). These different results may be due to the fact that not the leachates of *Eucalyptus* litter but the soil of different quality was used in the experiments. As del Moral & Muller (1970) showed, the allelopathic effect is dependent on the soil. In this experiment, *Eucalyptus* failed to inhibit annual herbs on sand whereas conditions for allelopathic interference seemed to be optimal on soils that were poorly drained, poorly aerated, shallow, and with high colloidal content. These factors permitted toxin concentrations to reach physiologically significant proportions.

Most of allelopathic studies are usually done in laboratory bioassay experiments focussing on the effects of leaf chemicals, volatile compounds, foliage decomposition and root exudates on such factors as seed germination and early seedling growth (Fang et al., 2009; Lisanetwork and Michelsen, 1993; Molina et al., 1991). However, these techniques hardly represent natural

ecological processes (May and Ash, 1990). According to Chu et al. (2014) and Jose et al. (2006), field experiments testing the allelopathic effects are rare and those focusing on eucalypt allelopathy on broad-leaved species are even rarer (Fang et al., 2009). Furthermore, allelopathic studies have been towards crops and weeds and not on relevant species usually associated with eucalypts in situ (Ahmed et al., 2008). Conclusions drawn from such studies may not be relevant in explaining allelopathic effects of eucalypt species.

***Eucalyptus* and the replacement of conventional forests**

Many native forests have been and are still being replaced by large-scale plantations of fast growing exotic tree species. In the process, some native ecosystems and indigenous species become extinct and others endangered and the ecosystem services provided by native forests are diminishing or even disappearing (Sangha and Jalota, 2005; Foroughbakhch et al., 2001). Continuous planting of eucalypts in pure stands for example, may cause accumulation of phytotoxins in soil which may lead to soil degradation and loss of productivity (El-Khawas and Shehata, 2005).

Sometimes eucalypts are accused for replacing natural forests. There are two ways in which natural forests can be replaced by eucalypts. One is by establishing plantation forests and the other is when eucalypts invade natural forests as aliens. In the first case, eucalypts do not deserve the blame because they are planted by man. For invasion as aliens, some eucalypt species such as *E. camaldulensis* and *E. globulus* were reported to have very high risk of invasion (Gordon et al., 2012). Stanturf et al. (2013) also report that *E. camaldulensis* has been a serious problem in South Africa. However, the potential invasiveness of eucalypts studied in the southern USA was observed to be very low, owing to the species' poor dispersal, small seeds with limited viability that require bare soil to germinate, and light demanding seedlings that do not grow successfully under closed forest or understory canopies (Lorentz et al., 2015; Stanturf et al., 2013). Booth (2012) observed that generally eucalypts have proved to have limited invasive potential for a number of reasons, including their poor dispersal capabilities. Relatively limited seed dispersal, high mortality of seedlings and lack of compatible ectomycorrhizal fungi were stated as three reasons making eucalypts unsuccessful invasive species (Rejmanek and Richardson, 2011). Calvino-Cancela and Rubido-Bara (2013) also observed that native vegetation in the Mediterranean region were resistant to invasion by eucalypts.

Keeping plantations away from water courses, maintaining clear fire breaks and interspersing eucalypt plantations with other intensive land uses are the suggested possible means to reduce the chances of spreading (Stanturf et al., 2013).

***Eucalyptus* and crop yield losses**

Eucalyptus species are blamed to exacerbate food insecurity problems through competing with crops and significantly reducing crop yields (Mukund and Palanisami, 2001). Significant depressions of crop yields have been reported in many areas when *Eucalyptus* trees were grown near crops. Even at very young age of two years, eucalypt seedlings reduced soybean

yield by 27% (Franchini et al., 2014). In Ethiopia for example, tef and wheat yield depressions occurred over the first 12 m from the tree line, with declines of 20 to 73% for tef and 20 to 51% for wheat, equivalent to yield losses of 4.4 to 26% and 4.5 to 10% per hectare respectively (Kidanu et al., 2005). The crop yield reductions observed when grown near eucalypt is a result of tree-crop competition for soil moisture (Kidanu et al., 2005), for soil nutrients (Harrison et al., 2000) and for light (Nissen et al., 1999). In our study (Chapter 3), soil moisture and soil N were significantly reduced in the crop field zone from 0-10.5 m from the tree-crop interface.

However, the way crop yield losses are usually presented in literature is not particularly informative since percentage reductions usually appear to be very large, while the losses actually apply to only small proportions of the cropland in question. Examining yields combining crop field portions of the same field, both the one affected and that not affected, may make the overall loss insignificant. An example is the maize experiment in this thesis (Chapter 3) where maize in the 10.5 m crop field strip next to eucalypt woodlots lost 80% of the grain yield since it produced an average yield of 0.5 t ha^{-1} in comparison with the yield in the open areas of 2.5 t ha^{-1} . A seemingly very high (80%) grain yield loss is not that big when it is spread over the whole field used for crop production. To illustrate this, consider one hectare square crop field adjoining a eucalypt woodlot along one side, making a tree-crop interface of 100 m long equivalent to the side of the crop field. Applying the results of our study, a 10.5 m crop field strip only (equal to 1050 m^2 or 0.105 ha ; or 10.5% of a hectare) next to the woodlot will produce 0.5 t ha^{-1} equivalent to 20% of the yield in the open areas (2.5 t ha^{-1}), losing 80% of the grain. The remaining 8950 m^2 or 0.895 ha (89.5% of a hectare) will produce 2.5 t ha^{-1} . The overall yield of this 1 ha farmland will be:

$$[(0.5 \text{ t ha}^{-1} \times 0.105 \text{ ha}) + (2.5 \text{ t ha}^{-1} \times 0.895)] = 2.29 \text{ t ha}^{-1}.$$

This is only 0.21 t ha^{-1} less than the yield that would be produced from one hectare in the absence of eucalypt-maize competition. The negative effect on the small area (10.5%) distributed over the whole (1 ha) area reduces the overall effect from 80% to only 8.4%. This however, applied to specific situations in the study area, may not present an insignificant loss. Let us consider the following three of possible scenarios basing on property ownership and land size: 1. When both the woodlot and the crop field belong to the same farmer and vice versa, 2. When the crop field is very small compared to the woodlot, and vice versa. When both the woodlot and crop field belong to the same farmer, losing 8.4% of the maize grain yield per hectare to increase the owner's woodlot productivity would actually be a benefit since woodlots earn farmers more income than maize (as Chapter 4). Country level average number of households with woodlots were reported to be about 42%, with the number increasing with altitude such that in the low altitude zone 31% of households had woodlots, mid altitude 43% and the high altitude 51% (Ndayambaje et al., 2014). *Eucalyptus* was the most dominant tree species occupying 90% of all the woodlots in each of the three altitude zones (Ndayambaje et al., 2014). This gives an impression that about 44 (90% of 49) to 62 (90% of 69) of all households may own both eucalypt woodlots and the crop fields on their own farms. The situation becomes an issue on the remaining 38-66% of the households which

do not own woodlots, in which case however small the grain yield loss may be, it is an absolute loss; and at the expense of a neighbour's woodlot (benefit).

For the small crop field scenario, the Rwandan household lands are generally very small, with an average of 0.7 ha (Ali et al., 2015). It is reported that 36% of the households own land of 0.11 ha on average (Warnest et al., 2011) and this may be divided into 4-5 small plots often in multiple locations (REMA, 2009). By taking the same setting of *Eucalyptus* woodlot and maize crop in example above, with half the crop field area of 0.5 ha (100 m × 50 m) for example, and the 100 m side adjoining a woodlot; average maize grain yield will be reduced to 1.04 t ha⁻¹. This yield is 10.5% less than half the yield obtained in 1 ha. In other words, by halving the crop field without altering the length of the tree-crop interface, the decline in grain yield is more than halved; implying that, the smaller the crop field, the greater the grain yield loss. Farmers owning 0.11 ha with a rectangular shape of 50 m × 22 m, with the longest (50 m) side being at tree-crop interface will produce yield (Y):

$$Y = 0.5 \times [(50 \times 11.5)/10,000] + (2.5 \times ((50 \times 10.5)/10,000)) = 0.16 \text{ t.}$$

If there was no tree-crop competition, the same 0.11 ha field would produce: $0.11 \times 2.5 = 0.275 \text{ t}$, which means a grain yield loss of about 42%. The last scenario where the woodlot becomes small, the edge effect generally will decrease as a result of the reduced length of the tree-crop interface. However, the benefit or loss pattern will remain the same as for scenario one based on the ownership of the two components of woodlot and crop field.

Basing on these estimates, the grain yield and corresponding losses can be extrapolated to country level by applying the situation to the available arable land area. It was shown in chapter 1 that the total arable land area in Rwanda is 1,369,576 ha. It was further estimated that the total eucalypt area in the country is 282,747 ha, equivalent to about 21% of the total arable land. With the average maize grain yield loss of 0.21 t ha⁻¹ recorded in Chapter 3, assuming an average of eucalypt woodlot-maize interface of 100 m ha⁻¹, the country level maize grain yield loss would be $282,747 \times 0.21$, equivalent to 59,376.9 t. The same observation applies, that, with the loss being experienced by woodlot owners would actually not be a loss; but the loss is felt by the 58% households who do not own woodlots.

To analyse the integration of eucalypt woodlot and maize on one farm, (Chapter 4) we considered the following. A household owning 1 ha to which both a *Eucalyptus* woodlot and maize are grown with the woodlot located at one corner of the hectare such that the woodlot adjoins the crop field along two of its sides and the other two sides are on the external perimeter and assumed not to affect the crops. We assume also that the woodlot sizes vary in proportions of the 1 ha, say from 15 × 15 m (225 m²) to 50 × 50 m (2,500 m²). This would lead to the area of maize field affected by the edge ranging from 0.042-0.105 ha (Fig. 6.7). The crop field area affected will increase linearly with the increasing tree-crop interface length and can be computed from equation 1 as:

$$A = 2L \times 10.5 \text{ (or } 21L)$$

where: A = the crop field prone to the edge effects (m^2) and L = the side length of a square woodlot adjoining the crop field (m). This 10.5 m is the length of the crop field affected by the edge perpendicular to the tree-crop interface. It is multiplied by 2 because two sides of the same woodlot are at the interface with maize.

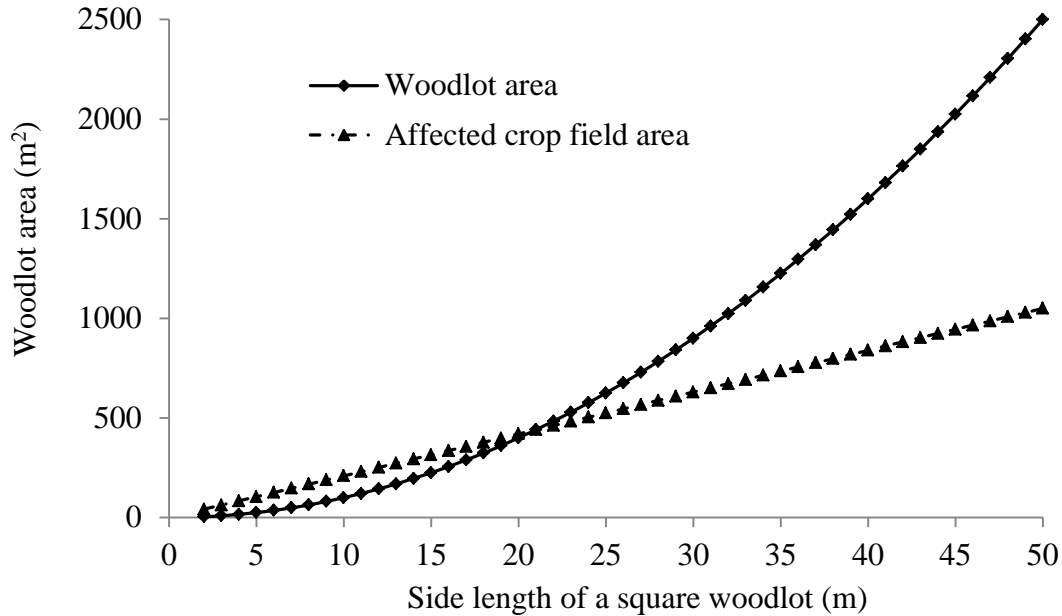


Figure 6.7. Area of the maize field experiencing woodlot (edge) effects in relation to woodlot size in a eucalypt woodlot-maize cropping system in southern Rwanda.

As earlier observed, many Rwandan farmers have small land holdings and the edge effects are more felt the smaller the crop field. The orientation of the crop field is also very important in smaller fields than in the bigger ones. This means that, if a rectangular crop field had its longer side adjoining the woodlot would be more affected than when its shorter side adjoined the woodlot. Consider a crop field of 0.4 ha laid such that 80 m adjoin a woodlot. The other side of 50 m will be affected by the edge up to 10.5 m from the tree-crop interface, making a total area affected to be $10.5 \text{ m} \times 80 \text{ m}$ ($= 924 \text{ m}^2$) or 21.3% of the 0.4 ha total crop area. If, on the other hand, the 50 m side adjoined the woodlot, the crop area affected by the edge would be $10.5 \text{ m} \times 50 \text{ m}$ ($= 525 \text{ m}^2$) or 13.1% of the 0.4 ha total crop area unaffected.

Woodlot trees are also affected by the edge as result of intraspecific competition, i.e., competition between trees. Trees in the eight outer tree rows making the woodlot edge were observed to be 17.8% bigger in DBH and to produce 34.5% more aboveground biomass than interior trees (Chapter 4). The woodlot area affected by the edge is derived as shown below (equation 2). Woodlots of $16 \times 16 \text{ m}$ will be completely affected by the edge and from those of $17 \times 17 \text{ m}$ and above, their proportion prone to edge effect will increase linearly with the size of the woodlot (Fig. 6.8).

$$\begin{aligned}
 A &= (L-2a)(L-2a) \\
 &= L^2 - 4La + 4a^2
 \end{aligned}$$

where A = the woodlot area prone to the edge effect (m^2), L = side length of a square woodlot (m), and a = the length of the portion of the woodlot affected by the edge, found to be 8 m (Chapter 4). To apply this formula, L must be $>2a$.

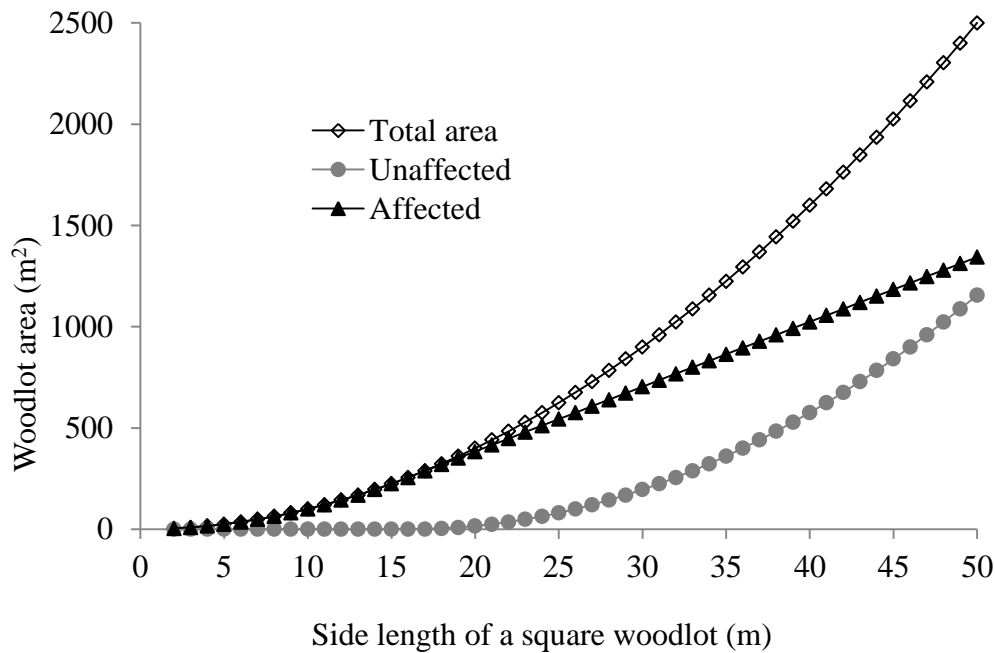


Figure 6.8. Total woodlot area in relation to its proportions that are prone- and not prone to the edge effects in a eucalypt woodlot-maize cropping system in southern Rwanda.

The estimated extra biomass obtainable from eucalypt woodlots would be computed as follows, using average aboveground biomass obtained in one hectare (reported in Chapter 4):

$$W \times A \times I \quad 3$$

where W = total above ground tree biomass (t ha^{-1}), A = country level eucalypt woodlot area (ha) and i = % increment in standing tree biomass due to edge, found to be 34.5% (Chapter 4). The average standing above ground tree biomass was found to be 204.5 t ha^{-1} (estimated from edge effect free woodlot area) and the woodlot area 282,747 ha and the increment due to the edge effect was 34.5%. This gives country level total biomass of 77,782,946 t containing extra total wood biomass (34.5%) equal to 19,951,759 t obtainable from eucalypt woodlots covering 21% of the arable land.

Other benefits of eucalypts

The value of eucalypt woodlots are sometimes overlooked probably due to ignorance. Examples cited here due to their important applicability to the eastern African region include prevention of run-off and soil loss and tree capacity to reduce CO_2 emissions and to enhance carbon sinks (Murthy et al., 2013). Soil erosion was reported to be a major concern for agricultural development in Rwanda (Bizoza et al., 2012). Annual rates of soil losses at

country level were estimated at 1.4 million tonnes, and 40% of Rwanda's land is classified by FAO as having a high risk of erosion (GoR, 2009). On-station soil loss reported by the national agricultural research institute (ISAR, now Rwanda Agriculture Board) as cited by Olson (1994) range from 35-246 t h⁻¹ per annum with most stations measuring over 100 t ha⁻¹ per year. On five of the seven research stations where erosion was measured, erosion would remove the fertile topsoil within 30 years if no anti-erosion techniques were used (König, 1994). Trees however have proved to be efficient tools to curb these problems.

Mulching by eucalypt leaves and branches is an important practice in Nyamagabe area, southern Rwanda where the mulch is primarily used to reduce soil loss through erosion by reducing kinetic energy of rain drops and nutrient enrichment through decomposition (Nzeyimana et al., 2013). A study in a valley region of western Himalaya showed a decrease in run-off and soil loss in contour cultivated maize by 27% and 45% compared to cultivated fallow (Narain et al., 1998). They further reported that *Eucalyptus* was more effective in controlling soil erosion than *Leucaena* trees both in alley cropping and in sole tree plots. In the study area, König (1992) had earlier reported a remarkable reduction in run-off and soil loss by using *Calliandra calothyrsus* and *Leucaena leucocephala* on a slope of 28%. Hedges and manure were observed to reduce runoff and erosion efficiently on the densely populated hill slopes of Rwanda (Roose and Ndayizigiye, 1997; Roose and Barthes, 2001).

If unchecked soil, erosion may cause more trouble as far as water supply is concerned. Siltation of rivers and other water bodies downstream may lower water quantities and quality significantly. Care must be put to the afforestation programmes regarding stand size and placement on the landscape to keep eucalypts far from riparian zones and near other water bodies, lakes and marshlands while curbing soil erosion and the associated soil losses.

Carbon sequestration as a feature of trees is very important in environmental conservation. Trees fix carbon in their biomass and at the same time enhance sequestration of atmospheric CO₂ in the soil. Soil organic matter provides a more lasting solution than just sequestering CO₂ in standing biomass (Gupta and Sharma, 2015). Creation and strengthening of carbon sinks in the soil were identified by the international panel on climate change (IPCC) as a clear option for enhancing the removal of CO₂ from the atmosphere and has recognises soil organic carbon pool as one of the five major carbon pools for land use, land use change in forestry (LULUCF) (paragraph 21, the annex to draft decision 16/CMP.1) (Anon, 2005). While soil contains an important pool of active carbon that plays a major role in the global carbon cycle (Melilo et al., 1995; Prentice et al., 2001), carbon in the soil is enhanced by trees. The build-up of each ton of soil organic matter removes 3.667 tons of CO₂ from the atmosphere (Bowen and Rovira, 1999).

Approximately 50% of tree biomass is carbon (Unwin and Kriedemann, 2000). Fast growing trees will therefore be more efficient fixers of carbon. *Eucalyptus* plantation was observed to fix more soil carbon compared to several tree species plantation including *Acacia catechu*, *Tectona grandis*, *Populus deltoids*, *Ailanthus excels*, *Haloptelia integrifolia*, *Pongamia pinnata* and a mixed plantation (Gupta and Sharma, 2015).

The effectiveness of agroforestry systems to store carbon will be affected by the environmental and socio-economic factors and in humid tropics, agroforestry systems have been reported to have the potential to sequester over 70 Mg ha⁻¹ in the top 20 cm of the soil (Mutuo et al., 2005; Newaj and Dhyani, 2008). Other factors include the species, geography, the structure and function of different components within the systems put into practice (Schroeder, 1993; Albrecht and Kandji, 2003). Agroforestry systems significantly accumulate C in living biomass and in the soil, demonstrating their potential to offer environmental service of C sequestration, and can contribute to reducing CO₂ emissions by avoiding burning of forest-based fuelwood and conserving soil (Murthy et al., 2013). *Eucalyptus* woodlots in the study area are likely to contribute significantly to C sequestration and related advantages due to their fast growth rates and greater numbers of trees per unit area compared to tree-crop mixtures in traditional agroforestry systems.

Trade-offs

Eucalyptus woodlots and crop production make the main components of plant production in Rwandan agriculture landscape. Due to the field setting, trees and crops compete, making what looks like monoculture tree stands and crop fields actually one system with the two being its components. Crop reduction resulting from tree-crop competition was seen to be insignificant, with an overall grain yield reduction of 8.4% (Chapter 3). Economic analysis showed that combined production of eucalypt woodlot and maize make a viable alternative compared to sole maize (Chapter 4). Positive economic benefits are known to occur when eucalypts are incorporated in the farming systems. This was reported in eastern Africa, in Ethiopia (Kebebew and Ayele, 2010; Kidanu et al., 2005; Jagger and Pender, 2000) and in Kenya (Peralta and Swinton, 2009). Kebebew and Ayele report a household income increase of up to 50% compared to the major food crop and further observe that by allocating only 12% of available land to eucalypt increased returns on the same land unit by 90%. They report a reduction in the returns from the land of 127 and 34% for teff and barley respectively when eucalypt contribution was not considered.

The findings reported in Chapter 4 are supported by observations in other places that tree component in agroforestry systems involving eucalypts adequately compensate for crop yield reductions and even generate additional income (Kidanu et al., 2005). In our case, not only tree-crop combinations but also even the additional wood increment obtained in the woodlots due to edge effects fully compensated for corresponding losses in maize revenue experienced due to woodlot tree competition. In addition to generating much income, *Eucalyptus* is less sensitive to changes in wages and has yields more than the other crops and can be highly profitable for smallholders (van Eijck et al., 2012). The income generated was significantly affected by the increase in woodlot area since the revenue from tree commodities was significantly greater than from the maize.

Water use rate of eucalypts observed in this study showed a small rate which is lower than the annual rainfall and the water use by important crops in the region (Chapter 5). Based on these, in addition to the dire need for firewood and wood for other uses by the farmers in the region, the integration rather than segregation of crop production and eucalypt woodlot is suggested a

suitable choice for farmers in the east African highland region which has reasonably high rainfall. This integration however would be more appropriate under two conditions: i) at the landscape level where eucalypts will play a significant role in soil erosion control, and ii) with farmers who have bigger land sizes where tree-crop competition is likely to be negligible. Negative effects on the environment are not evident which supports the conclusion.

Plantation planning and management

Eucalyptus plantations are now established on more than 20 M hectares globally (Carle et al., 2002; FAO, 2013). These plantations provide wood, environmental services and a source of renewable energy (Bauhus et al., 2010). In the eastern African region, the trend in eucalypt planting is also expected to continue and to expand with increasing population growth and technology. Rwandan government indicates its intension to expand eucalypt planting and states that approximately 11% of the total land area is potentially suitable for improved management of eucalypt woodlots and plantations especially in the northern and southern Provinces (MINITERE, 2014). This is necessary for both the supply of wood for firewood and charcoal supply, as well as other wood products and also for soil erosion control. As long as the planting is well planned, it would be appropriate to continue eucalypt planting. Although the water use rates observed in the current study are reasonably low and suggest no negative effects on the environment, some cautions may still be needed when planting eucalypts. Size and location of eucalypt plantations seem to be important aspects to be considered in afforestation programmes in the region.

Some features of Rwandan eucalypt plantings suggest they were well planned. The main Rwandan catchments from which major rivers originate, the Nyungwe and the Volcano forests, are both protected as national parks. Even the Gishwati and Mukura forests which had been seriously degraded in the past 2-3 decades are now being restored and a decision has been made to re-gazette and manage them as national parks to ensure their conservation. This status excludes any reforestation programmes using exotic tree species in these areas. Tree planting, if any, will be outside the national parks, restricted to the buffer zones. A similar protective effect was observed and suggestion made in Kerala, India, where eucalypts grown in Kanthalloor and the surrounding areas through which River Chengalar flowed from a natural spring uphill. Eucalypt water consumption did not seem to reduce stream flow (Kallarackal, 2010).

The small nature of eucalypt woodlots on Rwandan landscape is therefore appropriate. Plantation effects on the environment were reported to be directly related to plantation size with larger plantations causing more negative effects in Minas Gerais, Brazil (Ceccon, 2005). He concludes that eucalypt agroforestry appeared to be a promising alternative that could integrate timber and food production to meeting food demands while addressing wood production needs and, thus an important element in the plantation programs oriented towards small farm operations. In addition to plantation sizes, location is also an important factor as far as water use of trees is concerned.

Stands located near rivers are likely to affect stream flows more than those located far away, owing to the large size of individual trees and more abundant leaves with greater leaf areas (Dvorak et al., 2012). Negligible impacts are expected in catchments in which plantations are established on sites which are hydro-geologically isolated from streams (O'Loughlin and Nambiar, 2001). The severity of problems associated with water extraction are greater where plantations are large in size and cover most of the catchment area or in places with seasonally low rainfall. In this way, well planned agroforestry practice may provide an option of locating trees on sites where they can provide net hydrological benefits, such as controlling run off rates where this is a problem (Keenan et al. 2004).

Mixing *Eucalyptus* woodlots with native species woodlots or plantations could also be explored in terms of environmental conservation. Smethurst et al. (2015) observed that native forest reserves intercepted ground water moving laterally between eucalypt plantations and the stream. They observed that measured and simulated runoff were similarly small (5 and 3% respectively) but the simulated rate increased to 38% when native forest was removed. Appropriately planned agroforestry programmes would keep eucalypt stands far away from water sources where they can provide net environmental benefits, such as run off and soil erosion control (Keenan et al., 2004). Ferraz et al. (2013) also observed that a proportion of native forest plays an important role in reducing and regulating water use at landscape level and suggest a system of mosaic management to stabilise water flow across plantation landscapes.

Mixed (native and exotic species) plantations were suggested in order to solve nutrient mining by eucalypts and consequent soil degradation problems (Le Maitre et al., 2013). Resource use efficiency may increase with resource use (Binkley et al., 2004) and mixed species plantations may also use resources more efficiently through facilitation mechanisms (Kelty, 2006). An example is planting N₂-fixing tree species with a non-N₂-fixing tree species, which may enhance N soil availability and increase the growth of the non-N₂-fixing tree (Kaye et al., 2000; Richards et al., 2010). Several studies have shown that N-fixing species may have a positive effect on the overall productivity of mixed forest plantations (e.g., Richards et al., 2010; Forrester et al., 2010 Binkley et al., 2003). This kind of mixed species was also suggested as a means to enhance biodiversity conservation (Brokerhoff et al., 2013) among others.

Thinking across the east African region, so far, plantation water use in the region may not be expected to cause problems since they cover only a very small proportion of total land area. As seen earlier in Figure 5, the area of plantation forests (1.5 million hectares) in the eastern African states in 2011 was estimated to be only 0.3% of the total land area (507,839,000 ha) (Chamshama, 2011). With proper planning therefore, eucalypts are far from causing water use problems in the region. The region enjoys enough rainfall in most of the areas and it has been observed that in areas where annual rainfall exceeds annual potential evaporation, tree water use is limited by energy and not water deficit (Roberts et al., 2014). The east African region has some common issues including the shortage of wood for fuel and for other uses, high rates of deforestation as a result of high and increasing population. The demand for wood and other

non-timber forest products has exceed forest capacity to supply these resulting into the depletion of many natural forests (Burnett, 1985; Keerthisinghe, 1999). All these factors necessitate the increase of afforestation and reforestation rates in addition to agroforestry tree plantings so as to meet current and future wood demands which are expected to increase as well as protecting the environment.

Conclusions and recommendations

Detailed investigations done in this thesis on the role of *Eucalyptus* and their interaction with crops lead to the conclusion that the species are very useful agroforestry species for smallholder farmers despite the bad name the species have acquired. It is not possible to get an alternative species, being indigenous or exotic, that can equally provide benefits as does *Eucalyptus*. Owing to their competitive nature however, their integration in the spatially zoned agroforestry systems is recommended under two conditions: i) at landscape level where, in addition to the benefits farmers obtain from the woodlots, the latter will serve to conserve soils against erosion and landslides incumbent in the area; and ii) at farm level for farmers who own large land area, where crop yield losses due to tree-crop competition are likely to be negligible and fully compensated for by benefits from woodlot trees. Segregation or no woodlot planting at all will clearly be the option for farmers who own very small landholdings where crop yield losses are likely to be very significant. While no evidence of negative effects on catchment hydrology were observed in this study, some precautions which would equally be applied whenever planting any exotic tree species for plantation purposes still need to be taken and the following recommendations are suggested: 1. Plant and manage *Eucalyptus* woodlots outside riparian zones to avoid any possible impact on water sources 2. Maintain the current practice of not planting eucalypts in catchment areas 3. Maintain small woodlots to avoid possible impacts reported in literature caused by large plantation monocultures 4. Mix eucalypts with other species, preferably those capable of fixing nitrogen to improve soils and woodlot productivity as well as to allow for possible improvements of biodiversity 5. Make better species selection among eucalypts to allow for undergrowth development in the woodlots to improve soil erosion control especially in very steep sites and improve biodiversity. Finally, a holistic, integrated planning and management is needed, considering the environmental conservation by different stakeholders to achieve a more compatible agricultural production with the protection of ecosystems without impairing the livelihoods of the farmers. Currently, land use specific fields plan independent of the others which may be part of the cause of problems faced today.

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SUMMARY

Eucalyptus is the most planted *Genus* in Rwanda, with 69 species growing in Ruhande Arboretum and 10 species widely grown in plantations and woodlots countrywide. The species were introduced in the country in early 1900s by the colonial administration, to supply wood for fuel and for construction. Population pressure resulted in natural forest being greatly reduced in area, and so people have opted to grow trees on the farm to supply firewood and charcoal, the primary source of domestic energy for 90% of the population. Farmers obtain other wood and non-wood forest products from eucalypts and raise some income to sustain their families. Their intangible benefits such as soil erosion and landslide control are also very important for environmental conservation. *Eucalyptus* species are very competitive with crops and so farmers grow them in woodlots instead of intermixing them with annual crops in the fields. Farmers' eucalypt woodlots reflect the settlement pattern and thus are found scattered over most of the landscape, in a mosaic together with cropped fields. In such a setting the interaction between woodlot trees and annual crops is unavoidable.

In this thesis I characterise this type of agroforestry system in which eucalypt woodlots interact with annual crops, by investigating tree–crop interaction and the resultant effects on crop yield. The effects of edge on both the crop and woodlot trees were quantified and I evaluated the financial benefits of growing sole crops (annual or woodlot) versus their combination, and the trade-offs. In addition, water use of the woodlot trees was assessed, to evaluate the effects on environment at catchment level. An allometric relationship was used to estimate aboveground tree biomass using measurable tree parameters (DBH and height). The investigation was structured around four specific objectives: i) To develop a general equation for the estimation of aboveground tree biomass, applicable to both planted and coppice stands of *Eucalyptus saligna*; ii) To quantify the extent to which the competition for environmental resources manifests itself in maize grain yield and how this varies with distance from *Eucalyptus* woodlots along a line perpendicular to the tree–crop interface; iii) To determine the area of maize field/eucalypt woodlot affected by the edge, and the extent of maize grain loss or gain in aboveground biomass, and the trade-offs between the two components in an agroforestry system; and iv) To estimate the water use rate of eucalypt woodlots and compare this to the local precipitation to learn if eucalypts transpire more water than the rain supplies.

In chapter 2, allometric equations are developed for the estimation of aboveground tree biomass for both coppice and planted stands of *Eucalyptus saligna*. This is one of the most dominant tree species in southern Rwanda, where the study was done. DBH and total tree height were used as input variables to estimate total tree biomass and tree component (stem, branch, leaf and bark) biomass, and the best equations were selected. The equations can be used by different stakeholders to predict tree biomass for different purposes. In addition, they were used in other chapters of this thesis, as described below.

Chapter 3 is dedicated to investigating the interaction between eucalypt woodlots and maize (a staple crop in Rwanda) as an example field crop. An experiment was set up in three sites

where maize was grown adjacent to selected mature woodlots of *E. saligna*. Soil was characterised by measuring soil particle size distribution, soil nutrients (N, P and K) and soil pH at the beginning of the experiment. Environmental resources of soil water and solar radiation were assessed three times per season: the beginning, middle and end. This assessment was done at nine distances (1, 3, 4.5, 7.5, 10.5, 14, 18, 22.5 and 27.5 m) along a line perpendicular to the tree–crop interface. The experiment was repeated for three consecutive seasons: January–May 2007, September–December 2007 and January–May 2008. Maize growth and grain yield were assessed and related to the spatial variation of environmental resources in relation to distance from the tree–crop interface. In our experimental plot of $30 \times 12.75 \text{ m}^2$, a strip of $10.5 \text{ (width)} \times 12.5 \text{ (length)} \text{ m}^2$ crop field next to the woodlot showed a loss in grain yield of 80%. By increasing the crop field to $100 \times 100 \text{ m}^2$ (1ha) adjoining a woodlot along one side, the seemingly large grain yield loss will reduce to only 8.4% since the width of the crop field strip prone to the edge effects will remain constant.

Chapter 4 investigates the effect of the edge on both the maize crop and eucalypt woodlots. One hectare of land owned by a farmer was assigned to growing sole maize and sole woodlot and the benefits were compared with those obtained when both maize and woodlot were combined on the same 1 ha field at varying area proportions occupied by each. Maize and woodlot yields were quantified and an economic evaluation was made using a net present value (NPV) approach. A trade-off analysis showed that combining maize and firewood or pole or timber alternatives was more profitable than only growing maize. A further analysis was done to compare monetary loss from the maize loss due to competition with the monetary gain on the tree side obtained as a result of extra tree growth due to the edge effect. Trees in the outer 8 m zone of the woodlot were observed to grow 17.8% bigger in DBH and 34.5% more aboveground biomass. Combined maize–wood products were more profitable than the products from sole systems, and revenue from extra wood gains due to edge effects exceeded corresponding revenue losses in maize yield. However, land shortage, which is common in most households, may drive farmers to prioritise crop production to ensure food security. Again, since small cropped fields suffered more from tree–crop competition than bigger fields, farmers with large land holdings are likely to adopt eucalypt woodlot–maize agroforestry systems. Similarly, small woodlots are more prone to the edge effects, and the edge effect was felt throughout square woodlots with sides 16 m long.

Chapter 5 investigates the water use of eucalypt woodlots, using the tree DBH–leaf area index allometric relationship. This study was done in two catchments: Butare and Busoro in southern Rwanda. A monthly water balance was established. The woodlots had small coppice shoots ranging from 2–36 cm DBH and potential tree transpiration recorded was 3 mm d^{-1} . The annual potential tree transpiration was 10% below the local annual precipitation. Dry months had a water deficit, which could be covered by reductions in seasonal leaf area, stomatal closure and changes in soil water storage. A sensitivity analysis showed that 50% leaf area reduction corresponded to a 32.8 mm decline in potential tree transpiration. The deficit may not impact tree growth negatively, since in dry seasons tree growth. The moderate eucalypt water use rate observed in this study may be a function of tree size (which was

small) and low tree stocking, since such woodlots had less potential transpiration. The observed eucalypt water use rate is smaller than the range reported for eucalypts in Africa and was also smaller than that of key annual crops in the study area. Managing woodlots as short rotations and increasing initial tree spacing may contribute to resolving issues related to catchment hydrology associated with eucalypt plantations.

Detailed investigations done in this thesis on the role of *Eucalyptus* and their interaction with crops lead to the conclusion that the species are very useful agroforestry species for smallholder farmers, despite the bad name the Genus have acquired. The integration of *Eucalyptus* species into agroforestry systems is recommended, under two conditions: i) at landscape level where, in addition to financially benefitting farmers, the woodlots may serve to conserve soils against soil erosion and landslides; and ii) at farm level for farmers who own large areas of land, where total crop yield losses due to tree–crop competition are likely to be negligible. Segregation or no woodlots at all will clearly be the options for farmers who own very small areas of land; in their case, crop yield losses are likely to be appreciable and therefore likely to affect household food security.

Although no evidence of negative effects on the environment was observed in this study, some precautions (which would be applied whenever growing any exotic tree species) still need to be taken and the following recommendations are suggested: 1. Plant and manage *Eucalyptus* woodlots outside riparian zones to avoid possible impact on water sources 2. Maintain the current practice of not planting *Eucalyptus* trees in catchment areas in which the major rivers arise 3. Maintain small woodlots to avoid possible adverse impacts of large plantation monocultures reported in the literature 4. Mix *Eucalyptus* woodlots with other species (preferably N-fixing), to improve soils and woodlot productivity as well as to allow for possible improvement of biodiversity 5. Make a better species selection from among *Eucalyptus* species to allow for undergrowth development in the woodlots, which may also improve soil erosion control, especially on very steep sites, as well as biodiversity.

Finally, all-encompassing rather than field-specific, integrated environmental conservation planning and management by different stakeholders is needed, to develop agricultural production that is more compatible with the protection of ecosystems and will not impair farmers' livelihoods. Different natural resource management fields plan independently and lack of harmonised, multidisciplinary actions may be part of the cause of problems faced today.

PE&RC Training and Education Statement

With the training and education activities listed below the PhD candidate has complied with the requirements set by the C.T. de Wit Graduate School for Production Ecology and Resource Conservation (PE&RC) which comprises of a minimum total of 32 ECTS (= 22 weeks of activities)



Review of literature (6 ECTS)

- Tree-crop interactions and the use of eucalypts in agroforestry systems (2006)

Writing of project proposal (4.5 ECTS)

- The use of *Eucalyptus* in agroforestry systems of Southern Rwanda: to integrate or segregate? (2006)

Post-graduate courses (3.0 ECTS)

- Analysing farming systems and rural livelihoods in a changing world: vulnerability and adaptation competencies for integrated agricultural research; PE&RC/GERS-UZ (2008)

Laboratory training and working visits (3.0 ECTS)

- Soil and plant analysis; National University of Rwanda (NUR) (2007)

Deficiency, refresh, brush-up courses (7.5 ECTS)

- Research methods; ENP Group (2006)
- Basic statistics; PE&RC (2006)

Competence strengthening / skills courses (1.9 ECTS)

- PhD Competence assessment; WGS (2006)
- Information literacy and introduction to Endnote; WGS (2008)
- Mobilizing your scientific networking; WGS (2015)

PE&RC Annual meetings, seminars and the PE&RC weekend (1.5 ECTS)

- PE&RC Weekend first year (2006)
- PE&RC Day (2006)
- PE&RC Weekend final year (2015)

Discussion groups / local seminars / other scientific meetings (7.3 ECTS)

- Group discussions at FEM and FNP groups (2006, 2009, 2015)
- PhD Students discussion groups at the NUR (2007-2009)
- Annual national scientific meetings at the NUR (2007-2015)

International symposia, workshops and conferences (8 ECTS)

- International workshop on agroforestry and soil management; oral and three poster presentations; National University of Rwanda and Wageningen University, Butare, Rwanda (2008)
- First global workshop on improving forestry education in Africa; oral presentation; International Partnership for Forestry Education (IPFE) (2007)
- Workshop on biodiversity conservation in the Albertine Rift Region; oral presentation; Conservation Educators of the Albertine Rift (RNCEAR), Makerere University, Kampala, Uganda (2009)

Lecturing / supervision of Practical's / tutorials (13.2 ECTS)

- Multiple-purpose tree management (2006-2010)
- Agroforestry systems (2006-2010)

Supervision of MSc students

- Maize-*Eucalyptus* competition for soil water in Southern Rwanda
- Water consumption of *Eucalyptus* trees in Rwanda – an estimation based on leaf area
- Nutrient uptake by maize and *Eucalyptus* trees in an agroforestry system in Southern Rwanda
- Terrace stabilization: niche for producing *Alnus nepalensis* biomass as green manure and stakes for bean production in Gicumbi district, Rwanda

CURRICULUM VITAE

Canisius Patrick Mugunga was born in Kagera, Tanzania on 28 April 1964. After completing secondary education, he joined a forestry training institute in Olotonyi, Arusha, Tanzania, where he obtained a Diploma in forestry in 1990. He joined Sokoine University of Agriculture in Morogoro, Tanzania where he obtained his first degree in forestry in 1994. He was employed as an assistant research officer by “Institut des Sciences Agronomiques du Rwanda, (ISAR)” from 1995 to 2005. He undertook Master studies at Stellenbosch University in Cape Town, South Africa, where he obtained a master’s degree in forestry. During this period, he held different positions and responsibilities. He was the manager of a national tree seed centre and the National Designated Authority for the control of Forestry Reproductive Material moving in International Trade in the Organisation for Economic Cooperation and Development (OECD) Scheme from 1995-2004. From 2000-2004 he headed a Sida Salec sponsored National Plant Genetic Resources programme which worked in 10 eastern African countries. This post automatically rendered him two other designations namely: a National Focal Point for the International Plant Genetic Resources Institute (IPGRI) and country representative in the Eastern African Plant Genetic Resources Network (EAPGREN) and member of EAPGREN’s Regional Steering Committee. From February 2004 - March 2005, he was the head of Nyagatare research station in ISAR. From 2005 to date, he is a lecturer at the University of Rwanda in the department of Forestry and Nature Conservation. He has pursued a PhD in Agroforestry at Wageningen University from 2006 to 2016, a period during which he was also doing his teaching at the home university.

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