PNEUMATOPHORE HEIGHT AND DENSITY IN RELATION TO MICRO-TOPOGRAPHY IN THE GREY MANGROVE AVICENNIA MARINA

Farid Dahdouh-Guebas^{1,2,*}, James G. Kairo^{3,1}, Ruth De Bondt¹ and Nico Koedam¹

¹ Biocomplexity Research Focus, c/o Laboratory of General Botany and Nature Management,
Mangrove Management Group, Vrije Universiteit Brussel, Pleinlaan 2, B-1050 Brussel, Belgium

² Biocomplexity Research Focus, c/o Laboratoire de Botanique Systématique et de Phytosociologie,
Département de Biologie des Organismes, Université Libre de Bruxelles, ULB - Campus du Solbosch,
CP 169, Avenue Franklin D. Roosevelt 50, B-1050 Bruxelles, Belgium

³ Kenya Marine and Fisheries Research Institute, P.O. Box 81651, Mombasa, Kenya

(* Author for correspondence; e-mail: fdahdouh@ulb.ac.be)

Received 28 October 2006; accepted 11 June 2007.

ABSTRACT. — Mangroves are known to stabilise coastal sediments through their above-ground aerial root complex. The results presented in this paper suggest that the grey mangrove *Avicennia marina* has the ability to adapt its pneumatophores to micro-topographical irregularities in the otherwise regularly sloping intertidal zone. The difference in height above datum (and thus in hydrological regime) in this study was as little as 15 cm higher as opposed to that for the surrounding mangrove soil. Significantly higher pneumatophore densities and total pneumatophore lengths were observed in the centre of the landward depression, and significantly lower belowground pneumatophore length in the centre of the seaward depression. The mangrove's adaptations to these localised topographic differences are important in view of changes in intertidal hydrology, the latter being linked to changes in topography. We emphasize the need to consider the effect of topography in the intertidal environment of mangroves more closely in the future on different ecological scales (individual adaptation – regeneration of the entire population) and under different scenarios of change (direct anthropogenic impact – changes in sea level).

KEY WORDS. — Avicennia, aerial roots, Kenya, micro-topography.

INTRODUCTION

Mangrove species have developed many morphological and physiological adaptations that are essential for them to survive in the harsh intertidal zone in which they grow. Adaptations include aerial roots, salt balance, vivipary and hydrochory (Tomlinson 1986). Whereas most mangrove species display a choice for optimal conditions for their growth, others have developed an ability to withstand a wide range of global and local conditions (e.g., Ball 1988).

Of all mangroves, Avicennia marina (Forssk.) Vierh. has the largest latitudinal spread (cf. Spalding et al. 1997), and on a regional scale, it has been shown to adapt to varying climatic conditions by adjusting its phenology (Duke 1990). This species-specific plasticity has also been found on a much larger scale, as this species displays differential population structures at different latitudes (Osunkoya & Creese 1997). Even within a particular mangrove forest Avicennia marina has shown to be one of the most eurytopic species along environmental

Table 1. Overview of countries or regions where a 'double' or 'disjunct' zonation of *Avicennia marina* has been observed within sites with multiple mangrove species. The number of true mangrove species for the respective country is given according to SPALDING *et al.* (1997), but may differ on a local scale. For some countries more recent corrected species numbers have been given. This overview obviously excludes references to areas where *A. marina* occurs as the sole mangrove species (e.g., monospecific stands at latitudinal distribution limits).

Country or region	Number of true mangrove species	Reference(s)
Indo-West Pacific	up to 45	Macnae (1968)
East-Africa	11	Walter & Steiner (1936)
		Gallin et al. (1989), Dahdouh-Guebas et al.
Kenya	10	(1998), Matthus et al. (1999), Dahdouh-
•		GUEBAS et al. (2002a,b, 2004a,b)
India	28	Satyanarayana et al. (2002)
Malaysia	36	Watson (1928)
Australia, NT	37	Bunt (1996), Woodroffe & Grime (1999)
Australia, QLD	37	Macnae (1969), Bunt (1996), Clarke (2004)

gradients, which is particularly well illustrated by the appearance of a 'double' or 'disjunct' intertidal zonation in many parts of the world (Table 1). Such a zonation has been occasionally reported as observations (JOHNSTONE 1983, SMITH 1992, OCHIENG & ERFTEMEIJER 2002), but recently DAHDOUH-GUEBAS *et al.* (2004a) described *Avicennia marina*'s clear variation in morphological and genetic characteristics between landward and seaward zones.

Apart from effects induced by the natural mangroves environment. are additionally affected by direct and indirect human-induced stresses and disturbances, some effects of which are compounded in the intertidal zone (e.g., Alongi 2002, Dahdouh-Guebas et al. 2004b). In the light of the response of mangroves to global climatic change, and in particular of A. marina to sea-level rise, this preliminary study focuses on the adaptation of the pencil-root¹ complex of A. marina to changes in microtopography. DAHDOUH-GUEBAS et al. (2004a) demonstrated that root density and length were dependent on the intertidal zone in which the

tree was growing. The objective of the present study was to investigate whether very shallow and local micro-topographic depressions (0.2 – 0.4 m) have an effect on A. marina pneumatophore density or on their above- and belowground length. Our hypothesis was that there exists a trend in the density and the length of pneumatophores from the centre of a small depression (high/long) towards the edges (lower/shorter) in both landward and seaward A. marina zones.

MATERIAL AND METHODS

The study was conducted in the Kenyan mangrove area near Gazi (Fig. 1), for which the disjunct zonation of *Avicennia marina* has been described earlier (DAHDOUH-GUEBAS *et al.* 2002a,b, 2004a,b). The two *Avicennia* zones are located along the intertidal slope and are separated by approximately 100 m. Between these two *Avicennia* fringes, the mangrove vegetation is composed of other species, such as Rhizophoraceae. The spring tidal amplitude is about 3.5 m, and the seaward *Avicennia* zone is

¹ Mangrove aerial roots are commonly distinguished as prop or stilt roots (e.g., *Rhizophora* spp.), knee-roots (e.g., *Bruguiera* spp.), plank and buttress roots (e.g., *Xylocarpus* spp.), peg roots (e.g., *Sonneratia* spp.), and pencil roots (see TOMLINSON 1986). Pencil roots originate from the below-ground cable root and stick out vertically from the soil, and are a typical feature of *Avicennia* species (Fig. 1).

inundated twice daily (water column may be more than 2 m high), whereas the landward zone is inundated only during spring high tides (water column < 0.5 m). For a detailed description of the study site, including remotely sensed imagery and descriptive vegetation data, see DAHDOUH-GUEBAS et al. (2004a). A 10 m transect was laid across a natural topographic depression within the landward Avicennia marina vegetation zone. An 11 m transect was used in the seaward A. marina vegetation zone. The landward zone was composed entirely of A. marina. In the seaward zone A. marina was the dominant (87%) species with some Rhizophora mucronata Lamk. and Sonneratia alba J. Smith. Tree spread was approximately even in both zones. At 1 m intervals along each of the transects, 1 m² quadrates were established ($n_{\text{landward}} = 10$; $n_{\text{seaward}} = 11$), and within each quadrate, the number of pencil roots was counted and their above-ground length was recorded. In addition, four randomly chosen pneumatophores per quadrate were excavated down to the cable root in order to measure their below-ground-length. Since there was little variation in below-ground-length, the average of the four measurements was added to each of the above-ground-length measurements to obtain the total pneumatophore length. Spearman's rank correlation coefficient (r_s) was used as described by SOKAL & ROHLF (1981) in order to correlate height above datum (and thus relative depression depth) with the variables and parameters from the root complex. The height above datum was measured using a theodolite (Nikon, Auto level AE-3G). This was done at 1 m intervals within the depressions, and at 5 m intervals between the depressions, using a reference point or benchmark. This benchmark was the maximum water height above datum (highest water line), as predicted by tide tables available from the Kenya Port Authority. We are aware that tidal predictions may be off by a few centimetres, particularly in shallow bays, but we did not attempt to measure the actual hydro-graphic height, mainly because the objective of this study emphasized the existing relative differences in height and their effect on the vegetation. All elevations were expressed as 'meters above datum', and the error of the theodolite was experimentally recorded at 3 cm.

Redox potentials of the soil were measured with a platinum-Ag/AgCl redox electrode connected to a pH/mV/T-meter (P601, Eijkelkamp, Agrisearch Equipment), as described in MATTHIJS *et al.* (1999), in order to examine the effect of long inundation periods at the centre of the depressions.

RESULTS

Maximum depth of the topographic depression was 15 cm in the landward zone (3.4 m above datum; Fig. 2a). The highest density of pneumatophores in the landward depression was 2 500 m⁻² at the centre of the depression (Fig. 2c), but was only 200 m⁻² outside the pit. The redox potential of the interstitial water at the centre of the landward depression was -259 mV, while values outside the depression were less reduced (-55 mV and -178 mV). Total pneumatophore length increased towards the centre of the landward depression (Fig. 2a) and the relationship with height above datum was highly significant ($r_s = 0.71$; n = 10; P < 0.005). The below-ground length decreased significantly with a lower height above datum in the landward depression ($r_s = -0.86$; n = 10; P < 0.001; Fig. 2a).

Maximum depth of the topographic depression was 44 cm in the seaward zone (1.7 m above datum; Fig. 2b). However, there was a decrease in below-ground pneumatophore length with a lower height above datum ($r_s = -0.94$; n = 11; P < 0.001; Fig. 2b). There was no significant decrease in total pneumatophore length with a lower height above datum ($r_s = 0.44$; n = 11; n.s.). For the seaward depression, there seems to be no relationship between pneumatophore density and the position in the depression and thus height above datum (Fig. 2b,d). The redox potential of the interstitial water of the seaward depression was most reduced half-way down the depression slope (-352 mV and -386 mV), less reduced in the deepest quadrates (-296 mV and -306 mV) and least reduced at the top margins (-155 mV and -135 mV).

DISCUSSION

The discussion of our results is made within the seasonal conditions that prevailed during out field campaign (dry season). Although we are aware that our environmental measurements of the redox potential may be different in the wet season, or when multiple measurements are performed throughout the year, we have often observed that the more stressful conditions shaping the vegetation occurs in the dry season.

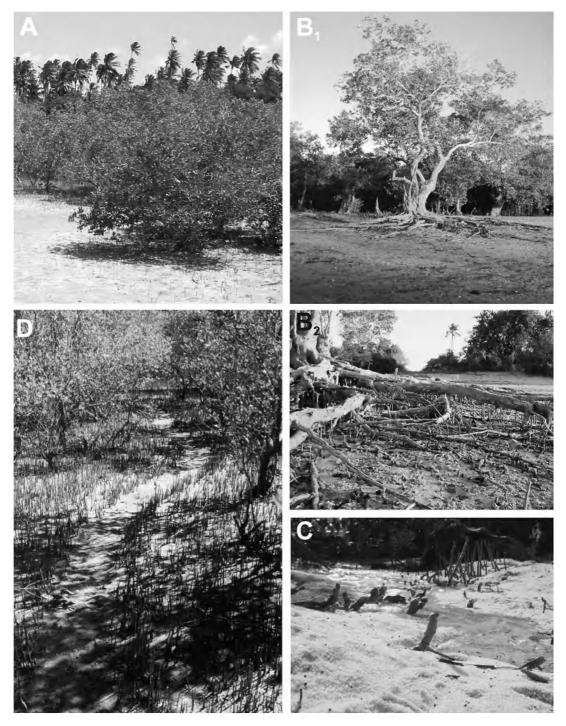


Fig. 1. (A) Physiognomy of a landward *Avicennia marina* fringe in Gazi, Kenya. (B) Overview (B1) and close-up (B2) of an *A. marina* tree uprooted by tidal and wave energy. (C) Close-up of *A. marina* pencil-roots and of the micro-topographic variations (note the pathway of the incoming tide). (C) Path amongst *A. marina* pneumatophores created by walking fisherfolk. (Photographs taken by Griet Neukermans and Farid Dahdouh-Guebas)

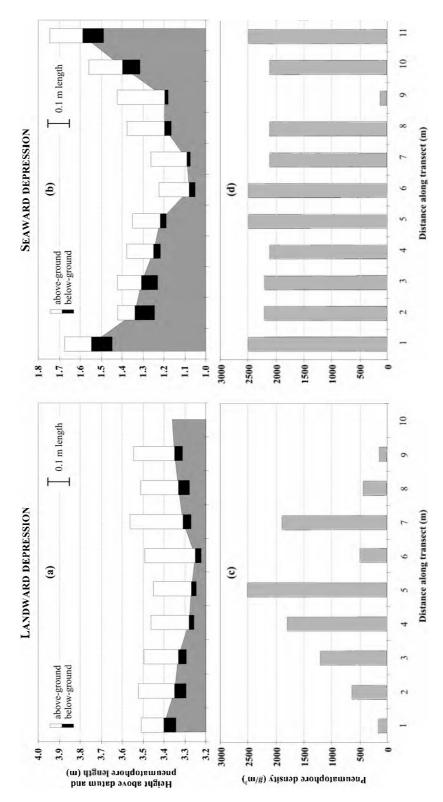


Fig. 2. (a-b) Micro-topography and above- and below-ground pneumatophore length, and (c-d) pneumatophore density for Avicennia marina sampled across a depression at 1 m intervals along a landward and a seaward mangrove transect

Observed pneumatophore density and length in the topographic depressions are in line with the hypothesis that pneumatophore density and length is higher for places with longer inundation periods (DAHDOUH-GUEBAS et al. 2004a). The hypothesis was confirmed for pneumatophore density and total and below-ground length in the landward depression. In addition, the present study revealed the sensitivity of the root complex of Avicennia marina with respect to micro-topographical settings. It is notable that even shallow depressions (0.2 m and 0.4 m for the landward and seaward depression, respectively) seem to trigger the plants to adjust their root complex significantly. However, further investigations in a variety of mangrove formations are necessary to validate these results.

Decreased below-ground pneumatophore length in the landward zone can be explained by the sandy substrate being easily removed by water, making the below-ground length appear shorter. In some landward areas outside the study area, the action of water has denuded entire root complexes of Avicennia marina (Fig. 1B). This removal of the substrate occurs in areas that display a sparse distribution of trees and roots, and where small waves (< 1 m length along the topographical gradient) occur at spring tide. Pneumatophore length and density being higher towards the centre of the depression may be indicative of root growth toward more oxygenated surface horizons (cf. DAHDOUH-GUEBAS et al. 2004a).

Pneumatophores in the seaward depression generally do not reach above the high water level. Tidal fluctuations of > 2 m in our sites cannot be overcome by the height of pencil roots of *Avicennia marina* (unlike peg roots in *Sonneratia* spp., pencil roots are thin throughout their length and tend to fall over as they grow tall). An increase in density would be beneficial, but of less importance in a depression, since the pneumatophores are already entirely exposed to water, and not covered by detritus.

Even though this study was conducted in a natural *A. marina*-dominated zone with an even spread of trees, local variations in pneumatophore density may be due to tree density

(both A. marina and other species) and cable root length (distance to tree). However, at the place of the depressions no other species were present. The pneumatophore density or length in a large majority of quadrates studied consistently displayed the hypothesized trend in both depressions.

The results suggest that the grey mangrove Avicennia marina has the ability to adapt its root complex to micro-topographical irregularities in the otherwise regularly sloping intertidal zone. Although this study was carried out in Kenya, a review of recent literature indicates that it may apply to other mangroves dominated by A. marina worldwide, where pneumatophore density, pneumatophore density range or microtopography display similarities in absolute or relative values (Table 2). Yet, differences in root density as a result of micro-topographic differences have never been reported before. The hydrological regime in this study was as little as 15 cm higher than that for the surrounding mangrove soil. The mangrove species' adaptation to this localised topographic difference is important in view of changes in intertidal hydrodynamics, the latter of which is linked to changes in topography. Apart from individual adaptation, the effects on the regeneration of the entire populations also should be considered, as substratum heterogeneity has been found to influence the recruitment of A. marina (MINCHIN-TON 2001). Changing root complexes may alter their ability to entangle mangrove propagules directly, or indirectly through woody debris (STIEGLITZ & RIDD 2001, KRAUSS et al. 2005).

From a human impact or management perspective, the observed results are of importance and suggest that trees will react to the creation of topographical depressions or pneumatophore destruction, both of which may occur when people make their way through the mangrove on foot (Fig. 1D). Although we do not know at what time scale the difference in pneumatophore density and length have developed, or to which extent it can be found in other mangrove formations, it is evident that the influence of microtopography merits closer attention in the future, as it may be significant on other scales of the

Table 2. Recent peer-reviewed papers referring to variations in pneumatophore density for mangrove species or to the effects of micro-topographical differences (*A. mar = Avicennia marina*; *B. gym = Bruguiera gymnorrhiza*; *R. api = Rhizophora apiculata*; *S. alb = Sonneratia alba*; *X. gra = Xylocarpus granatum*; n.a. = not available).

Height above datum	Pneumatophore density (number/m²)	Species	World region	Reference
1 to 3 m n.a. n.a. n.a. n.a. n.a. 6 to 13 m n.a. n.a. n.a. n.a. n.a. n.a.	4 - 1950 80 - 180 56 - 1168 200 ca. 65 - 80 24 - 347 382 - 718 389 - 1040 ca. 20 - 85 50 - 381 ca. 40 - 250 80 - 200 n.a.	A. mar A. mar B. gym R. api S. alb X. gra	Kenya Mozambique Pakistan QLD, Australia QLD, Australia QLD, Australia NSW, Australia NSW, Australia NSW, Australia NSW, Australia New Zealand New Zealand Micronesia	Dahdouh-Guebas et al. (2004a) Macia et al. (2003) Saifullah & Elahi (1992) Laegdsgaard & Johnson (2001) Skilleter & Warren (2000) Harty & Cheng (2003) Bishop et al. (2007) Melville & Pulkownik (2007) Kelaher et al. (1998) Burchett et al. (1999) Young & Harvey (1996) Alfaro (2006) Fujimoto et al. (1995)

mangrove ecosystem as well (e.g., vegetation structure, dispersion processes). In addition, the strength and density of the root complex of mangrove trees is worth focusing on in the light of their ability to protect coastal areas from human, meteorological and oceanographical hazards (BADOLA & HUSSAIN 2005, DAHDOUH-GUEBAS et al. 2005a,b). Not only for Avicennia spp., but also for other mangrove species there is a lack of studies relating micro-topography (rather than intertidal position and intertidal height above datum) to vegetation characteristics. We recommend that more research be focused on the interrelationships between hydrodynamics, topography and vegetation, particularly the effects of water on the mangroves' root complex, and the effect of the mangrove root complex on the water currents and on a wide range of water-related impacts, among which storm surges, sea-level rise, daily tidal action, heavy El-Niño rains and tsunamis.

ACKNOWLEDGEMENTS

The first author was a Postdoctoral Researcher from the Science Foundation in Flanders (FWO-Vlaanderen). The research and its presentation was

also financed by the European Commission (Contract No. TS3*-CT92-0114), the Fund for Collective Fundamental Research (FKFO) and with a specialisation fellowship of the Institute for the Promotion of Innovation by Science and Technology in Flanders (IWT). We thank S. Matthijs, Y. Samyn, K. Millis, J. Mwachirumu, D. Van Speybroeck, J.F. Tack, P. Polk, E. Okemwa, G. Ngari, M. Kodjo and families of J. Seys, P. Reyniers and L.S. Ba'alawy for fieldwork, logistic and socio-cultural assistance. We also thank anonymous reviewers and Editor Patrick Endels for their critical and constructive comments.

REFERENCES

ALFARO A.C., 2006. — Benthic macro-invertebrate community composition within mangrove/seagrass estuary in northern New Zealand. *Estuar: Coast. Shelf Sci.* **66**: 9-110.

ALONGI D.M., 2002. — Present state and future of the world's mangrove forests. *Environm. Conserv.* **29**: 331-349.

BADOLA R. & HUSSAIN S.A., 2005. — Valuing ecosystem functions: an empirical study on the storm protection function of Bhitarkanika mangrove ecosystem, India. *Environm. Conserv.* **32**: 85–92.

BALL M.C., 1988. — Salinity tolerance in the mangroves, *Aegiceras corniculatum* and *Avicennia*

- marina. I. Water use in relation to growth, carbon partitioning and salt balance. Austr. J. Plant Physiol. 15: 447–464.
- BISHOP M.J., KELAHER B.P., SHARP S.M. & VERHOEVEN M., 2007. Dispersal of the estuarine gastropod *Pyrazus ebeninus* is only weakly influenced by pneumatophore density. *J. Exp. Marine Biol. Ecol.* **340**: 62–69.
- Bunt J.S., 1996. Mangrove zonation: an examination of seventeen riverine estuaries in tropical Australia. *Ann. Bot.* **78**: 333-341.
- BURCHETT M.D., PULKOWNIK A., GRANT C. & MACFAR-LANE G., 1999. — Rehabilitation of saline wetlands, Olympics 2000 site, Sydney (Australia) – I: Management strategies based on ecological needs assessment. *Marine Pollu. Bull.* 37: 515-525.
- CLARKE P.J., 2004. Effects of experimental canopy gaps on mangrove recruitment: lack of habitat partitioning may explain stand dominance. *J. Ecol.* **92**: 203-213.
- Dahdouh-Guebas F., Verneirt M., Tack J.F., Van Speybroeck D. & Koedam N., 1998. Propagule predators in Kenyan mangroves and their possible effect on regeneration. <u>Marine Freshw. Res.</u> **49**: 345-350.
- Dahdouh-Guebas F., Kairo J.G., Jayatissa L.P., Cannicci S. & Koedam N., 2002a. An ordination study to view vegetation structure dynamics in disturbed and undisturbed mangrove forests in Kenya and Sri Lanka. *Plant Ecol.* **161**: 123-135.
- Dahdouh-Guebas F., Verneirt M., Cannicci S., Kairo J.G., Tack J.F. & Koedam N., 2002b. An exploratory study on grapsid crab zonation in Kenyan mangroves. *Wetlands Ecol. Manage.* 10: 179-187.
- Dahdouh-Guebas F., De Bondt R., Abeysinghe P.D., Kairo J.G., Cannicci S., Triest L. & Koedam N., 2004a. Comparative study of the disjunct zonation pattern of the grey mangrove *Avicennia marina* (Forsk.) Vierh. in Gazi Bay (Kenya). *Bull. Marine Sci.* 74: 237-252.
- DAHDOUH-GUEBAS F., VAN POTTELBERGH I., KAIRO J.G., CANNICCI S. & KOEDAM N., 2004b. Human-impacted mangroves in Gazi (Kenya): predicting future vegetation based on retrospective remote sensing, social surveys, and distribution of trees. *Marine Ecol. Progr. Ser.* 272: 77-92.
- Dahdouh-Guebas F., Hettiarachchi S., Lo Seen D., Batelaan O., Sooriyarachchi S., Jayatissa L.P. & Koedam N., 2005a. Transitions in ancient inland freshwater resource management in Sri

- Lanka affect biota and human populations in and around coastal lagoons. *Curr. Biol.* **15**: 579-586.
- Dahdouh-Guebas F., Jayatissa L.P., Di Nitto D., Bosire J.O., Lo Seen D. & Koedam N., 2005b. How effective were mangroves as a defence against the recent tsunami? *Curr. Biol.* 15: R443-447.
- Duke N.C., 1990. Phenological trends with latitude in the mangrove tree *Avicennia marina*. *J. Ecol.* **78**: 113-133.
- FUJIMOTO K., TABUCHI R., MORI T. & MUROFUSHI T., 1995. Site environments and stand structure of the mangrove forest on Pohnpei Island, Micronesia. *Jap. Agric. Res. Quart.* **29:** 275-284.
- GALLIN E., COPPEJANS E. & BEECKMAN H., 1989. The mangrove vegetation of Gazi bay (Kenya). *Bull. Soc. Rov. Bot. Belg.* **122**: 197-207.
- HARTY C. & CHENG D., 2003. Ecological assessment and strategies for the management of mangroves in Brisbane Water—Gosford, New South Wales, Australia. *Landsc. Urban Plann.* **62**: 219–240.
- JOHNSTONE I.M., 1983. Succession in zoned mangrove communities: where is the climax ? In: Teas H.J. (ed.), Tasks for vegetation vcience 8, pp. 131-139. Dr. W. Junk Publishers, The Hague, The Netherlands.
- Kelaher B.P., Chapman M.G. & Underwood A.J., 1998. Changes in benthic assemblages near boardwalks in temperate urban mangrove forests. *J. Exp. Marine Biol. Ecol.* **228**: 291-307.
- Krauss K.W., Doyle T.W., Twilley R.R., Smith III T.J., Whelan K.R.T. & Sullivan J.K., 2005. Woody debris in the mangrove forests of South Florida. *Biotropica* 37: 9-15.
- LAEGDSGAARD P. & JOHNSON C., 2001. Why do juvenile fish utilise mangrove habitats? <u>J. Exp.</u>

 Marine Biol. Ecol. **257**; 229-253.
- MACIA A., ABRANTES K.G.S. & PAULA J., 2003. Thorn fish *Terapon jarbua* (Forskål) predation on juvenile white shrimp *Penaeus indicus* H. Milne Edwards and brown shrimp *Metapenaeus monoceros* (Fabricius): the effect of turbidity, prey density, substrate type and pneumatophore density. *J. Exp. Marine Biol. Ecol.* **291**: 29–56.
- MACNAE W., 1968. A general account of the fauna and flora of mangrove swamps and forests in the Indo-West-Pacific region. *Adv. Marine Biol.* **6**: 73-270.
- MACNAE W., 1969. Zonation within mangroves associated with estuaries in north Queensland. *In:* LAUFF G.H. (ed.), *Estuaries*, pp. 432-441. American Association for the Advancement of Science, Washington D.C., USA.

- MATTHUS S., TACK J., VAN SPEYBROECK D. & KOEDAM N., 1999. Mangrove species zonation and soil redox state, sulphide concentration and salinity in Gazi Bay (Kenya), a preliminary study. <u>Mangroves and Salt Marshes</u> 3: 243-249.
- MELVILLE F. & PULKOWNIK A., 2007. Seasonal and spatial variation in the distribution of mangrove macroalgae in the Clyde River, Australia. *Estu. Coast. Shelf Sci.* 71: 683-690.
- MINCHINTON T.E., 2001. Canopy and substratum heterogeneity influence recruitments of the mangrove *Avicennia marina*. <u>J. Ecol. 89</u>: 888-902.
- OCHIENG C.A. & ERFTEMEIJER P.L.A., 2002. Phenology, litterfall and nutrient resorption in *Avicennia marina* (Forssk.) Vierh in Gazi Bay, Kenya. *Trees* **16**: 167-171.
- OSUNKOYA O.O. & CREESE R.G., 1997. Population structure, spatial pattern and seedling establishment of the grey mangrove *Avicennia marina* var. *australasica*, in New Zealand. *Austr. J. Bot.* 45: 707-725.
- SAIFULLAH S.M. & ELAHI E., 1992. Pneumatophore density and size in mangroves of Karachi, Pakistan. *Pak. J. Bot.* **24**: 5-10.
- Satyanarayana B., Raman A.V., Dehairs F., Kalavati C. & Chandramohan P., 2002. Mangrove floristic and zonation patterns of Coringa, Kakinada Bay, East Coast of India. <u>Wetlands Ecol. Manage</u>. 10: 25-39.
- SKILLETER G.A. & WARREN S., 2000. Effects of habitat modification in mangroves on the structure of

- mollusc and crab assemblages. <u>J. Exp. Marine</u> Biol. Ecol. **244**: 107-129.
- SMITH III T.J., 1992. Forest structure. *In:* ROBERT-SON A.I. & ALONGI D.M. (eds.), *Tropical man*grove ecosystems, pp. 101-136. American Geophysical Union, Washington D.C., USA.
- SOKAL R.R. & ROHLF F.J., 1981. Biometry. The principals and practice of statistics in biological research. W.H. Freeman & Co, New York, USA.
- Spalding M., Blasco F. & Field C., 1997. World mangrove atlas: 178 p. The International Society for Mangrove Ecosystems, Okinawa, Japan.
- STIEGLITZ T. & RIDD P.V., 2001. Trapping of mangrove propagules due to density-driven secondary circulation in the Normanby River estuary, NE Australia. *Marine Ecol. Progr. Ser.* 211: 131-142.
- Tomlinson P.B., 1986. *The botany of mangroves*: 419 p. Cambridge University Press, Cambridge, UK.
- WALTER H. & STEINER M., 1936. Die Ökologie der Ost-Afrikanischen Mangroven. Zeitschrift für Botanik 30: 65-193.
- WATSON J.G., 1928. Mangrove forests of the Malay Peninsula. *Malay Forest Records* **6**: 1-275.
- Woodroffe C.D. & Grime D., 1999. Storm impact and evolution of a mangrove-fringed chenier plain, Shoal Bay, Darwin, Australia. *Marine Geol.* **159**: 303-321.
- Young B.M. & Harvey E.L., 1996. A apatial analysis of the relationship between mangrove (*Avicennia marina* var. *australasica*) physiognomy and sediment accretion in the Hauraki Plains, New Zealand. *Estu. Coast. Shelf Sci.* **42**: 231-246.