An overview of the terrestrial carboncycle from a global modelling perspective

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Figure from Ciais et al. (2013)





$d[CO_2]/dt = E_{FF} + E_{LUC} - S_0 - S_L$

Figure from Ciais et al. (2013)



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Basic elements of the terrestrial carbon cycle:



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Non-exhaustive overview of available measurements for terrestrial carbon cycle

Flux towers

- GPP
- NEE
- Total respiration

Spatial coverage: point Frequency: <1 sec



Satellites

- NDVI
- Canopy height
- GPP (derived, e.g. from NDVI)

Spatial coverage: regional/ global Frequency: days-weeks

Inventories

- Biomass
- Growth rates
- Allometry

Spatial coverage: plotlandscape Frequency: years





Typically we use process-based models to study the terrestrial biosphere.

The main models in use today have evolved from a variety of backgrounds, which make each differentially suited to the study of different aspects of the biosphere.



Terrestrial biosphere models must span a range of spatial scales...



Terrestrial biosphere models must span a range of spatial scales...



... and also timescales



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Representations of key ecosystem processes in global terrestrial models

- Boundary conditions
- Primary production
- Respiration
- Structures and pools
- Species and vegetation dynamics
- Managed landcovers

Projecting the global carbon cycle - examples

Boundary conditions

The key boundary conditions for terrestrial biosphere models are usually atmospheric:

- Incoming short-wave radiation
- Surface/Air temperature (sometimes surface temp. is calculated explicitly from other variables)
- Precipitation
- Atmospheric CO₂ mixing ratio
- Incoming long-wave radiation
- Humidity
- Wind-speed
- Nitrogen/Phosphorus deposition

Others are:

• Soil parameters

- Vegetation type maps (not in DGVMs)
- Landcover/land-use/management variables

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Come from either historical reconstructions, or based on future scenarios (based on assumed human trajectories of e.g. emissions, management)



Photosynthesis is modelled as minimum of two limiting rates:



Arneth et al., 2014, Slide A. Arneth

Ecological Climatology, Chapter 9, © G. Bonan (2002)

$$6CO_2 + 6H_2O + hv \rightarrow C_6H_{12}O_6 + 6O_2 \qquad \lambda = 400-700 \text{ nm}$$

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Primary production



Guanter et al. (2014)

Autotrophic respiration

Plant respiration usually divided into growth and maintenance respiration Example functions from LPJ/LPJ-GUESS:



Growth:

$$R_{\rm g} = 0.25 \times (GPP - R_{\rm leaf} + R_{sapwood} + R_{root})$$

Further reading: Thornley and Cannell (2000), Annals of Botany

Autotrophic respiration

Plant respiration usually divided into growth and maintenance respiration Example functions from LPJ/LPJ-GUESS:

$$R_{\text{leaf}} = r \cdot \frac{C_{\text{leaf}}}{cn_{\text{leaf}}} \phi \cdot g(T)$$

$$R_{\text{sapwood}} = r \cdot \frac{C_{\text{sapwood}}}{cn_{\text{sapwood}}} g(T)$$

BUT: Hugh simplification of underlying biochemistry. Acclimatisation could moderate response in warm future climates. Model appears to fail under drought conditions.

Rate is tissue dependent – related to nitrogen content

$$g(T) = \exp\left[308.56 \cdot \left(\frac{1}{56.02} - \frac{1}{(T+46.02)}\right)\right]$$

Strong dependence on temperature

Growth:

$$R_{\rm g} = 0.25 \text{ X } (GPP - R_{\rm leaf} + R_{sapwood} + R_{root})$$

Further reading: Thornley and Cannell (2000), Annals of Botany

Heterotrophic respiration

Respiration by decomposers in the soil.

Typically global models use simple lifetime functions modified by temperature and moisture:



Heavily over-simplified in most global models. Is actually a function of a whole range of factors – bacteria, substrate, moisture, N, temperature. See, e.g. Koven et al. (2013)

Net ecosystem flux balance



Bottom figure Sitch et al. (2003)

Model structure

We model the effect of these fluxes on terrestrial carbon storage using the concept of pools.



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Model structure

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Vegetation pool may be one average individual, or a range of different individuals of different age and species, depending on the model



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Prentice et al. (2007)



Smith et al. (2014)

Plants can also die. Typical death mechanisms related to resource availability or physiological limits include:

- Bioclimatic limits
- Negative productivity
- Growth efficiency threshold (biomass increase per unit leaf area)
- Maximum age
- Background rate
- Shading/competition (mortality increases with canopy cover)

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Plants can also die due to ecosystem disturbances, e.g.

- Fire
- Wind-throw
- Insect attack

This is a major uncertainty in the understanding of ecosystem response to future environments. A doubling of mortality rates leads to a large drop in vegetation carbon stocks.

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Species composition

Huge range of species in reality. We have neither computational capacity or data to capture these in large-scale models. Typical approach is to classify species into PFTs.

Distinguished by, e.g. **Plant physiology** (C3/C4 photosynthesis) **Phenology** (Evergreen/Deciduous) **Physiognomy** (Woody/Herbaceous) **Bioclimatic limits** (Cold/Heat tolerance)





Species composition



Until recently, ESMs and DGVMs concentrated on PNV. But managed system differ fundamentally in many respects:



Models are now being expanded to account for the range of processes existing in managed systems, e.g. agriculture.

Productivity of croplands can be very different



Extraction of global croplands from chlorophyll fluoresence data shows that GPP can be much higher than indicated by models which do not account for management (Guanter et al., 2014)

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However, these productivity increases may not propagate to increases in terrestrial carbon stocks because of processes such as harvest and tillage

Change in soil carbon stocks after complete conversion from natural vegetation (global mean values):



Change in terrestrial C accumulation: (LPJ-GUESS, forced by MPI-ESM-LR following RCP 8.5)



Together agricultural processes can make a huge difference to projections of global carbon uptake





Projecting the global carbon cycle: Nutrient limitation

Nutrients other than nitrogen may also be important.



Goll et al. (2012)

Spatial projections very different between models

Difference 1860 to 2099. Driven with climate-carbon-cycle coupling.

Figures from Sitch et al. (2008)

Projecting the global carbon cycle: Mortality

Amazon dieback?

Difference 1860 to 2099. Driven with climate-carbon-cycle coupling.

Figures from Sitch et al. (2008)

Projecting the global carbon cycle: Mortality

Or robust tropical forests?

TRIFFID model forced with climates from a range of GCMs. Only one climate realisation resulted in a loss of carbon.

Summary and implications

- A lack of appropriate observations means that we must rely on models to understand the overall role of the land biosphere in the carbon cycle.
- These models attempt to capture the key processes, but in order to be applied at the global scale they adopt highly simplified process representations
- Models generally project a strong take up of carbon by the terrestrial biosphere under environmental change. However, they don't universally agree on this, and there are many uncertainties, relating to both explicitly resolved processes and to missing processes (e.g. peatlands).
- As you will see in the coming talks, these changes in natural vegetation properties under environmental change mean that calculations of human impacts based on the current land system may not hold in the future.
 - E.g. The tropical forest may be a much large C store in the future, making it even more important than it is now for keeping C out of the atmosphere. Or it may dieback anyway, meaning that todays deforestation emissions would occur naturally in the future anyway.

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Projecting the global carbon cycle: Nutrient limitation

Must take results from models without nutrient constraints with a pinch of salt...

Hungate et al. (Science, 2003)

Projecting the global carbon cycle: Nutrient limitation

FACE experiments differ their long-term response to elevated [CO₂] Must take results from models without nutrient constraints with a pinch of salt...

Projecting the global carbon cycle: Feedbacks

Biogeochemical feedbacks

Many other feedbacks exist Further reading: Arneth et al. (2010)