Including more plasticity of functional traits in a DGVM

Functional diversity in DGVMs: The PFT problem

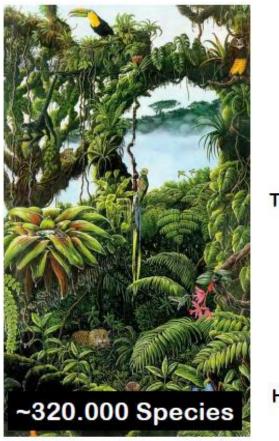
- Plant Functional Types (PFTs): represent whole biomes e.g. *"*tropical broadleaved evergreen tree"
- Fixed parameter settings (e.g. leaf longevity)

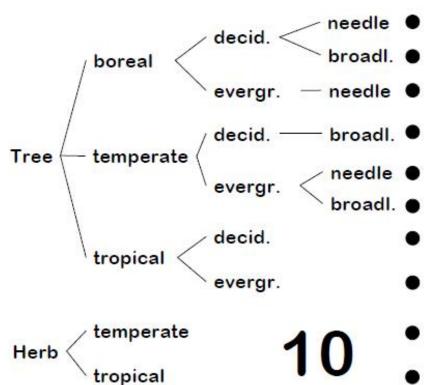
same plant on biome level

no adaptation to environment

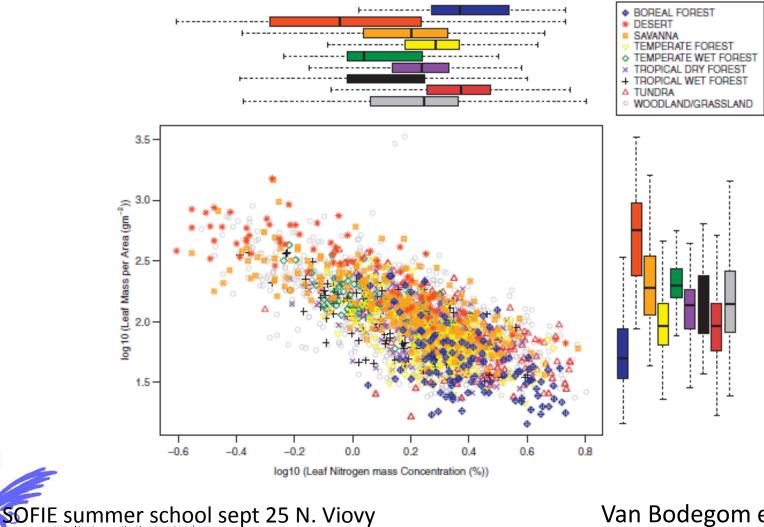
no biodiversity effects on ecosystem processes

A view of diversity in DGVMs





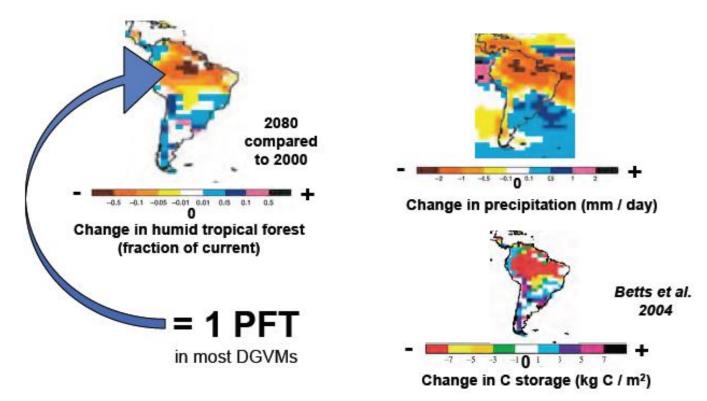
Trait variation is only partly captured by classification in biomes/PFTs



Van Bodegom et al. 2012 GEB

An example of PFT limitation

• Dieback of amazonian forest



first CLIMATE/CARBON coupled simulation show a total dieback of the amazonian Forest in 2080 with a high feedback of climate SOFIE summer school sept 25 N. Viovy

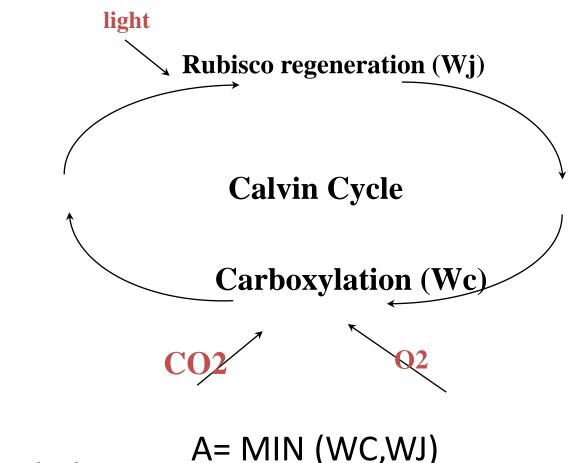
How to introduce more plasticity in DGVMs?

- 1. define empirical relationships between traits and environement (Verheijen et al. 2013)
- Use ecological concepts of optimality (e.g coordination *(Chen et al. 1993)* Use tradeoff between traits *(Wright et al. 2004)*

First attempt to use these concepts in ORCHIDEE DGVM

The coordinated photosynthesis

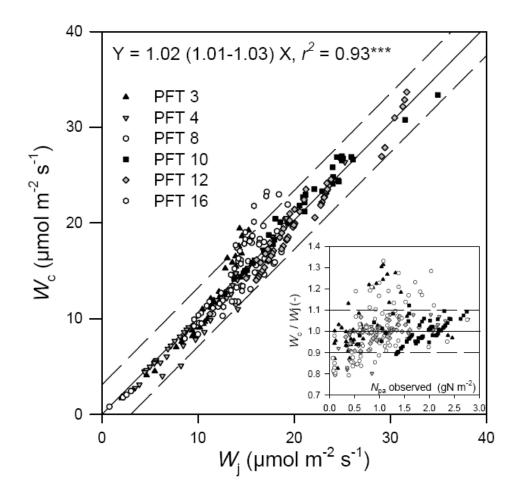
• Reminder on photosynthesis:



- Wc is proportional to rubisco concentration (i.e Vcmax)
 And rubisco is the main source of N
 (Vcmax=k3*N)
- Wj is proportional to energy (light available)
- ➔ If N is too low, Wc is the main limiting factor and A is not optimal
- ➔ If N is sufficiently high Wj become the limiting factor and then additional N does not increase A and additional N intake has a cost !

So there is an optimum leaf N (Vcmax) concentration that plant will try to reach for which Wc=Wj

This is indeed observed in the reality



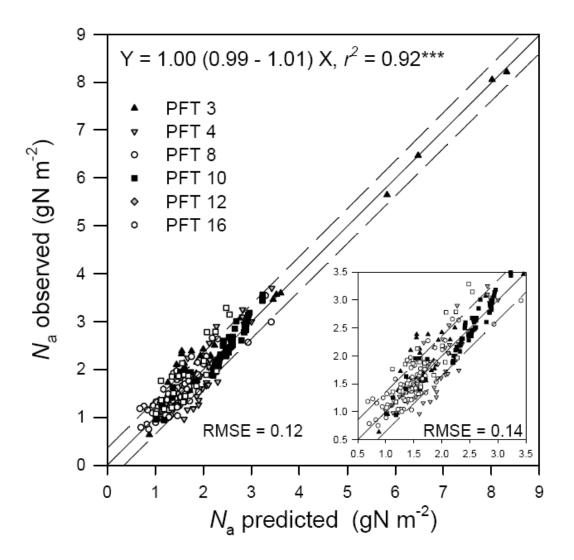
Synthesis from try database (Maire et al. 2010)

• From inversion of the Farqhuar model we can estimate the N of coordinated photosynthesis:

$$N_{\text{pac}} = \frac{4 \cdot \alpha \cdot \text{PPFD}}{k_3^{\text{ac}}} \cdot \left(\left(\frac{C_i + k_2}{\left(4 \cdot C_i + 8 \cdot \Gamma^*\right) \cdot \Phi_{V_{\text{c_{max}}}}} \right)^2 - \left(\frac{1}{J_{\text{fac}}^{\text{atc}} \cdot \Phi_{J_{\text{max}}}} \right)^2 \right)^{1/2}$$

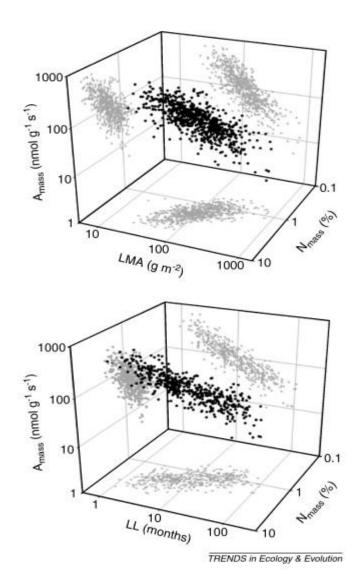
. č.

It gives satisfying results

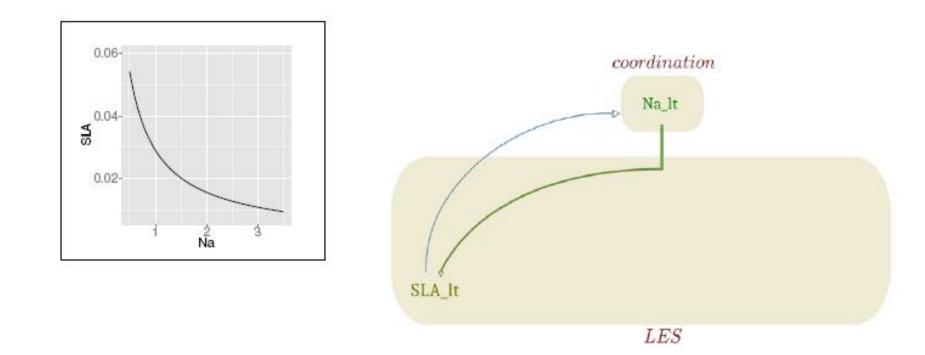


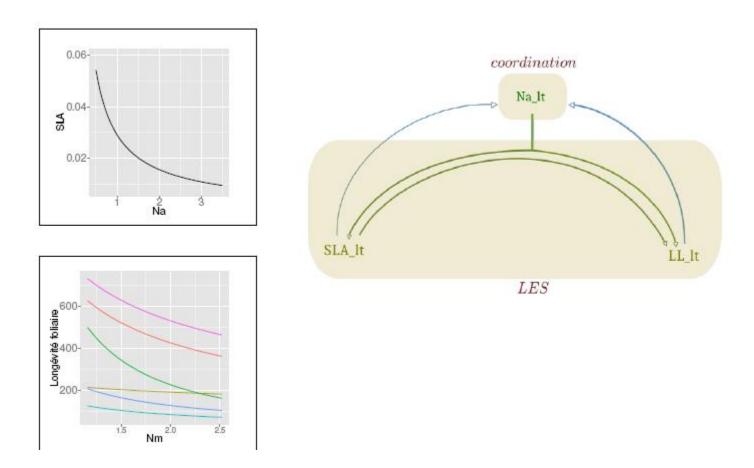
Coupling with the Leaf economic spectrum

(Wright et al. 2004) showed that there is a correlation between 3 main parameters: Na, SLA, leaf longevity.

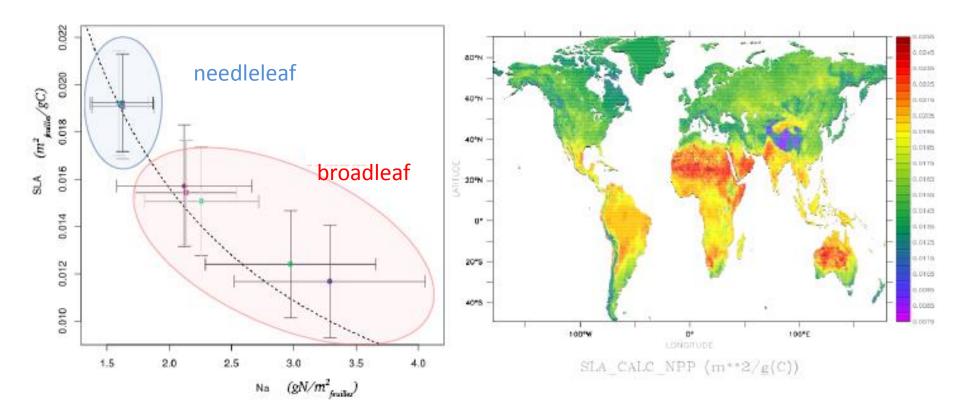


Algorithm



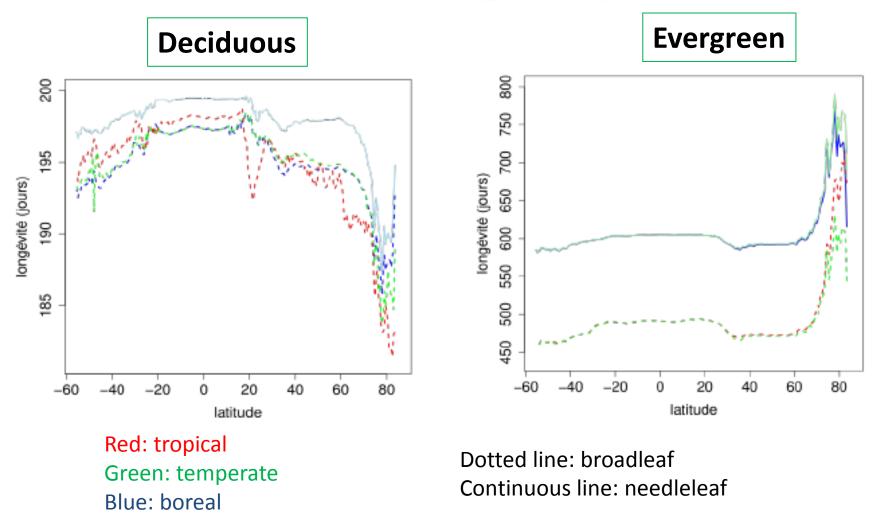


Simulated SLA

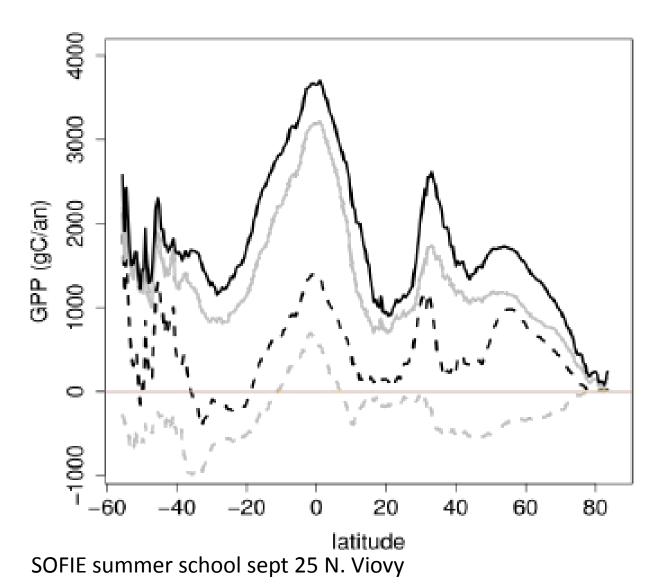


Good agreement between simulated and observed relationship

Leaf longevity



Difference in simulated GPP (with standard version)



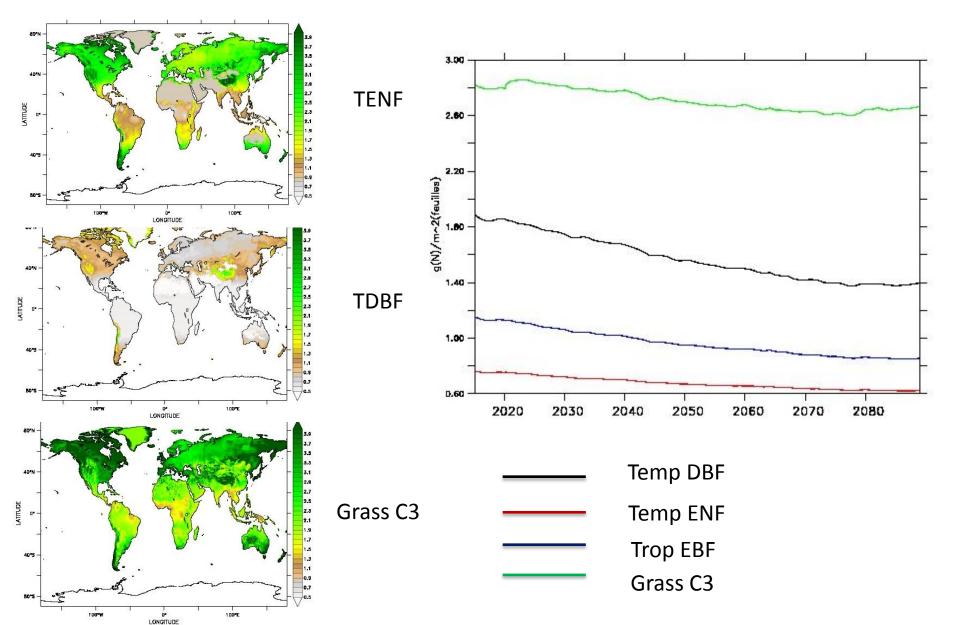
Dotted: broadleaf Cont.: needle leaf Black: sempervirens Grey: deciduous

Coherent within the PFT envoloppe. Divergence outside

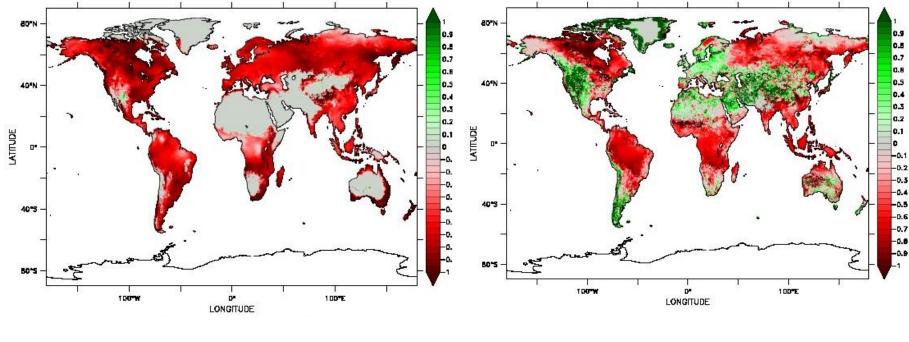
Impact of climate change on predicted traits

Simulation with ISIMIP scenario IPSL rcp8.5

Impact on coordinated Nitrogen



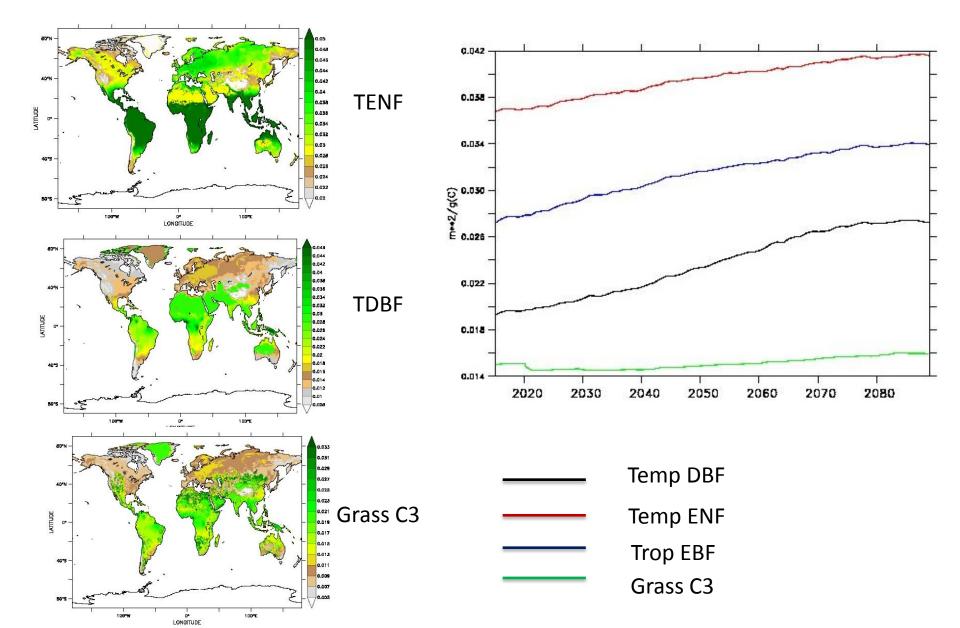
Spatial differences of N between 2100 and 2010



Temp DBF

Grass C3

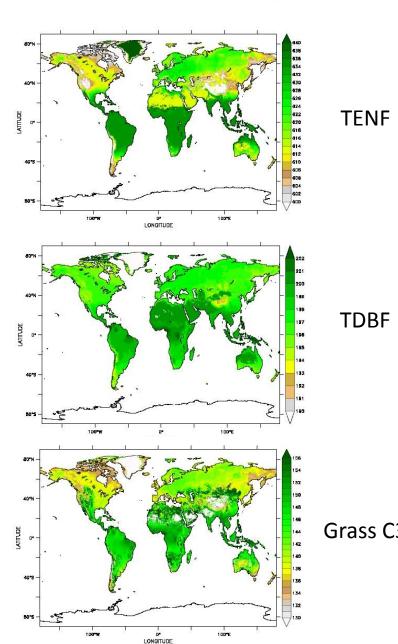
Impact on simulated SLA

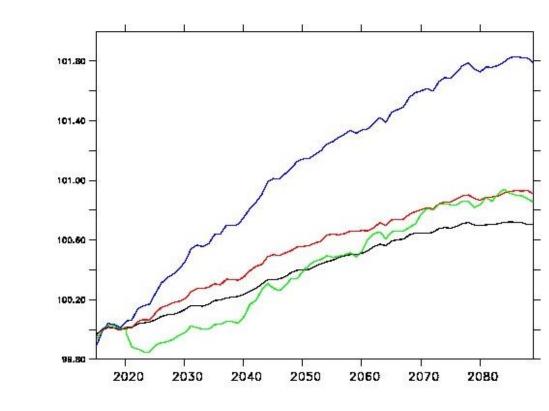


Impact on leaf longevity

TENF

TDBF





3	

Temp DBF Temp ENF Trop EBF

Grass C3

conclusion

- Preliminary results but:
- Simulated N/SLA/LL is coherent with expected variation between PFTs
- ➔ simulated GPP is correct where PFT is present But diverge outside of climate enveloppe
- Decreasing N with climate change (CO2 effect ?)
- But different response for different PFTs.