ORIGINAL PAPER

Structure, composition and diversity of plant communities in FSC-certified, selectively logged forests of different ages compared to primary rain forest

Arbainsyah · H. H. de Iongh · W. Kustiawan · G. R. de Snoo

Received: 8 January 2014/Revised: 3 May 2014/Accepted: 28 May 2014/ Published online: 18 June 2014 © Springer Science+Business Media Dordrecht 2014

Abstract The impact of logging on plant communities was studied in forest that has been logged selectively 1, 5 and 10 years previously (following a certified procedure): diversity was compared with that of primary rain forest in the Berau region of East Kalimantan, Indonesia. Four sets of 20 transects located within an area of 6 ha were sampled for all trees, saplings and seedlings, and records were made of topographic position, structure, composition and species diversity. There was a high level of floristic similarity between primary forests at the study sites compared to primary forest elsewhere in Kalimantan. The impact of logging is therefore likely to be the most important factor determining any differences between the plant communities of the selectively logged and primary forest sites. We found differences in species composition and abundance of most plants between selectively logged and primary forest. Overall, stem densities of trees in the primary forest were higher than in the three selectively logged forest sites. Stem densities of saplings were equivalent in all four forests. Seedling stem densities were higher in the forest site logged 10 years previously than in primary forest. Our results showed that the forests logged selectively under certified regimes still have a high plant diversity, possibly indicating that biodiversity values may be conserved by following certification procedures.

Keywords Sustainable forest management · Selective logging · Species diversity · Forest structure · Tropical rain forest

Communicated by Peter Ashton.

e-mail: arbainsyah.ins@gmail.com; arbainsyah@cml.leidenuniv.nl

W. Kustiawan Universitas Mulawarman, Kampus Gunung Kelua, Jl. Kuaro I/5, PO-Box 1068, Samarinda, Kalimantan Timur, Indonesia

Arbainsyah $(\boxtimes) \cdot H$. H. de Iongh \cdot G. R. de Snoo Institute of Environmental Science (CML), Leiden University, Einsteinweg 2, PO-Box 9518, 2300 RA Leiden, The Netherlands

Introduction

Tropical rain forests are recognized for their high biological diversity and their ecosystem services (Richards 1952; Whitmore 1984; Sheil and van Heist 2000; Jennings et al. 2001). Large parts of East Kalimantan are now covered by forests that are degraded as a result of fire and logging (Slik et al. 2002; van Nieuwstadt 2002; Meijaard et al. 2005; Eichhorn et al. 2006). Forest certification [Lembaga Ekolabel Indonesia and the Forest Stewardship Council (FSC)] has been introduced in Indonesia since several decades. The impact of FSC-certified logging on biodiversity has rarely been quantified, however (van Kuijk et al. 2009). There is a need to develop suitable biological indicators of sustainable forest management at the forest management unit level (Ghazoul and Hellier 2000; de Iongh and van Weerd 2006; de Iongh and Persoon 2010).

Commercial logging leads to fragmentation and degradation of the remaining tropical rain forests (Kartawinata 1977; Skole and Tucker 1993; Parthasarathy 1999), and results in many processes negatively affecting populations of plants and animals. When basic biological characteristics of the commercial species are considered in timber harvesting prescriptions, mixed dipterocarp forests appear capable of sustained timber yield in combination with habitat conservation. The Indonesian selective logging system allow selective logging intensity of ≥ 8 trees/ha associated with a felling cycle of 40–60 years depending on site conditions (Sist et al. 2003; van Kuijk et al. 2009). It has been more than 10 years since parts of the forest were selectively logged in the initial exploitation period in the 2,000 s (Kuswandari 2004). Intermediate disturbance hypothesis is one of the most frequently suggested non-equilibrium explanations for intervene maintain of species diversity in all communities (Connell 1978; Wilson 1990; Roxburgh et al. 2004).

Tree mortality in the understorey of logged forest is at least 2–3 times lowers than in the forest overstorey, and mostly occurs near and on skid trails (Webb 1998; Woods 1989; Pinard et al. 2000; Slik et al. 2002). In addition, some light-demanding, non-pioneer species may exhibit higher growth rates after logging. The increased light levels in the understorey of logged forests result in the rapid growth of many herbaceous and woody pioneer species (Woods 1989; Fredericksen and Mostacedo 2000). Trees make up only a part of the tropical rain forest ecosystem; herbs, shrubs, ferns and lianas generally constitute a large component of total plant diversity (Eichhorn et al. 2006; Yassir et al. 2010). To evaluate its biodiversity it is very important to know the vegetation composition of a forest type, from canopy to forest floor including trees, climbers (liana and rattan), non-rattan (Palmae), herbs, shrubs, etc., all of which are genetic resources for plant species within the forest.

Many impact of logging have been studied; tree mortality in the forest overstorey (diameter at breast hight (dbh) ≥ 10 cm) (Slik et al. 2002; van Nieuwstadt 2002), the mortality of canopy trees due to edge effects (Laurance et al. 2000), recruitment failure resulting from over-predation of seeds (Curran et al. 1999; Eichhorn et al. 2006), reduced seedling establishment and plant growth (Slik 2001; Bruna et al. 2002; Bruna 2003), local extinction of plants (Benitez-Malvido and Martinez-Ramos 2003), decline in butterfly abundance and/or diversity (Cleary 2002), decline in bird abundance and/or diversity (Boulinier et al. 2001; Beier et al. 2002; Slik and van Balen 2006), and decreased pollination (Ashworth et al. 2004). Logging also often leads to an increase in local human populations and to increased accessibility of the forest (Kartawinata and Vayda 1984), which in turn results in increased illegal logging and hunting and a decrease in biodiversity of remaining forest fragments (Laurance 1998; Hartshorn and Bynum 2001; Curran et al.

2004). The final outcome may be local mass extinctions of species as has been recently documented for Singapore (Brook et al. 2003). Because tropical rain forests harbour most of the world's biodiversity, tropical deforestation has become the major cause of global species extinctions (Pimm and Raven 2000).

The main goal of our research is to quantify the impact of FSC certification on botanic diversity and forest structure of tropical lowland forest in Borneo. Here we present the results of a detailed study of selectively logged forests (FSC certified) and primary rain forest site in the Indonesian province of East Kalimantan, including all terrestrial vascular plants. The three logged forest plots had been logged 1, 5 and 10 years ago. We analysed the structure and composition of forest plots under different logging regimes by assigning species to life forms that can be readily applied in the field (e.g., Eichhorn et al. 2006). We assessed the impact of selective logging at the landscape level to ensure that our plant diversity assessment was representative for the large scale at which disturbance by logging activities occurs. The numbers were expected to reflect the scale and severity of the disturbances taking place in a large forest area (Primack and Lee 1991; Davies et al. 1998; Slik et al. 2002), and as such could be useful to estimate the impact of logging on future plant diversity. Finally, we address the question what is the vegetation structure and composition in under sustainable forest management, FSC-certified selectively logged forests of different ages in comparison to primary forest?

Materials and methods

Study area

The study area is located in a lowland forest within the forest concession of PT. Hutansanggam Labanan Lestari (HLL) Labanan, East Kalimantan. The largest share of the new company area belong to the state-owned logging company of PT. Inhutani I, in Berau district, in the northeastern part of the Indonesian province of East Kalimantan (Fig. 1). The elevation range at the study area is 25–140 m above sea level. The topography consists of a rolling hilly landscape with shallow valleys and gullies, the highest elevation being 140 m. The soils consist of loamy clay and sandy soils with a top soil layer of approximately 5–10 cm (Mantel et al. 2002).

Sites were established in primary forest (1 site) and selectively logged forest (3 sites) (Fig. 1). In these plots, three groups of plants (trees, saplings and seedlings) were systematically recorded along a line transect of 10×300 m. In total, 20 transects were sampled, 5 in primary forest and 5 in each of the 3 selectively logged over forest. We divided each line transect into 30 plots of 10×10 m (a total of 150 plots in each site) to measure all trees with a dbh ≥ 10 cm (dbh 130 m above ground level or, if buttresses are present, 30 cm above buttresses) using the circumference method. Within each plot, a subplot of 5×5 m for saplings and 2×2 m for seedlings were established and measured (number of individuals per species and cover estimate). These measurement quadrats for seedlings and saplings were positioned alternately to the left and right of transect centre lines at intervals of 100 m, resulting in 15 subplots per site. Plants were sampled and identified, i.e. whenever a fertile plant, labeled (vouchers stored in the Herbarium Wanariset Samboja, Indonesia). The field work was done by the principle author, together with field assistants.



Fig. 1 Map of East Kalimantan with the location of study areas P1 plots: primary forest site, L1 plots: forest site logged 1 year ago (2011), L2 plots: forest site logged 5 years ago (2007), L3 plots: forest site logged 10 years ago (2003)

Life forms

In order to provide a detailed description of the structure and composition of the terrestrial plant community in the four research sites, all species were assigned to life forms and taxonomic criteria (Table 1). The criteria used were chosen in such a way that they provided maximum information about the forest structure and composition while still being applicable for para-taxonomists in comparative studies in East Kalimantan and elsewhere in the tropics (Eichhorn et al. 2006). To enhance compatibility with growth forms that were used in similar studies in the past, species were first assigned to three major groups: trees, saplings and seedlings. Several life forms were distinguished within these three major groups, based on taxonomic criteria and growth form. Throughout this study, each plant species was assigned referred to one of the following three life forms.

- (1) *Trees* defined as non-climbing woody species of which the mature individuals had a stem diameter ≥ 10 cm.
- (2) Saplings defined as all herbaceous species, non-climbing woody species and climbing woody species of which the mature individuals had a stem diameter ≤ 10 cm and were on average more than 1.5 m tall.
- (3) Seedlings defined as all herbaceous species, non-climbing woody species and climbing woody species of which the mature individuals were on average less than 1.5 m tall.

2449

Life form	Taxa	Growth form and size
Trees (woody non-climbers w	with stem diameter ≥ 10 cm)	
Palms-trees	Palmae	Woody non climbers, height > 1.3 m
Dicots-trees	Dicotyledonae	Woody non climbers, height > 1.3 m
Saplings (herbs, shrubs, clim	bers, woody non-climbers with	diameter < 10 cm)
Monocots-other herbs	Monocotyledonae	Herbaceous non climbers, height > 1.5 m
Dicots-trees	Dicotyledonae	Woody non climbers, height > 1.5 m
Dicots-lianas	Dicotyledonae	Climber, height > 1.5 m
Dicots-shrubs	Dicotyledonae	Woody non climbers, with many branches from the ground, height > 1.5 m
Seedlings (herbs, shrubs, clir	nbers, woody non-climbers < 1 .	5 height
Palms-lianas (rottans)	Palmae	Climber, height < 1.5 m
Palms-palmlets	Palmae	Woody non-climber, height < 1.5
Monocots-small lianas	Monocotyledonae	Climber, height < 1.5 m
Monocots-other herbs	Monocotyledonae	Herbaceous non climbers
Monocots-grass-like	Graminae + Cyperaceae	Herbaceous, leaves linear
Dicots-small treelets	Dicotyledonae	Woody non-climber, height < 1.5
Dicots-small lianas	Dicotyledonae	Climber, height < 1.5 m
Dicots-small shrubs	Dicotyledonae	Woody non-climber, with many branches from the ground, height < 1.5
Ferns-small lianas	Filicopsida	Climbers
Ferns-herbs	Filicopsida	Herbaceous non climbers

Table 1 List of the life forms used in this study and the taxa, growth form and size class they represent

Data analyses and statistics

All data analyses were performed with Microsoft Excel and SPSS 13.0 software to calculate standard deviation for the estimated average number of stems. Tree species diversity of the plots was compared between of selectively logged forests, and primary forest. These comparisons were made to compensate for differences in sample sites between of selectively logged forests in comparison to primary forest. Post hoc comparisons between of selectively logged forests, and primary forest were made using the Fisher's Least Significant test (one-way ANOVA based on ln transformed data, with Bonferroni multiple comparison test). The stages in evaluating the data were as follows:

- (1) Counting the number of stems of all trees, saplings and seedlings found in each transect.
- (2) Calculating Species Diversity Index (H), Evenness Index (E) and Dominance Index (C).

Species diversity analysis was done with the Shannon Diversity Index, with the Jost (2006) formula as follows:

Diversity Index
$$H = \sum_{S=1}^{S} p_i \ln p_i$$
,

where H is the Shannon Diversity Index, S is the total number of species in the community, p_i is the proportion of abundance of the i species. p_i is calculated by dividing the number of species i by the total number of all species.

Then to determine the Evenness Index (E) the Pielou Evenness Index (Ludwig and Reynold 1988) formula was used:

Evenness Index (E) =
$$\sum H/\ln(S)$$
,

where E is the Pielou Evenness Index, ln is the normal logarithm, S number of species. The Dominance Index (C) was determined using the formula: Dominance Index (C) = \sum (ni/N)²

Dominance Index (C) =
$$\sum (ni/N)^2$$
,

where C is the Dominance Index, ni is the number of individuals of a certain species, N is the total number of individuals of all species.

(3) Calculating the estimated average number of stems (N) and stems per hectare (q) for each class (trees, saplings and seedlings) Ludwig and Reynold (1988):

$$q = \frac{\sum_{i=1}^{n} Y_i}{\sum_{i=1}^{n} X_i},$$

where q is the average number of stems (N) per hectare, Yi is the Number of stems (N) per hectare of given transect, Xi is the area of a given transect.

(4) Calculating the Importance Value index (I.V.) for each level/strata. The formula used in calculating I.V. was the quadrate method (Mueller-Dombois and Ellenberg 1974). The (I.V.) of species is defined as sum of its relative density (RD), relative dominance (Rd) and relative frequency (RF) (I.V.) = RD + RF + Rd), which are calculated using the following equations:

$$\begin{aligned} \text{Density} \left(D \right) &= \frac{\text{Number of individuals of a species}}{\text{Area of all sample units}} \\ \text{Relative Density} \left(\text{RD} \right) &= \frac{\text{Number of individuals of a species}}{\text{Density for all species}} \times 100 \% \\ \text{Frequency} \left(\text{F} \right) &= \frac{\text{Number of quadrats containing a certain species}}{\text{Total number of quadrats}} \\ \text{Relative Frequency} \left(\text{RF} \right) &= \frac{\text{Frequency of a certain species}}{\text{Total number of species}} \times 100 \% \\ \text{Dominance} \left(\text{d} \right) &= \frac{\text{Basal area of a species}}{\text{Area of all sample units}} \\ \text{Relative Dominance} \left(\text{Rd} \right) &= \frac{\text{Dominance of one species}}{\text{Dominance of all species}} \times 100 \% \end{aligned}$$

The Importance Value index for trees and saplings was calculated based on the formula:

Importance Value index (I.V.) = RD + RF + Rd

For seedling levels, the species importance value index was calculated using the formula:

Importance Value index (I.V.) = RD + RF

Results

Composition and biodiversity in selectively logged and primary forests

Tree diversity was higher in forest site logged selectively 1 year ago, where H > 4.5, but the dominance index and number of stems were almost as high in the primary forest site (Table 2). This high dominance index is indicated by the very abundant tree species of *Hopea semicuneata*, with an I.V. score of 43.4 % (Table 6). In contrast to the tree diversity index, the highest evenness index was found in the forest site logged 1 year ago, with a total of 156 species encountered (Table 3).

Sapling diversity was higher in the primary forest site compared to selectively logged forest sites, but the dominance index of the forest site logged 5 years ago was lower than that of the other forest sites (Table 2). This high dominance is indicated by regenerating species of *Madhuca malaccensis*, with an I.V. score of 25.5 %, two times more dominant than other species (Table 7). In contrast to the saplings diversity index, the highest evenness index was found in primary forest with a total number of 97 species encountered (Table 4).

Seedling diversity was high in some of the selectively logged forest sites, in the forest site logged 1 year ago, but due to the dominance index was found in primary forest site (Table 2). This dominance was indicated by the very abundant species of *H. semicuneata*, with an I.V. score of 24.6 %, which is also the highest in the regeneration of saplings and trees (Tables 7 and 8). In contrast to the seedlings diversity index, the highest evenness index was found in the forest site logged 1 year ago, with a total number of 95 species encountered (Table 5).

Index	Primary forest	Selectively log	ged forest	
		1 year ago	5 years ago	10 years ago
Tree (1.5 ha)				
Diversity (H)	4.259	4.509	4.339	4.260
Dominance (C)	0.033	0.019	0.023	0.022
Evenness (E)	0.853	0.893	0.879	0.888
Number of stems	612	492	501	558
Sapling (0.375 ha)				
Diversity (H)	4.352	4.175	4.095	4.150
Dominance (C)	0.015	0.020	0.024	0.023
Evenness (E)	0.951	0.937	0.922	0.918
Number of stems	245	247	242	244
Seedling (0.06 ha)				
Diversity (H)	4.067	4.319	4.090	4.217
Dominance (C)	0.030	0.017	0.025	0.020
Evenness (E)	0.908	0.948	0.913	0.930
Number of stems	328	306	303	346

Table 2 Comparison between the diversity index (H), dominance index (C), evenness index (E) and number of stems for all trees (dbh ≥ 10 cm) per 1.5 ha, all saplings (dbh < 10 cm) per 0.375 ha and all seedlings in ground cover per 0.06 ha in primary forest site and three selectively logged forest: logged 1 year ago, logged 5 years ago and logged 10 years ago

	-		-	/	0		/	Ì				
Trees	Abundance si	tems (per 100 1	m ²)		Species per si	ubplot (10×10)	0 m)		Total spec	cies numb	er	
	Primary	Selectively lc	ogged forest		Primary	Selectively lo	gged forest		Primary	Selective	ly logged f	orest
	lorest	1 year ago	5 years ago	10 years ago	lorest	1 year ago	5 years ago	10 year ago	IOTESI	1 year ago	5 years ago	10 years ago
Life form												
Palms- trees	0.00 ± 0.00	0.05 ± 0.29	0.00 ± 0.00	0.02 ± 0.25	0.00 ± 0.00	0.04 ± 0.19	0.00 ± 0.00	0.01 ± 0.08	0	1	0	1
Dicots- trees	4.08 ± 1.97	3.23 ± 2.13	3.34 ± 2.08	3.70 ± 1.86	3.70 ± 1.78	3.30 ± 1.67	3.19 ± 1.79	3.49 ± 1.62	147	155	139	120
Total	4.08 ± 1.97	3.28 ± 2.15	3.34 ± 2.08	3.72 ± 1.87	3.70 ± 1.78	3.33 ± 1.70	3.19 ± 1.79	3.49 ± 1.62	147	156	139	121
Abundance the total ol ago, in ple correction	e expressed as constructed species of $10 \times 10 \text{ r}$ for multiple test	densities of ster numbers in all n, in total of 1 sts)	ms exceeding 1 subplot togethe 1.5 ha. Bold av	.3 height. Spec er in primary fo erages for sele	ies richness at rest site and thr ctively logged	the subplot scal ree selectively l forest sites dif	le expressed as ogged forest: la fer significantl	species numbe ogged 1 year ag y from those o	er per subp go, logged (of the prim	lot and at 5 years ag ary forest	the landsca o and logge site (with	pe scale as d 10 years Bonferroni

Table 3 Comparison between the abundance and species richness (average \pm standard deviation) for all trees (dbh ≥ 10 cm) life form and forest sites

Saplings	Abundance ster	ms (per 25 m^2)			Species per sul	bplot (5 \times 5 m)	(Total species nu	umber		
	Primary	Selectively log:	ged forest		Primary	Selectively lo	gged forest		Primary forest	Selective	ely logged	forest
	Iorest	1 year ago	5 years ago	10 years ago	Iorest	1 year ago	5 years ago	10 years ago		l year ago	5 years ago	10 years ago
Life form												
Monocots-herbs	0.00 ± 0.00	0.13 ± 0.52	0.00 ± 0.00	0.00 ± 0.00	0.00 ± 0.00	0.07 ± 0.26	0.00 ± 0.00	0.00 ± 0.00	0	1	0	0
Dicots-treelets	16.27 ± 4.11	15.80 ± 6.19	15.40 ± 5.26	15.60 ± 3.58	12.93 ± 3.01	9.40 ± 3.14	11.93 ± 4.20	10.40 ± 4.10	96	83	84	89
Dicots-lianas	0.00 ± 0.00	0.40 ± 1.55	0.00 ± 0.00	0.33 ± 1.29	0.00 ± 0.00	0.07 ± 0.26	0.00 ± 0.00	0.13 ± 0.52	0	1	0	7
Dicots-shrubs	0.06 ± 0.26	0.13 ± 0.35	0.73 ± 1.16	0.33 ± 0.72	0.07 ± 0.26	0.13 ± 0.35	0.40 ± 0.51	0.20 ± 0.41	1	1	1	1
Таха												
Monocots	0.00 ± 0.00	0.13 ± 0.52	0.00 ± 0.00	0.00 ± 0.00	0.00 ± 0.00	0.07 ± 0.26	0.00 ± 0.00	0.00 ± 0.00	0	1	0	0
Dicots	16.33 ± 4.06	16.34 ± 6.35	16.13 ± 5.88	16.27 ± 4.37	13.00 ± 2.98	9.60 ± 3.18	12.27 ± 4.40	10.73 ± 4.04	97	85	85	92
Total	16.33 ± 4.07	16.47 ± 6.50	16.13 ± 5.88	16.27 ± 4.37	13.00 ± 2.98	9.67 ± 3.13	12.27 ± 4.40	10.73 ± 4.04	97	86	85	92
Abundance express numbers of species 0.375 ha. Bold ave	sed as densities c in all subplots to rages for selectiv	of stems exceedin ogether in prima vely logged fores	$ng \ge 1.5$ height. ry forest site and st sites differ sign	Species richness 1 three selectively nificantly from th	t at the subplot s y logged forest: hose of the prim	scale expressed logged 1 year <i>i</i> nary forest site of	as species numb ago, logged 5 yei (with Bonferroni	ber per subplot a ars ago and logg	nd at the landsca ed 10 years ago, multiple tests)	pe scale a in plot of	s the total 5 x 5 m, i	observed n total of

Table 4 Comparison between the abundance and species richness (average \pm standard deviation) for all saplings (dbh < 10 cm) life form and forest sites

ŝ
site
est
for
and
E
for
life
ß
dlir
see
all
for
(u
atio
evis
1 de
larc
anc
E st
ы Ц
srag
ave
ss (
nne
ricł
es
Seci
l st
and
JCe
ıdaı
pun
e a
th
eer
etw
n b
iso
par
uo
Ŭ
e S
lde
Ë

Seedlings	Abundance ster	ns (per 4 m ²)			Species per sub	plot $(2 \times 2 m)$			Total spec	ies number		
	Primary	Selectively logg	ted forest		Primary	Selectively logg	ged forest		Primary	Selective	ily logged fi	orest
	101031	1 year ago	5 year ago	10 year ago	101051	1 year ago	5 year ago	10 year ago	101631	1 year ago	5 years ago	10 years ago
Life form												
Palms-lianas (rottans)	0.80 ± 1.21	0.27 ± 0.46	0.13 ± 0.35	0.13 ± 0.35	0.53 ± 0.74	0.27 ± 0.46	0.13 ± 0.35	0.13 ± 0.35	6	7	7	7
Palms-small trees	0.13 ± 0.52	0.07 ± 0.26	0.00 ± 0.00	0.00 ± 0.00	0.07 ± 0.26	0.07 ± 0.26	0.00 ± 0.00	0.00 ± 0.00	1	1	0	0
Monocots- small lianas	0.07 ± 0.26	0.07 ± 0.26	0.00 ± 0.00	0.00 ± 0.00	0.07 ± 0.26	0.07 ± 0.26	0.00 ± 0.00	0.00 ± 0.00	1	1	0	0
Monocots- herbs	0.07 ± 0.26	0.33 ± 0.72	0.67 ± 1.84	0.13 ± 0.35	0.07 ± 0.26	0.20 ± 0.41	0.20 ± 0.41	0.13 ± 0.35	1	7	7	7
Monocots- grass	0.20 ± 0.56	0.53 ± 1.19	0.00 ± 0.00	0.13 ± 0.52	0.13 ± 0.35	0.27 ± 0.59	0.00 ± 0.00	0.07 ± 0.26	6	2	0	1
Dicots-small trees	18.80 ± 9.22	12.87 ± 6.86	15.80 ± 8.05	18.60 ± 9.91	8.33 ± 2.32	6.87 ± 3.16	8.13 ± 2.64	8.40 ± 3.58	69	67	65	75
Dicots-small lianas	1.67 ± 1.95	3.33 ± 3.54	1.93 ± 1.39	0.80 ± 1.26	1.00 ± 1.00	1.47 ± 0.99	1.47 ± 1.13	0.47 ± 0.74	6	13	15	9
Dicots-shrubs	0.07 ± 0.26	0.47 ± 0.92	0.13 ± 0.52	0.27 ± 0.46	0.07 ± 0.26	0.20 ± 0.41	0.07 ± 0.26	0.27 ± 0.46	1	3	1	1
Ferns-small lianas	0.07 ± 0.26	0.60 ± 1.59	1.47 ± 3.09	1.20 ± 2.78	0.07 ± 0.26	0.13 ± 0.35	0.20 ± 0.41	0.40 ± 0.83	1	1	7	4
Ferns-herbs	0.00 ± 0.00	1.67 ± 2.23	0.07 ± 0.26	1.80 ± 4.18	0.00 ± 0.00	0.60 ± 0.74	0.07 ± 0.26	0.27 ± 0.59	0	5	1	2
Ferns-small trees	0.00 ± 0.00	0.20 ± 0.41	0.00 ± 0.00	0.00 ± 0.00	0.00 ± 0.00	0.20 ± 0.41	0.00 ± 0.00	0.00 ± 0.00	0	1	0	0
Таха												
Monocots	1.27 ± 1.39	1.27 ± 1.28	0.80 ± 1.86	0.40 ± 0.74	0.87 ± 0.83	0.87 ± 0.74	0.33 ± 0.62	0.33 ± 0.62	6	8	4	5
Dicots	20.53 ± 8.83	16.67 ± 8.80	17.87 ± 8.24	19.67 ± 9.82	9.40 ± 2.47	8.40 ± 3.91	9.67 ± 2.58	9.13 ± 3.64	78	83	81	82
Ferns	0.07 ± 0.26	2.47 ± 2.83	1.53 ± 3.20	3.00 ± 6.89	0.07 ± 0.26	1.07 ± 0.96	0.27 ± 0.59	0.67 ± 1.40	1	4	3	6

continued
S
le
Lab

Seedlings	Abundance ste	times (per 4 m^2)			Species per sul	pplot (2 \times 2 m)			Total speci	es number		
	Primary	Selectively log	ged forest		Primary	Selectively log	ged forest		Primary	Selective	ly logged f	orest
	IOrest	1 year ago	5 year ago	10 year ago	lorest	1 year ago	5 year ago	10 year ago	Iorest	1 year ago	5 years ago	10 years ago
Total	21.87 ± 8.84	20.40 ± 9.82	20.20 ± 6.46	23.07 ± 11.20	10.33 ± 2.64	10.33 ± 4.27	10.27 ± 2.34	10.13 ± 3.09	88	95	88	93
Abundance expre species numbers i Bold averages for	ssed as densities of all subplot toget selectively logge	of stems percent ther in primary f	age ground cover orest site and thre ffer significantly	:. Species richnes: se selectively logg from those of the	s at the subplot s ged forest: logge e primary forest	scale expressed a d 1 year ago, log site (with Bonfe	s species numbe ged 5 years ago rroni correction	ar per subplot an and logged 10 y for multiple test	d at the lands ears ago, in p s)	scape scale plot of 2 ×	e as the tota 2, in total	l observed of 0.06 ha.

Abundance and composition of three major groups of species

Tree densities were significantly lower in the forest sites logged selectively 1 and 5 years ago than in the primary forest site, but tree densities in the forest site logged 10 years ago were similar to those of the primary forest site (Table 3). Dicot trees were clearly the dominant tree type, as they accounted for ca. 99 % of the stems in all four forest sites. However, palm trees still exceeded densities of 3 stems ha⁻¹ in at least one of the four forest sites.

There were no significant differences in sapling densities among all four forests, but due to the dicot shrubs, the total number of stems in the forest site logged 5 years ago was higher than in the other three forest sites (Table 4). When the dicot shrubs were excluded from the analysis, stem density in the selectively logged forest, namely of the forest site 5 years ago was 242 stems ha⁻¹ this is the lowest when compared to primary forest site, the sites logged 1 year ago and 10 years ago with stem density of 245, 247 and 244 stems ha⁻¹.

There were no significant differences between seedlings, which were in general equally abundant in all four forest sites. Due to the presence of small dicot lianas, the total number of stems in forest sites logged selectively 1 and 10 years ago was significantly greater than in the other forest sites (Table 5). Palm lianas were clearly the most abundant life form in primary forest, while small fern lianas and fern herbs were both very abundant in selectively logged forest sites.

Small trees, small lianas, herbs and shrubs all contributed importantly to overall seedling densities, but there were pronounced differences between these growth forms with respect to the forest type (Fig. 2, Table 8). Densities of small tree seedlings were highest in the primary forest site and lowest in the forest sites 1 and 5 years ago with the forest site logged 10 years ago being more or less intermediate. When compared to small liana seedlings, the forest site 1 year ago had value twice that of the primary forest site (Fig. 2). Densities of both herb and shrub seedlings were almost three times higher in the logged forest sites compared to the primary forest site (Fig. 2; Table 5).

Abundance and composition of different types of trees

Dicot trees were significantly less abundant in the forest sites logged 1 and 5 years ago than in the primary forest site, but dicot tree densities in the forest site logged 10 years ago were similar to those of the primary forest site (Tables 2 and 3). In the primary forest site,



the dominant families were Dipterocarpaceae, Euphorbiaceae, Caesalpiniaceae, Burseraceae and Sapotaceae (Table 6). Species that contributed greatly to the dominance of these families were *H. semicuneata* and *Dipterocarpus lowii* (both Dipterocarpaceae), *Chaetocarpus castanocarpus* (Euphorbiaceae), *Cynometra elmeri* (Caesalpiniaceae) and *Palaquium stenophyllum* (Sapotaceae).

The abundance of tree species of *Hopea* was highest in the primary forest site (Table 6), only Dipterocarpaceae were more dominant in the forest site logged 10 years ago than in the primary forest site, while other dominant families of the three selectively logged forest sites were much less abundant compared with the primary forest site (Fig. 3). *Hopea cernua* and *Hopea pachycarpa* were especially abundant in the three selectively logged forest sites (Table 9). Other species that were abundant in the logged forest sites were *Syzygium tawahense* (Myrtaceae), *Shorea parvifolia* (Dipterocarpaceae), *Gironniera nervosa* (Ulmaceae), *Palaquium calophyllum* (Sapotaceae) and *Neoscortechinia kingii* (Euphorbiaceae).

Species that were only abundant in the forest site 10 years ago were Allanthospermum borneensis (Simaroubaceae), Canarium denticulatum (Burseraceae), C. castanocarpus and Macaranga gigantea (both Euphorbiaceae), Gluta renghas (Anacardiaceae), M. malaccensis (Sapotaceae), Myristica villosa (Myristicaceae), Scaphium macropodum (Sterculiaceae), S. parvifolia, Shorea inappendiculata and Vatica nitens (all Dipterocarpaceae). All these species were absent or rare in the forest sites logged 1 and 5 years ago.

No significant differences between forest sites were observed in other types of palm trees. Palm trees were only abundant in the forest sites logged 1 and 10 years ago. Most stems were *Oncosperma horridum* (Table 9). Even though this species has a multi-stemmed growth form, total stem densities of this species were rather low, particularly in the forest site logged 10 years ago. Stem clusters of this species were present in only seven plots of the forest site logged 1 year ago and five plots of the one forest site logged 10 years ago.

Monocot trees were only represented by species of *O. horridum*. This species occurred in very low densities in the four forest sites and was never observed in the forest site logged 5 years ago or in the primary forest site. In contrast to this, there were 7 stems presence among the 150 plots of the forest site logged 1 year ago, making it the dominant monocot tree of this forest type (Table 3).

Abundance and composition of different types of saplings

Dicot saplings had a similar abundance in all four forest sites, with stem densities of around 240 stems ha⁻¹ (Tables 2 and 4). In the primary forest site, the dominant families were Euphorbiaceae, Dipterocarpaceae, Myristicaceae, Ebenaceae, Sapotaceae, and Polygalaceae (Table 7). Species that contributed greatly to the dominance of these families were *H. semicuneata* (Dipterocarpaceae), *Knema laurina* (Myristicaceae), *Croton argy-ratus, Koilodepas brevipes* and *Cleistanthus erycibifolius* (all Euphorbiaceae), *Xanthophyllum obscurum* (Polygalaceae), *M. malaccensis* (Sapotaceae) and *Diospyros curanii* (Ebenaceae).

Generally the number of families of saplings in the 1 year ago logged site was about twice the number in the other sites (Table 7). However the differences between forest sites were less pronounced for the dominant dicot tree families (Table 6). In the selectively logged forest sites, and particularly in the one logged 1 year ago, species of *Macaranga*, *Glochidion* and *Shorea* were more abundant than in the primary forest site. Apparently *Melastoma malabathricum* and *Glochidion arborescens* had ecological characteristics

Table 6 The ten most abundant fi	families,	genera	nera and species of trees in the primary forest site and three selectively logged forest: logged 1 year ago, logged 5 years ago	so and
logged 10 years ago				
Trees-Primary forest	Abun	%	% Selectively logged forest	

Tree	ss-Primary forest	Abun	%	Selectively logged fore:	st							
				1 year ago	Abun	%	5 years ago	Abun	%	10 years ago	Abun	%
Fam	ilies											
-	Dipterocarpaceae (6, 22)	123	73.7	Dipterocarpaceae (6, 20)	98	61.6	Dipterocarpaceae (6, 25)	92	63.2	Dipterocarpaceae (6, 30)	138	86.7
7	Euphorbiaceae (10, 13)	51	21.8	Myrtaceae (1, 6)	29	29.2	Sapotaceae (2, 6)	69	37.3	Sapotaceae (3, 5)	72	29.2
3	Caesalpiniaceae (5, 5)	34	17.7	Lauraceae (6, 7)	41	28.7	Myrtaceae (1, 4)	45	28.8	Euphorbiaceae (7, 9)	49	25.0
4	Burseraceae (2, 5)	33	16.2	Euphorbiaceae 10, 15)	45	23.1	Euphorbiaceae (8, 13)	50	25.0	Simaroubaceae (2, 2)	48	23.2
5	Sapotaceae (2, 6)	32	15.2	Myristicaceae (4, 5)	43	21.0	Burseraceae (3, 6)	22	13.0	Myrtaceae (1, 4)	28	13.9
9	Verbenaceae (1, 2)	39	14.7	Ebenaceae (1, 5)	24	13.5	Lauraceae (3, 3)	21	12.1	Burseraceae (3, 4)	23	13.2
٢	Flacourtiaceae (1, 1)	19	13.2	Caesalpiniaceae (5, 7)	14	10.8	Myristicaceae (4, 8)	25	11.8	Myristicaceae (4, 6)	23	12.0
×	Myrtaceae (2, 4)	23	12.3	Burseraceae (3, 6)	18	10.5	Sterculiaceae (3, 5)	11	10.7	Caesalpiniaceae (3, 3)	14	9.7
6	Ebenaceae (1, 5)	29	11.7	Sapotaceae (2, 7)	22	10.5	Caesalpiniaceae (3, 5)	14	10.4	Anacardiaceae (3, 3)	15	9.3
10	Guttiferae (3, 3)	19	8.6	Annonaceae (6, 8)	18	8.7	Anacardiaceae (3, 5)	13	8.3	Sterculiaceae (3, 4)	14	9.2
Gen	era											
1	Hopea (Dipt, 2)	78	44.5	Shorea (Dipt, 12)	70	45.3	Shorea (Dipt, 14)	38	33.6	Shorea (Dipt, 18)	78	51.4
7	Dipterocarpus (Dipt, 4)	15	14.9	Syzygium (Myrt, 6)	29	28.8	Syzygium (Myrt, 4)	45	28.4	Madhuca (Sapo, 2)	60	22.8
ю	Teijsmanniodendron (Verb, 2)	39	14.4	Eusideroxylon (Laur, 1)	25	17.9	Madhuca (Sapo, 3)	56	26.2	Allanthospermum (Sima, 1)	47	21.8
4	Hydnocarpus (Flac, 1)	19	13.1	Diospyros (Eben, 5)	24	13.1	Hopea (Dipt, 2)	24	13.2	Dipterocarpus (Dipt, 7)	22	15.9
5	Shorea (Dipt, 12)	22	11.9	Palaquium (Sapo, 4)	19	8.7	Palaquium (Sapo, 3)	13	11.8	Syzygium (Myrt, 4)	28	13.5
9	Diospyros (Eben, 5)	29	11.4	Hopea (Dipt, 2)	11	8.0	Vatica (Dipt, 3)	19	10.3	Vatica (Dipt, 2)	19	9.5
٢	Syzygium (Myrt, 3)	22	11.4	Myristica (Myri, 2)	14	7.8	Alseodaphne (Laur, 1)	14	8.4	Chaetocarpus (Euph, 1)	12	8.7

continued
9
e
ab
r .

1 al												
Tre	es-Primary forest	Abun	%	Selectively logged forest								
				1 year ago	Abun	%	5 years ago	Abun	%	10 years ago	Abun	%
~	Canarium (Burs, 3)	20	10.3	Knema (Myri, 1)	16	7.6	Horsfieldia (Myri, 2)	15	7.2	Canarium (Burs, 2)	14	7.4
6	Chaetocarpus (Euph, 1)	19	9.6	Artocarpus (Mora, 2)	11	6.7	Barringtonia (Lecy, 2)	15	7.0	Gluta (Anac, 1)	11	7.1
10 Spe	Palaquium (Sapo, 2) cies	13	7.9	Gymnacranthera (Myri, 1)	12	6.4	Gluta (Anac, 3)	10	6.6	Scaphium (Ster, 1)	10	6.8
Г	Hopea semicuneata	76	43.4	Syzygium tawahense	15	19.1	Syzygium tawahense	42	26.5	Madhuca malaccensis	58	21.6
0	Hydnocarpus polypetala	19	13.1	Eusideroxylon zwageri	25	17.7	Madhuca malaccensis	46	21.1	Allanthospermum borneensis	47	21.5
3	Teijsmanniodendron coriaceum	32	11.5	Shorea parvifolia	24	15.3	Hopea cerma	21	11.1	Syzygium tawahense	21	9.8
4	Chaetocarpus castanocarpus	19	9.6	Shorea pinanga	17	8.5	Alseodaphne elmeri	14	8.4	Shorea parvifolia	12	9.2
2	Syzygium tawahense	16	8.8	Knema laurina	16	7.5	Palaquium stenophyllum	٢	6.6	Shorea inappendiculata	10	9.1
9	Dipterocarpus lowii	6	8.3	Palaquium calophyllum	16	7.2	Horsfieldia polyspherula	14	6.6	Chaetocarpus castanocarpus	12	8.6
2	Cynometra elmeri	16	7.7	Syzygium caudatilimba	8	6.4	Neoscortechinia kingii	11	6.4	Vatica nitens	13	7.1
×	Palaquium stenophyllum	12	7.2	Gymnacranthera farquhariana	12	6.3	Shorea pinanga	9	6.3	Gluta renghas	11	7.0
6	Diospyros curranii	16	6.6	Hopea dryobalanoides	6	6.1	Lithocarpus coopertus	6	5.8	Scaphium macropodum	10	6.8
10	Chionanthus sp.1	15	5.9	Diospyros borneensis	11	6.1	Drypetes kikir	11	5.8	Myristica villosa	12	6.5
In c	volumns after each taxon the	le corresp	onding	abundance (Abun), expresse	d as the	densit	ies of stems exceeding	≤ hdbh	10 cm)	in height 1.3 m, and the I	percenta	ge of



Fig. 3 Stem density (**a**) and total observed species number (**b**) in four forest sites of the 10 most speciesrich tree family of primary forest in Labanan, PT. Hutansanggam Labanan Lestari. *Solid bars*: primary forest site; cross-hatched bars: selectively logged forest; forest site logged 1 year ago; *dotted bars*: forest site logged 5 years ago; *open bars*: forest site logged 10 years ago

similar to species such as *Macaranga hypoleuca*, as it was fast-growing and particularly abundant in the selectively logged forest sites absent in the primary forest site (Table 7).

Dicot shrubs were the only life form more abundant in the forest site logged 5 years ago than in the other forest sites (Table 4). Dicot treelets were more abundance in the primary forest site than in the three logged ones. Typically, the dominant species were different in all four forest sites (Table 7). Some dicot liana species were abundant in the selectively logged forest sites, while being almost absent in the primary forest site. Species of *Uncaria* were only present in the forest sites 1 and 10 years ago (Table 9). Monocot herbs were less abundant than dicot herbs in the forest site logged 1 year ago and were absent in the three other forest sites (Table 4). Monocots were mainly represented by *Costus speciosus* (Table 9).

Abundance and composition of different types of seedlings

Dicot seedlings were more abundant in primary forest site than in the selectively logged forest sites, but total stem densities were highest in the forest site 10 years ago (Tables 2 and 5). In the primary forest site, the dominant families were Dipterocarpaceae, Euphorbiaceae,

Sapl	lings-Primary forest	Abun	%	Selectively logged forest								
				1 year ago	Abun	%	5 years ago	Abun	%	10 years ago	Abun	%
Fam	ilies											
-	Euphorbiaceae (14, 16)	43	28.2	Euphorbiaceae (8, 13)	93	71.4	Dipterocarpaceae (5, 12)	38	42.7	Euphorbiaceae (7, 13)	35	40.8
7	Dipterocarpaceae (4, 9)	24	18.8	Dipterocarpaceae (2, 7)	24	23.7	Euphorbiaceae (9, 11)	43	41.2	Dipterocarpaceae (5, 13)	35	36.8
3	Myristicaceae (2, 3)	10	15.0	Myristicaceae (3, 4)	10	19.1	Sapotaceae (1, 3)	26	31.4	Melastomataceae (3, 5)	31	26.7
4	Ebenaceae (1, 4)	12	14.1	Myrtaceae (1, 4)	11	16.5	Burseraceae (2, 4)	13	18.7	Sapotaceae (2, 3)	15	22.7
5	Sapotaceae (2, 4)	11	14.0	Rubiaceae (5, 5)	17	15.3	Rubiaceae (4, 4)	15	17.7	Myrtaceae (1, 1)	11	19.0
9	Polygalaceae (1, 2)	6	13.5	Annonaceae (2, 4)	7	12.7	Lauraceae (4, 4)	12	15.1	Rubiaceae (7, 8)	18	16.3
2	Annonaceae (6, 6)	16	12.5	Ebenaceae (1, 3)	6	12.4	Myrtaceae (1, 2)	10	15.0	Ulmaceae (1, 1)	7	13.5
8	Myrtaceae (1, 4)	11	9.9	Tiliaceae (2, 2)	9	12.0	Annonaceae (4, 4)	6	10.9	Polygalaceae (1, 2)	8	13.1
6	Burseraceae (2, 3)	7	9.6	Sapotaceae (1, 2)	9	10.5	Polygalaceae (1, 3)	5	10.8	Anacardiaceae (3, 3)	10	10.7
10	Meliaceae (1, 3)	7	8.2	Papilionaceae (1, 1)	5	8.3	Fagaceae (2, 2)	4	10.6	Burseraceae (3, 5)	6	9.8
Gen	era											
1	Diospyros (Eben, 4)	12	15.3	Macaranga (Euph, 3)	37	24.9	Madhuca (Sapo, 3)	26	30.2	Shorea (Dipt, 7)	25	26.9
5	Xanthophyllum (Poly, 2)	6	13.7	Glochidion (Euph, 2)	32	23.0	Shorea (Dipt, 6)	21	23.4	Macaranga (Euph, 5)	12	19.4
Э	Knema (Myri, 2)	8	11.7	Shorea (Dipt, 6)	23	20.8	Vatica (Dipt, 3)	12	17.0	Melastoma (Mela, 1)	27	19.2
4	Syzygium (Myrt, 4)	11	11.0	Syzygium (Myrt, 4)	11	15.4	Syzygium (Myrt, 2)	10	14.0	Syzygium (Myrt, 1)	11	18.0
5	Hopea (Dipt, 1)	10	10.2	Diospyros (Eben, 3)	6	11.4	Canarium (Burs, 2)	8	13.1	Madhuca (Sapo, 2)	12	18.0
9	Madhuca (Sapo, 2)	9	10.1	Knema (Myri, 2)	4	11.3	Koilodepas (Euph, 1)	15	12.4	Gironniera (Ulma, 1)	٢	12.8
7	Croton (Euph, 2)	9	9.8	Koilodepas (Euph, 1)	10	10.1	Cleistanthus (Euph, 1)	10	11.9	Xanthophyllum (Poly, 2)	8	12.3
8	Cleistanthus (Euph, 2)	7	9.1	Palaquium (Sapo, 2)	9	9.7	Ixora (Rubi, 1)	11	10.2	Barringtonia (Lecy, 1)	ю	9.3
6	Fordia (Papi, 1)	11	9.1	Baccaurea (Euph, 2)	4	9.2	Xanthophyllum (Poly, 3)	5	10.1	Fordia (Papi, 1)	7	7.6
10	Chionanthus (Olea, 2)	10	8.4	Polyalthia (Anno, 3)	5	8.9	Alseodaphne (Laur, 1)	9	8.3	Mallotus (Euph, 2)	9	7.6

cars ago	pun %	10 years ago	Abun	%
dhuca malaccensis 2	2 25.5	Melastoma malabathricum	27	19.1
ygium tawahense	9 12.4	Syzygium tawahense	11	17.8
ilodepas brevipes 1	5 12.2	Madhuca malaccensis	11	16.5
istanthus 1 rycibifolius	0 11.7	Gironniera nervosa	٢	12.6
narium denticulatum	6 10.8	Barringtonia macrostachya	б	9.2
ra fucosa 1	1 10.1	Xanthophyllum obscurum	5	9.1
tica nitens	4 8.4	Shorea macroptera	10	8.0
eodaphne elmeri	6 8.2	Fordia splendidissima	٢	7.5
tica oblongifolia	7 7.3	Macaranga hypoleuca	5	7.3
rdia splendidissima	6.9 6.9	Macaranga gigantea	б	6.6
$\begin{array}{rrrr} ana & z & 0.0 & Va. \\ 6 & 6.3 & Fo. \\ 1), \text{ expressed as the densi} \\ 10 \text{ cm in height } \geq 1.5 \text{ m} \end{array}$	ma 2 0.0 Varies correction operations 6 6.3 Fordia splendidissima 1 , expressed as the densities of stems exceeding d 10 cm in height $> 1.5 m (%)$. In parentheses after f	c_{11} c_{22} c_{23} c_{33} c_{41} c_{41} c_{42} c_{41} c_{42} c_{43} c_{41} c_{42} c_{43} c_{43} c_{43} c_{44} c	ana = 2 0.0 Varies operation $b = 0.0$ Varies operation $b = 0.0$ Varies operation $b = 0.3$ Fordia splendidissima $b = 0.9$ Macaranga gigantea b), expressed as the densities of stems exceeding dbh < 10 cm in height > 1.5 m (%). In parentheses after families the numbers of observed genera ar	ana 2 0.0 varied consignation b 0.2 varied approximation b 0.3 Fordia splendidissima 6 6.9 Macaranga signatea 3 0.1, expressed as the densities of stems exceeding dbh < 10 cm in height > 1.5 m, and the percentage 10 cm in height > 1.5 m (%). In parentheses after families the numbers of observed genera and specie:

Seedlin	ngs-Primary forest	Abun	%	Selectively logged forest								
				1 year ago	Abun	%	5 years ago	Abun	%	10 years ago	Abun	%
Famili	es											
1 D	hipterocarpaceae (4, 7)	92	37.7	Euphorbiaceae (8, 9)	42	21.9	Dipterocarpaceae (5, 11)	50	23.5	Dipterocarpaceae (5, 19)	65	30.2
2 E	uphorbiaceae (7, 9)	47	24.7	Dipterocarpaceae (3, 5)	34	17.0	Euphorbiaceae (10, 11)	40	23.4	Melastomataceae (3, 3)	39	14.5
3 A	nnonaceae (4, 4)	20	11.3	Rubiaceae (8, 9)	22	15.3	Sapotaceae (2, 3)	29	15.8	Euphorbiaceae (11, 11)	17	12.2
4 P	'almae (4, 4)	14	9.5	Melastomataceae (5, 5)	24	12.3	Papilionaceae (1, 1)	22	14.3	Papilionaceae (1, 1)	17	11.4
5 E	benaceae (1, 4)	18	9.2	Lauraceae (6, 6)	12	9.1	Myrtaceae (1, 2)	22	13.5	Dryopteridaceae (1, 1)	25	9.7
6 N	felastomataceae (2, 5)	11	8.5	Dryopteridaceae (1, 1)	13	8.7	Guttiferae (2, 2)	12	9.4	Sapotaceae (1, 1)	15	8.4
7	leaceae (1, 2)	11	7.8	Sapotaceae (3, 3)	16	8.2	Connaraceae (3, 3)	6	6.9	Guttiferae (3, 3)	11	8.1
8 P	olygalaceae (1, 3)	8	6.1	Selaginellaceae (1, 1)	12	7.6	Melastomataceae (5, 6)	6	6.9	Rubiaceae (3, 3)	8	8.0
9 R	ubiaceae (4, 4)	7	5.8	Myristicaceae (3, 3)	10	6.2	Burseraceae (2, 3)	8	6.5	Oleaceae (1, 1)	15	7.6
10 S.	ymplocaceae (1, 1)	7	5.8	Anacardiaceae (5, 5)	7	6.0	Schizaeaceae (1, 1)	13	5.9	Anacardiaceae (3, 3)	12	6.7
Genera	I											
1 H	lopea (Dipt, 1)	68	24.6	Shorea (Dipt, 3)	32	15.1	Fordia (Papi, 1)	22	13.3	Shorea (Dipt, 10)	38	19.3
2 C	roton (Euph, 2)	22	11.9	Macaranga (Euph, 2)	20	9.8	Syzygium (Myrt, 2)	22	12.6	Melastoma (Mela, 1)	35	11.5
о С	iospyros (Eben, 4)	18	8.7	Nephrolepis (Dryo, 1)	13	8.2	Madhuca (Sapo, 2)	25	12.3	Fordia (Papi, 1)	17	10.4
4 X	oilodepas (Euph, 1)	12	7.6	Selaginella (Sela, 1)	12	7.2	Koilodepas (Euph, 1)	17	10.3	Nephrolepis (Dryo, 1)	25	9.3
5 C	hionanthus (Olea, 2)	11	7.2	Koilodepas (Euph, 1)	11	6.2	Calophyllum (Gutt, 1)	11	8.3	Madhuca (Sapo, 1)	15	7.8
6 D	aemonorops (Palm, 1)	×	5.7	Combretum (Comb, 1)	11	4.9	Shorea (Dipt, 5)	13	7.0	Chionanthus (Olea, 1)	15	7.1
7 P	ternandra (Mela, 3)	×	5.7	Diospyros (Eben, 2)	5	4.3	Vatica (Dipt, 2)	14	9.9	Dipterocarpus (Dipt, 4)	10	7.0
8 X	(anthophyllum (Poly, 3)	×	5.7	Lygodium (Schi, 1)	6	4.3	Hopea (Dipt, 1)	16	6.0	Lygodium (Schi, 2)	15	5.7
S S	ymplocos (Symp, 1)	7	5.4	Pternandra (Mela, 1)	8	3.9	Lygodium (Schi, 1)	13	5.6	Hopea (Dipt, 1)	6	5.4

See	llings-Primary forest	Abun	%	Selectively logged fores	ţ							
				1 year ago	Abun	%	5 years ago	Abun	%	10 years ago	Abun	%
10	Shorea (Dipt, 2)	11	5.3	Pleiocarpidia (Rubi, 1)	5	3.6	Dacryodes (Burs, 2)	7	5.0	Syzygium (Myrt, 2)	6	5.4
Spe	cies											
1	Hopea semicuneata	68	24.6	Shorea parvifolia	17	9.4	Fordia splendidissima	22	13.1	Melastoma malabathricum	35	11.4
6	Croton argyratus	21	11.6	Macaranga hypoleuca	17	8.8	Madhuca malaccensis	24	11.8	Fordia splendidissima	17	10.2
Э	Koilodepas brevipes	12	7.5	Nephrolepis bisserata	13	8.1	Syzygium tawahense	20	11.8	Nephrolepis bisserata	25	9.2
4	Chionanthus sp.1	6	6.0	Selaginella caulescens	12	7.1	Koilodepas brevipes	17	10.2	Madhuca malaccensis	15	7.6
5	Daemonorops sabut	8	5.7	Koilodepas brevipes	11	6.2	Calophyllum gracilipes	11	8.2	Chionanthus sp.1	15	7.0
9	Diospyros macrophylla	14	5.6	Combretum nigricans	11	4.9	Vatica nitens	13	6.2	Shorea macroptera	12	5.4
2	Symplocos crassipes	٢	5.4	Shorea leprosula	12	4.6	Hopea cernua	16	5.9	Hopea cernua	6	5.2
8	Uvaria elmeri	٢	4.1	Lygodium circinatum	6	4.2	Lygodium circinatum	13	5.6	Lygodium circinatum	13	5.1
6	Anaxagorea javanica	4	3.8	Pternandra rostrata	8	3.9	Gluta renghas	٢	4.9	Syzygium tawahense	7	4.7
10	Dipterocarpus acutangulus	8	3.7	Bauhinia diptera	٢	3.6	Alpinia galanga	6	4.3	Calophyllum gracilipes	9	4.4
In c	olumns after each taxon the co	orrespon	iding at	bundance (Abun), express	ed as the	e dens	ities of stems exceeding po	er subple	$t 2 \times 10^{-1}$	2 m, and the percentage of im	portant	value

index of all stems of seedlings exceeding in subplot 2 × 2 m (%). In parentheses after families the numbers of observed genera and species and after genera the corresponding family and the number of observed species

Table 8 continued

Family	Genus	Species	Life form	NO	NI	N2	N3	$\mathbf{S0}$	$\mathbf{S1}$	S2	S3
Trees (1.5 ha)											
Dipterocarpaceae	Hopea	сетиа	Dicot tree	0	0	21	10	0	0	15	6
Dipterocarpaceae	Hopea	pachycarpa	Dicot tree	2	2	3	0	2	2	3	0
Palmae	On cosperma	horidum	Palm tree	0	7	0	3	0	5	0	-
Saplings (0.375 ha)											
Euphorbiaceae	Macaranga	hypoleuca	Dicot tree	0	24	3	5	0	4	3	-
Rubiaceae	Uncaria	borneensis	Dicot liana	0	0	0	2	0	0	0	-
Rubiaceae	Uncaria	calophylla	Dicot liana	0	0	0	3	0	0	0	-
Rubiaceae	Uncaria	cordata	Dicot liana	0	9	0	0	0	1	0	0
Zingiberaceae	Costus	speciosus	Monocot herb	0	2	0	0	0	1	0	0
Seedlings (0.06 ha)											
Cyperaceae	Mapania	latifolia	Monocot grass	1	5	0	0	1	7	0	0
Cyperaceae	Scleria	terrestris	Monocot grass	2	3	0	0	1	2	0	0
Dipterocarpaceae	Hopea	semicuneata	Dicot tree	68	0	0	0	9	0	0	0
Ebenaceae	Diospyros	macrophylla	Dicot tree	14	0	0	0	2	0	0	0
Euphorbiaceae	Macaranga	hypoleuca	Dicot tree	0	17	1	1	0	5	1	-
Ophioglossaceae	Helminthostachys	zeylanica	Fern tree	0	3	0	0	0	3	0	0
Selaginellaceae	Selaginella	caulescens	Fern herb	0	12	0	0	0	5	0	0

 $\stackrel{{}_{\scriptstyle{\frown}}}{\underline{\bigcirc}}$ Springer

Annonaceae, Palmae and Ebenaceae (Table 8). The main representing these families were *H. semicuneata* and *Dipterocarpus acutangulus* (both Dipterocarpaceae), *C. argyratus* and *K. brevipes* (both Euphorbiaceae), *Uvaria elmeri* (Annonaceae), *Daemonorops sabut* (Palmae) and *Diospyros macrophylla* (Ebenaceae).

There were significantly more small dicot liana seedlings in the forest sites logged 1 and 10 years ago, compared with the primary forest site (Table 5). Small dicot lianas were the only of species *Combretum nigricans* (Table 8) in the 2×2 m subplots. This species was very abundant in the forest site logged 1 year ago but was absent in the primary forest site. Seedlings of the species of *Strychnos axillaris* was present in the selectively logged forest sites, but were not observed in the primary forest site. Small dicot trees were more abundant in the primary forest site but not in the three logged ones (Table 5).

Dicot shrubs were more abundant in the forest site logged 1 year ago than in the other forest sites (Table 5). Dicot species were typically shrubs belonging to the genera of the Rubiaceae: *Ixora* and *Psychotria*. The first genus consists of species that were mainly restricted to the primary forest site, while *Psychotria* sp.1 was most abundant in the forest site logged 1 year ago.

Monocot herbs were abundant in the forest site logged 5 years ago (Table 5) and this was mainly due to extensive ground cover of *Alpinia galanga* (Table 8), which was absent or rare in the other three forests.

Grass-like monocots were more abundant in the forest site logged 1 year ago (Table 5), mainly due to plentiful ground cover of *Mapania latifolia* and *Scleria terrestris* (Table 9). Both species were absent in two other logged forest sites.

Both palm lianas (rattans) and small palm trees were more abundant in the primary forest site than in the three selectively logged forest sites (Table 5). The climbing palm D. *sabut*, was the only climber that was very abundant in the primary forest site (Table 8). Other species were rare in all four forest sites.

Small liana ferns were more abundant in the forest site logged 5 year ago than in the other forest sites (Table 5). Two genus *Lygodium* and *Stenochlaena*, were abundant in the forest site logged 5 years ago, while in the other three forest sites these genera were absent or rare. These genera often form a very tangled mat at places where the small pioneer trees and small lianas are absent or rare.

Both herbaceous and small tree ferns were abundant in selectively logged forest sites, but were absent in the primary forest site (Table 5). Small tree ferns were represented solely by *Helmintostachys zeylanica*, a species that was only observed in the forest site logged 1 year ago. Of the same life form, the species *Selaginella caulescens* was also very abundant in the forest site logged 1 year ago, but absent in the three other forest sites (Table 9).

Discussion

Changes in forest structure and composition after logging compared to primary forest

Biodiversity levels indicate the stability of a forest community: the higher the levels of biodiversity, the more stable the community (Richards 1964; Whitmore 1990). In our study we compared forest sites which were logged 1, 5 and 10 years ago and a primary forest site. Our study covered canopy and forest floor vegetation, trees, saplings and seedlings, climbing trees (liana and rattan), non-rattan palms, herbs, epiphytes, and mosses.

Since we included 10 years ago logged forests, a before-, between- and after treatment study was not possible. We also did not include a heavily logged forest site, because these were not available within the forest concession. The distance of 50 m between the five transects per site is limited (risk of pseudo replication), but this distance has been used in other studies and is proposed as a standard by several authors (Slik et al. 2002). The present comparison of the four forest sites: three forest sites logged selectively 1, 5 and 10 years ago (certified logging) and a primary forest site provides important information on the status of logged lowland rain forest in East Kalimantan. The logging procedures of our study sites are described in detail (Meijaard et al. 2005). The present study showed that overall tree densities were significantly higher in the forest site of primary forest than in the sites 1 and 5 years ago, but approximately similar to that in the forest site logged 10 years ago (Table 3). The number of tall trees (dbh ≥ 10 cm) is very similar in the three logged forest sites of Labanan (492-558 stems ha⁻¹), suggesting that the impact of selective logging was similar for the three sites (Table 2). Nevertheless, we found small though significant differences in stem densities in the forest sites logged 1 and 5 years ago compared with the primary forest site (Table 3). In addition, typical sapling and seedling life forms, such as the fast growing species *M. hypoleuca* were very abundant in the forest site logged 1 year ago less abundant in the forest sites 5 and 10 years ago, and absent in the primary forest site (Table 9). Macaranga hypoleuca is a very common and characteristic pioneer tree species in most of Southeast Asia and especially in East Kalimantan (Primack and Lee 1991; Davies et al. 1998; Slik et al. 2000, 2002; Eichhorn et al. 2006). Without information on species composition and diversity, a comparison of the vegetation structure in the four forests sites therefore suggested that selective low impact logging largely compensated for the strong negative impact of initial logging. However, we found that only a small number of pioneer tree species accounted for the high small tree densities in the forest site 10 years ago (Table 5). Our result therefore show that abundance of tree species regeneration, as was observed at several sites in East Kalimantan (e.g. Siegert et al. 2001; Slik et al. 2002; Yassir et al. 2010), does not in itself, ensure recovery of the forest's original botanic diversity. Information on species composition is needed to know how many species of trees and other plants are able to recover in the selectively logged forest sites (Tables 3, 4 and 5).

The total number of saplings species were belonged to one or a few different species in the selectively logged forest sites (Table 4); this means that new trees of sapling which entered the overstorey (dbh \geq 10 cm), whereas the in-growth of seedlings in the selectively logged forests consisted of many species (Table 5); this was apparently due to more light on the open places to grow the pioneer species (Arbainsyah, pers. obs.). This finding doesn't contradict the generally held view that pioneer species occur only after disturbance when the light or temperature levels are raised substantially (Bazzaz and Pickett 1980; Uhl and Clark 1983; Swaine and Whitmore 1988; Vazquez-Yanes and Orozco-Segovia 1993; Eichhorn et al. 2006). After selective logging of the forest, many small pioneer tree seedlings were therefore likely to be available for tree establishment and this would explain why in a forest site, which was logged previously, seedlings could become dominant. This explanation implies that after selectively logging, there will not be a permanent deforestation at Labanan, as ingrowth of seedlings of small trees in the topsoil were most abundant in the forest site logged 10 years previously (Fig. 2).

10 years after selective logging, the sapling and tree densities were still high in the forest sites, but rather low in the forest sites logged 1 and 5 years ago, compared with the primary forest site. Apart from the differences in tree densities, there was also a difference in tree composition among the selectively logged forest sites (1, 5 and 10 years ago). The



Fig. 4 Diameter at breast height (average + standard deviation) of four species abundant in three logged forest sites (*open bars*) and of four species abundant only in forest site logged 10 years ago (*solid bars*)

most abundant tree species in the forest site logged 10 years ago were *M. malaccensis*, *A. borneensis* and *S. tawahense* (Table 6). *S. tawahense* also belongs to the highest 10 most abundant tree species of forest sites logged 1 and 5 years ago (Table 6; Fig. 4).

The impact of selective logging on plant diversity

Comparison of the plant diversity in different habitats shows that the overall impact of logging on plant species richness is highly dependent on the scale of assessment. We realize that the maximum number of 156 tree species we found in our study does represent of the tree species diversity in Borneo (Raes 2009). Since most species in hyperdiverse rainforests occur in low densities, their response to logging can not be assessed with small sampling plots only. So we suggest that our study at best, gives an indication of an impact of logging on the more common tree species. We also realize that the tree category >10 cm dbh will include a wide range of diameter classes, which may have changed following logging. Therefore we recommend to include more diverse diameter categories in a follow up study (Tables 3, 4 and 5). Logging in a sense is equivalent to gap formation in the forest canopy, but at a much larger scale and will surely alter the nature of the original forest (Kartawinata 1977; Eichhorn et al. 2006). Similar scale-dependent effects of disturbance on species diversity have also been reported by other studies, for example on the abundance of bark beetles in pine forests of Finland (Peltonen et al. 1998). In tropical rain forests, the impact of logging on species richness and evenness in butterflies has been shown to be highly scale-dependent as well (Hammer and Hill 2000; Cleary 2002; Summerville and Crist 2002). As a result, numbers of trees and small dicot trees were always higher in the primary forest site (Tables 3 and 6), this was mainly due to the abundant regeneration of invasive species such as *H. semicuneata* (Tables 7 and 8), this species is not logged and the mother tree was absent in the three logged forest sites (Table 9). For forest regeneration it is important that the high richness of saplings were found in the forest site logged 1 year ago (Table 4). When taxonomic composition was compared species diversity was higher in the in the forest site logged 1 year ago than in the primary forest site (Table 3). Our study showed that the abundance of Caesalpiniaceae was considerably decreased in the selectively logged forest sites relative to the primary forest site, whereas the abundance of Dipterocarpaceae, Sapotaceae and Euphorbiaceae increased in the selectively logged forest sites (Fig. 3). The latter family is typical fast-growing pioneer taxa throughout the tropics (Turner 2001) and are of little economic interest. Our results show that, despite attaining a height comparable to that of the primary forest site, selectively logged forest sites have different plant taxa (Tables 3, 4 and 5).

Total numbers of trees accounted for a higher overall plant diversity in primary forest sites, while the number of tree species was higher in the forest site logged 1 year ago (Table 3). Saplings had a more or less similar diversity in all forest sites, while the number of sapling species was higher in the primary forest site (Table 4). While numbers of ingrowth of seedlings was higher in the forest site logged 10 years ago, the number of seedling species was higher in the forest site logged 1 year ago (Table 5).

Conclusions

Overall, our study has revealed a rich natural vegetation on Borneo and major differences in the vegetation structure, composition and in plant diversity between selectively logged forest sites and a primary forest site. We have confirmed the importance of distinguishing between area of primary forest site and selectively logged forest sites for documenting and interpreting plant species richness for sustainable forest in tropical rain forest. Selectively logged rain forest of this study still showed high regenerating diversity of plant.

Logging practices in selectively logged forest with normal management operations have not resulted in a high deforestation of the study sites. The tree numbers still recovered with abundant regeneration. The numbers of tree species composition were clearly affected neither and increased nor decreased within forest sites logged 5–10 years ago.

Composition of saplings species were fewer within in selectively logged forest sites, in the 5–10 years ago-logged sites than in primary forest site. This indicates that the selectively logged forests have more or less the same value in evenness number species. These saplings will therefore slowly form a lower and lower proportion of all saplings present in selectively logged forest sites. That means that the impact of logging results in the same proportion of the total number of saplings in all selectively logged forest sites in near future.

The ingrowth of seedlings of the *M. hypoleuca* was found to be independent of the light availability in the forest understorey. Instead the number of this species depended strongly on the presence of mature parent trees species in and around the forest sites. This caused one species (*M. hypoleuca*) to be the most dominant species in the 1 year ago logged forest site. In the 5 and 10 year ago forest sites *M. hypoleuca* seedlings were probably replaced by competition with seedlings from other species. However, apart from seedling numbers, did depend strongly on the light availability in the forest understorey.

The diversity index used as indicators the stability regeneration for all growth stage of forest community showed that were still floristically very diverse and indicated that the selectively logged forest sites effected the abundance of species rather than species richness itself. This renders the selectively logged forest still valuable for conservation, especially since the studied forests were in selectively logged forest sites and tree species diversity to be higher in diversity of plant.

Acknowledgments We would like to thank Tien Wahyuni (B2PD, Samarinda), for the information of PhD Louwes fellowship to study in University of Leiden, the Netherlands. We would also like to thank Irsal Yasman, Joni Mujiono, Pudja Satata, Director of the PT. Inhutani and Sewoko Priyoyudoko, Rajudin Abdul Rahman, Director of the PT. Hutansanggam Labanan Lestari, for their support and permission to use the field station in Labanan. The head of BPTKSDA, Nur Sumedi, The head of Herbarium Wanariset Samboja, Kade Sidiyasa, Zainal Arifin are thanked for their help with plant identifications and for the opportunity to use the material housed in the herbarium. We would also like to acknowledge M.C. Roos, ter Stage H (NHN, Leiden), C.J.M (Kees) Muster, M. (Merlijn) van Weerd (CML, Leiden), K.A.O. Eichhorn (Bosflora, Utrecht), Wawan Gunawan, Ishak Yassir, Tri Atmoko (BPTKSDA, Samboja) and Amiril Saridan (B2PD, Samarinda) for their many discussion on methodology and statistical analysis. The fieldwork would have been impossible without the help of many people from Berau and Samboja who assisted him. We would especially like to mention Pujiansyah, Anto, Mahmud, Sugito and Prapto for their great assistance in the field. This study was supported under the umbrella of a LOUWES fellowship grant.

References

- Ashworth L, Aquilar R, Galetto L, Aizen MA (2004) Why do pollination generalist and specialist plant species show similar reproductive susceptibility to habitat fragmentation? J Ecol 92:717–719
- Bazzaz FA, Pickett STA (1980) Physiological ecology of tropical succession: a comparative review. Annu Rev Ecol Syst 11:287–310
- Beier P, van Drielen M, Kankam BO (2002) Avifaunal collapse in West African forest fragments. Conserv Biol 16:1097–1111
- Benitez-Malvido J, Martinez-Ramos M (2003) Impact of forest fragmentation on understory plant species richness in Amazonia. Conserv Biol 17:389–400
- Boulinier T, Nichols JD, Hines JE, Sauer JR, Flather CH, Pollock KH (2001) Forest fragmentation and bird community dynamics: inference at regional scales. J Ecol 82:1159–1169
- Brook BW, Sodhi NS, Ng PKL (2003) Catastrophic extinctions follow deforestation in Singapore. Nature 424:420–423
- Brown S, Lugo AE (1990) Tropical secondary forests. J Trop Ecol 6:1-32
- Bruna EM (2003) Are plant populations in fragmented habitats recruitment limited? Test with an Amazonian herb. Ecology 84:932–947
- Bruna EM, Nardy O, Strauss SY, Harrison S (2002) Experimental assessment of *Heliconia acuminata* growth in a fragmented Amazonian landscape. J Ecol 90:639–649
- Cleary DFR (2002) Biodiversity and environmental change in the rainforests of Borneo. PhD thesis, University of Amsterdam
- Connell JH (1978) Diversity in tropical rain forests and coral reefs. Science 199:1302–1310
- Curran LM, Caniago I, Paoli GD, Astianti D, Kusneti M, Leighton M, Nirarita CE, Haeruman H (1999) Impact of El Niño and logging on canopy tree recruitment in Borneo. Science 286:2184–2188
- Curran LM, Trigg SN, McDonald AK, Astiani D, Hardiono YM, Siregar P, Caniago I, Kasischke E (2004) Lowland forest loss in protected areas of Indonesian Borneo. Science 303:1000–1003
- Davies SJ, Palmiotto PA, Ashton PS, Lee HS, Lafrankie JV (1998) Comparative ecology of 11 sympatric species of Macaranga in Borneo: tree distribution in relation to horizontal and vertical resource heterogeneity. J Ecol 86:662–673
- de Iongh HH, Persoon G (2010) Monitoring the impact of certification. ETFRN News 51:48-50
- de Iongh HH, van Weerd M (2006) The use of avian guilds for the monitoring of tropical forest disturbance by logging. Tropenbos, Wageningen
- Eichhorn KAO, Eichhorn LS, Arbainsyah du Pon I (2006) Plant diversity after rain-forest fires in Borneo. Structure, composition and diversity of plant communities in burnt and unburnt rain forest. Blumea J Plant Taxon Plant Geogr 18:37–64
- Fredericksen TS, Mostacedo B (2000) Regeneration of timber species following selection logging in a Bolivian tropical dry forest. For Ecol Manag 131:47–55
- Ghazoul J, Hellier A (2000) Setting limits to ecological indicators of sustainable tropical forestry. Int For Rev 2:243–253
- Hammer KC, Hill JK (2000) Scale-dependent effects of habitat disturbance on species richness in tropical forests. Conserv Biol 14:1435–1440
- Hartshorn G, Bynum N (2001) Tropical forest synergies. Nature 404:493-495
- Jennings SB, Brown ND, Bshier DH, Whitmore TC, Loves do CA (2001) Ecology provides a pragmatic solution to the maintenance of genetic diversity in sustainably managed tropical rain forests. For Ecol Manag 154:1–10
- Jost L (2006) Entrophy and diversity. Oikos 113:363-375
- Kartawinata K (1977) Biological changes after logging in lowland Dipterocarp forest. Herbarium Bogoriensis, Lembaga Biologi Nasional, LIPI, Bogor, Indonesia

- Kartawinata K, Vayda AP (1984) Forest conversion in East Kalimantan, Indonesia: the activities and impact of timber companies, shifting cultivators, migrant pepper-farmers, and others. In: Castri FD, Baker FWG, Hadley M (eds) Ecology in practice, part 1: Ecosystem management. Ticooly International Publications, Dublin, pp 99–126
- Kuswandari R (2004) Assessment of different methods for measuring the sustainable of forest management. International Institute for Geo-Information Science and Earth Observation, Enschede
- Laurance WF (1998) A crisis in the making: responses of Amazonian forests to land use and climate change. Trends Ecol Evol 13:411–415
- Laurance WF, Delamonica P, Laurance SG, Vasconcelos HL, Lovejoy TE (2000) Rainforest fragmentation kills big trees. Nature 404:836
- Ludwig JA, Reynold (1988) Statistical ecology. Wiley Interscience Publ. John Wiley and Sons, Toronto
- Mantel S, Tyrie GR, Oosterman A (2002) Exploring sustainable land use options for district planning in the Berau regency, Indonesia. International Soil Reference and Information Center, Wageningen
- Meijaard E, Sheil D, Nasi R, Augeri D, Iskandar B, Rosenbaum D, Setyawati T, Lammertink M, Rachmatika I, Wong A, Soehartono T, Stanley S, O'Brien T (2005) Life after logging. Reconciling wildlife conservation and production forestry in Indonesia Borneo. CIFOR. Indonesia
- Mueller-Dombois D, Ellenberg H (1974) Aims and methods of vegetation ecology. John Wiley and Sons, New York
- Parthasarathy N (1999) Tree diversity and distribution in undisturbed and human impacted sites of tropical wet evergreen forest in southern Wastern Ghats, India. Biodivers Conserv 8:1365–1381
- Peltonen M, Heliovaara K, Vaisanen R, Keronen J (1998) Bark beetles diversity at different spatial scales. Ecography 21:510–517
- Pimm SL, Raven P (2000) Extinction by numbers. Nature 403:843-845
- Pinard MA, Barker MG, Tay J (2000) Soil disturbance and post-logging forest recovery on bulldozer paths in Sabah, Malaysia. For Ecol Manag 130:213–225
- Primack RB, Lee HS (1991) Population dynamics of pioneer (Macaranga) trees and understorey (Mallotus) trees (Euphorbiaceae) in primary and selectively logged Bornean rain forests. J Trop Ecol 7:439–458
- Raes N (2009) Borneo. A quantitative analysis of botanical richness, endemicity and floristic regions based on herbarium records. PhD thesis, Nationaal Herbarium Nederland, University Leiden Branch
- Richards PW (1952) The tropical rain forest. An ecologically study. Cambridge University Press, Cambridge
- Richards PW (1964) The tropical rain forest. Cambridge University, New York
- Roxburgh SH, Shea K, Wilson JB (2004) The intermediate disturbance hypothesis: patch dynamics and mechanisms of species coexistence. The ecological society of America. Ecology 8(2):359–371
- Sheil D, van Heist M (2000) Ecology for tropical forest management. Int For Rev 2:261–270
- Sidiyasa K, Zakaria, Ramses I (2006) The forests of Setulang and Sengayan in Malinau, East Kalimantan: their potential and the identification of steps for their protection and sustainable management, Bogor, Indonesia: Center for International Forestry Research (CIFOR)
- Siegert F, Rűcker G, Hinrich A, Hoffmann AA (2001) Increased damage from fire in logged forest during droughts caused by El-Niño. Nature 414:437–440
- Sist P, Fimbel L, Sheil D, Nasi R, Chevallier MH (2003) Towards sustainable management of mixed dipterocarps forests of Southeast Asia: moving beyond minimum diameter cutting limits. Environ Conserv 30(4):364–374
- Skole D, Tucker C (1993) Tropical deforestation and habitat fragmentation in the Amazon: satellite data from 1978 to 1988. Science 260:1905–1910
- Slik JWF (2001) Macaranga and Mallotus (Euphorbiaceae) as indicators in the lowland dipterocarp forests of East Kalimantan. PhD dissertation. Tropenbos-Kalimantan Series 4. Wageningen: Tropenbos
- Slik JWF, van Balen S (2006) Bird community changes in response to single and repeated fires in a lowland tropical rainforest of eastern Borneo. Biodivers Conserv 15:4425–4451
- Slik JWF, Priyono, van Welzen PC (2000) Key to the Macaranga and Mallotus species (Euphorbiaceae) of East Kalimantan (Indonesia). The Gardens' Bulletin Singapore 52:11–87
- Slik JWF, Verburg RW, Keβler PJA (2002) Effects of fire and selective logging on the tree species composition of lowland dipterocarp forest in East Kalimantan, Indonesia. Biodivers Conserv 11:85–98
- Summerville KS, Crist TO (2002) Effects of timber harvest on forest Lepidoptera: community, guild, and species responses. Ecol Appl 12:820–835
- Swaine MD, Whitmore TC (1988) On the definition of ecological species groups in tropical rain forests. Vegetation 75:81–86
- Turner IM (2001) The ecology of trees in the tropical rainforest. Cambridge University Press, Cambridge Uhl C, Clark K (1983) Seed ecology of selected Amazon basin successional species. Bot Gaz 144:407–419

- van Kuijk M, Putz FE, Zagt RJ (2009) Effect of forest certification on biodiversity. Wageningen: Tropenbos International, p. 94. www.tropenbos.org/image/Tropenbos/publications_TBI_certification_and_biodiversity. pdf
- van Nieuwstadt MGL (2002) Trial fire. Postfire development of a tropical dipterocarp forest. PhD thesis, Utrecht University, the Netherlands
- Vazquez-Yanes C, Orozco-Segovia A (1993) Patterns of seed longevity and germination in the tropical rain forest. Annu Rev Ecol Syst 24:69–87
- Webb EL (1998) Gap-phase regeneration in selectively logged lowland swamp forest, Northeastern Costa Rica. J Trop Ecol 14:247–260

Whitmore TC (1984) Tropical rain forest of the far east, 2nd edn. Clarendon Press, Oxford

Whitmore TC (1990) Tropical rain forest. An introduction. Clarendon Press, Oxford

- Wilson JB (1990) Mechanisms of species coexistence: twelve explanations for Hutchinson's "paradox of the plankton": evidence from New Zealand plant communities. N Z J Ecol 13:17–42
- Woods P (1989) Effects of logging, drought, and fire on structure and composition of tropical forests in Sabah, Malaysia. Biotropica 21(4):290–298
- Yassir I, van der Kamp J, Buurman P (2010) Secondary succession after fire in Imperata grasslands of East Kalimantan, Indonesia. Agric Ecosyst Environ 137:172–182