

A Patchy Growth via Successive and Simultaneous Cambia: Key to Success of the Most Widespread Mangrove Species *Avicennia marina*?

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- **Background and Aims** Secondary growth via successive cambia has been intriguing researchers for decades. Insight into the mechanism of growth layer formation is, however, limited to the cellular level. The present study aims to clarify secondary growth via successive cambia in the mangrove species *Avicennia marina* on a macroscopic level, addressing the formation of the growth layer network as a whole. In addition, previously suggested effects of salinity on growth layer formation were reconsidered.
- **Methods** A 1-year cambial marking experiment was performed on 80 trees from eight sites in two mangrove forests in Kenya. Environmental (soil water salinity and nutrients, soil texture, inundation frequency) and tree characteristics (diameter, height, leaf area index) were recorded for each site. Both groups of variables were analysed in relation to annual number of growth layers, annual radial increment and average growth layer width of stem discs.
- **Key Results** Between trees of the same site, the number of growth layers formed during the 1-year study period varied from only part of a growth layer up to four growth layers, and was highly correlated to the corresponding radial increment (0–5 mm year⁻¹), even along the different sides of asymmetric stem discs. The radial increment was unrelated to salinity, but the growth layer width decreased with increasing salinity and decreasing tree height.
- **Conclusions** A patchy growth mechanism was proposed, with an optimal growth at distinct moments in time at different positions around the stem circumference. This strategy creates the opportunity to form several growth layers simultaneously, as observed in 14 % of the studied trees, which may optimize tree growth under favourable conditions. Strong evidence was provided for a mainly endogenous trigger controlling cambium differentiation, with an additional influence of current environmental conditions in a trade-off between hydraulic efficiency and mechanical stability.

Key words: *Avicennia marina*, cambial marking, mangrove, phloem, salinity, secondary growth, successive cambia, xylem.

INTRODUCTION

One of the curious mechanisms of plant growth is secondary wood formation via successive cambia (Wheeler *et al.*, 1989). This means that each growth layer is formed by a new cambium and hence is composed of both xylem and phloem tissue. The cambial variant is especially present in lianas (e.g. Nair, 1993; van Veenendaal and den Outer, 1993; Carlquist, 1996, 1999a; Jacques and De Franceschi, 2007), but is also reported in woody shrub and tree species, with descriptions of the wood anatomy (McDonald, 1992; Carlquist and Gowans, 1995; Carlquist, 1996, 1999b, 2002, 2003a, b) and wood formation (Wheat, 1977; Zamski, 1979, 1981; Fahn and Zimmermann, 1982; Nair and Mohan Ram, 1990; Terrazas, 1991; den Outer and van Veenendaal, 1995; Rajput and Rao, 1999; Carlquist, 2004; Schmitz *et al.*, 2007). However, it remains unclear if the growth layers, originating from the combined action of the several cambia, are formed periodically or continuously, and which factors trigger the formation of a new growth layer.

The mangrove genus *Avicennia* belongs to this exclusive group of trees showing secondary growth via successive cambia (Schmitz *et al.*, 2007). Early studies speculated on the periodicity of the conspicuous pattern of light- and dark-coloured bands resulting from the alternating xylem and phloem tissue (Baker, 1915; Chapman, 1944, 1947; Gill, 1971). A preliminary study on six *Avicennia marina* trees in Gazi Bay, Kenya showed the non-annual nature of their growth layers (Schmitz *et al.*, 2007). However, this study suggests that within one site a constant number of growth layers may form per year. While only half a growth layer was formed in a site of high salinity, an average of three growth layers was formed in a site with relatively lower salinity (Schmitz *et al.*, 2007). Therefore, the formation of growth layers seems to be influenced by local environmental factors such as soil water salinity rather than by seasonal climate. In addition, the proportion of phloem tissue increased slightly in parallel with the salinity of the site (Schmitz *et al.*, 2007). In agreement with these findings, Carlquist (2001) and Fahn and Shchori (1967) assume that dispersed phloem tissue over the entire stem offers a functional advantage to trees growing under xeric conditions.

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Avicennia tree species tolerate a wide range of soil water salinity levels, related to different inundation frequencies and evaporation of the substrate (Matthijs *et al.*, 1999; Dahdouh-Guebas *et al.*, 2004; Ye *et al.*, 2005; Naidoo, 2006; Sobrado and Ewe, 2006). To study the formation of growth layers, a technique with cellular resolution is offered by mechanically wounding the cambium, leaving a datable scar (Kuroda and Shimaji, 1984). Thanks to the distinct time mark, the cambial mark, the presence of clear tree rings becomes superfluous in studies of wood formation, making the method especially useful for the study of tropical tree species (Détienne, 1989; Nobuchi *et al.*, 1995; Sass *et al.*, 1995; Jalil *et al.*, 1998; Bauch and Dünisch, 2000; Ohashi *et al.*, 2001; Heinrich and Banks, 2002), including mangrove trees (Shiokura, 1989; Verheyden *et al.*, 2004; Schmitz *et al.*, 2007).

In the present study, the cambium of 80 *A. marina* trees was wounded. The trees were sampled from seven sites in Gazi Bay and one more distant location in Dabaso, both situated in Kenya. The aim was to clarify the secondary growth mechanism of *A. marina* characterized by

successive cambia. This study tested the hypotheses that (a) growth layer formation is periodic, with the number of growth layers formed per year depending on the soil water salinity of the site; and (b) the width of the growth layers decreases with increasing soil water salinity.

MATERIALS AND METHODS

Study sites and sample collection

Wood samples were collected in Gazi and Dabaso, two locations along the Kenyan coast. The mangrove forest of Gazi Bay (39°30'E, 4°25'S) covers about 710 ha (UNEP, 2001) and is situated approx. 50 km south of Mombasa. Sampling was performed in seven sites (Table 1), differing in local environmental conditions (Table 2). For comparison, an additional set of trees was sampled in Dabaso (39°21'E, 03°59'S), in the 1500 ha large mangrove forest of Mida Creek situated about 100 km north of Mombasa (Gang and Agatsiva, 1992). In the rainy season of 2005 and 2006 (May–June) and the dry season of 2006 and

TABLE 1. Studied stem discs and outermost wood blocks of *Avicennia marina* with the corresponding tree characteristics for the eight study sites at two locations in Kenya, Gazi and Dabaso

Location	Accession no.		Tree height (m)	R ₁₃₀ * (cm)	R _{base} † (cm)
	Discs	Samples			
Gazi					
Site 1	Tw58910–13, Tw58927–31	T72, T76, T85	11 (7–14)	4 (1–13)	43 (20–100)
Site 2	Tw58905–09, Tw58926, Tw58988	T40–41	3 (2–4)	2 (1–3)	9 (5–17)
Site 3	Tw58916–18, Tw58933–35	T26–28	2 (2–4)	1 (1–3)	7 (3–11)
Site 4	Tw58943–48, Tw58987	T14–16	6 (4–8)	3 (2–5)	17 (5–54)
Site 5	Tw58936–42	T45, T51, T56	6 (6–6)	3 (1–5)	7 (3–10)
Site 6	Tw58900–04, Tw58932	T35, T37	4 (2–5)	3 (1–4)	6 (6–14)
Site 7	Tw58920–25, Tw58989	T19–22	6 (3–10)	4 (1–9)	22 (4–44)
Dabaso					
Site 8	–	Tw58950–61	10 (6–14)	11 (6–16)	36 (16–100)

Values are means with ranges in parentheses.

* Stem radius at 130 cm height.

† Stem radius at the base of the tree.

TABLE 2. Stand characteristics of seven study sites in Gazi Bay and one study site in Dabaso, Kenya

Location	Soil water nutrients (10 ⁻³ μmol cm ⁻³ soil)				Soil texture	Soil water salinity (‰)		
	NO ₃ ⁻	NH ₄ ⁺	P	LAI		Minimum	Maximum	Inundation class
Gazi								
Site 1	0.5 (0.2–0.6)	3 (0–5)	1.4 (0.9–1.8)	1.42	Silty clay	20.9	46.0	1
Site 2	0.3 (0.2–0.5)	*	0 (0–0)	0.25	Sandy loam	40.0	70.0	2
Site 3	0.15 (0.10–0.24)	*	0.2 (0.1–0.3)	0.23	Loamy sand	38.0	86.0	3
Site 4	0.3 (0.0–0.4)	1 (0–4)	2 (0–4)	1.29	Loamy sand–sandy loam	5.0	68.2	3
Site 5	2 (0–6)	10 (7–14)	0.1 (0–0.3)	1.18	Clay loam	40.0	79.9	3
Site 6	1 (0–4)	0.5 (0.1–1.1)	0.1 (0–0.3)	0.62	Sandy loam	10.0	90.0	4
Site 7	23 (0–68)	*	2 (0–4)	1.73	Loamy sand–sandy loam	10.0	48.0	4
Dabaso								
Site 8	*	*	*	*	*	33.0	72.2	3

Soil water for nutrient and salinity analyses was taken at about 25 cm depth. Inundation classes are according to Tomlinson (1994). Values are means with range in parentheses.

* No data records.

2007 (February–March) soil water was collected at approx. 25 cm depth with a punctured plastic tube connected to a vacuum pump. At each site, 1–3 salinity measurements were carried out with a hand-held refractometer (ATAGO, Tokyo, Japan). One day after spring tide in February 2007, soil water was collected in triplicate at the seven study sites in Gazi Bay. From these water samples, NO_3^- , NH_4^+ and P concentrations were measured by standard procedures (APHA-AWWA-WEF, 1995) and expressed relative to soil water content, which was calculated from the fresh and oven-dry weight. Soil texture was determined by standard field characterization methods (GLOBE, 2005). The height above datum was measured with tracing paper, placed at each site before high tide, and the corresponding inundation classes were calculated. According to Tomlinson (1994), inundation classes one, two, three and four correspond to an area being inundated by 100–76, 75–51, 50–26 and 25–5 %, respectively, of the high tides. The leaf area index (LAI), integrated over the zenith angle 0–75°, was calculated from hemispherical images using the software program Gap Light Analyzer v. 2.0 (Simon Fraser University, British Columbia and the Institute of Ecosystem Studies, New York).

The cambial marking was performed with a surgical needle of 1.2 mm diameter (Verheyden *et al.*, 2004) in May 2005 (wet season). In view of a potential time mark of the dry season by changes in wood anatomy, the trees were marked a second time in February 2006 (dry season). For two trees (Tw58955, -57) the cambium of both the stem and one branch was wounded. Of these 80 trees, 49 were felled in June 2006 and samples are now part of the Xylarium of the Royal Museum for Central Africa, Tervuren, Belgium (discs, Table 1). From the remaining 31 trees, wood samples were taken with a handsaw of the outermost wood comprising the cambial mark, and 19 of them were subsequently stored in FAA (formalin–acetic acid–alcohol). Before felling or sampling of the trees, the stem circumference at 130 cm height and the stem diameter at the base were measured. The height was calculated trigonometrically.

Climate description

The climate along the Kenyan coast is characterized by a bimodal distribution of precipitation. A distinct dry season (January–February) is followed by a long (April–July) and a short rainy season (October–November; Fig. 1). During the wet season, the rivers Mkurumuji and Kidogweni provide an important freshwater source for the mangroves of Gazi Bay (Kitheka, 1997). The average temperature at the Kenyan coast ranges from 22 to 30 °C, with a mean relative humidity of 65–81 % (annual averages of minima and maxima for Mombasa for the period 1972–2001, data from the Kenyan Meteorological Department, Mombasa, Kenya).

Sample preparation and microscopic analysis

In this study, the following terms and definitions are used: *phloem band* designates a zone of phloem strands united in a band of parenchyma tissue and bordered on

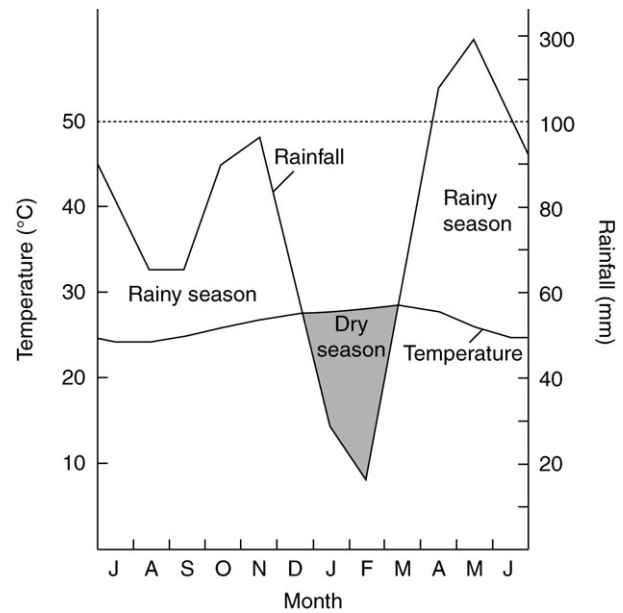


FIG. 1. Climate diagram of Mombasa (39°36'E, 4°0'S) adapted from Lieth *et al.* (1999), showing the long (April–July) and short (October–November) rainy season and one distinct dry season (January–February). The precipitation axis is reduced to one-tenth scale above the dotted horizontal line.

the outside by a layer of sclereids; *xylem band* designates the zone in between two phloem bands; and *growth layer* designates one ontogenetic unit of phloem and xylem (Figs 2 and 3).

To investigate the periodicity of growth layer formation, the number of growth layers formed in the period from 17–26 May 2005 to 9 June–9 July 2006 was counted from 79 cambially marked trees. Wood discs and wood samples were prepared as described by Schmitz *et al.* (2007). In brief, wood discs were sanded with a series of sandpaper from 100 to 1200 grit, while wood samples were dehydrated in an ethanol series and embedded in PEG1500. Transverse sections were made with a sliding microtome and double stained with safranin–fast green. The number of growth layers from the cambial mark onwards was counted using an Olympus BX 60 microscope. This was done at both sides of the wound and the growth layer count was averaged (Figs 2B, C and 3A). The growth layer network of one stem, Tw57798 (see also Schmitz *et al.*, 2007), was visualized with a CT scan (Fig. 4; CT-scan Brilliance 64 slice, Philips, The Netherlands).

In addition, the increment during the period of the cambial marking experiment was measured with the image analysis software AnalySIS Pro 3.2 (Soft Imaging System GmbH, Münster, Germany) via a camera connected to a microscope (Olympus BX60). The increment was measured at about 2 mm distance from the abnormal growth at the wound, which is the furthest position showing a cambial mark. For trees showing a double wound, resulting from two simultaneously dividing cambia, the increment was corrected for the part of the xylem tissue of the outermost growth layer already formed at the time of wounding (Figs 2C and 3A).

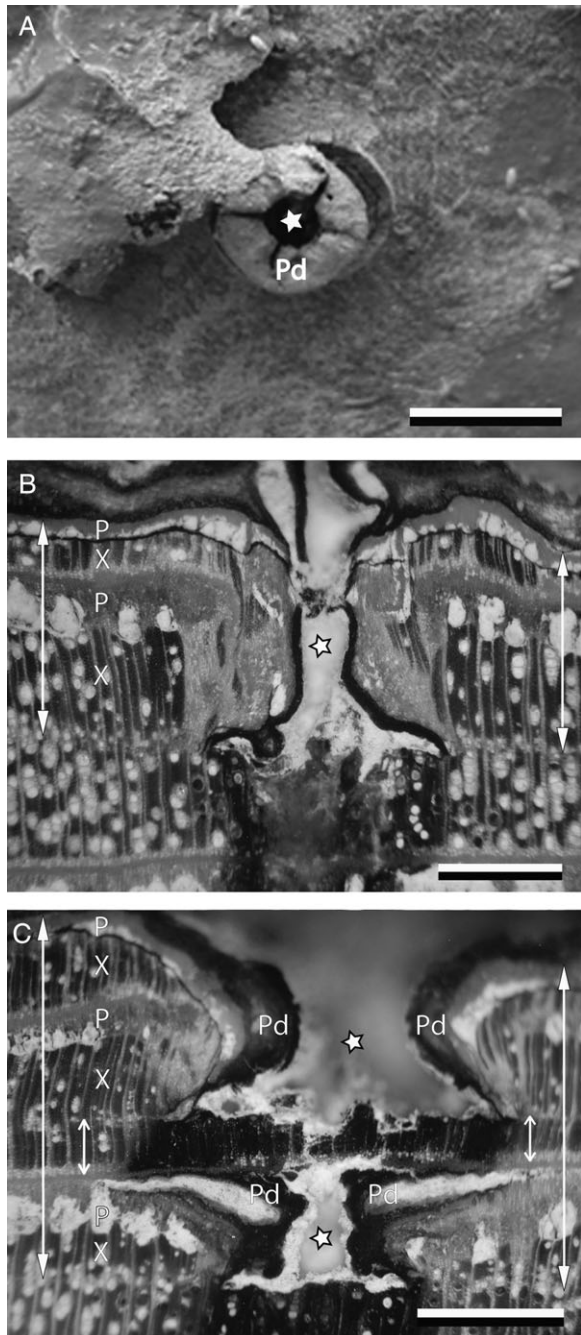


FIG. 2. Macro- and microscopic characteristics of *Avicennia marina* stem wood after wounding the cambium with a needle of 1.2 mm diameter. (A) Pinhole encircled by wound periderm, scale bar = 1 cm. (B, C) Detail of sanded stem discs showing (B) a decreased vessel density and vessel diameter in the zone around the pinhole, marking the time of cambial wounding, and (C) a double cambial wound illustrating the simultaneous formation of the two growth layers. Scale bars = 1 mm. P, phloem band; Pd, periderm; X, xylem band. Arrows, radial increment from May 2005 to June 2006. Small arrows, part of the growth layer already formed at the time of cambial wounding. Asterisks, pinhole.

On each of 48 stem discs, the width of the xylem and phloem bands of each growth layer was measured along three transects from pith to bark at the maximum, minimum and medium sized disc radius. Because of the

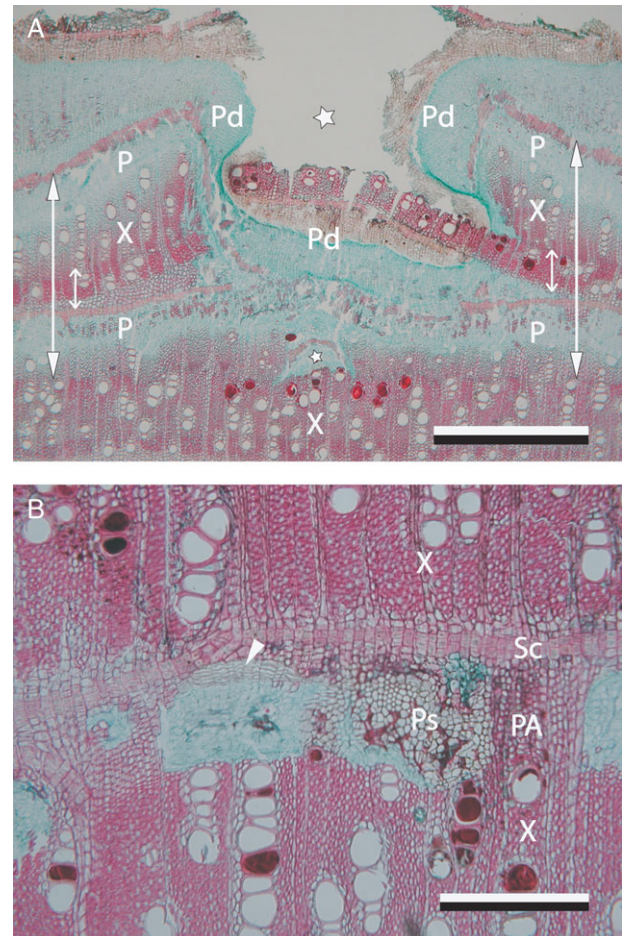


FIG. 3. Transverse microsections of the stem wood of *Avicennia marina* after wounding the cambium with a needle. (A) Microscopic wood structure after wounding of the two outermost cambia. Scale bar = 1 mm. (B) Detail of two phloem strands located inward from the pinhole showing a coating of the cells and an induced meristem (arrowhead) in the right and left phloem strand, respectively. Scale bar = 500 μm . P, phloem band; PA, parenchyma; Pd, periderm; Ps, phloem strand; Sc, sclereids; X, xylem band. Arrows, radial increment from February 2006 to June 2006. Small arrows, part of the growth layer already formed at the time of cambial marking. Asterisks, pinhole.

networking pattern of the growth layers, a pencil line was drawn and all growth layers crossing the line were measured.

Statistics

To quantify relationships between different tree and growth layer characteristics, Pearson correlation coefficients were calculated. Simple linear regressions were used to study relationships in growth increment with environmental and tree characteristics. One-way analyses of variance (ANOVAs) were used to test (a) the difference in radial increment between the different study sites and (b) the relationship between the average ring width on different sides of asymmetric stem discs and corresponding radii. A *t*-test for independent variables was performed to compare the growth increment at opposite radii of

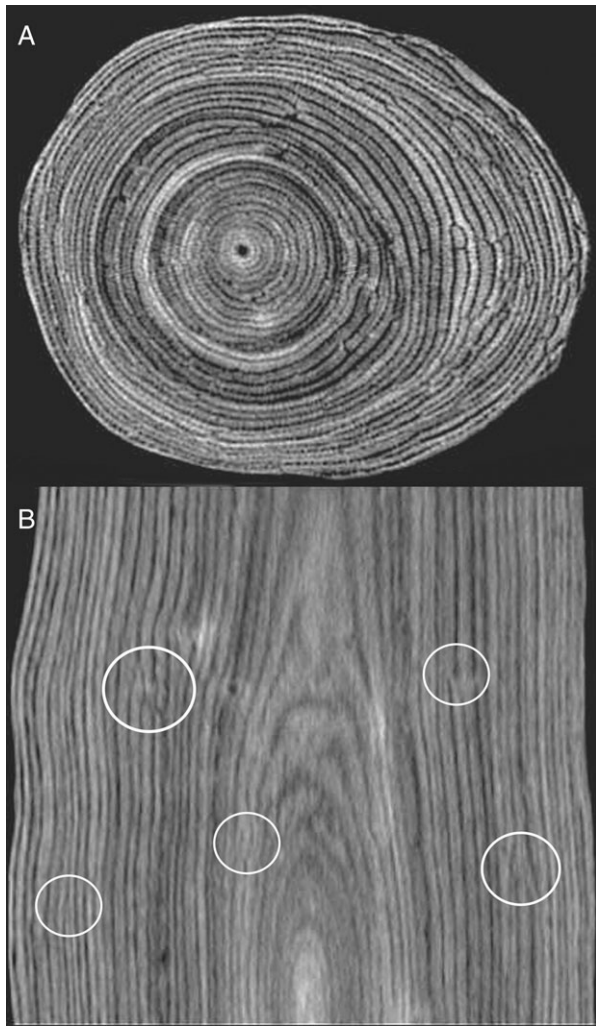


FIG. 4. Networking pattern of the growth layers in the (A) transverse and (B) longitudinal plane as visualized by a CT scan of a tree stem portion of 7 cm diameter (Tw57798, *Tervuren xylarium*). Circles indicate the positions of branching phloem bands.

asymmetric trees and between trees with and without signs of die-back (with separate variance estimates). To comply with the assumptions of a normal distribution of the data and homogenous variances, data were transformed with either logarithmic or square root functions. If transformations failed, Spearman R correlation coefficients were calculated (STATISTICA, StatSoft Inc.).

RESULTS

Wood anatomical description of the cambial mark

On a macroscopic level, the pinhole was regularly encircled by an extra periderm (Fig. 2A). Microscopically, the position of the cambium at the time of wounding could be determined from a variable combination of wood anatomical changes (Figs 2B and C, and 3). In agreement with earlier reports, the time mark was formed by an initial parenchyma band in 53% of the trees ($n = 80$) (Shiokura, 1989; Nobuchi *et al.*, 1995; Ohashi *et al.*, 2001; Stobbe

et al., 2002) and relatively smaller vessels (Nobuchi *et al.*, 1995; Schmitt *et al.*, 2000; Ohashi *et al.*, 2001) at a lower density (Carlquist, 2001) in an area of about 1 mm^2 on both sides of the pinhole. The canal created by wounding was closed by periderm (Carlquist, 2001; Stobbe *et al.*, 2002), with an extra internal wound periderm in 14% of the studied trees ($n = 79$) showing a double wound (Figs 2C and 3A). Regularly, a coating that stained pink with safranin was observed in the phloem strands inward from the canal caused by wounding (Fig. 3B). Suberin has been mentioned to coat parenchyma tissue of the reaction zone (Schmitt and Liese, 1993). It is known as a barrier-forming agent protecting against invading pathogens (Franke and Schreiber, 2007) and necrosis of the deeper lying tissues (Schmitt and Liese, 1993). In addition, in 53% ($n = 19$) of the microsections, the phloem strands showed a meristematic zone (Fig. 3B), corresponding to the finding in poplar of secondary phloem cells dedifferentiating into meristematic cambium cells (Frankenstein *et al.*, 2005; Frankenstein, 2006). Finally, a dark zone of protection wood (Carlquist, 2001) was observed inward from the wound canal (Fig. 2B, C) with partly occluded vessels and fibres (Schmitt *et al.*, 2000).

Annual increment

From only a tenth of a growth layer up to four completed growth layers was formed in the period of the cambial marking experiment from May 2005 to June 2006 (Fig. 5, Table 3). The number of growth layers formed during the 1-year study period was highly correlated with the corresponding radial increment ($r^2 = 0.73$, $n = 73$, $P < 0.0001$). Also at the level of the entire stem disc, the number of growth layers showed a strong correlation with its radius ($r^2 = 0.89$, $n = 141$, $P < 0.0001$). In agreement with this high correlation, the difference between the maximum and minimum disc radius of asymmetrical stem discs was correlated with the corresponding difference in number of growth layers ($r^2 = 0.37$, $n = 42$, $P < 0.0001$). Growth layer width increased from a minimum, through a medium to a maximum disc radius ($F = 50.79$, $n = 2663$, $P < 0.0001$; linear trend analysis: $F = 101.57$, $n = 2663$, $P < 0.0001$).

No significant difference was observed in the number of growth layers formed in the 1-year study period between the eight study sites (Fig. 5B; $F = 1.48$, $n = 79$, $P = \text{ns}$). By analysing different environmental factors separately, characterizing the different study sites, no or only very weak relationships were found with the radial increment of the study period (Table 4). Also stem radius at the sampling height of the trees and tree height showed a weak but significant correlation with the radial increment measured from the cambial markings onwards (Table 4). The growth rate during the 1-year study period was significantly larger in trees showing a double cambial mark, resulting from the additional radial increment of the last but one growth layer (Figs 2C and 3A; $t = -3.2$, $n_1 = 62$ and $n_2 = 11$, $P < 0.01$). The cambium of this inner growth layer produced on average $249 \pm 105 \mu\text{m}$ ($n = 10$) of xylem tissue during the 1-year study period (Table 3). Within trees Tw58955 and Tw58957, radial growth was

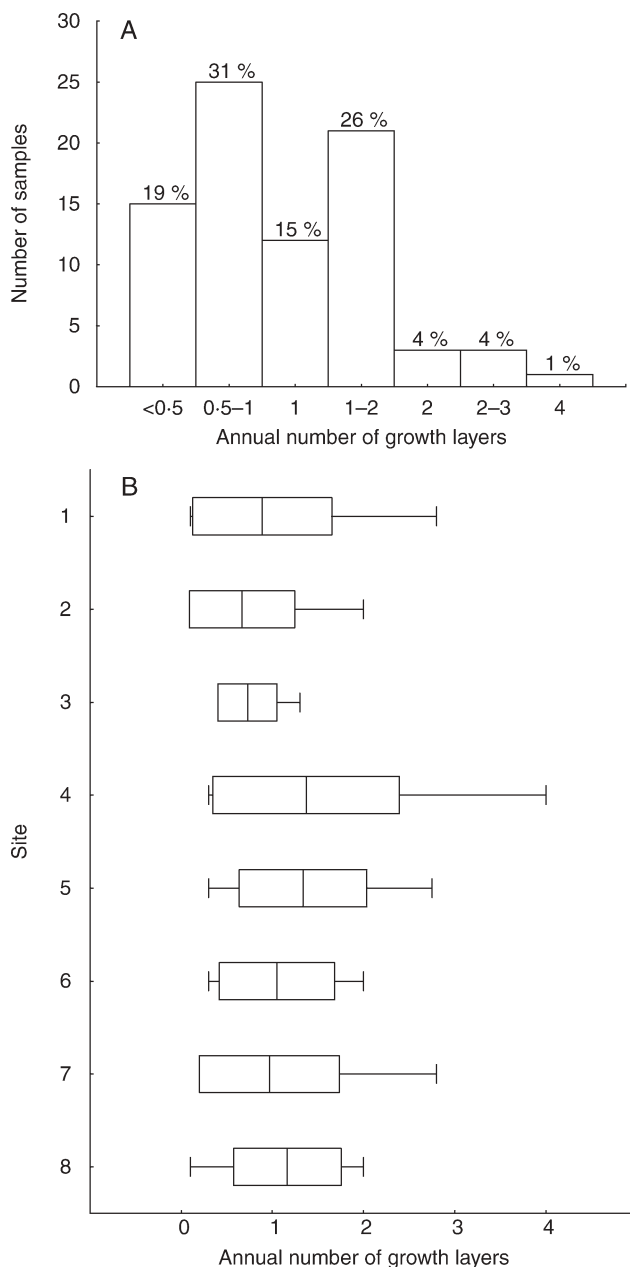


FIG. 5. Number of growth layers of 79 *Avicennia marina* trees formed in the period from May 2005 to June 2006 (A), distribution of the annual growth layer number over the eight study sites (B) (see also Table 2). Line, mean; box, s.d.; whiskers, min.-max.

370 and 1065 % higher, respectively, in branches compared with stems. Trees without dead leaves or branches had a significantly larger growth increment compared with trees with dead leaves and/or branches dispersed in the crown ($t = -3.68$, $n_1 = 15$ and $n_2 = 57$, $P < 0.01$). The growth from the cambial mark in February 2006, the middle of the dry season, until the felling in June 2006, the end of the long rainy season, was mostly small and restricted to the zone around the wound, hindering relevant measurements and study of potential wood anatomical changes in the dry season.

TABLE 3. Characteristics of the secondary growth of *Avicennia marina* averaged over the 1-year study period between May 2005 and June 2006 ($n = 80$) or averaged over the entire stem disc ($n = 49$)

Growth characteristic	Mean \pm s.d.	Minimum	Maximum
1-year study period			
Growth rate (mm year^{-1})	1 ± 1	0	5
Included growth* ($\mu\text{m year}^{-1}$)	249 ± 105	64	376
Growth layers (nr year^{-1})	1.0 ± 0.7	0.1	4
Disc average			
$D(M - m)$ (nr)	4 ± 3	0	12
Growth layer width (μm)	898 ± 137	402	1564
Xylem width (μm)	657 ± 107	257	1198
Phloem width (μm)	241 ± 35	117	403
% Phloem growth layer ⁻¹	28 ± 2	20	44

$D(M - m)$, difference in number of growth layers between the minimum and maximum disc radius; nr, number of growth layers.

* Growth of the last but one growth layer ($n = 10$, see Figs 2C and 3A).

TABLE 4. Results of simple linear regressions for the May 2005–June 2006 growth increment of *Avicennia marina* trees of different size and growing under different environmental conditions (see Table 1)

Factor	F-value	P-value	n	$r^{2\ddagger}$
Environment				
Minimum salinity	3.00	ns	73	0.03
Maximum salinity	0.04	ns	73	-0.01
Inundation class	1.47	ns	73	0.01
$[\text{NO}_3^-]$	1.11	ns	62	0.00
$[\text{NH}_4^+]$	0.02	ns	37	-0.03
[P]	1.09	ns	54	0.00
LAI	6.05	<0.05	62	0.08
Tree				
R_{130}^*	16.87	<0.001	73	0.18
R_b^\dagger	0.34	ns	73	-0.01
Height	9.13	<0.01	73	0.10

Data were transformed with a logarithmic function.

* Stem radius at 130 cm height.

† Stem radius at the base.

‡ Adjusted correlation coefficient.

Growth layer characteristics

The 2005–2006 growth increment was significantly correlated with the average growth layer width of a stem disc ($r^2 = 0.31$, $t = 4.37$, $n = 44$, $P < 0.0001$). The average growth layer width was strongly correlated with tree height ($r^2 = 0.43$, $F = 36.35$, $n = 47$, $P < 0.0001$) and weakly but significantly with soil water salinity, showing a higher correlation with maximum salinity ($r^2 = 0.16$, $F = 9.60$, $n = 47$, $P < 0.01$) than with minimum salinity ($r^2 = 0.10$, $F = 5.97$, $n = 47$, $P < 0.05$). No correlation was observed between growth layer width and age, represented by growth layer number, with the first growth layer at the pith having number 1 and the last growth layer at the bark having the highest number (partial correlation, controlling for tree: $r^2 = 0$, $t = 0.0067$, $n = 2663$,

$P = \text{ns}$). The width of a xylem band showed a strong positive correlation with the phloem band width of the same growth layer (Table 3, $r^2 = 0.53$, $t = 54.78$, $n = 2663$, $P < 0.0001$). However, the increase in phloem width was only 22 % of the corresponding xylem width, resulting in a negative relationship between the percentage of phloem and the growth layer width (Spearman $r^2 = 0.14$, $n = 2663$, $t = -21.26$, $P < 0.0001$) with a lower limit at 20 % phloem per growth layer (Table 3).

DISCUSSION

The patchy growth hypothesis

The first hypothesis of a periodic growth with the number of growth layers produced during the year depending on site conditions was rejected. Instead, a patchy growth mechanism is proposed based on circumstantial evidence: (a) a network pattern of growth layers (Fig. 4); (b) two simultaneously forming growth layers without the occurrence of wood cracks or crushed phloem tissue indicating the absence of internal strains (Figs 2C and 3A); (c) a remarkable variation in growth layer formation within the different study sites (Fig. 5B); and (d) an extensive variation of the radial increment, seemingly independent of any of the environmental factors considered (Tables 3 and 4). The patchy growth hypothesis states a basic growth rate all around the tree, with a more vigorous growth at one or several positions around the stem circumference. The side of active growth changes with time to end up with a tree of regular circumference. A patchy growth would clarify the three-dimensional phloem network, first reconstructed from serial branch microsections by Zamski (1979) and here for the first time visualized *in situ* from a mature tree trunk. Not only around (Fig. 4A) but also along the stem (Fig. 4B), growth patches instead of rings were observed. Nair and Mohan Ram (1990) propose a similar explanation for the growth layer network of *Dalbergia paniculata*, resembling that of *A. marina*. They suggest it could be due to the outermost parenchyma cells that do not invariably become meristematic. More recently, discontinuous new cambia have been observed in some species of Amaranthaceae (Rajput and Rao, 2000) and a liana species of the Menispermaceae (Jacques and De Franceschi, 2007).

The earlier suggestion of a simultaneous development of two growth layers (Schmitz *et al.*, 2007) was illustrated by a double cambial mark (Figs 2C and 3A). Cracks in the lignified part of the outermost growth layer can be overcome by collapse of phloem cells (Carlquist, 2007). Supporting the patchy growth hypothesis, this was not observed in the present study nor in *A. resinifera* and *A. germinans* in which at least the four outer phloem bands were uncrushed (Zamski, 1979). Moreover, the finding of 14 % ($n = 79$) of the trees with more than one active cambium could be an underestimation. The depth of wounding was not standardized because the depth needed to wound both cambia varies with the width of the outer growth layer. Recently, more than one active cambium was observed in the herbaceous and shrubby plants of Aizoaceae (Carlquist, 2007), questioning the existence of two mechanisms for successive

cambia formation. Unlike *Avicennia* (Zamski, 1979) and some other species (Esau, 1969), Carlquist (1999a, b, 2003a, 2004) found in members of the Caryophyllales successive cambia originating from one lateral meristem in the cortex that also produces the rays and the tissue in between the phloem strands. This would be hindered by the partially formed outer growth layer in between the lateral meristem and the inner forming growth layer. Moreover, the patchy differentiation of new cambia makes the existence of one circular meristem, giving rise to the vascular cambia, unlikely in *A. marina*.

Secondary growth in discrete units would explain the variation in growth between *A. marina* trees (Fig. 5). On the side of wounding, the cambial activity and the duration of active growth during the study period would determine the number of growth layers produced and the annual radial increment. Therefore, radial increment was not a reflection of past environmental conditions (Table 4). However, the total increase in stem width is expected to be influenced by external factors such as soil water nutrients (Feller *et al.*, 2003; Lovelock *et al.*, 2004), salinity (Lin and Sternberg, 1992; Tuffers *et al.*, 2001; Ball, 2002) and inundation periods (Worbes, 1989; Callado *et al.*, 2001; Menezes *et al.*, 2003; Verheyden, 2004). In favour of this idea, the average growth layer width of stem discs did show a decrease with increasing salinity. Despite the underestimation of total secondary growth, radial increment mirrored differences in leaf area related to crown die-back or to site-specific LAI (Table 4). Although only slightly, compared with tree species from Cameroon (Worbes *et al.*, 2003), tree size (Table 4) was also reflected in *A. marina*'s growth rate for 2005–2006 as supported by the overall positive effect of tree height on mangrove biomass production (Saenger and Snedaker, 1993). The absence of a relationship between radial increment during the study period and basal diameter (Table 4) agrees with earlier findings of tree growth being determined by tree structural characters rather than by age (e.g. Pélissier and Pascal, 2000; Sterck *et al.*, 2001; Worbes *et al.*, 2003; King *et al.*, 2005).

Additional support for the patchy growth is given by the slower radial growth, on average $0.57 \text{ mm year}^{-1}$ (see also Fig. 5B), at site 5 during the previous study (Schmitz *et al.*, 2007). The longer study period of 2.5 years in combination with a cambial mark at a single position around the stem would have increased the underestimation of the total increase in stem width if growth takes place at discrete and changing positions around the stem. Therefore, when comparing the growth rate of *A. marina* with that of other tree species, one has to be cautious. The measured growth rate concerns the radial increment on the side where wounding occurred. Depending on the number of positions around the stem circumference active during the study period, this value might underestimate the total increase in stem width. On the other side, growth layers growing at the time of wounding have already produced (part of) their phloem band, which might have resulted in an overestimation of the growth rate. Among mangrove species, the present findings on radial growth corresponded quantitatively with the radial growth of *Rhizophora mucronata* growing in the same forest in Gazi Bay, $<0.5\text{--}4.81 \text{ mm year}^{-1}$ (Verheyden,

2004), and of *Heritiera fomes* and *Sonneratia apetala* from Bangladesh, 1.4 ± 0.5 and 2 ± 1 mm year⁻¹, respectively (Md Q Chowdhury, Wageningen University, The Netherlands, unpubl. res.). Shiokura (1989) and Thampanya (2006) mention a higher growth rate of 3.8–5.7 mm year⁻¹ for *Avicennia* species from Japan and Thailand. A soil water salinity level rarely exceeding 40 ‰ (see also Table 2) in these countries' mangrove forests (Wakushima *et al.*, 1994a, b) might be responsible for the inconsistency. In India, three *A. officinalis* trees even had a diameter growth of 25.3 mm year⁻¹ (Baker, 1915). The growth layers there are suggested to be monthly (Baker, 1915) and thus are approx. 1 mm wide, as are those of *A. marina* (Table 3) and of *A. germinans* from Japan (Sun and Suzuki, 2000).

What triggers cambium differentiation?

The second hypothesis of the growth layer width being determined by soil water salinity was only partially accepted. The similar width of the growth layers of different *Avicennia* species together with the influence of radial growth on the formation of a new growth layer is in favour of a mainly endogenous control of cambium differentiation. An additional external influence of tree height and soil water salinity was recorded. Also, in other species, stem radius and number of growth layers are highly related (Ogden, 1981; Ruthrof *et al.*, 2003), but in *A. marina* the relationship applies even along the different radii of a single asymmetric stem disc. Moreover, the radial increment of 2005–2006 was only slightly related to the width of the corresponding growth layers as compared with trees with one vascular cambium. A strictly controlled growth layer formation results in a regular dispersion of the phloem tissue over the entire stem disc. Within one growth layer, the ratio of xylem to phloem tissue is lower compared with trees with conventional secondary growth (ratio: 2–9; Table 3) (Artschwager, 1950; Waisel *et al.*, 1966; Rajput and Rao, 2000). Besides, they accumulate only a limited number of phloem increments, making the total phloem fraction of an *Avicennia* stem considerably higher (Esau, 1969). The functional advantage of such a three-dimensional phloem network lies in its proposed role in embolism repair and water storage (Mauseth and Plemons-Rodriguez, 1997; Salleo *et al.*, 2004, 2006; Scholz *et al.*, 2007), supported by the decreasing growth layer width with increasing soil water salinity. The smaller the growth layers are, the higher is the percentage of phloem tissue and the lower the fraction of thick-walled xylem fibres, since the increase in phloem tissue was only 22 % of the xylem production. Therefore, the disappearance of growth layers at the smaller radius of asymmetric stems, instead of becoming smaller, might ensure the mechanical stability of the tree. The finding of wider growth layers in larger trees confirms this idea though no age trend was observed. Nevertheless, the considerable variation in growth layer width within a tree, with an average range of 402 ± 92 to 1564 ± 354 μm ($n = 47$), underscored the existence of an external control on the development of a new cambium superimposed on a genetically defined basic growth layer width.

In conclusion, a patchy growth via successive cambia may offer a functional advantage under xeric and fluctuating environmental conditions as occur in the mangrove environment. It is proposed that growth layer width is partly predetermined by mature tree height and partly controlled by the micro-environmental conditions via a trade-off between hydraulic safety and mechanical stability. Optimal growth conditions could prematurely stimulate the differentiation of a new cambium, resulting in two simultaneously forming growth layers. In this way, beneficial growth conditions could be exploited, which is supported by the significantly larger increment of the trees showing a double wound. To test this hypothesis and the proposed patchy growth, future studies should focus on controlled eco-physiological experiments and address the cambial activity on an intra-annual level, considering intra-tree variations and the relationship with tree phenology. Further insight into the functioning of successive cambia in different tree species and the link with environmental conditions is needed to reveal this growth strategy as a functional adaptation or a mere structural oddity.

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