

Old-growth forests as global carbon sinks

Sebastiaan Luyssaert^{1,2}, E. -Detlef Schulze³, Annett Börner³, Alexander Knohl⁴, Dominik Hessenmöller³, Beverly E. Law², Philippe Ciais⁵ & John Grace⁶

Old-growth forests remove carbon dioxide from the atmosphere^{1,2} at rates that vary with climate and nitrogen deposition³. The sequestered carbon dioxide is stored in live woody tissues and slowly decomposing organic matter in litter and soil⁴. Old-growth forests therefore serve as a global carbon dioxide sink, but they are not protected by international treaties, because it is generally thought that ageing forests cease to accumulate carbon^{5,6}. Here we report a search of literature and databases for forest carbon-flux estimates. We find that in forests between 15 and 800 years of age, net ecosystem productivity (the net carbon balance of the forest including soils) is usually positive. Our results demonstrate that old-growth forests can continue to accumulate carbon, contrary to the long-standing view that they are carbon neutral. Over 30 per cent of the global forest area is unmanaged primary forest, and this area contains the remaining old-growth forests⁷. Half of the primary forests (6×10^8 hectares) are located in the boreal and temperate regions of the Northern Hemisphere. On the basis of our analysis, these forests alone sequester about 1.3 ± 0.5 gigatonnes of carbon per year. Thus, our findings suggest that 15 per cent of the global forest area, which is currently not considered when offsetting increasing atmospheric carbon dioxide concentrations, provides at least 10 per cent of the global net ecosystem productivity⁸. Old-growth forests accumulate carbon for centuries and contain large quantities of it. We expect, however, that much of this carbon, even soil carbon⁹, will move back to the atmosphere if these forests are disturbed.

We conducted a literature search to test the hypothesis that old-growth forests continue to accumulate atmospheric carbon dioxide (CO_2). Site-level estimates of the annual sums of carbon-cycle components were compiled, including those of biometry-based net primary production (NPP), eddy-covariance or biometry-based net ecosystem production (NEP) and chamber-based heterotrophic respiration. The data set was completed with site information related to stand characteristics, standing biomass and stand age. Data were compiled from 519 plot studies that reported one or more components of the carbon cycle. The studies involved boreal ($\sim 30\%$) and temperate ($\sim 70\%$) forests and represented the full range of conditions of such forests, excluding those subjected to experimental treatments such as fertilization and irrigation (Supplementary Information, section 1.1). Tropical forests were excluded from the analysis because only 12 sites were found for which NEP and age estimates are available.

The NEP is the net carbon balance of the forest as a whole, and is the difference between CO_2 uptake by assimilation and losses through plant and soil respiration. On the basis of our global data set we find that in forests between 15 and 800 years old, the NEP is usually positive; that is, the forests are CO_2 sinks (Fig. 1a). The maximum probabilities of finding a single forest to be a source of carbon at 60, 180 and 300 years of age are 0.20, 0.25 and 0.35, respectively. However, the probability of finding an ensemble of ten old-growth forests that are carbon neutral is negligible

(Supplementary Fig. 1). In the small number of case studies on the effect of age on the carbon balance of forests, several have demonstrated some age-related decline in NEP but very few have shown old forests to be sources^{1,2,10–13}. Our NEP estimates suggest that forests 200 years old and above sequester on average $2.4 \pm 0.8 \text{ tC ha}^{-1} \text{ yr}^{-1}$ (tC , tonnes of carbon; Fig. 1a). In our model (Supplementary Information, section 1.3), we find that old-growth forests accumulate $0.4 \pm 0.1 \text{ tC ha}^{-1} \text{ yr}^{-1}$ in their stem biomass and $0.7 \pm 0.2 \text{ tC ha}^{-1} \text{ yr}^{-1}$ in coarse woody debris, which implies that about $1.3 \pm 0.8 \text{ tC ha}^{-1} \text{ yr}^{-1}$ of the sequestered carbon is contained in roots and soil organic matter.

The commonly accepted and long-standing view that old-growth forests are carbon neutral (that is, that photosynthesis is balanced by respiration) was advanced in ref. 6 and was originally based on ten years' worth of data from a single site⁵. It is supported by the observed decline of stand-level NPP with age in plantations^{14,15}, but is not apparent in some ecoregions¹⁶. Yet a decline in NPP is commonly assumed in ecosystem models (Supplementary Information, section 1.4). Moreover, it has led to the view that old-growth forests are redundant in the global carbon cycle.

If, however, the hypothesis of carbon neutrality⁶ were correct, the expected probabilities of observing a sink or source would be equal and around one-half, the average sink strength for a random ensemble of forests 200 years old and above would be zero and the mean CO_2 release from heterotrophic respiration would equal the mean CO_2 sequestration through NPP (thus, the ratio of heterotrophic respiration to NPP would be approximately one). However, we observe this ratio to be well below one on average (Fig. 1b) and not to increase with age. Hence, all three quantitative tests fail to support the hypothesis of carbon neutrality. The currently available data consistently indicate that carbon accumulation continues in forests that are centuries old.

In fact, young forests rather than old-growth forests are very often conspicuous sources of CO_2 (Fig. 1a) because the creation of new forests (whether naturally or by humans) frequently follows disturbance to soil and the previous vegetation, resulting in a decomposition rate of coarse woody debris, litter and soil organic matter (measured as heterotrophic respiration) that exceeds the NPP of the regrowth^{2,17–22} (Fig. 1b).

The scatter in the relationship between NPP and age is considerable, but given the climatic, edaphic and biological diversity of the observations in combination with differences in disturbance histories, this is to be expected. There is some degree of age-related decline in NPP beyond 80 years of age (Fig. 1c), and temperate and boreal forests both show a consistent pattern of declining NPP beyond an early maximum (Supplementary Fig. 2a) when analysed separately. The decline in NPP could be partly attributed to the presence or absence of management (Supplementary Fig. 2b). However, we expect that this decline is not strictly a management effect, but a

¹Department of Biology, University of Antwerp, 2610 Wilrijk, Belgium. ²College of Forestry, Oregon State University, Corvallis, Oregon 97331-5752, USA. ³Max-Planck Institute for Biogeochemistry, 07701 Jena, Germany. ⁴ETH Zürich, Institute of Plant Sciences, CH-8092 Zürich, Switzerland. ⁵Laboratoire des Sciences du Climat et de l'Environnement, IPSL-LSCE, CEA-CNRS-UVSQ, 91191 Gif sur Yvette Cedex, France. ⁶School of GeoSciences, The University of Edinburgh, Edinburgh EH9 3JN, UK.

reflection of differences in disturbance history between managed and unmanaged forests.

Consistent with earlier studies², biomass continues to increase for centuries irrespective of whether forests are boreal or temperate (Supplementary Fig. 3). In the course of succession, plants compete for resources and self-thinning²³ (or thinning by humans in the case of managed forests) occurs (Fig. 2), so the older stands contain a relatively small number of individuals, although of course these trees tend to be large. Obviously biomass cannot accumulate forever. Our data (Supplementary Fig. 3) suggest a possible upper limit somewhere between 500 and 700 tC ha⁻¹ (equivalent to 1,400 to 1,800

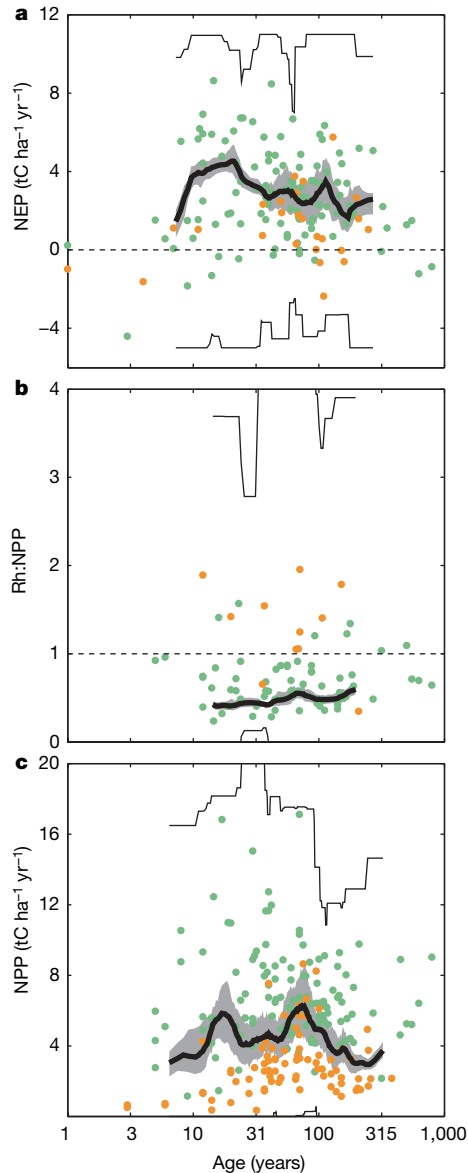


Figure 1 | Changes in carbon fluxes as a function of age. **a**, Observed NEP versus age; positive values indicate carbon sinks and negative values indicate carbon sources. **b**, Observed ratio of heterotrophic respiration (Rh) to NPP versus age; Rh:NPP < 1 indicates a carbon sink. **c**, Observed NPP versus age. It appears that temperate and boreal forests both show a pattern of declining NPP. Most probably, the late-successional increase in NPP is caused by the combination of data from different climate regions or the combination of disturbance regimes (Supplementary Fig. 2a, b). In each panel, the green dots show observations of temperate forests, the orange dots show observations of boreal forests, the thick black line shows the weighted mean within a moving window of 15 observations, the grey area around this line shows the 95% confidence interval of the weighted mean and the thin black lines delineate the 95% confidence interval (where visible) of the individual flux observations.

cubic metres of wood per hectare); these high-biomass forests were located in the Pacific Northwest USA¹⁶.

We speculate that when high above-ground biomass is reached, individual trees are lost because of lightning, insects, fungal attacks of the heartwood by wood-decomposers, or trees becoming unstable in strong wind because the roots can no longer anchor them. If old-growth forests reach high above-ground biomass and lose individuals owing to competition or small-scale disturbances, there is generally new recruitment or an abundant second canopy layer waiting in the shade of the upper canopy to take over and maintain productivity.

Although tree mortality is a relatively rapid event (instantaneous to several years long), decomposition of tree stems can take decades. Therefore, the CO₂ release from the decomposition of dead wood adds to the atmospheric carbon pool over decades, whereas natural regeneration or in-growth occurs on a much shorter timescale. Thus, old-growth forest stands with tree losses do not necessarily become carbon sources, as has been observed in even-aged plantations (that is, where trees are all of the same age). We recognize that self-thinning theory was originally developed and validated for even-aged single-species stands; however, it has been shown to hold for uneven-aged multi-species plant communities (Supplementary Information, section 1.3). In reasonable agreement with our observations (Fig. 1b), self-thinning theory predicts that the ratio between heterotrophic respiration and NPP is constant and around 0.65 ± 0.02 (indicating a carbon sink; Supplementary Fig. 4), as long as stand density is driven by small-scale, rather than stand-replacing, disturbances. Old stands, with sufficiently high densities (that is, through development of a multilayer canopy structure) are thus expected to maintain biomass accumulation for centuries. Hence, we postulate that biomass accumulation and decline are largely driven by stand structure.

A stand must be spared for centuries from stand-replacing disturbances (such as fires, insect outbreaks, wind-throw and avalanches) in order to accumulate sufficient aboveground biomass to become old growth. Because the cumulative probability of disturbances is higher in stands with high above-ground biomass, old stands are rarer than young stands, even in unmanaged landscapes. At the landscape level, we expect a mosaic of forests characterized by different times since the last stand-replacing disturbance²⁴. Despite differences in age and density, these forests are, however, expected to follow the same relationship between biomass and density (Fig. 2).

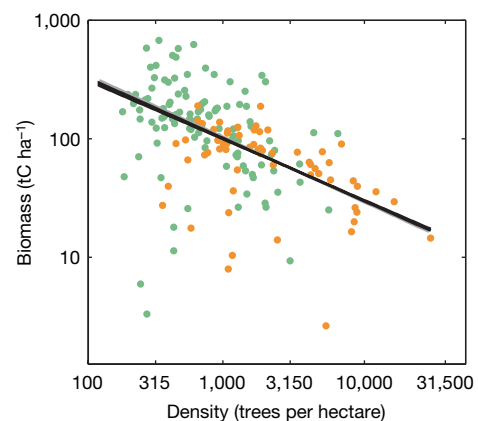


Figure 2 | Biomass accumulation as a function of stand density. Each data point represents a different forest, many of which have different growing conditions and tree species. Not all growing conditions and species compositions allow for the accumulation of the global maximum observed biomass. Self-thinning, the process of density-dependent mortality, is shown (solid line, of slope γ) as the relationship between the logarithm of above-ground biomass and the logarithm of stand density according to ref. 23 ($\gamma = -0.51 \pm 0.08$, $r^2 = 0.25$, $P < 0.01$). The green dots show observations of temperate forests, the orange dots show observations of boreal forests and the grey area (which is barely wider than the solid line) shows the 95% confidence interval of the median.

Under the Kyoto Protocol (<http://unfccc.int/resource/docs/convkp/kpeng.pdf>) only anthropogenic effects on ecosystems are considered (Article 2 of the Framework Convention on Climate Change (<http://unfccc.int/resource/docs/convkp/conveng.pdf>); Supplementary Fig. 5) and the accounting for changes in carbon stock by afforestation, reforestation and deforestations is mandatory (Article 3.3), operating from a base line of 1990. Leaving forests intact was not perceived as an anthropogenic activity. In addition, the potential consequences of excluding old-growth forests from national carbon budgets and from the Kyoto Protocol were downplayed in the carbon-neutrality hypothesis⁶. However, over 30% (1.3×10^9 ha) of the global forest area is classified⁷ by the Food and Agriculture Organization of the United Nations as primary forest, and this area contains the world's remaining old-growth forests. Half (0.6×10^9 ha) of the primary forests are located in the boreal and temperate regions of the Northern Hemisphere. On the basis of our analysis, we expect that these forests alone sequester at least $1.3 \pm 0.5 \text{ GtC yr}^{-1}$. Hence, 15% of the global forest surface, which is currently not being considered for offsetting increasing atmospheric CO₂ concentrations, is responsible for at least 10% of the global NEP⁸. Sporadic disturbances will interrupt carbon accumulation, implying that net biome productivity²⁵ will be lower, but it will remain positive as demonstrated by the accumulation of carbon in soils^{4,26}, coarse woody debris and charcoal^{27,28}.

The present paper shows that old-growth forests are usually carbon sinks. Because old-growth forests steadily accumulate carbon for centuries, they contain vast quantities of it. They will lose much of this carbon to the atmosphere if they are disturbed, so carbon-accounting rules for forests should give credit for leaving old-growth forest intact.

METHODS SUMMARY

We conducted a literature and database search to determine the fate of the carbon sequestered in forests. Observation-based estimates were compiled for carbon-cycle components, including biometry-based NPP, eddy-covariance or biometry-based NEP and chamber-based heterotrophic respiration²⁹. The data set was extended with site information related to stand characteristics, standing biomass and stand age. In general, uncertainties in flux estimates were not reported in the literature. Therefore, we estimated the total uncertainty for every component flux contained in the data set using a consistent framework based on expert judgment (Supplementary Information, section 1.2). The uncertainty framework in our database was designed to account for differences in data quality between sites due to length of time series, methodology and conceptual difficulties (that is, gap filling and dark respiration). Also, an uncertainty of 20% was assigned to the biomass, age and density estimates. These uncertainties were propagated through the statistical analyses by means of random realizations based on Monte Carlo principles. Within each of the 1,000 random realizations, normally distributed random errors, based on the uncertainty framework of our database, were added to the observed fluxes. Therefore, all results that are based on flux data are reported as the weighted mean and the 95% confidence interval of the probability distribution.

Despite the climatic, edaphic and biological diversity of our observations, above-ground biomass was observed to be related to stand density in the way described by self-thinning theory²³. Although, this theory was initially developed for even-aged single-species plant communities, we applied it to our data (Supplementary Information, section 1.3) to determine the components of the flux-computed NEP, specifically the above-ground biomass, woody debris and soil sequestration. Furthermore, self-thinning theory was used to calculate the theoretical ratio of heterotrophic respiration to NPP and compare it with the observed ratio in support of the hypothesis that biomass accumulation and decline are largely driven by stand structure.

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- Carey, E. V., Sala, A., Keane, R. & Callaway, R. M. Are old forests underestimated as global carbon sinks? *Glob. Change Biol.* **7**, 339–344 (2001).
- Pregitzer, K. S. & Euskirchen, E. S. Carbon cycling and storage in world forests: biome patterns related to forest age. *Glob. Change Biol.* **10**, 2052–2077 (2004).
- Magnani, F. *et al.* The human footprint in the carbon cycle of temperate and boreal forests. *Nature* **447**, 848–850 (2007).
- Zhou, G. Y. *et al.* Old-growth forests can accumulate carbon in soils. *Science* **314**, 1417 (2006).
- Kira, T. & Siddei, T. Primary production and turnover of organic matter in different forest ecosystems of the western Pacific. *Jpn. J. Ecol.* **17**, 70–87 (1967).

- Odom, E. P. The strategy of ecosystem development. *Science* **164**, 262–270 (1969).
- FAO. *Global Forest Resources Assessment 2005. Progress towards sustainable forest management*. Forestry Paper 147 (Food and Agriculture Organization of the United Nations, 2006).
- Bolin, B. *et al.* in *IPCC, Land Use, Land-Use Change, and Forestry. A Special Report of the IPCC* (eds Watson, R. T. *et al.*) 23–51 (Cambridge Univ. Press, 2000).
- Fontaine, S. *et al.* Stability of organic carbon in deep soil layers controlled by fresh carbon supply. *Nature* **450**, 277–280 (2007).
- Acker, S. A., Halpern, C. B., Harmon, M. E. & Dyrness, C. T. Trends in bole biomass accumulation, net primary production and tree mortality in *Pseudotsuga menziesii* forests of contrasting age. *Tree Physiol.* **22**, 213–217 (2002).
- Knohl, A., Schulze, E. D., Kolle, O. & Buchmann, N. Large carbon uptake by an unmanaged 250-year-old deciduous forest in Central Germany. *Agric. For. Meteorol.* **118**, 151–167 (2003).
- Law, B. E. *et al.* Changes in carbon storage and fluxes in a chronosequence of ponderosa pine. *Glob. Change Biol.* **9**, 510–524 (2003).
- Desai, A. R. *et al.* Comparing net ecosystem exchange of carbon dioxide between an old-growth and mature forest in the upper Midwest, USA. *Agric. For. Meteorol.* **128**, 33–55 (2005).
- Gower, S. T., McMurtrie, R. E. & Murty, D. Aboveground net primary production decline with stand age: Potential causes. *Trends Ecol. Evol.* **11**, 378–382 (1996).
- Binkley, D. *et al.* Age-related decline in forest ecosystem growth: an individual-tree, stand-structure hypothesis. *Ecosystems* **5**, 58–67 (2002).
- Van Tuyl, S., Law, B. E., Turner, D. P. & Gitelman, A. I. Variability in net primary production and carbon storage in biomass across Oregon forests—an assessment integrating data from forest inventories, intensive sites, and remote sensing. *For. Ecol. Manage.* **209**, 273–291 (2005).
- Harmon, M. E., Ferrell, W. K. & Franklin, J. F. Effects on carbon storage of conversion of old-growth forests to young forests. *Science* **247**, 699–702 (1990).
- Janish, J. E. & Harmon, M. E. Successional changes in live and dead wood carbon stores: implications for net ecosystem productivity. *Tree Physiol.* **22**, 77–89 (2002).
- Wirth, C., Czimczik, C. I. & Schulze, E. D. Beyond annual budgets: carbon flux at different temporal scales in fire-prone Siberian Scots pine forests. *Tellus* **54**, 611–630 (2002).
- Knohl, A. *et al.* Carbon dioxide exchange of a Russian boreal forest after disturbance by wind throw. *Glob. Change Biol.* **8**, 231–246 (2002).
- Kowalski, A. S. *et al.* Paired comparisons of carbon exchange between undisturbed and regenerating stands in four managed forests in Europe. *Glob. Change Biol.* **10**, 1707–1723 (2004).
- Irvine, J., Law, B. E. & Hibbard, K. A. Postfire carbon pools and fluxes in semiarid ponderosa pine in Central Oregon. *Glob. Change Biol.* **13**, 1748–1760 (2007).
- Yoda, K., Kira, T., Ogawa, H. & Hozumi, K. Self-thinning in overcrowded pure stands under cultivated and natural conditions. *J. Biol. Osaka City Univ.* **14**, 107–132 (1963).
- Mladenoff, D. J., White, M. A., Pastor, J. & Crow, T. R. Comparing spatial pattern in unaltered old-growth and disturbed forest landscapes. *Ecol. Appl.* **3**, 294–306 (1993).
- Chapin, F. S. *et al.* Reconciling carbon-cycle concepts, terminology and methodology. *Ecosystems* **9**, 1041–1050 (2005).
- Ciais, P. *et al.* in *The Carbon Balance of Forest Biomes* (eds Griffith, H. & Jarvis, P.) 109–150 (Taylor and Francis, 2005).
- Jomura, M. *et al.* The carbon budget of coarse woody debris in a temperate broad-leaved secondary forest in Japan. *Tellus B* **59**, 211–222 (2007).
- Preston, C. M. & Schmidt, M. W. I. Black (pyrogenic) carbon: a synthesis of current knowledge and uncertainties with special consideration of boreal regions. *Biogeosciences* **3**, 397–420 (2006).
- Luyssaert, S. *et al.* CO₂-balance of boreal, temperate and tropical forest derived from a global database. *Glob. Change Biol.* **13**, 2509–2537 (2007).

Supplementary Information is linked to the online version of the paper at www.nature.com/nature.

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