

Effects of irrigation on litterfall, fine root biomass and production in a semideciduous lowland forest in Panama

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Abstract

The effects of irrigation on fine root biomass, root production and litterfall were measured at the community level, in a semideciduous lowland forest in Panama. Biomass of roots less than 2 mm in dia. in the first 10 cm of the soil (measured with soil cores), was higher in irrigated (1.80 Mg ha⁻¹) than in non-irrigated plots (1.24 Mg ha⁻¹). During the dry season, productivity of roots (measured with ingrowth cylinders filled with root-free soil), was higher in irrigated (1.6 g m⁻² day⁻¹) than in control plots (0.3 g m⁻² day⁻¹). In control plots, root productivity was highly seasonal. Maximum root growth into the root-free soil, occurred during the transitions from dry to wet, and from wet to dry season, possibly as a response to water and/or nutrient pulses. Litterfall was not significantly different between irrigated (3.8 g m⁻² day⁻¹) and control plots (3.7 g m⁻² day⁻¹). The results of this study show that root-productivity is limited by the water supply during the dry season, and that water by itself, is not a limiting factor for community-level litter production.

Introduction

Rainfall in many tropical forests is seasonal, with dry seasons varying from a few weeks to several months (Beard, 1944, 1955; Richards, 1952). With increasing drought, forest physiognomy change gradually from lowland rain forest (*sensu stricto*) to evergreen seasonal forest, to semi-evergreen seasonal forest, and to deciduous forest (Beard, 1944, 1955). Along this humidity gradient, the occurrence of deciduous tree species increases and thus the seasonallity in litter production, with leaf fall peaking during the dry seasons (Folster and de las Salas, 1976; Franken et al., 1979; Frankie et al., 1974; Haines and Foster, 1977; Klinge and Rodriguez, 1968; Ogawa, 1978; Schaik, 1986).

Leaf flush often peaks soon after the rains begin (Leigh and Windsor, 1982), and trunk growth is more pronounced during the wet season (Daubenmire, 1972; Raich and Borchert, 1982). On Barro Colorado Island, where the dry season usually begins in December and ends in April or May, leaf fall increases at the beginning of the dry season with maximum rates in January (Dietrich et al., 1982; Wright and Cornejo, 1990b). As the dry season progresses, leaf fall rates decline slowly dropping to a minimum with the onset of rains. Low leaf fall rates are characteristic of the wet season, and may be interrupted by a small increase during the 'veranillo' (shorter dry season) during August and September (Wright and Cornejo, 1990b). The number of deciduous trees producing young leaves peaks in May, while evergreen trees peak during the wet season (September), at the end of the year, and after the onset of rains (Leigh and Windsor, 1982). Furthermore, a survey of saplings of canopy and understory species, showed that leaf production occurs mostly during the transitions from wet to dry and from dry to wet seasons (Aide, 1988).

In contrast to the response of the above-ground biomass to changes in water supply, there is little information on the effects of the dry season on the below-ground ecosystem, particularly on fine-root biomass and production. In spite of the great im-

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portance of fine roots on forest productivity (Persson, 1979; Vogt et al., 1981; Gower et al. 1992), fine root biomass has been measured in only a few tropical forests (i.e. Klinge and Herrera, 1978; Sanford, 1989a,b; Stark and Spratt, 1977), and fine-root productivity in even fewer sites (Cavelier, 1989; Cuevas and Medina, 1983, 1988; Jordan and Escalante, 1980; Sanford, 1985). Responses of roots to tree falls (Sanford, 1990) and mineral nutrient amendments have been studied recently (Cavelier, 1989; Gower and Vitousek, 1989).

In this paper we present the results of an irrigation experiment in a semideciduous lowland forest in central Panama. We investigate whether seasonal changes in soil water content affect productivity at the community level and whether there are effects on the patterns of allocation of resources above- and belowground. If water is a limiting factor, as suggested by seasonal differences in rainfall and soil water potential, irrigation should have a direct and positive effect on the biomass and productivity of fine roots, and indirectly on fine litterfall through higher water and/or nutrient consumption by the roots.

Methods

Study site

Soil irrigation was carried out in a semideciduous lowland forest at an altitude of 60m on Barro Colorado Island (BCI). This forest is composed of a mixture of deciduous and evergreen species (Croat, 1978; Foster and Brokaw, 1982). The number of deciduous trees bearing young leaves peak in May and June and is lowest from October to February. Canopy evergreen trees show a continuous change in foliage, with a minimum during the leaf flush of deciduous canopy trees (Leigh and Windsor, 1982). In all, the canopy is never entirely deciduous. This forest experiences a marked seasonal variation in water supply. The annual rainfall on BCI averages 2612 mm (1925-1986; Windsor, 1990) with a median length of the dry season of 17 weeks between mid December-early January and late April-early May. During this part of the year, panevaporation exceeds rainfall and soils become dry and hard. Surface soil moisture content declines to 27% (wet weight) during the dry season, in contrast to 45% during the wet season (Windsor, 1990). This forest has remained without human intervention for more than 500 years (Piperno, 1990). Detailed description

of the flora, structure and ecology of this forest can be found in Croat (1978), Leigh et al. (1982) and Windsor (1990).

Experimental design

Four 2.25-ha plots were located in Poacher's Peninsula. The distance between plots averaged over 50 m. Two of these plots were irrigated and two served as controls. Water was added to the irrigated plots by 160 sprinklers placed 1.8 m above the ground at 15.3 m intervals in an hexagonal array. Thus, each point in the forest would receive water from three or more sources, except plot borders and where water was intercepted by vegetation. During a typical week in the dry season, each irrigated plot received 135 metric tons of water during 1.5 h of irrigation between 11:00 and 14:00 h on each of five days. This is equivalent to a daily rainfall at the forest floor of 6 mm. Mean soil water potential was maintained at or slightly above field capacity (-0.04 MPa) throughout the dry seasons of 1986-1990.

Water availability

Rainfall was measured daily at the site with a metric rain gauge (All Weather Rain Gauge). Soil water content was measured at eight randomly located stations in the interior of each plot (> 15 m from edge). Tensiometers (Soil Moisture Equipment Corp., 2725A), soil psychrometers (Wescor PCT55) and a hydroprobe (Campbell Nuclear Pacific, Model 503) were used to monitor soil water content as described elsewhere (Wright and Cornejo, 1990a).

Root measurements

Standing root biomass was measured with ten soil cores per plot on 14–15 December 1988 at the end of the rainy season, and on 29–30 March 1989 at the end of the dry season. Root biomass was sampled by driving into the soil a sharp-edge steel corer with an internal diameter at the hardened cutting edge of 5.6 cm. The corer had an internal PVC pipe, 7.5 cm in diameter, divided longitudinally in two halves to allow the cores to be removed (Ford and Deans, 1977). The soil corer was driven to a depth of -10 cm. Each soil core was placed in a plastic bag, labeled and stored at 3 °C until soil washing could take place, usually within 30 days. The samples were rinsed with tap water over a 500 μ m sieve to loosen the soil and facilitate root sorting. Root fragments which were mixed with organic

matter (and not easily identifiable as roots), were not collected. Roots were sorted by diameter (Very Fine Roots, VFR=<1 mm; Fine Roots, FR=1-2 mm and Medium Roots, MR=2-5 mm). After sorting, roots were dried at 60 °C for 48 h, weighed to the nearest 0.0001 g.

Root growth was measured by means of ingrowth cylinders (Cuevas and Medina, 1983, 1988; Lund et al., 1970; Persson, 1979). Each ingrowth cylinder (IC) was made with 30 mm² plastic mesh, 9 cm in diameter and 25 cm tall. The cylinders were placed in holes created with a Dutch auger. The holes were placed in random positions, at least 15 m from the plot edges and each point at least 10 m from all others. Each hole was filled with root free soil. This soil was obtained by drying (60 °C for 48 h), crushing and removing the roots with forceps, from soil collected at the site. IC were collected by carefully cutting the roots that had penetrated the cylinder, and then lifting out the column. Four series of 5 ingrowth cylinders were inserted in each plot, and the location of all 20 IC within the plot, was selected at random. The first series (dry season) was installed on 11 February and collected on 10 April 1988. The second series (transition dry-wet season) was installed on 5 May and collected 5 July 1988. The third series (wet season) was installed 22 September and collected 23 November 1988. The fourth series (transition wet-dry season) was installed on 7 December and collected on 14 February 1989. The contents of the cylinder were processed as described for the soil cores.

Leaf fall

Leaf fall was measured with 15 litter traps, located in random positions in each plot. The surface area of each trap was 0.25 m^2 . Traps were constructed with PVC tubing and plastic screening (1.2 mm mesh), and mounted 40 cm above the soil. Traps were emptied every seven days. Plant material was dried to a constant weight at 60 °C and weighed to the nearest 0.1 g. In this paper we present total weekly litterfall from 7 December 1987 through 29 May 1989.

Calculations and statistical analysis

Two-way ANOVAs were performed for standing root biomass and root ingrowth. Treatment and seasons were fixed main effects. Plots were a random factor nested within treatment. Levels of treatment were irrigated and control. For the analysis of standing root biomass, levels of season were December 1988 (end



ROOT DJAMETER

₽R

(1-2 mm)

VFR

(<1mm)

Figure 1. Standing root biomass for Very-line (<1 mm in diameter), fine (1–2 mm) and medium roots (2–5 mm) at the end of the wet season of 1988 (December) and at the end of the dry season of 1989 (March). (\square) = root biomass in control plots and (\square) = root biomass in the irrigated plots. (*) represent significant differences (p<0.05) between treatments.

of wet season) and March 1989 (late dry season). For standing root biomass, analyses were performed separately for untransformed biomass of VFR (<1 mm) and FR (1-2 mm). For root ingrowth, analyses were performed for logarithmically transformed biomass of roots < 2 mm in diameter. The transformation was necessary to achieve homoscedastic residuals.

Annual root production was estimated using rates of root ingrowth. Because the IC were in the field for less than 12 months, the productivity measured during the four seasons (dry season = 60 days, dry-wet season transition = 61, wet season = 61 and wet–dry season transition = 68 days) were extrapolated assuming that the seasons were 81, 61, 155, and 68 days in duration, respectively.

Results

Standing root biomass and root ingrowth

Biomass (measured with soil cores) of VFR and FR was significantly higher in irrigated than in control plots (Figure 1). Neither season nor the season-treatment interaction were significant for either VFR (F=0.81, $\alpha = 0.05$; F = 0.65, $\alpha = 0.05$, respect-

MR

(2-5 mm)



Figure 2. Root biomass in the ingrowth cylinders at the end of a 60 day period during the dry, dry-wet, wet and wet-dry seasons of 1988 and 1989. In the upper panel (A) is the biomass of roots less than 1 mm in diameter, in the second panel (B) is the biomass of roots 1-2 mm in diameter, and in the third panel (C) is the biomass of all roots inside the cylinders. (\Box) = Irrigated plots and (\mathbb{S}) = control plots. (*) represent significant differences (p < 0.05) between treatments.

ively) or FR (F=0.31, $\alpha = 0.05$; F = 0.92, $\alpha = 0.05$, respectively).

There was a season × treatment interaction in the productivity of VFR (measured with IC) during the dry season (Figure 2). During the wet season and seasonal transitions, root productivity was not significantly different between irrigated and control plots (p>0.20). Annual root production (of roots <2 mm), estimated from root ingrowth, was not significantly different between irrigated (4.32 Mg ha⁻¹ year⁻¹) and control plots (3.52 Mg ha⁻¹ year⁻¹). Thus, statistically significant dry-season increase under irrigation is offset by non-significant treatment differences in the opposite direction in the wet season and in the transitions between seasons. As a result, year-long estimates of productivity do not show a treatment effect.

Litterfall

Between 7 December 1987 and 29 May 1989, mean daily litterfall was not significantly different between irrigated (3.8 g m⁻² day⁻¹) and control plots (3.7 g m⁻² day⁻¹) as shown in Figure 3. Litterfall increased during the dry season (7.7 g m⁻² day⁻¹ in January 1988) and later decreased to a minimum in mid October (about 2.0 g m⁻² day⁻¹) in both treatments.

Discussion

Treatment effects

The large treatment difference for VFR and FR standing biomass in December 1988 and March 1989, is probably the result of small cumulative increments in biomass under irrigation during the three previous dry seasons. This conclusion is supported by the results of the ingrowth cylinder experiments, where fine-root productivity was significantly higher in irrigated plots only during the dry season.

Fine-root production, measured with ingrowth cylinders, was significantly higher in irrigated than in control plots only during the dry season (Figure 2). Roots grew more in irrigated cylinders, as a response to higher water availability (a direct effect of irrigation), and probably also as a response to softer soils. Soil strength, measured with a soil penetrometer, decreased from 3.5 kg cm⁻² (+/-0.30; n=15) in dry soils (27% water content at the end of the 1992 dry season), to 1.8 kg cm⁻² (+/-0.13; n=15) in saturated soils (45% water content). Because soils are very hard during the dry season, fine-root elongation and exploration are likely to be reduced. Furthermore, soils crack as a result of low water content (around 27% on a percentage wet weight; Windsor, 1990) and high clay content (Cavelier, 1992), probably killing some of the fine-roots and making it difficult for other roots to grow. These cracks usually open at the beginning of the dry season and deepen and widen as the dry season progresses (Cavelier, 1992).

In irrigation and fertilization experiments in temperate conifer forests (*Pseudostuga menziesii* var. *glauca*), fine root production (< 2 mm) decreased in comparison with control plots (Gower et al. 1992). While fertilization with N + P in a nearby semideciduous forest at Gigante península in Panama also resulted in reduction in fine root biomass and production (Cavelier, 1989), irrigation at Poacher's Península on Barro Colorado Island, resulted in only a seasonal



Figure 3. Rainfall (mm), soil water potential (MPa) and fine litter fall (g m⁻² day⁻¹) between December 1988 and March 1989. In the bottom panel, weekly rainfall is presented by the histogram, mean soil water potential in irrigated plots is represented by the solid line at the top of the panel (above -0.04 MPa), and mean soil water potential and standard errors for control plots are represented by the closed circles and error bars (n = 16). Soil water potentials are at 25 cm depths. In the top panel, weekly mean fine litter fall for irrigated and control plots is represented by the solid and broken lines, respectively.

(i.e. dry season) increase in both very fine root biomass (< 1 mm) and production. The increase in fine root biomass and production in irrigated plots during the dry season, may be the response to the amelioration of a seasonal limiting factor (water availability), that drives most of the phenological responses in this forest type. Fine root biomass and production, are likely to decrease year round, in non-seasonal ecosystems with relatively low water availability. Suppression of the dry season at the soil level, had no effect on litterfall, an important component of the above-ground productivity (see also Wright, 1991; Wright and Cornejo, 1990a,b). This result contrasts with the increases in new twigs and foliage in irrigated plots of *Pseudostuga* menziesii var. glauca (Gower et al. 1992). It is possible that irrigation during the dry season had a positive effect on other components of forest productivity not measured in this study, like small branches and total height of trees.

In contrast to the lack of effect of irrigation on litterfall of trees and lianas, irrigation had a weak effect on the phenology of some shrubs (Mulkey et al., 1991; Wright, 1991) and strong effects on photosynthesis, growth and survivorship of herbs and seedlings (Fisher et al., 1991; Mulkey et al., 1991). Changes in relative humidity, and thus in vapour pressure deficits, are probably the cause for changes in stomatal conductance of plants close to the ground but not in the canopy (Sternberg et al., 1989). Indeed, carbon isotopic ratios of leaves from control and irrigated plots were different for understory saplings but not for canopy trees of the same species (Sternberg et al., 1989).

Seasonal variations in root biomass

Strong seasonal variation was measured in fine-root production. In control plots, productivity was high during seasonal transitions and low both during the dry and wet seasons (Figure 2). The peak in root biomass

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in the transition from dry to wet season, was marked by heavy rains (140 mm in May in contrast to 10 mm in April 1988) and the largest amount of leaf litter on the forest (Leigh and Windsor, 1982). This peak in fine-root biomass can be suppressed by the addition of a N+P fertilizer (Cavelier, 1989). Indeed, at a site 1 km from this study, the dry-wet season transition peak in fine-root growth was suppressed by the addition, in three doses, of 300 Kg N ha⁻¹ y⁻¹ and 100 Kg P ha⁻¹ y⁻¹ (Cavelier, 1989). Thus, this peak was probably a response to increasing nutrient availability, rather than to water.

The high productivity during the transition from wet to dry season is more difficult to explain because there are no peaks in N or P availability (Yavitt and Wright, 1996). It is likely that the roots responded to the increase in aeration that results from the decreasing soil water content, or to the build up in exchangeable cations such as K, Ca and Mg (Yavitt and Wright 1996). This peak in root biomass was not observed during the fertilization experiment at Gigante Peninsula (Cavelier, 1989).

While the low root production during the dry season is probably related to low soil water content and hard soils, low wet season productivity is probably related to very high soil water content and a lack of nutrient pulses. A second and smaller peak in fine-root biomass was detected in the same forest type (Gigante Peninsula, part of Barro Colorado Natural Monument), several weeks before the wet season series of ingrowth cylinders was placed at Poacher's Peninsula (Cavelier, 1989). This second peak in fine-root biomass, apparently missed by the ingrowth cylinders of the present study, coincides with the peak in leaf flush of evergreen trees (Leigh and Windsor, 1982).

Thus, seasonality of fine-root production is related to water and mineral pulses and also to the growth of the above-ground biomass. Relationships between fine-root productivity and environmental and biological variables, is also known in temperate forests. For instance, in a 23-y old *Abies amabilis* forest in the north-west USA, conifer fine root biomass peaks in spring and autumn. The spring peak correlates well with the onset of water percolating into the soil from the snow pack (Grier et al., 1981; Vogt et al., 1981).

The experimental results presented here show that water supplementation has a positive effect on one of the below-ground components of productivity, but not on litterfall. In the long term, increases in belowground biomass should have a positive effect on the water and nutrient balance of the trees, that should be reflected in the above-ground biomass and production. Apparently, four years of irrigation and a marginal increase in root biomass and productivity are not enough to stimulate growth that can be measured at the community level. Further research is needed at the species level to study the timing of events in the root and shoot biomass, and the effect of the addition of water and mineral nutrients.

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References

- Aide T M 1988 Herbivory as a selective agent on the timing of leaf production in a tropical understory community. Nature 336, 574– 575.
- Beard J S 1944 Climax vegetation in tropical America. Ecology 25, 127-158.
- Beard JS 1955 The classification of tropical American vegetation types. Ecology 36, 89–100.
- Cavelier J 1989 Root biomass, production and the effect of fertilization in two tropical rain forests. Ph.D. thesis, University of Cambridge, England. 116 p.
- Cavelier J 1992 Fine-root biomass and soil properties in a semideciduous and a lower montane rain forest in Panama. Plant Soil 142, 187–201.
- Croat T B 1978 Flora of Barro Colorado Island. Stanford University Press, Stanford, CA. 943 p.
- Cuevas E and Medina E 1983 Root production and organic matter decomposition in a tierra firme forest of the upper Rio Negro basis. *In* Wurzecoekologie und ihre Nutzanwendung. Int Symp Gumpenstein. Eds W Bohm L Kutshera and E Lichtenegger. pp. 653–666. Irdning, FRG.
- Cuevas E and Medina E 1988 Nutrient dynamics within amazonian forests. II. Fine root growth, nutrient availability and leaf litter decomposition. Oecologia 76, 222–235.
- Daubenmire R 1972 Phenology and other characteristics of tropical semideciduous forest in north-west Costa Rica. J. Ecol. 60, 147– 170.
- Dietrich W E, Windsor D M and Dune T 1982 Geology, climate, and hydrology of Barro Colorado Island. *In* The Ecology of a Tropical Forest. Seasonal Rhythms and Long Term Changes. Eds E G Leigh Jr, A S Rand and D M Windsor. pp. 21–46 Smithsonian Institution Press, Washington, DC.
- Fisher B, Howe H F and Wright S J 1991 Survival and growth of Virola surinamensis seedlings; water augmentation in gap and understory. Oecologia 86, 292–297.

- Folster H and de las Salas G 1976 Litterfall and mineralization in three tropical evergreen forest stands. Colombia. Acta Científica Venezolana 27, 196–202.
- Ford E D and Deans J S 1977 Growth of a Sitka spruce plantation: spatial distribution and seasonal fluctuations of lengths, weights and carbohydrate concentrations of fine roots. Plant Soil 47, 463– 485.
- Foster R B and Brokaw N V L 1982 Structure and history of the vegetation of Barro Colorado Island. In The Ecology of a Tropical Forest. Seasonal Rhythms and Long Term Changes. Eds E G Leigh Jr, A S Rand and D M Windsor. pp 67–82. Smithsonian Institution Press, Washington, DC.
- Franken M, Immler U and Klinge H 1979 Litterfall in inundation, riverine and terra firme forests on central Amazonia. Trop. Ecol. 60, 147–170.
- Frankie G W, Baker H G and Opler P A 1974 Comparative phenological studies of trees in tropical wet and dry forests in the lowlands of Costa Rica. J. Ecol. 62, 881–919.
- Gower S T, Vogt K A and Grier C C 1992 Carbon dynamics of Rocky mountain Douglas-fir: influence of water and nutrient ' availability. Ecol. Monogr. 62, 43–65.
- Gower S T and Vitousek P M 1989 Effects of nutrient amendments on fine root biomass in a primary successional forest on Hawaii. Oecologia 81, 566–568.
- Grier C C, Vogt K A, Keyes M R and Edmonds R L 1981 Biomass distribution and above- and below-ground production in a young and mature *Abies anabilis* zone ecosystems of the Washington Cascades, Can. J. For. Res. 11, 155–167.
- Haines B and Foster R B 1977 Energy flow through litter in a Panamanian forest. J. Ecol 65, 147-155.
- Jordan C and Escalante G 1980 Root productivity in an Amazonian forest. Ecology 61, 14–18.
- Klinge H and Herrera R 1978 Biomass studies in Amazon Caatinga forest in southern Venezuela. 1. Standing crop of composite root mass in selected stands. Trop. Ecol. 19, 93–110.
- Klinge H and Rodriguez W 1968 Litter production in an area of Amazonian terra firme forest. Part I. Litter-fall, organic carbon and total nitrogen contents of litter. Amazoniana 1, 287–302.
- Leigh E G Jr and Windsor D M 1982 Forest production regulation of primary consumers on Barro Colorado Island. *In* The Ecology of a Tropical Forest. Seasonal Rhythms and Long Term Changes. Eds E G Leigh Jr, A S Rand and D M Windsor. pp 111–122. Smithsonian Institution Press, Washington, DC.
- Leigh E G Jr, Rand A S and Windsor D M (Eds) 1982 The Ecology of a Tropical Forest. Seasonal Rhythms and Long Term Changes. Smithsonian Institution Press, Washington, D.C.
- Lund Z F, Pearson RW and Buchanan GA 1970 An implanted soil mass technique to study herbicide effects on root growth. Weed Science 18, 279–281.
- Mulkey S S, Wright S J and Smith A P 1991 Drought acclimation of an understory shrub (*Psychotria limonensis*; Rubiaceae) in a seasonally dry tropical forest in Panama. Amer. J. Bot. 78, 579– 587.
- Mulkey S S, Smith A P and Wright S J 1991 Comparative life history and physiology of two understory neotropical herbs. Oecologia 88, 263–273.

- Ogawa H 1978 Litter production and carbon cycling in Pasoh Forest, Malaysia. Malayan Nature Journal 30, 367–373.
- Persson H 1979 Fine-root production, mortality and decomposition in forest ecosystems. Vegetatio 41, 101–109.
- Piperno D 1990 Fitolitos, arqueología y cambios prehistoricos de la vegetación en un lote de cincuenta hectárcas de la isla de Barro Colorado. Ecología de un Bosque Tropical. Ciclos estacionales y cambios a largo plazo. Eds E.G. Leigh Jr, A S Rand and D M Windsor. pp 153–156. Smithsonian Institution Press, Washington, DC.
- Raich PB and Borchert R 1982 Phenology and ecophysiology of the tropical tree, *Tabebuia neochrysantha* (Bignoniaceae). Ecology 63, 294–299.
- Richards P W 1952 The Tropical Rain Forest. Cambridge University Press, Cambridge, England. 450 pp.
- Sanford RL Jr 1985 Root ecology and successional Amazon forests. Ph.D. thesis, University of California, Berkely, U.S.A. 135 pp.
- Sanford R L Jr 1989a Fine root biomass under a tropical forest light gap opening in Costa Rica. J. Trop. Ecol. 5, 153–159.
- Sanford R L Jr 1989b Root systems of three adjacent old growth Amazon forests and associated transition zones, J. Trop. For. Science 1, 268–279.
- Sanford R L Jr 1990 Fine root biomass under light gap openings in an Amazon rain forest. Oecologia 83, 541-545.
- Schaik C P Van 1986 Phenological changes in a Sumatran rain forest. J. Trop. Ecol. 2, 327–347.
- Stark N and Spratt M 1977 Root biomass and nutrient storage in rainforest oxisols near San Carlos de Rio Negro. Trop. Ecol. 18, 1–9.
- Sternberg L S L, Mulkey S S and Wright S J 1989 Ecological interpretation of leaf carbon isotope ratios: influence of respired carbon dioxide. Ecology 70, 1317–1324.
- Vogt K A, Edmonds R L and Grier C C 1981 Seasonal changes in biomass and vertical distribution of mycorrhizal and fibroustextured conifer fine roots in 23- and 180-year-old subalpine *Abies amabilis* stands. Can. J. For. Res. 11, 223–229.
- Windsor D M 1990 Climate and moisture variability in a tropical rain forest, long term records for Barro Colorado Island, Panama. Smithsonian Contributions to Earth Sciences. 145 pp.
- Wright S J and Cornejo F H 1990a Seasonal drought and leaf fall in a tropical forest. Ecology 71, 1165–1175.
- Wright S J and Cornejo F H 1990b Seasonal drought and the timing of flowering and leaf fall in a tropical forest. *In* The Reproductive Biology of Tropical Forests Plants. Man and the Biosphere Series, UNESCO. Eds K Bawa and M Hadley. pp 49–61. Paris and Parthenon Publishing, Carnforth, England.
- Wright S J 1991 Seasonal drought and the phenology of understory shrubs in a tropical moist forest. Ecology 72, 1643–1657.
- Yavitt J B and Wright S J 1996 Temporal patterns of soil nutrients in panamanian moist forest revealed by ion exchange resin and experimental irrigation. Plant Soil 183, 117–129

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