

# Root symbioses as belowground C pumps: a mitigation strategy against rising CO<sub>2</sub> levels

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## Abstract

Mycorrhizal and N-fixing root symbioses evolved at two points in the past when global CO<sub>2</sub> was highest, consistent with the high demand these symbioses place on host C. Trees hosting both mycorrhiza and N-fixing bacteria are able to fix more atmospheric CO<sub>2</sub> and grow at faster rates than non-symbiotic plants, or plants with only mycorrhiza. We argue that on the basis of this improved C capture, N-fixing trees act as C-pumps, sequestering C and locking it in biomass, thus, if properly managed, can contribute significantly towards the mitigation of rising CO<sub>2</sub> levels.

**Citation:** Mortimer PE, Hammond J, Hyde KD, Gui H, Xu J. 2021. Root symbioses as belowground C pumps: a mitigation strategy against rising CO<sub>2</sub> levels. *Circular Agricultural Systems* 1: 9 <https://doi.org/10.48130/CAS-2021-0009>

## Introduction

Around 80% of plant species form symbiotic relationships with mycorrhizal fungi, which assist the plant in accessing soil nutrients beyond their roots<sup>[1]</sup>. A second, less widespread symbiosis between plant roots and N-fixing bacteria (symbiotic nitrogen fixation – SNF), allows for the further provision of N to the host plant. In return, host plants provide the symbionts with photosynthetically derived carbohydrates<sup>[2]</sup>. The improved nutrition from these symbioses enables plants to maintain higher photosynthetic rates, thus fixing more atmospheric CO<sub>2</sub>. Furthermore, if both symbioses occur in the same root system, there is an enhanced synergistic effect on plant productivity and symbiont performance<sup>[3]</sup>.

This article presents the case that the co-evolution of these root symbioses contributed towards the decline in atmospheric CO<sub>2</sub> levels at two points in the Earth's history. The potential for harnessing these symbioses to address current planetary system challenges are discussed.

## Evolution of root symbioses linked to global CO<sub>2</sub> levels

During the Phanerozoic Eon, there were two peaks in temperature and CO<sub>2</sub> concentrations (Fig. 1). The first peak in CO<sub>2</sub> occurred between the Cambrian and Devonian Periods (540–420 million years ago (Ma)), coinciding with the earliest mycorrhizal records<sup>[4,5]</sup>. The second was during the late Jurassic and early Cretaceous Periods (150–100 Ma), at approximately the same time when the first precursors of symbiotic N-fixation evolved (ca. 100 Ma) (Fig. 1)<sup>[6]</sup>.

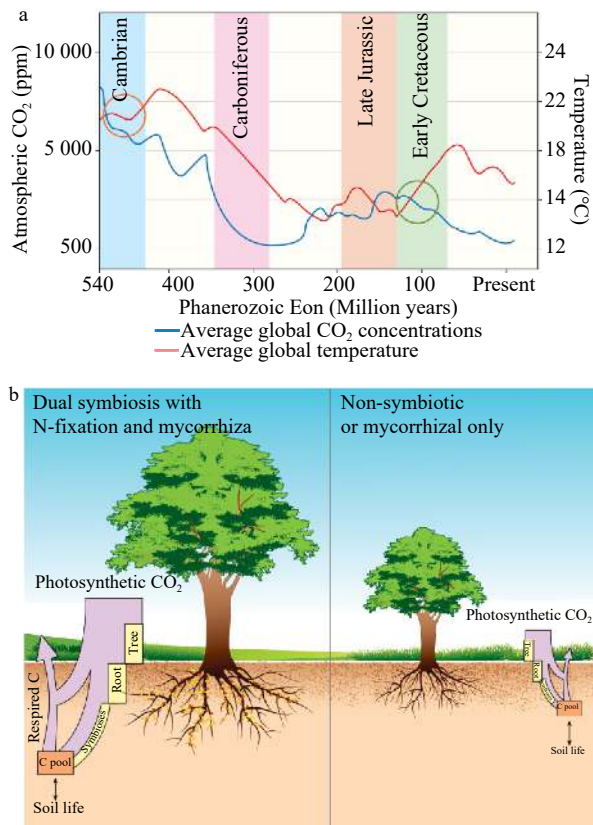
These symbioses evolved during periods of elevated atmospheric CO<sub>2</sub> concentrations, when plants could easily provide the additional C for the symbionts. During the late

Devonian Period, trees and larger plants evolved and began to dominate the landscape<sup>[7]</sup>. There is evidence that mycorrhizal symbioses were critical to this process<sup>[8]</sup>. During the following Carboniferous Period (ca. 360–300 Ma), forest systems became widespread and atmospheric CO<sub>2</sub> declined to around 300 ppm (Fig. 1a). Average temperatures also declined by about 10 °C (Fig. 1a). This climatic change led to the Carboniferous Rainforest Collapse<sup>[9]</sup>, a period when many forest systems died, depositing large amounts of C belowground.

With the decline of forest systems, global CO<sub>2</sub> levels began to rise, possibly because Agaricomycetes, the first organisms capable of degrading lignin and releasing CO<sub>2</sub>, had evolved<sup>[10]</sup>. In addition to high CO<sub>2</sub> levels, temperatures rose to levels more than double those experienced today (Fig. 1a). The first evidence of SNF coincides with high temperatures and CO<sub>2</sub> levels around 100 Ma. During this period, plants with N-fixing symbioses proliferated and colonized various landscapes, acting as pioneer species in soils with low N<sup>[11]</sup>. These changes were followed by another decline in global CO<sub>2</sub> levels that continued to decline until the Quaternary Period (Fig. 1a).

## Root symbioses as carbon pumps

Plants produce organic C compounds by fixing atmospheric CO<sub>2</sub> through photosynthesis. This C is used for the plant's own growth; an additional 25% can be consumed by mycorrhiza<sup>[12]</sup> and a further 28% by N-fixing bacteria<sup>[13,14]</sup>. To maintain these symbioses, the photosynthetic rates of dual inoculated host plants can increase by over 50%<sup>[14]</sup>. These figures only represent the C directly attributed to growth and maintenance of the symbionts. Increased C-flow through root exudates and increased plant biomass are difficult to



**Fig. 1** Atmospheric carbon dioxide (CO<sub>2</sub>, ppm) and temperature (°C) levels during the Phanerozoic Eon (540 Ma to present) (a). Data adapted from [5], [20], [21]. The highlighted portions indicate relevant periods in the past: light blue indicates the Devonian Period, purple the Carboniferous Period, light orange the late Jurassic Period, and light green the early Cretaceous Period. The proposed periods for the development of the mycorrhizal symbioses are circled in orange and that of symbiotic N-fixation in green. The illustrative description of the C-pumps model proposed for N-fixing trees shows the relative movement of C through these systems (b). The relative proportion of C captured via photosynthesis is represented by the purple box, with as much as 50% more C fixed in dual symbiotic plants compared to non-symbiotic plants<sup>[10]</sup>. The yellow boxes represent the relative amounts of C stored in biomass, with N-fixing trees growing as much as nine times faster than their non-N-fixing counterparts<sup>[20]</sup>.

accurately quantify but are significant additional contributions to C sequestration. Plants with both symbioses are therefore greater C sinks, moving a higher percentage of C belowground, and acquiring more C for growth (Fig. 1b). There are also beneficial downstream effects: N-fixing plants can cause improved soil nutrition and growth of surrounding plants<sup>[15]</sup>.

### Implications for forestry and agriculture in the Anthropocene

The Anthropocene describes the period of recent history in which humankind's impact on the earth has been globally significant. The Anthropocene is characterised by rising temperatures, erratic weather events, biodiversity loss,

chemical pollution, forest loss, and high-input agricultural systems<sup>[16]</sup>. In 'On the Possibilities of a Charming Anthropocene', Buck (2015) highlights the need for a shift in focus from the negative impacts of this period towards the possibilities for stimulating positive changes<sup>[17]</sup>. We posit the increased usage of plants with both forms of root symbioses as one such positive intervention, with particular potential in tropical regions.

The tropics is home to about 68% of legume species, which comprise the vast majority of plants capable of SNF; and more than 50% of all biological N-fixation takes place in the tropics<sup>[18,19]</sup>. Furthermore, N is often the most limiting factor for plant growth, and therefore C sequestration, in the tropics. Batterman et al (2013) showed that in tropical forests N-fixing trees can accumulate C nine times faster than non N-fixing trees, and that younger forest systems have much higher C and N accumulation rates<sup>[20]</sup>. Thus an ideal means of landscape restoration and C sequestration would be to plant N-fixing tree species in disturbed and/or agricultural landscapes within tropical areas.

However, we could go further than simply modifying natural regeneration to favor SNF plant species. We could use new forest restoration technologies, such as superior genetic materials, pre-inoculation of saplings with symbionts, rotational forest management with sustainable timber harvest (for C storage), and superior agroecology and agroforestry systems<sup>[21]</sup>.

The projections indicate that CO<sub>2</sub> levels will continue to rise, temperature will increase, and the tropics will become drier. Evidence from past geological time periods suggest that under these conditions, we can expect the rate of C accumulation by SNF trees to increase<sup>[22]</sup>. The current direction of global climatic change provides a positive feedback for N-fixing trees, and their role in C sequestration could offer a useful buffering effect.

Improved understanding and management of N-fixing trees will enhance the contribution of agriculture and forestry to climate change mitigation, ecosystem resilience, food security and environmental sustainability. Future agricultural systems that encourage the planting of trees and diversification of agricultural landscapes will increase sustainability and productivity. We therefore propose harnessing the C pump and ecosystem enhancement capacities of N-fixing trees as a means of taking us forward into a new, positive era.

### ACKNOWLEDGMENTS

This work was generously supported by the Key Project from the Ministry of Sciences and Technology of China (No: 2017YFC0505101).

### Conflict of interest

The authors declare that they have no conflict of interest.

### Dates

Received 24 May 2021; Accepted 25 May 2021; Published online 29 June 2021

## REFERENCES

1. Ahmad P, Prasad MNV. 2012. *Environmental adaptations and stress tolerance of plants in the era of climate change*. New York: Springer New York. xvi, 515 pp. <https://doi.org/10.1007/978-1-4614-0815-4>
2. Puri A, Padda KP, Chanway CP. 2020. Can naturally-occurring endophytic nitrogen-fixing bacteria of hybrid white spruce sustain boreal forest tree growth on extremely nutrient-poor soils? *Soil Biology & Biochemistry* 140:107642
3. Zhu C, Tian G, Luo G, Kong Y, Guo J, et al. 2018. N-fertilizer-driven association between the arbuscular mycorrhizal fungal community and diazotrophic community impacts wheat yield. *Agriculture Ecosystems & Environment* 254:191–201
4. Brundrett MC. 2002. Coevolution of roots and mycorrhizas of land plants. *New Phytologist* 154:275–304
5. Orme DA, Surpless KDJG. 2019. The birth of a forearc: The basal Great Valley Group, California, USA: REPLY. *Geology* 47:e494
6. Werner GDA, Cornwell WK, Sprent JI, Kattge J, Kiers ET. 2014. A single evolutionary innovation drives the deep evolution of symbiotic N<sub>2</sub>-fixation in angiosperms. *Nature Communications* 5:4087
7. Algeo TJ, Scheckler SE. 1998. Terrestrial-marine teleconnections in the Devonian: links between the evolution of land plants, weathering processes, and marine anoxic events. *Philosophical Transactions of the Royal Society of London Series B: Biological Sciences* 353:113–30
8. Humphreys CP, Franks PJ, Rees M, Bidartondo MI, Leake JR, et al. 2010. Mutualistic mycorrhiza-like symbiosis in the most ancient group of land plants. *Nature Communications* 1:103
9. Dunne EM, et al. 2018. Diversity change during the rise of tetrapods and the impact of the 'Carboniferous rainforest collapse'. *Proceedings of the Royal Society B: Biological Sciences* 285:20172730
10. Floudas D, Binder M, Riley R, Barry K, Blanchette RA, et al. 2012. The Paleozoic origin of enzymatic lignin decomposition reconstructed from 31 fungal genomes. *Science* 336:1715–19
11. Manzanedo RD, HilleRisLambers J, Rademacher TT, Pederson N. 2020. Evidence of unprecedented rise in growth synchrony from global tree ring records. *Nature Ecology & Evolution* 4:1622–29
12. Terrer C, Vicca S, Hungate BA, Phillips RP, Prentice IC. 2016. Mycorrhizal association as a primary control of the CO<sub>2</sub> fertilization effect. *Science* 353:72–74
13. Chen Y, Yu Z, Wang J, Zhang X. 2014. Allocation of photosynthetic carbon to nodules of soybean in three geographically different Mollisols. *European Journal of Soil Biology* 62:60–65
14. Kaschuk G, Kuypers TW, Leffelaar PA, Hungria M, Giller KE. 2009. Are the rates of photosynthesis stimulated by the carbon sink strength of rhizobial and arbuscular mycorrhizal symbioses? *Soil Biology and Biochemistry* 41:1233–44
15. Zhao M, Zhao J, Yuan J, Hale L, Wen T, et al. 2021. Root exudates drive soil-microbe-nutrient feedbacks in response to plant growth. *Plant, Cell & Environment* 44:613–28
16. Rull V. 2016. The 'Anthropocene': A requiem, for the Geologic Time Scale? *Quaternary Geochronology* 36:76–77
17. Buck HJ. 2015. On the possibilities of a charming Anthropocene. *Annals of the Association of American Geographers* 105:369–77
18. Daryanto S, Wang L, Jacinthe PA. 2015. Global synthesis of drought effects on food legume production. *PLoS One* 10:e0127401
19. Cleveland CC, Townsend AR, Schimel DS, Fisher H, Howarth RW, et al. 1999. Global patterns of terrestrial biological nitrogen (N<sub>2</sub>) fixation in natural ecosystems. *Global Biogeochemical Cycles* 13:623–45
20. Batterman SA, Hedin LO, van Breugel M, Ransijn J, Craven DJ, et al. 2013. Key role of symbiotic dinitrogen fixation in tropical forest secondary succession. *Nature* 502:224–27
21. Boinot S, Fried G, Storkey J, Metcalfe H, Barkaoui K, et al. 2019. Alley cropping agroforestry systems: Reservoirs for weeds or refugia for plant diversity? *Agriculture, Ecosystems & Environment* 284:106584
22. Laganière J, Angers DA, Paré D. 2010. Carbon accumulation in agricultural soils after afforestation: a meta-analysis. *Global Change Biology* 16:439–53



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