TROPICAL FIRE ECOLOGY

Climate Change, Land Use, and Ecosystem Dynamics





Tropical ecosystems constitute about half of the planet's fire-affected surface and are the location of the majority of all fires. *Tropical Fire Ecology* gives an extensive explanation of both the historic and current situations, describing the fire ecology of tropical ecosystems from around the globe. Regional chapters address a set of common subjects, including the causes of fire, typical fire behavior, and elements of the fire regime. They also study the impacts of human land use, landscape fragmentation and climate change on the fire environment, and the challenges of fire management in these ecosystems. The common set of topics provides consistency, enables comparisons and syntheses of fire effects in disparate tropical ecosystems, and facilitates a comprehensive understanding of fire's place in tropical ecology.

Tropical Fire Ecology uniquely provides a global synthesis and overview of fire in tropical ecosystems, and fills a large void in our current understanding of how fire affects terrestrial biota.

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Fires in Australia's tropical savannas: Interactions with biodiversity, global warming, and exotic biota

1. Malcolm Gill, Richard J. Williams, and John C.Z. Woinarski

5.1 ABSTRACT

The vast tropical savanna region of northern Australia is generally flat, vegetated by native species and sparsely populated by people. The climate is typified as monsoonal, ith contrasting wet and dry seasons. Temporal and spatial patterns of rainfall, cattle crazing, and fires help shape the vegetation of the region. Eucalyptus spp. dominate the tree cover while grasses are a feature of woodland understories. Grasses fuel fires which occur mostly in the dry season. Fire regimes (including between-fire interval, re intensity, and season of occurrence) affect the relationships between the woody and grassy components of the vegetation. Fires, at a given point in the landscape, can vecur with mean intervals as short as about two years but mean intervals vary widely across the region. Fire intensity varies seasonally: it can arbitrate between outcomes such as tree survival, shoot survival, or tree death. Introduced grasses, especially gamba grass (Andropogon gayanus), can alter fire regimes to the extent that the structure and composition of plant communities is strongly affected. Near major towns and cities, increased fire intensities due to gamba grass increases the fire threat to houses. Fire regimes, whether or not influenced by exotic species, affect land management for broad-scale pastoralism and biodiversity conservation, and influence carbon emissions and storage.

5.2 INTRODUCTION

This chapter describes the pattern and management of fire in the tropical savannas of northern Australia. This region has characteristics shared with strongly seasonal tropical areas elsewhere in the world, but has a distinct ecological context. Fire is a pervasive management issue in the region and the management of fire affects, or is affected by, land use, biodiversity conservation, the maintenance of indigenous

culture, the spread of exotic plants and animals, and global climate change. Across most of the region, fire regimes have changed over the 150 years or so since European settlement.

Reflecting the main arena of research activity, our focus in this chapter is on the central third of the tropical savanna area, locally known as the "Top End" of the Northern Territory (Figure 5.1). However, consideration of the region as a whole is possible for some topics and research in areas outside the Top End are not neglected.

5.2.1 The region

The island continent of Australia is generally flat. hot, and dry. but has great variety in climate, soils, vegetation, and topography. Its northern quarter—an area of 1.9 million km² (Russell-Smith et al., 2003b)—is characterized by a monsoonal (strongly seasonal wet—dry) climate and tropical savanna vegetation. The circumscription and naming of this area has varied: here, the definition based on bioregional boundaries defined in the interim biogeographic regions of Australia (Thackway and Cresswell 1995), and its broad subdivisions ("provinces") are used (Russell-Smith et al., 2003a) and shown in Figure 5.1. The whole area is referred to as northern Australia or the Australian tropical savanna, reflecting local usage.

This region spans parts of three Australian political jurisdictions (Figure 5.1 inset), with variable legislative, land use, and management contexts. Unlike most of the World's other tropical savanna regions, it is sparsely populated by people: excluding a handful of larger towns, the average population density is less than 0.1 person/km² (Woinarski et al., 2007). Australia's tropical savannas are managed by both indigenous people (who own 20% of it) and by relatively recent settlers of predominantly European origin. Cattle-based pastoralism is by far the most wide-spread land use, extending over about 70% of the savannas (Woinarski et al., 2007); it is more intensive (i.e., with much smaller properties) in the eastern and southeastern parts of the region, compared with the west and northwest. The iconic 2-million-hectare World Heritage Kakadu National Park is found in the savanna region along with a number of other national parks, totaling about 6% of the region's area (Woinarski et al., 2007). There are much smaller areas devoted to military training, horticulture, mining, and other industries (Russell-Smith et al., 2003a).

Climate is a defining feature of the region. but also shows ecologically important variation. Rainfall is strongly seasonal, with a distinct short and hot summer wet season and a long warm dry season (April–November in the Top End): seasonality is least marked in the southeast. Total rainfall varies from >2.000 mm yr⁻¹ in some near-coastal areas to less than 400 mm at the fringe of the vast arid zone of Australia's interior and central west (Gill *et al.*, 1990: Williams *et al.*, 1997a; Russell-Smith *et al.*, 2003a). There is considerable variability in the duration, intensity, and the timing of the onset of the wet season (Ridpath, 1983; Holland, 1986; Cook and Heerdegen, 2001).

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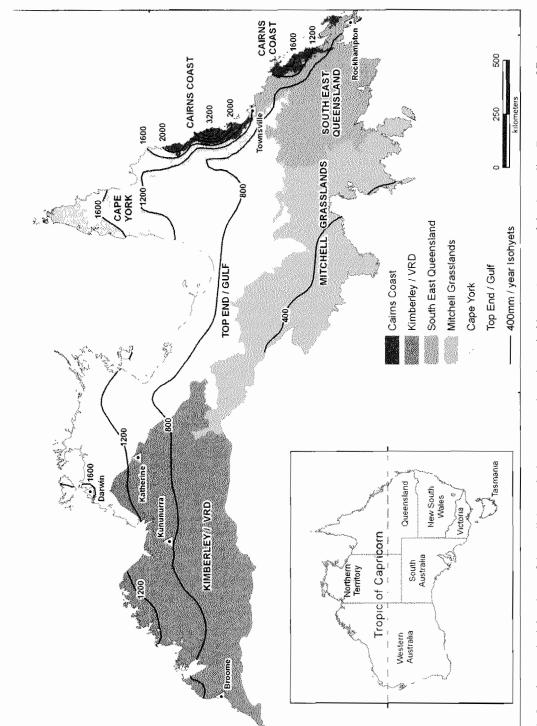
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Figure 5.1. Interim provincial regions of Australia's tropical savanna region—in capital letters (courtesy of the Australian Department of Environment and Water Resources, Canberra. ACT, Australia; after Thackway and Cresswell, 1995, see also Russell-Smith et al., 2003a), national context (inset), and rainfall isohyets in millimeters.

The period of change from the dry to the wet season is often characterized by highly localized storm activity, with lightning strikes being a major ignition source for fires. Particularly in the wet season, coastal areas are subjected to frequent destructive cyclones.

The passage of the dry season is marked by increasing depth of drought and the presence of the southeast Trade Winds; it is conveniently subdivided into "early" and "late". May and June are typical early dry-season months in the Top End (R.J. Williams *et al.*, 2003a) and Russell-Smith *et al.* (2003a) defined the early dry season there as the period up until the end of July. The rest of the dry season, August to November, is regarded as the late dry season (Hill *et al.*, 2006). Cook and Heerdegen (2001), with an eye to biological responses, defined a "rainy season" and a "wet season" based on differing probabilities of 10-day dry spells. Seasons could also be defined in terms of ecological indicators (Lewis, 1985). Crowley and Garnett (2000) defined "storm-burns" as those being undertaken soon after the first heavy rains of the wet season.

Native vegetation is extensive across the savanna region and is a highly significant asset for biodiversity conservation (Woinarski et al., 2007). Apart from some parts of Queensland (e.g. see Tassicker et al., 2006), there has been relatively little intensive clearing for agriculture; forestry is rare. As a consequence, the savannas are relatively intact structurally, and "relatively little transformed by modern development" but despite this "the savanna landscapes of Australia are in flux" (Whitehead et al., 2005) with increasing pressure for substantial modification and resource use. Naturalized exotic plants have established and spread in some areas, strongly affecting fire regimes and their outcomes (Douglas and Setterfield, 2005); feral cats, horses, pigs, and donkeys are widespread (Strahan, 1995).

The dominant, savanna, vegetation of northern Australia comprises extensive grasslands, grassy forests, and woodlands, with far smaller areas of other structural types (Fox et al., 2001; Williams et al., 2002; Russell-Smith et al., 2003b), notably including heathlands (mostly on skeletal sandy soils in rugged rocky areas) and closed forest (rainforest), typically occurring as small patches in relatively fire-protected areas or in sites with seasonally persistent water supply, especially along the coastlines and in the immediate hinterland (Russell-Smith et al., 2003a). While these more restricted vegetation types are not strictly tropical savannas, some consideration is given to them here as their ecology, conservation, and management is intimately associated with the savannas that typically surround them.

Forests and woodlands are typically 10 m to 20 m in height, occasionally reaching 30 m (Gillison and Walker. 1981). Tree height and density tend to decrease with decreasing rainfall: for example, along the aridity gradient south from Darwin (Williams et al., 1996). Eucalypts, trees of the genera Eucalyptus and Corymbia, family Myrtaceae, and especially the Eucalyptus subgenera Blakella and Eudesmia (Gill et al., 1985, p. 29) feature as dominants to a grassy understory. The presence of these eucalypt subgenera in the north contrasts with the predominance of the speciesrich subgenera Monocalyptus and Symphyomyrtus in the southern part of Australia (Gill et al., 1985, p. 29). Other common tree genera include the pantropical Erythrophleum and Terminalia, the native conifer Callitris, and Myrtaceae like

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Melaleuca. There is a wide range of other tree and shrub genera present including Acacia, Alphitonia. Brachychiton, Calytrix. Cochlospermum, Grevillea, Petalostigma, and Planchonia.

Grasses dominate the ground stratum of savannas. Common dominant or subdominant grass genera include Aristida, Astrebla, Bothriochloa, Chrysopogon, Dicanthium, Eriachne, Heteropogon, Imperata, Plectrachne, Schizachyrium, Sarga (formerly Sorghum). Sporobolus, Triodia. and Xerochloa (Tothill and Gillies, 1992). Kimberley/Victoria River province has widespread Triodia and Plectrachne (spinifex); the western part of the Top End Gulf province is dominated by annual Sarga; the eastern Top End Gulf Province has widespread Aristida—Bothriochloa and Dicanthium pastures; the Mitchell Grasslands province has widespread Astrebla (Mitchell Grass) (Tothill and Gillies, 1992). Schizachyrium and perennial Sarga pastures are common in Cape York while Heteropogon pastures are common in southeast Queensland (Weston, 1988).

While grass predominates as the understory throughout the savanna region, litter from the trees may be substantial in quantity in some places and times, the late dry season being associated with increased litter fall and litter fuel load (Gill *et al.*, 1996; Williams *et al.*, 1997b). In the wet season the grasses are green but with the approach of the long dry season, grasses start to cure (i.e., shoots die) in a rapid mottled wave from the south to the north and towards the northern coastlines (S. Berry, unpublished data). With the arrival of the early wet-season convective storms, and, later the monsoon proper, a wave of greening up spreads from the north to the south. The monthly raingreen (mostly grass) component of the vegetation follows monthly rainfall (Spessa *et al.*, 2005).

Thus the period in which fuels are predisposed to burn is longest in the south and shortest in the north, although greater than about 4 months in all regions. The length of the growing season may be seen as setting the grassy fuel load while the length of the dry season—the non-growing season (Walker 1981)—may be seen as setting the length of the potential "fire season". However, fires are also possible in drier parts of the wet season (Williams and Lane. 1999: Crowley and Garnett. 2000).

5.2.2 Fires in the region

In this section, the broad characteristics of fire in this region are described. In subsequent sections, more detailed attention is given to some of the factors that underlie or define these characteristics and we consider in more detail the management and consequences of particular fire regimes. Management considerations include the manner in which fires affect the interrelationships between woody plants and grasses, the interactions between global warming effects and atmospheric emissions, the conservation of biodiversity in pastoral systems and in reserves, and the emergent issue of the protection of human life and property, particularly in districts with a transition from rural to urban.

Like tropical savannas in monsoonal climates elsewhere, fires are a pervasive feature of northern Australia. Fires are frequent and extensive especially in the north of the region; individual fires can cover an area of over a million hectares (Luke and

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McArthur, 1978, p. 260); the region has been characterized as "perhaps the most extensive and flammable ecosystem in the world" (Liedloff and Cook, 2007). Fires affect all landscapes and land tenures—pastoral properties, conservation lands, indigenous freehold land, mining leases. and, increasingly, the urban–savanna interface (Dyer et al., 2001; Williams et al., 2002).

Without ignition there is no fire. In the early dry season, all fires are caused by humans while in the late dry season lightning can be a major source of ignition. Early dry-season fires include those deliberately lit with the aim of mitigating the effects (or reducing the likelihood) of intense late dry-season fires, some of which are caused by arson and some by lightning strikes; no systematic data are known regarding causes of fire starts across northern Australia, other than those for central Cape York Peninsula (Crowley and Garnett, 2000). Early dry-season fires typically go out at night but the chances of overnight burning (and hence greater fire extent) increase as the dry season progresses. There are a number of reasons for this including increasingly drier conditions in which fuel is more continuous, fewer dews, and increasing wind speeds (Gill et al.. 1996). The control of prescribed fires, then, becomes increasingly problematic as the dry season advances. Hence, particularly in the Top End province (Figure 5.1), early dry-season fires are typified by short-lived humaninitiated fires burning grassy fuels in relatively mild weather, whereas late dry-season fires are of generally higher intensity and capable of burning day and night in grass-litter fuels under drier and windier conditions.

5.3 FIRE BEHAVIOR, FIRE WEATHER

"Fire behavior" is represented by a range of attributes varying from the rate of spread of the fire burning with the wind to the rate of the setting of new fires downwind as a result of firebrands arising from the "mother" fire. Besides rates of spread, fire behavior variables include temperature (Weber et al., 1995), intensity (Byram, 1959), duration of burning (flaming and smoldering), and various dimensions of flames (height, length, depth), and flame angles. Fire behavior can also be seen to be the way in which fires interact with tree stems (Tunstall et al., 1976). Rates of spread of fires burning with the wind are the primary object of measurement and prediction. In conjunction with "fuel loading", rates of spread contribute to fire intensity (Byram, 1959) which, in turn, may be correlated with flame dimensions and immediate effects of fires like the scorch height and char height of leaves (Williams et al., 1998). Australian researchers have traditionally adopted an empirical approach to fire prediction (e.g., Cheney et al., 1993), an approach that is particularly advantageous where there are large tracts of similar fire-prone vegetation.

The only formal fire-modeling program conducted in Australia's tropical savanna region has been that of Cheney et al. (1993) in the Top End province. These authors examined rate of spread of fire burning with the wind in manipulated and control swards of cured (i.e., with dead shoot systems) *Themeda triandra* and *Eriachne burkittii*. They found that neither the grass species nor the fuel load above about 1 tha⁻¹ significantly influenced the rate of fire spread, but wind speed and

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moisture content were important. Their research formed the basis of a rate-of-spread predictive system that required added input variables for its practical application, namely curing (CSIRO, 1997a; Cheney *et al.*, 1998) and slope (CSIRO, 1997a). This northern Australian spread meter applies to open forest, woodland, and open grassland (CSIRO, 1997b).

Rates of spread of fires burning with the wind on level ground form the basis of fire weather forecasts and fire danger warnings. Australia's Bureau of Meteorology uses the modified McArthur Grassland Fire Danger Index Mark 4 (Purton, 1982), or GFDI_4, for such forecasts and warnings in northern Australia. McArthur's GFDI (McArthur, 1966), and Purton's (1982) modification of it, have a scale of 1 to 100; 100 represents the "worst possible" weather condition. The indices are grouped as ratings in a generally logarithmic fashion with "extreme" being the range from 50 to 100. The modified McArthur GFDI 4 has inputs of temperature, relative humidity, wind speed, grass curing, and fine-fuel load (Purton, 1982). Weather inputs are provided from widely distributed weather stations in the savanna region while estimates of curing and fuel loading (from "very sparse" to "heavy"-Purton. 1982) are provided by bushfire agencies. Across the Northern Territory and northern Queensland at least, indices in the "extreme" range occur (Bureau of Meteorology, pers. commun.), thereby predicting possible extreme rates of spread, but the frequency of "extreme" values and historically peak values have not been reported. Tapper et al. (1993) document the synoptic conditions for high fire danger in the Northern Territory. In coastal Darwin, and in its hinterland at Jabiru, the GFDI (McArthur, 1966) rises during the dry season and peaks in September (Gill et al., 1987, 1990).

The effects of individual fires often depend on fire intensity. Williams *et al.* (1998), measuring fire intensity in their plots at Kapalga in the Top End province, found a mean of 2,100 kW m⁻¹ for early dry-season fires and 7,700 kW m⁻¹ for late dry-season fires with a peak intensity of 18,000 kW m⁻¹. In this province, fuel loads up to 13 t ha⁻¹ have been measured in native ecosystems but 2 t ha⁻¹ to 5 t ha⁻¹ are more typical, perhaps due to frequent burning (Williams *et al.*, 1998). Bowman *et al.*'s (2007) study of grass biomass across the northern parts of the Northern Territory and Western Australia found sampled grassy fuel loads in savannas dominated by *E. tetrodonta* to vary from 0.1 t ha⁻¹ to 5.8 t ha⁻¹.

In Australian savannas, fuel loads are generally higher in the wetter north of the region and lower in the south in accord with trends in annual rainfall, but are also influenced by fuel type and fire history. Williams *et al.* (2002) report unpublished data of G. Cook for an "open-forest savanna" in the Top End in which fuel loads increased rapidly in the first 4 or 5 years after fire to reach a quasi-equilibrium value of between 8 t ha⁻¹ and 9 t ha⁻¹. Large local variations may be expected due to the variety of circumstances in which various fuels occur. Locally, distinguishing features may be lowland (e.g., floodplain or swamp) vs. upland (e.g., plateau sites, sometimes very rocky); grass species composition (including the productive exotic grasses, native annuals, and native perennials); and, early vs. late dry season, the fuels in the early dry season being mainly grassy and only partly cured if perennial, whereas those of the late dry season comprise cured grasses supplemented by litter from semi-deciduous eucalypts and other woody species.

Sec. 5.4]

Fire behavior models need a weather and landscape context for their use—which may be actual or virtual. In the latter case the fire prediction system of Cheney et al. (1998) has been incorporated into a fire effects savanna model called "FLAMES" (Liedloff and Cook, 2007). FLAMES allows for the modeling of fire effects using multiple fires with different properties in different seasons—in short, the fire regime.

5.4 FIRE REGIMES

Fires spread across landscapes forming burnt patches of many magnitudes (Gill et al., 2003). As these patches age, further fires overlay them to various extents thereby diminishing their area. Thus, a time sequence of patch size starting with the frequency distribution of the newest (less than 1-year-old) patches, is transformed into successive frequency distributions of ever older, smaller, patches. In a similar way, frequency distributions of areas with the same between-fire interval can be constructed, the mean area of the patches declining as the interval increases (Gill et al., 2003).

Mean "interval between fires" is the reciprocal of the average proportion of the landscape burnt per year. The interval between fires is an important component of the fire regime along with type of fire, season of fire, and fire intensity (Gill, 1975). Peat fires in *Melaleuca* swamps (Innis, 1994). then, may be contrasted with the more common above-ground fires. Fire intensity varies with diurnal, seasonal topographic and fuel load variations (R.J. Williams *et al.*, 2003a). Seasonal variation in fire occurrence in any one location can be extensive, as has been mentioned above, while variation in interval between fires is also large and is considered below.

Intervals between fires can be determined using a series of yearly maps of burnt areas, the maps often being derived from satellite imagery. Analysis of five years of NOAA-AVHRR satellite imagery across northern Australia suggested that there were two distinct groups of vegetation with respect to the mean proportion burnt per year (Spessa et al., 2005):

- (i) low open woodland, open forest, open woodland, woodland, and low woodland—with 23% to 34% burnt per year; and,
- (ii) open heathland and closed [or "rain"] forest—the remaining two vegetation types—with 8.1% and 8.5% burnt per year, respectively.

These proportions translate to mean intervals between fires of 4.4 to 2.9 years for the first group and 12.3 and 11.8 years for the second. A statistical model examining the proportion burnt in relation to "raingreen" and "evergreen" components of the vegetation suggested that the proportions burnt per year can be greater than 0.54 per year (a mean interval of approximately 2 years).

Using a similar 5-year set of NOAA-AVHRR data, Spessa *et al.* (2005) and Russell-Smith *et al.* (2003a) examined a wider area—the entire tropical savanna region (Figure 5.1)—and estimated the average percentage of burnt area according

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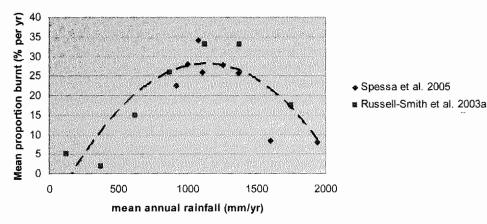
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Figure 5.2. Mean NOAA-AVHRR proportion burned per year as a function of interpolated mean annual rainfall (based on five years of data only): combined results of Spessa *et al.* (2005) and Russell-Smith *et al.* (2003a) for the "wet-dry tropics" and the more extensive "tropical savannas". respectively for an almost identical period. The baseline for Spessa *et al.*, (2005) was the mean annual rainfall for vegetation types (including closed forest—rainforest); that of Russell-Smith *et al.* (2003a) for rainfall classes, the top class (>1,500 mm) being taken as 1,750 mm yr⁻¹. The trend line is a quadratic function.

to rainfall category rather than mean rainfall for vegetation categories. The results are plotted in Figure 5.2 with a trendline for the combined data. The trendline suggests that the mean proportion burnt is greatest—and, therefore, the mean interval between fires is least—when mean annual rainfall is about 1,100 mm. These data are spatially averaged so cannot be expected to reflect local circumstances; they are also derived for areas of various size and for a short period, and they may provide only a coarse index of the actual extent of land cover being burned. They necessarily gloss over the effects of other variables like topography and land use. Across the vast area of northern Australia, there is substantial variability in the seasonal patterning and extent of fire, associated largely with differences in land use and intensity, topography, and climate.

At the regional scale of their study. Russell-Smith *et al.* (2003a) calculated that about 77% of the savanna region that burned did so in the late dry season. This proportion can be very variable. For example, while more than 50% of Pine Creek, Central Arnhem, and Daly Basin bioregions (all in the Top End province, Figure 5.1) was burnt by late dry-season fires in four years of a nine-year study period (1997–2005), this was the case for only one (North Kimberley) of the remaining 19 bioregions that comprise the tropical savanna region (G. Crowley, pers. commun.). For 12 of these bioregions, the mean rate was less than 25%, and for five bioregions the rate was less than or equal to 1% (G. Crowley, pers. commun.). There are various reasons for this variability, one of which is that there can be a tradeoff between the proportion of landscape burning in the early dry season with that in the late dry season (as in lowland Kakadu National Park during the 15-year study period of Gill *t al.*, 2000).

The seasonality of fires affects the scale of patterning of burnt areas. For sandstone heath vegetation, Price et al. (2003) found that individual early dry-season fires burnt, on average, 64% of the vegetation within their perimeter, whereas this proportion averaged 84% for fires in the late dry season. Similarly, R.J. Williams et al. (2003a) reported that the burned percentage of their transects in experiments in mesic savanna increased from zero to over 90% as fire intensities rose up to, then above, 2,000 kW m⁻¹.

Local studies of fire regimes have been conducted across the region using historical accounts (e.g., Stanton, 1992) and Landsat imagery (e.g., Gill et al., 2000; references in Russell-Smith et al., 2003a; Felderhof and Gillieson, 2006), for example. These studies are more appropriate to guiding fire management (Felderhof and Gillieson, 2006) than the region-wide studies but the latter provide an overview not possible from local studies. The foci of local studies have usually been on between-fire interval and season.

Fire regimes have changed in the last 150 years or so (e.g., Stanton, 1992; Russell-Smith et al., 2003a) in association with the general replacement of indigenous land management by pastoralism, changes in human dispersion, changed fuel conditions, and the spread of exotic invasive grasses. Global warming can be expected to produce further changes and attempts to reduce emissions of greenhouse gases such as those from fires (see below): thus, with frequent fires and a vast area, increasing attention is being paid to the greenhouse gas implications of management (Gill et al., 1990: Russell-Smith et al., 2007). Like other savannas of the world, those in northern Australia generate the majority of such emissions that result from biomass burning in the continent (Meyer, 2004; Russell-Smith et al., 2007).

A living legacy of changes in fire regimes can be found in landscapes in the form of increased tree and shrub density. The relationships between woody plants, grasses, and fires is important in relation to the cattle industry, biodiversity conservation, and global warming.

5.5 RELATIONSHIPS BETWEEN FIRE, WOODY PLANTS, AND GRASSES

As with tropical savannas throughout the world (Scholes and Archer, 1977), in northern Australia the tree-grass relationship is driven primarily by rainfall (Liedloff and Cook, 2007) and soil type; the general lack of trees on the Mitchell Grass (Astrebla) grasslands can be attributed to the clay soils there (Figure 5.1; Thackway and Cresswell, 1995). However, in this section, emphasis is placed on the dynamics of woody plants locally and the impact of fire regimes, a secondary driver of the grass-tree balance. Where woody plants become more abundant, the grass component of the community can be expected to diminish.

Tree survival in a fire is a function of fire intensity, the height of the tree, and its bud-protective mechanisms (Gill, 1995). In the Top End, the study of the effect of fires on trees has largely been directed towards the ubiquitous eucalypts (Williams *et al.*, 1999) and the sporadically distributed conifer *Callitris intratropica*. Seedlings and

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saplings of Callitris are readily killed by fire, and adults may be readily killed if the crowns are completely scorched (browned). If scorched trees of Callitris have no canopy-stored seed then local extinction can occur as this seed is the only source of regeneration; the species is an obligate seeder (Yates and Russell-Smith, 2003). Fires have led to the depletion of Callitris in various parts of northern Australia (e.g., Yates and Russell-Smith, 2003), probably because of changes in fire regimes associated with the arrival of Europeans (Bowman and Panton, 1993).

Eucalypt trees behave quite differently from Callitris in the face of fire. They often recover after crown-scorching fires and even after above-ground stem death. Williams et al. (1999) found stem death to increase quite markedly with increasing intensity and whole tree death to increase in the same manner but at a much reduced rate; smaller and larger tree stems were the most affected in their model. In Queensland, P.R. Williams et al. (2003) found that eucalypt saplings were suppressed by frequent fire (<4-year interval). In the simulations of Liedloff and Cook (2007), frequent fire, in the early or late dry season, affected the size distribution of stems by continually removing recruits and killing larger stems.

Woody plant thickening (typically an increase in the density of woody stems) has become an issue on pastoral properties across the savannas (Crowley et al., 2004, p. 36)—including parts of the Kimberley (Craig, 1997), the Top End (McGuffog and Starr, 1997; Dyer and Stafford Smith. 2003). central Queensland (Landsberg, 1997; Burrows et al., 2002), Gulf country (Grice, 2006) and Cape York Peninsula (Neldner et al., 1997; Crowley and Garnett, 1998). This may be detrimental to pastoral productivity and lead to decline of grassland ecosystems and the biota that depend upon them (Stanton, 1992; Neldner et al., 1997; Crowley and Garnett, 1998; Crowley, 2001; Crowley et al., 2004), but conversely may benefit bird species diversity following partial vegetation clearance (Tassicker et al., 2006) and can increase carbon storage in woody material, thereby benefiting users of a carbon-trading scheme (Ockwell and Lovett, 2005).

The complex dynamic of the tree-grass relationship is influenced differentially across the region by variations within grazing (Sharp and Whittaker, 2003) and fire regimes, by climatic variation and more globally by changes in greenhouse gas concentrations (Bond et al., 2003). Periods of above-average rainfall may trigger exceptional episodes of woody plant recruitment (Crowley, 1995, p. 13; Fensham and Fairfax, 2003), while extended periods of below-average rainfall may trigger exceptional rates of tree mortality across broad areas (Fensham and Holman, 1999). With very low fire frequency or fire exclusion, woody thickening can occur Bowman et al., 1988; Crowley and Garnett, 1998; Russell-Smith et al., 2003c).

Grass is a major fuel of the savanna but also the major food of cattle generally Bos indicus) on pastoral properties. The pastoralist is faced with a choice of using grass for cattle or grass for fire (Liedloff et al., 2001; Dyer and Stafford Smith, 2003) but the choice can be pre-empted by an unplanned fire. Without fire, there is the chance of an increasing density of native or exotic woody plants which then deplete pastures. With fire, pastures are temporarily removed but woody plants re suppressed; this is a simplistic characterization of the situation as fires have a ariety of advantages to the pastoralist as well as disadvantages (Leigh and Noble, 1981; Dyer and Stafford-Smith. 2003). The management key here is the choice of the appropriate fire regime for various parts of a pastoral property for different purposes using fires of different intensities, times of year, and intervals (Leigh and Noble, 1981; Liedloff *et al.*, 2001; Dyer and Stafford Smith, 2003; Crowley *et al.*, 2004, pp. 18-20).

The interactions between rainfall, livestock grazing, and fire regimes as they affect temporal aspects of woody plant–grass relationships is a theme that extends beyond savannas. Multiple stable states are possible according to the models of Janssen *et al.* (2004) who applied them to sheep country in New South Wales, Australia. These authors showed the importance of rainfall variability to their results. an outcome also emphasized by Liedloff and Cook (2007) who concluded, for tree density across a rainfall gradient: "While fire is an important factor in northern Australia, the model indicates its effects on tree populations at a sub-continental scale are secondary to the impact of rainfall variability."

Changing climates and vegetation, along with land use intensification, may cause shifts in carbon stocks and fluxes. Cattle grazing and its intensification may have increasing importance given the methane content of emissions from ruminants (Blaxter and Clapperton, 1965; Howden and Reyenga; 1999), while the creation of a market for carbon "has the potential to drive land use away from fire-assisted pastoralism toward sustainable forestry and environmental conservation" in northern Australia (Ockwell and Lovett, 2005). Thus, the management of greenhouse gases and biodiversity conservation in Australian savannas can be considered to be a cross-tenure issue of increasing importance.

5.6 FIRE, GLOBAL WARMING, AND GREENHOUSE GASES

5.6.1 Predicted changes in climate

Rising concentrations of greenhouse gases such as CO_2 . CO, CH_4 , N_2O , and NO_x in the atmosphere (greenhouse gas emissions) cause increased planetary warming and other atmospheric changes which, in turn, affect nearly all aspects of life on Earth (IPCC, 2007). Predictions of changes in climate as a result of human-mediated emissions in the decades ahead can be made by using global climate models assuming various emission scenarios and by extrapolating current trends.

The regional climate of northern Australia has changed since the beginning of the 20th century (http://www.bom.gov.au/cgi-bin/silo/reg/cli_chg/trendmaps.cgi), and further changes are predicted as a consequence of global climate change (Pearce et al.. 2007). Since about 1950, northwestern Australia has become wetter by about 5 mm per year, but there have been declines in rainfall over much of the drier inland savannas of the Northern Territory and Queensland. Mean annual temperature has been generally rising in the same period but not so in the Kimberley region of Western Australia. Pan evaporation has generally declined over much of the savanna biome since 1970 and this has considerable significance for fire behavior as it appears to be due to a reduction in wind speed (Roderick et al., 2007).

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With due caveats for imprecision and regional variation, predicted climate changes for northern Australia by 2030, as a consequence of greenhouse gas-induced global warming, are for an increase in temperature of about 1°C and an increase in the number of days on which temperatures exceed 35°C in major centers like Darwin, Broome, and Cairns (Pearce et al., 2007). Predicted changes in annual rainfall are either nil or show small decreases of about 1% to 3%. Other changes predicted by Pearce et al. (2007) are that evapotranspiration will increase and relative humidity will show little change or record a small decrease. Predictions with respect to the southeast Trade Winds, which predominate during the dry season (southern winter and spring), are either for no change or increases of 2% to 5%. There is a likelihood of an increase in incidence of severe cyclones (Pearce et al., 2007). Lightning ignition of fires may increase (Goldammer and Price, 1998).

Predictions made by extrapolating current climatic trends and those made by most global climate models are presently inconsistent for parts of northern Australia but the situation may be resolved for the northwest, at least for rainfall, by the inclusion of Asian aerosols in the models (Rotstayn et al., 2007).

Climate change alone may or may not cause major changes to the fire regime of the savannas but rising human populations, patterns of human ignitions of fires, land use, grazing intensification, and the spread of exotic plants such as gamba grass (Andropogon gayanus) (Rossiter et al., 2003), are likely to do so.

5.6.2 Management of emissions

Managing savannas for carbon raises a number of questions. How best can we manage grass—use it as fuel for fires, or for cattle fodder, or just let it decompose (Gill et al., 1990)? Would some forms of management allow grass to be replaced by woody vegetation which maximizes carbon storage and offsets emissions?

In the Top End (Figure 5.1) and some other parts of northern Australia, fires in the late dry season may be extensive and of high intensity, resulting in widespread consumption of fuel and the potential death of trees (Williams et al., 1999). In comparison, fires in the early dry season may be of relatively low intensity and consume less fuel. Greater tree death means less carbon sequestration and more carbon in the atmosphere; alternatively, tree thickening creates a gain in sequestration.

A short-term perspective on losses and gains through burning is apparent from the Kapalga experiments (Andersen et al., 2003b) in the Top End province where tree death from high-intensity fires increased the net emissions per fire by 15-44 times compared with the emissions from the burning of fine fuels by low-intensity fire Cook, 2003). Similarly, estimates using a number of methods, suggest that the videspread Eucalyptus miniata-E. tetrodonta savannas of higher rainfall areas of northern Australia are a carbon sink, even when burnt by early dry-season fires Williams et al., 2004; Beringer et al., 2007). Sequestration capacity can be decreased by repeated late dry-season fires and, under these circumstances, fires can be a net source of carbon to the atmosphere (Cook et al., 2005).

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Management of emissions 5.6.2

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If cattle consumed the grass, fire-caused emissions might be depleted due to fuel reduction and lower fire intensities. However, northern Australia's 14 million head of cattle (Petherick, 2005) emit copious quantities of methane (e.g., Blaxter and Clapperton, 1965; Howden and Reyenga, 1999; NGGIC, 2005), which has a greenhouse gas effect much more potent than carbon dioxide (Hansen et al., 2007). With less grass and more trees, and therefore more carbon sequestration (Williams et al., 2004), there may be fewer cattle, and fewer cattle—in general terms only—may mean less income for pastoralists.

Cattle properties have benefited from the introduction of productive exotic grasses but the escape and naturalization of such grasses has raised the potential for the loss of tree-stored carbon and increased emissions from burning, particularly with gamba grass. On pastoral properties gamba grass fuels may be reduced by grazing but on properties devoted to other land uses reduction by grazing does not occur. Gamba grass can reach 4 m in height and have a standing biomass of up to 17 t/ha—which is up to seven times that of the local grasses (Rossiter et al.. 2003). Unpublished data show fuel loads can reach 30 t/ha (N. Rossiter, pers. commun.). Measurements suggest that this exotic grass has the potential to raise the fire intensity to at least eight times that of fires in native grasses in the early dry season (Rossiter et al., 2003). Invasion of this grass initiates the grass—fire cycle in which fires are more intense, thereby decreasing tree canopy cover and increasing the grass to tree ratio until grass dominates the system (see Rossiter et al., 2003). Thus, reducing the extent of gamba grass will bring greenhouse gas abatement benefits.

Given the complex interactions between grazing, fire, and greenhouse accounting, especially with respect to methane, further investigation of interactions between pasture production, grazing, fires, and tree demographics is urgently needed for this region; a cue may be taken from Moore *et al.* (2001) who simulated burning and grazing effects on a semi-arid rangeland, but outside the region of interest; the system could be managed as a source (with livestock grazing) or a sink (especially without grazing or burning) for emissions.

5.7 BIODIVERSITY AND FIRE REGIMES

The effects of fire regimes on indigenous biodiversity may be seen in two contexts: in non-park landscapes generally (where fire regimes interact with cattle-grazing regimes); and, in the conservation—reserve context. In Section 5.7.1, examples of the interaction of plant and animal species and grazing and fire regimes are described In Section 5.7.2, some examples of responses of biodiversity to fire regimes in conservation reserves are considered.

5.7.1 Biodiversity, fire, and grazing regimes

In concert with the gradual transitions of dominant grass species across the longitudinal expanse of the northern Australian savannas, there is a muted change

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the nge in native animal species (Woinarski, 1992). The most notable directional trends are associated with the rainfall gradient (and tree density) from relatively high on the north coast to relatively low in the inland south (Woinarski et al., 1999a).

There are clear indications that there have been, and continue to be, substantial changes in the faunal biodiversity of northern Australia since European settlement of the area about 150 years ago. Unsurprisingly—given that fire and pastoral impacts are concentrated particularly in the grass layer—change appears most pronounced in granivorous (seed-eating) birds and in the assemblage of small native terrestrial mammals. Animal species that have particular dependencies on one or a few grass species for at least part of the year have been particularly prone to decline. Both fire regimes and grazing regimes affect the species composition of the grass layer and thereby affect the native animals that depend on them.

The grazing regime may be defined in parallel with the fire regime (Gill, 1975) and have as its component variables the type of herbivore (e.g., grazer/browser or particular species), the season of feeding, the intensity of feeding, and the intervals between feeding at points on the ground (Gill, 2008). To the extent that the grazing regime affects the fuel, and thus fire behavior, there is an interaction between the grazing regime and the fire regime. In a semi-arid fringe area of northern Australia, Suijdendorp (1981) studied the interactions of grazing by sheep and fire. He found that pastorally inspired winter burning had a detrimental effect on Triodia pastures out that summer burning with deferred grazing was best for them. There were interactions between the effects of fire and grazing regimes on vegetation composition that favored sheep on the one hand and native herbivores (in this case euros, *Macropus robustus*) on the other.

In eastern Queensland, Themeda-dominated pastures have been replaced in many areas by Heteropogon-dominated pastures (Shaw and Bisset, 1955; Shaw, 1957; Groves and Williams, 1981; Crowley and Garnett, 1998). Themeda is palatable, and is replaced by Heteropogon contortus under heavy grazing. However, even without livestock grazing, Walker et al. (1983) found that the proportions of Themeda and Heteropogon varied in response to the season of burning.

The strongest evidence of broad-scale changes in native fauna due, at least in ourt, to fire and grazing comes from the detailed reporting of bird observations over about 60 years by Barnard (1925, 1934) at a single pastoral property in central Queensland, with a further assessment of the bird fauna at that location about years later (Woinarski and Catterall, 2004). Together these observations constitute a chronicle spanning about 120 years since the introduction of pastoralism this area. Of 192 bird species with adequate information, 18 species disappeared from the property over this period, a further 68 species declined, 13 species ncreased, and 11 species were new colonists. The species that suffered local loss er decline included many that favored dense grassy understories, and some that wored fine-scale vegetation mosaics including both open and relatively dense inderstories.

Franklin (1999) provided evidence for the above pattern of change to be prevalent across northern Australia by means of an analysis of historical changes n the reporting rates of granivorous bird species. Over the historical span of his study (150 years), he found declines in 12 of the 49 species considered and increases in three species. One species was "probably extinct" while two species were considered to be "critically endangered". Declines were most marked in locations exposed to the longest history of, and most intensive, pastoralism, and in environments with little topographic relief (Franklin *et al.*, 2005).

A strong role for grazing and, interrelatedly, fire, in structuring faunal communities of the tropical savannas of northern Australia has been demonstrated in a range of experimental (e.g., Woinarski and Ash, 2002) and correlative studies (e.g., Kutt and Woinarski, 2007). In experimental studies with no pastoral factor, long-term imposition of different fire regimes produced substantial divergence in fauna (Woinarski, 1990; Woinarski et al., 1999b. 2004; Pardon et al., 2003; Corbett et al., 2003; Andersen et al., 2006), typically with the most notable polarization in faunal communities being between long-unburnt treatments and those burnt frequently by intense late dry-season fires.

Research on individual fauna species has revealed much of the important detail that underpins community-based responses to fire regimes, and demonstrates that there may be highly idiosyncratic—but ecologically pivotal—relationships between individual animal species and components of fire regimes. For example, the ecology and persistence of some animal species may be linked irrevocably with one grass species, or a small subset of such species. The endangered golden-shouldered parrot (Psephotus chrysopterygius) of Cape York Peninsula is one such case (Crowley et al... 2004). For part of the year (the early wet season), the principal seeds available in the landscape for this granivorous bird are those from cockatoo grass (Alloteropsis semialata). Grazing and uprooting by cattle and pigs, respectively, reduced the seed production of this grass species (Crowley and Garnett, 2001: Crowley et al., 2004). Where pastoral managers have lit fires after early storms at the start of the wet scason, seed production may be delayed by a month. Thus, where this "storm burning" is practiced patchily, there can be a landscape-wide expansion in the temporal availability of these seeds, and hence a direct benefit to the parrots (Crowley and Garnett, 2001; Crowley et al., 2004). Similar circumstances have been reported for other declining granivorous birds (Woinarski et al., 2005). While the seasonal timing of burning may influence the timing of seed production, the overall abundance of cockatoo grass may also be affected by fire interval as shown by its substantial increase following five years of fire exclusion at Kapalga (Top End province) (R.J. Williams et al., 2003b) and in the absence of grazing by livestock.

Other animal species may be keyed into the spatial patterning of fire events. The threatened partridge pigeon (Geophaps smithii), a ground-dwelling granivore, requires both unburnt areas (for providing shelter from predators for its rudimentary ground nests) and burnt areas (to enhance access to the fallen seeds on which it feeds (Fraser et al., 2003); and, its survival and reproductive output will be maximized where these are both available simultaneously within its territory—that is, where fireare of relatively low intensity and provide a fine-scale (I ha to 10 ha) mosaic of burnt and unburnt patches. A similar preference for heterogeneity has been demonstrated and modeled for the iconic frilled lizard (Chlamydosaurus kingii), where survival is typically low in unburnt areas (because dense grasses reduce foraging ability) and in

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period soon after intense late-season fires (when the fires themselves may kill tards) (Griffiths and Christian, 1996).

Biodiversity conservation in reserves

be savanna region of northern Australia contains many areas set aside for nature asservation. These areas have regional, national, and international significance as include World Heritage Areas like Kakadu National Park (Top End) and anululu National Park (Kimberley).

Two major experimental programs have been conducted in, or near, Kakadu Monal Park (Top End): the Kapalga experiment (Andersen et al., 2003b) and the Manmarlary experiment (Bowman et al., 1988; Russell-Smith et al., 2003c). Here we utch the findings of these studies, introduce fire-related concerns associated with roduced species, and note the effects of fire regimes.

Over the two to three decades of the Munmarlary experiments, metronomic mual burning maintained the dominance of annual Sarga—a major fuel species the Top End—while, in the absence of fire, annuals, including Sarga, declined; canwhile, the perennial grass Eriachne triseta, and shrub cover, increased (Russellwith et al., 2003c). The small size of plots in the Munmarlary experiment (1 ha) coluded their general utility for the study of faunal responses, but the unburnt plots prorted more bird species associated with a relatively dense woody understory ser (Woinarski, 1990).

The Kapalga experiment was larger in spatial scale, but conducted over a catively brief period (5 years) (Andersen et al., 2003b). In the ground stratum of e woodlands there, plant species composition was not affected by variation in the regime but was strongly correlated with wet-season rainfall and the cover of the anual Sarga (R.J. Williams et al., 2003b). Annual Sarga can be locally eliminated by es in the early wet season, soon after its germination (Williams and Lane, 1999), a at used by managers to reduce fuel loads in strategic areas, thus assisting with the ntrol of late dry-season fires.

In the Kapalga experiment, small mammals in general increased in the unburnt eatment compared with both early dry-season and late dry-season fire treatments relatively few species of frogs (1 of 11), lizards (5 of 16), or birds (5 of 25) were feeted by experimental treatment (Corbett et al., 2003). Those animal species that ere affected by fire treatment did so in a variety of ways such as by population aduction or increase under late dry-season treatment, or reduction or increase in aburnt relative to fire treatments (Corbett et al., 2003). Such variation is to be spected and highlights the importance of: (i) conserving the species rather than aximizing populations; and, (ii) varying fire regimes within limits appropriate to anagement aims (Gill, 2008). Some species, such as the northern brown bandicoot (woodon macrourus), a small fossorial marsupial, decreased under all fire treatments, solibly because it is reliant on a mosaic of burnt and unburnt patches that is finer called than that produced in the 15 km² to 20 km² landscapes of the fire treatments in rs experiment (Pardon et al., 2003).

2005).

At Boggy Plains, a wetland in Kakadu National Park, the native grass *Hymenachne acutigluma* proliferated when populations of feral Asian water buffalo (*Bubalus bubalis*) were removed (Robinson and Whitehead, 2003). In a program to restore pre-buffalo assemblages of plant and animal species to the wetland, low-intensity fires were ignited twice within a short period in the late dry season (CSIRO, 2005; Davidson, 2005). This caused a marked change in plant species composition in the wetland, from the *Hymenachne*-dominated one to one with a variety of native plant species including red lily (*Nelumbo* sp.) and other water lilies, wild rice

(Oryza meridionalis), and water chestnut (Eleocharis dulcis) (CSIRO, 2005; Davidson.

A number of vigorous grasses have been introduced into northern Australia to increase the productivity of pastures for cattle; besides gamba grass (mentioned above), these include para grass (*Urochloa mutica*), and mission grasses (*Pennisetum* spp.). Para grass has spread to wetlands near Kakadu National Park on the Mary River (Ferdinands *et al.*, 2005). This species can completely exclude all species of native vascular plants, including the native wild rice and native birds. Ferdinands *et al.* (2005) point out that wild rice is a critical resource for the magpie goose (*Anseranas semipalmata*), the most characteristic bird species of floodplain areas, because it produces 'large and abundant carbohydrate and protein-rich seeds' which are a preferred food source for their newly hatched young.

Individual species form communities which vary in structure according to the stature and density of the constituent plants. Fire regimes affect both the structure and the composition of savanna communities as noted above but, embedded within these communities, are closed forests (rainforests) with apparently different responses. These embedded communities (often in particular and limited topographic or other positions that provide some shelter from fire) have substantial conservation value, especially in national parks. In parts of the savanna that now have frequent intense fire, these communities, such as monsoon rainforests and heathlands, may be shrinking (Russell-Smith and Bowman, 1992). Conversely, there is recent evidence of widespread expansion of at least some rainforest types (a spatial counterpart to the vegetation thickening described above). While there are substrate limitations to expansion, "low fire disturbance" and favorable moisture conditions (Russell-Smith et al., 2004a, b) and/or changes in greenhouse gases (Banfai and Bowman, 2006) may be implicated. However, the impacts of fire regimes on the rainforest boundary are complex (Williams, 2000) but dramatically altered when invasion by highly flammable Lantana camara occurs as it has in north Queensland (Fensham et al., 1994).

Scientists working in savanna conservation reserves highlight the need for adaptive management (Andersen et al., 2003a: Russell-Smith et al., 2003c). Russell-Smith et al. (2003c) suggest further that adaptive management involving collaborative partnerships between managers and scientists is desirable; they question the value of experimental studies using deterministic fire regimes because fire regimes "vary idiosyncratically over multiple time and spatial scales." Such variation for between-fire intervals is reflected in the randomness of fire occurrence at a point in the landscape as in Kakadu National Park (Gill et al., 2000). Incorporating randomness as a formal treatment in experimental programs constitutes a major challenge.

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PROTECTION OF HUMAN LIFE AND PROPERTY

Until very recently, fires in northern Australia have been regarded as posing a very low risk to human life and infrastructure; hence, the management of fire has been restricted largely to pastoralists who want to maintain good habitat for their animals, to managers of conservation lands who attempt to use fire to maintain an optimal mix of habitats for biodiversity, and to indigenous people who use fire according to various cultural imperatives. The spread of vigorous exotic pasture grasses has meant that fire is becoming a problem for the society of northern Australians more generally, even the suburban or semi-suburban householder.

Attention in this section is focused on the area around the village of Batchelor, about 70 km south-southeast of Darwin, in the Top End province. The area consists of rocky hills and sometimes extensive flats all covered with shrubby or grassy woodland. Relatively small holdings (0.5 ha) are a feature of the landscape, and there are several villages in the area, but holdings may be as large as 6,000 ha. Mango orchards are sometimes established where there are deeper soils: Asian buffalo or cattle are raised on some properties while other properties are undeveloped as far as farming or horticulture is concerned: the latter properties retain the appearance of natural woodland although houses and other buildings have been erected in clearings.

In 1980 gamba grass was introduced into the area as a pasture grass. In the late dry season, peaking in this area in August-September, fire weather can reach "extreme" on the McArthur Mark 4 grassland meter: Trade Winds may reach speeds of 35 km/hr, with gusts 50% higher (Northern Territory Branch, Commonwealth Bureau of Meteorology, pers. commun.). Air temperature in the shade may be 37°C or more at 3 PM while relative humidity can drop to 20% (Northern Territory Branch of the Bureau of Meteorology, pers. commun.). In the evening a strong sea breeze may travel through the area from the northwest—as opposed to the daytime Trade Winds from the southeast thereby complicating fire control: the duration of the sea breeze extends as the dry season progresses.

With extreme weather and high fuel loads due to gamba grass (see above), fire suppression is sometimes impossible and losses of buildings including a house have occurred. Flame heights of up to 15 meters have been observed with spot fires occurring up to 1 km ahead of the fire front, and a maximum rate of spread of a hire noted as being 5 km in 25 minutes, in 2004. These fires are dangerous for firefighters and residents alike. Filters on vehicles are readily clogged in thick smoke and the vehicles then disabled, rendering their occupants in peril.

Fuel managers aim to establish buffer areas for fire spread by burning the gamba grass early in the dry season. Fire suppression capacity at prescribed fires is ramped up as the dry season progresses because the chances of fire escape increase concomitantly. The first prescribed fire occurs while the plants are carrying dead leaves below the mostly green foliage; a follow-up fire in the same season may be

The information in this section, unless otherwise noted, was supplied by the regional management officer of NT Bushfires, John Whatley, and his informed team-especially Sue amb and Lee Humphris.

needed to burn regrowth or the dead fuel created by the first fire. Heavy grazing, early dry-season cutting and harvesting, and protracted use of herbicides may reduce grassy fuel loads to low levels. Gamba grass may burn at any time of the year, weather permitting; however, fire spread in the wet season may be more likely if fuel has accumulated over at least a 2-year period.

In this area, gamba grass is spreading rapidly. At the same time, the human population and numbers of dwellings in the area is growing. Few people are aware of the seriousness of the fire issue because it is a relatively new one and people moving into the area may come from areas without this grass. Dense stands of the grass occur in the vicinity of the city of Darwin and its satellite Palmerston in the Northern Territory and also in Queensland. Further serious consequences of fires in gamba grass seem inevitable.

With large fuel loads and intense fires, the smoke volume arising exacerbates an already high smoke load in the regional atmosphere during the dry season, the adverse air quality increasing hospital admissions for asthma sufferers (Johnston *et al.*, 2002).

5.9 DISCUSSION AND CONCLUSIONS

Fire regimes, and the environments they interact with, vary spatially at different scales. Fire interval, season of occurrence, and fire intensity all vary in ecologically significant ways in northern Australia. Peat fires may have local significance. Fire patchiness can be important at the event level—such as in the grazing-burning interaction on grasses—but the effects of patch size distributions of land with specific between-fire intervals and times since fire, as shown by Gill *et al.* (2003), has not been explored; landscape dynamics of fire-prone environments may be better understood when the dynamics of patches with different between-fire intervals (and other components of the fire regime) are also understood.

Fire regimes have changed over the course of the last century, and continue to change. This change has led to some significant environmental change, including, in some places, woody plant thickening but, in other places, decline of woody species like *Callitris intratropica*. With increasing intensification of the cattle industry, global warming, and further spread of exotic plants, perhaps even more rapid change is to be expected. These changes will affect the species composition of pastures and vegetation generally and, in turn, affect populations of dependent fauna. There is an obvious challenge to conservation managers as a result.

With the uncertainties of the future, the continuance and increasing the resolution and verification of fire mapping (Pereira, 2003) is essential. Enhancement of the monitoring of fires by remote sensing of fire intensity within and between events, or routine recording of the distribution of char and scorch heights in woody vegetation is possible but universally overlooked to date.

Fire is one of several key event-based drivers of ecosystems across northern Australia. If we are to have an understanding of management issues such as global warming, cattle husbandry (rather than "hunting"— Lewis, 1985), conservation of a

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hern lobal of a Istinctive biodiversity and human health and safety, we need to improve and regrate knowledge of fire regimes, herbivory, climate and resource economics. Ashieving such understanding remains a challenge despite the many gains in rowledge so briefly summarized in this chapter.

5.10 ACKNOWLEDGMENTS

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