



*The Abdus Salam
International Centre for Theoretical Physics*



2022-4

Workshop on Theoretical Ecology and Global Change

2 - 18 March 2009

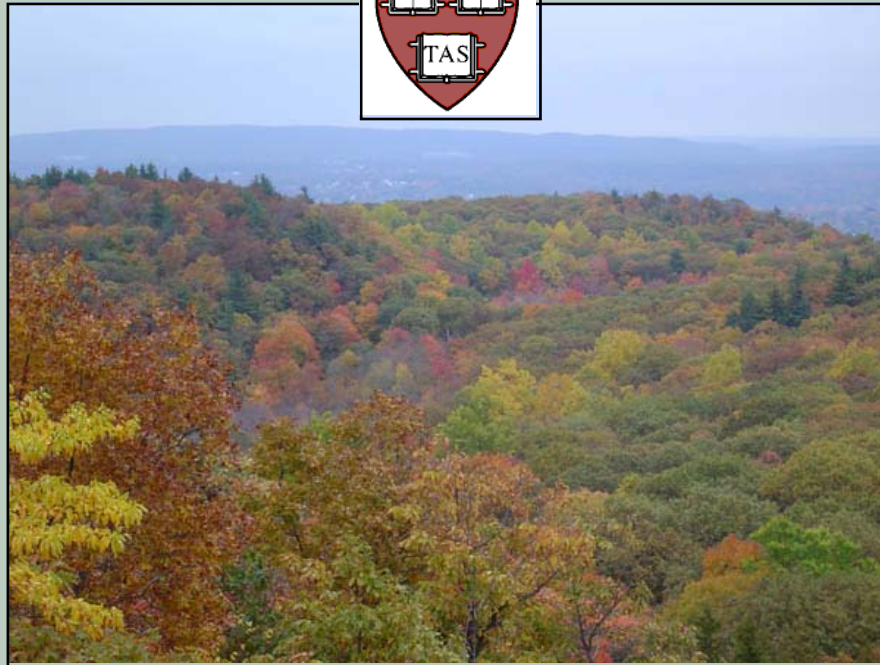
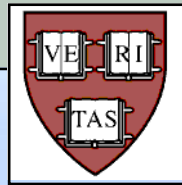
Developing a Predictive Science of the Biosphere

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Developing a predictive science of the biosphere

Paul R. Moorcroft

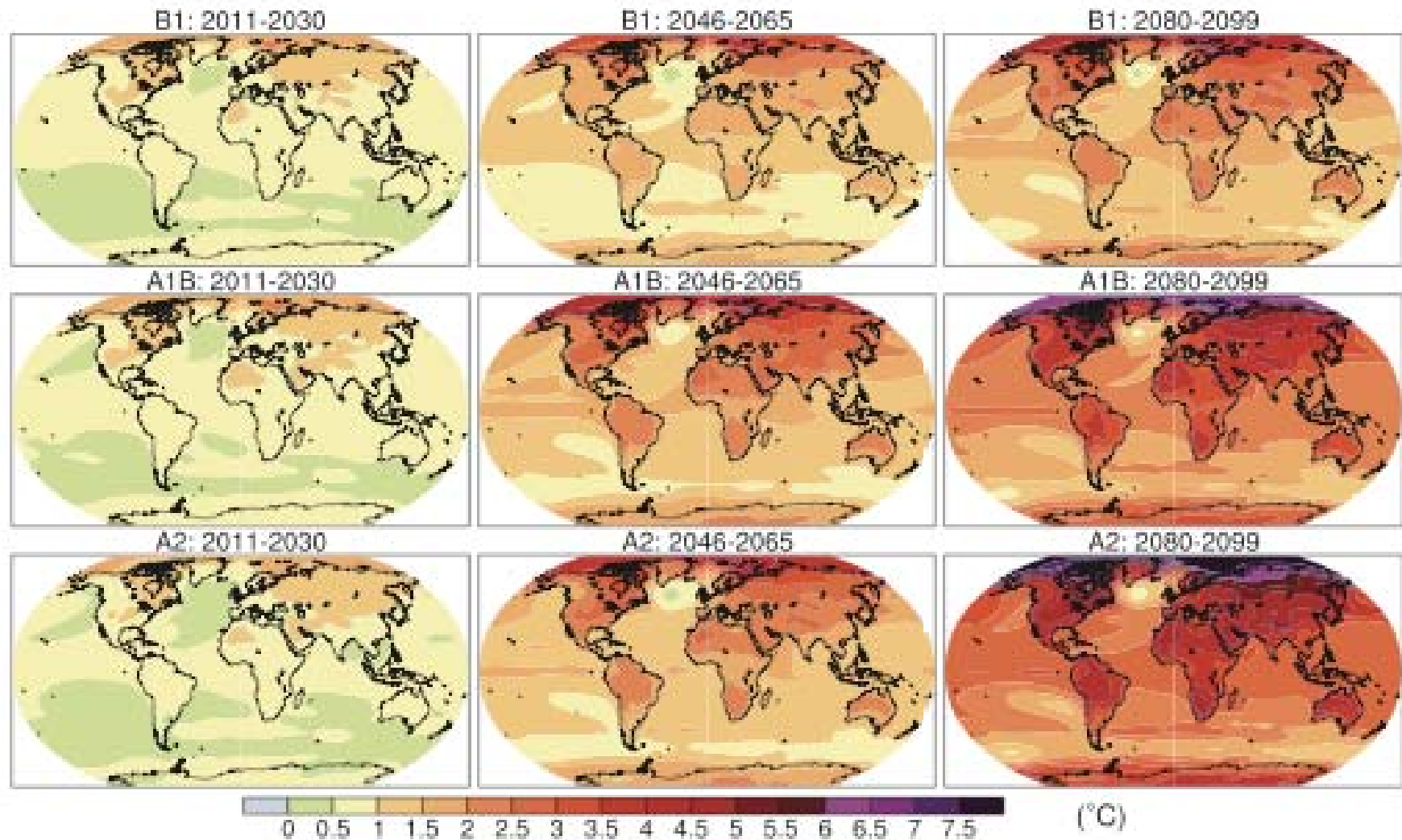
Harvard University

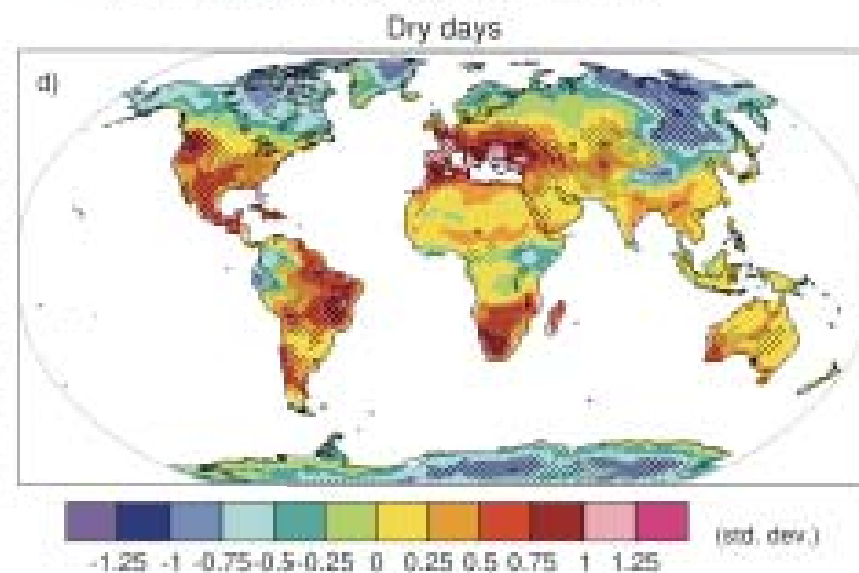
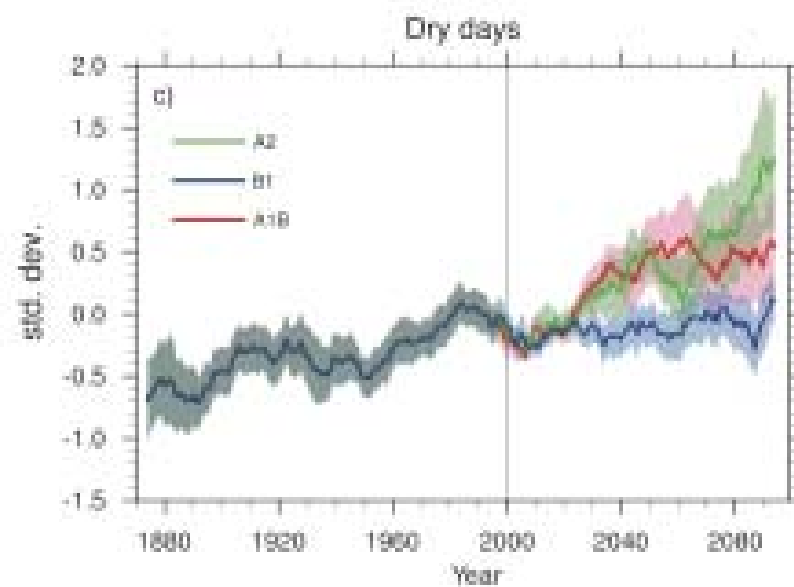
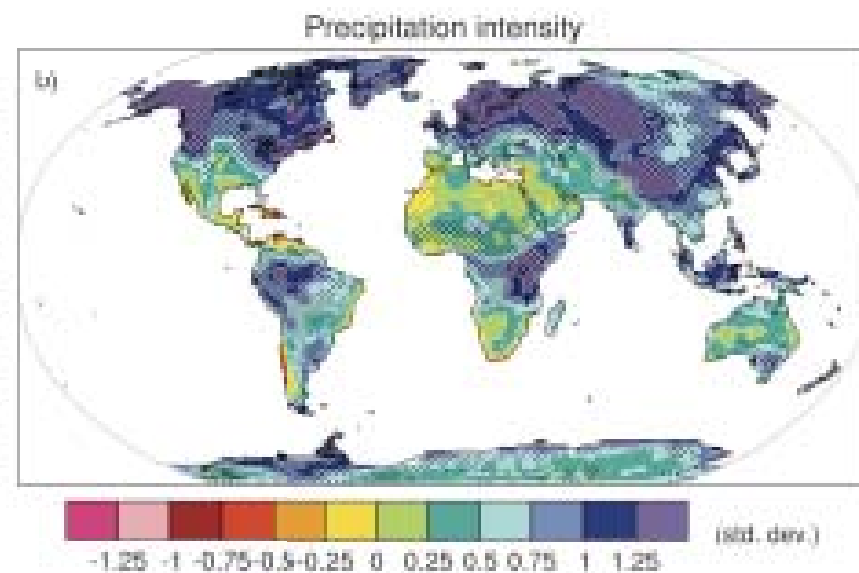
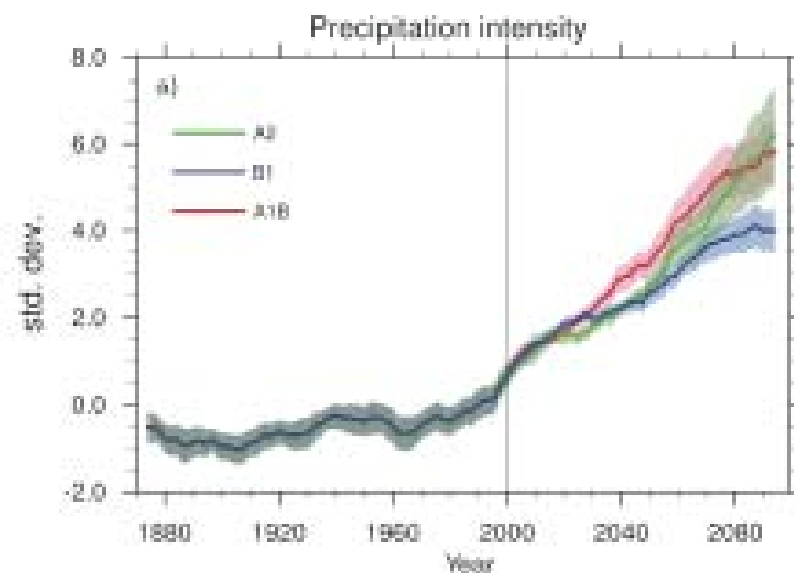


Overview

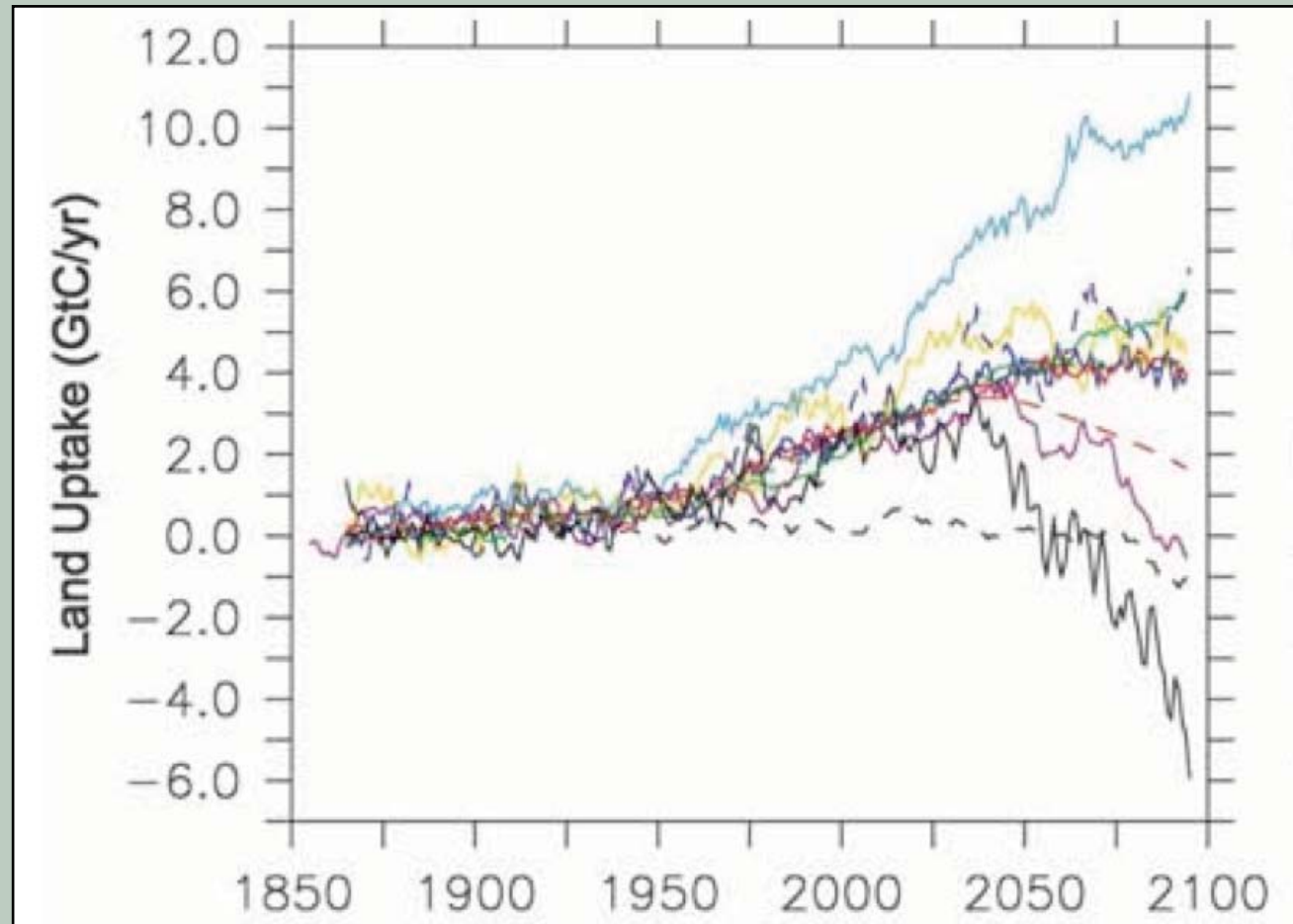
- Background
- Approaches to modeling the terrestrial biosphere
- Structured ecosystem models

Predicted temperature changes over the coming century





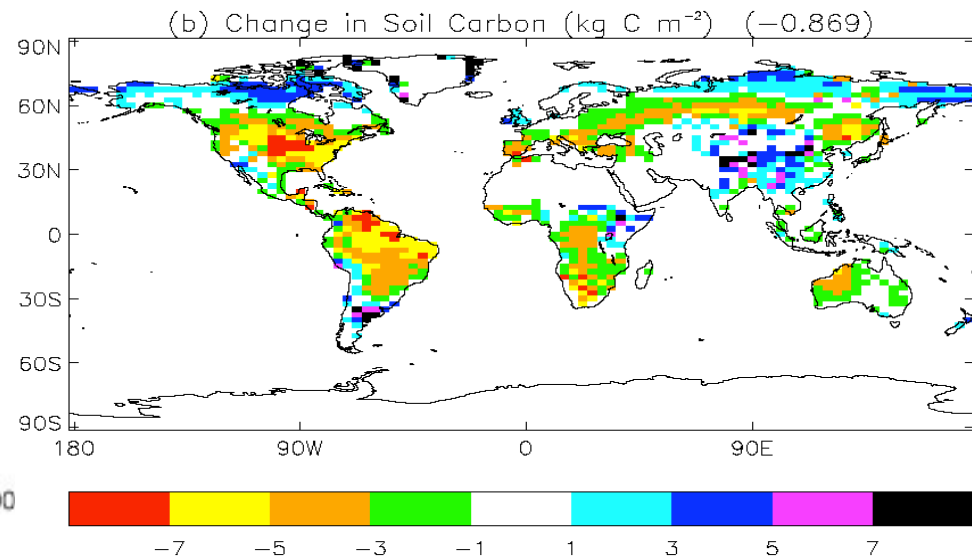
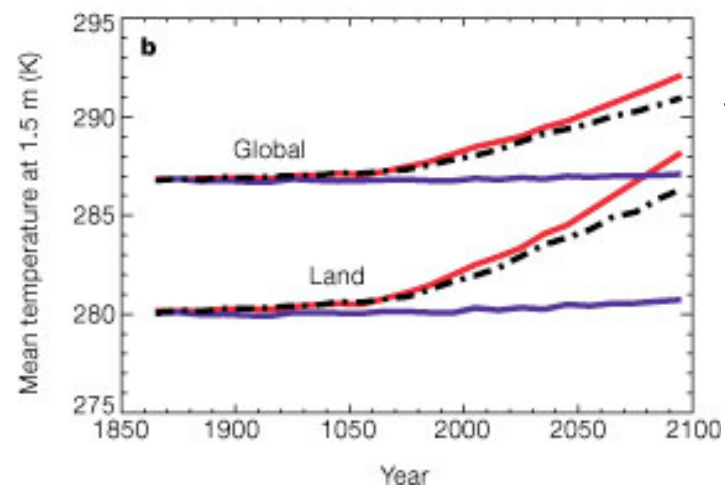
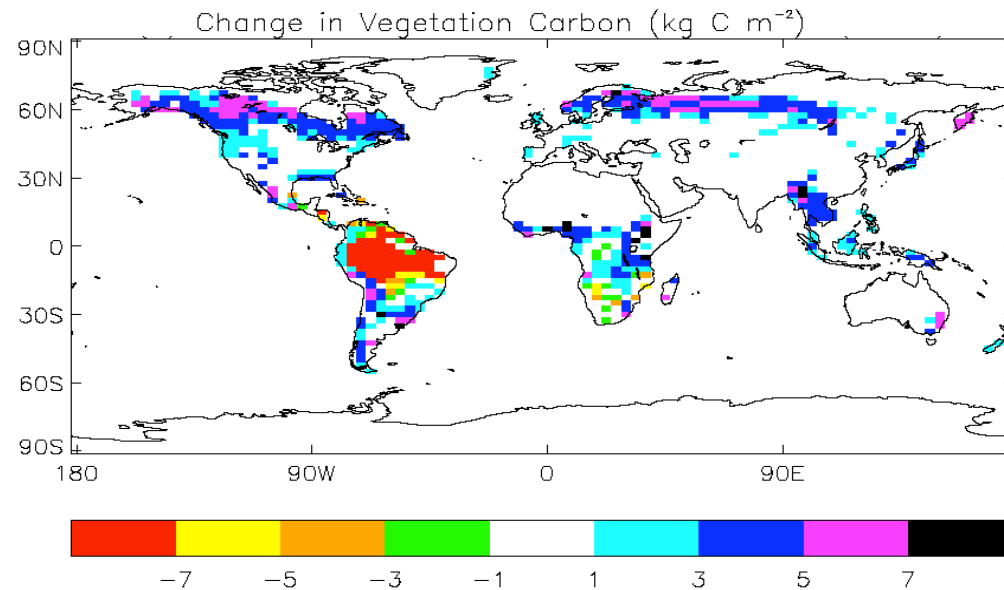
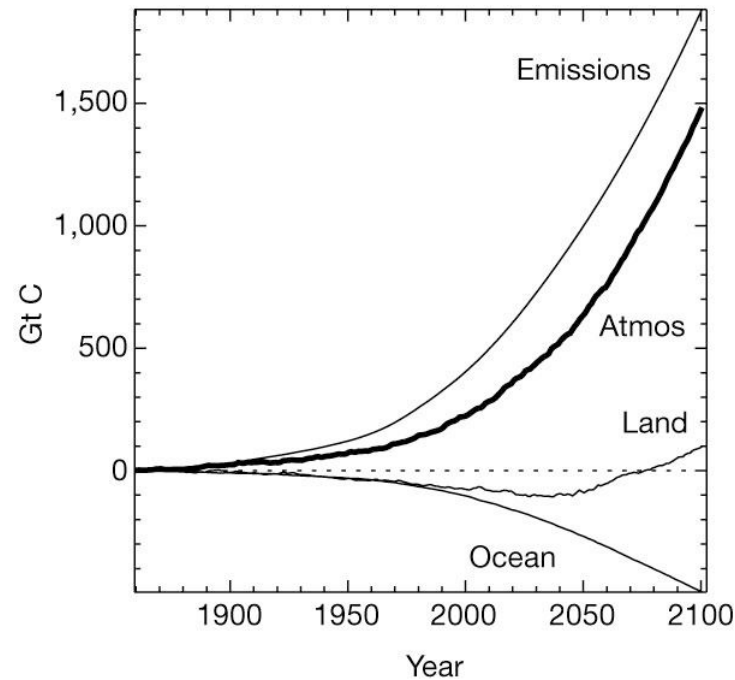
Predictions for the long-term responses of terrestrial ecosystems to climate change (IPCC 2007)



HadCM3LC (solid black), IPSL-CM2C (solid red), IPSL-CM4-LOOP (solid yellow), CSM-1 (solid green), MPI (solid dark blue), LLNL (solid light blue), FRCGC (solid purple), UMD (dash black), UVic-2.7 (dash red), CLIMBER (dash green), and BERN-CC (dash blue).

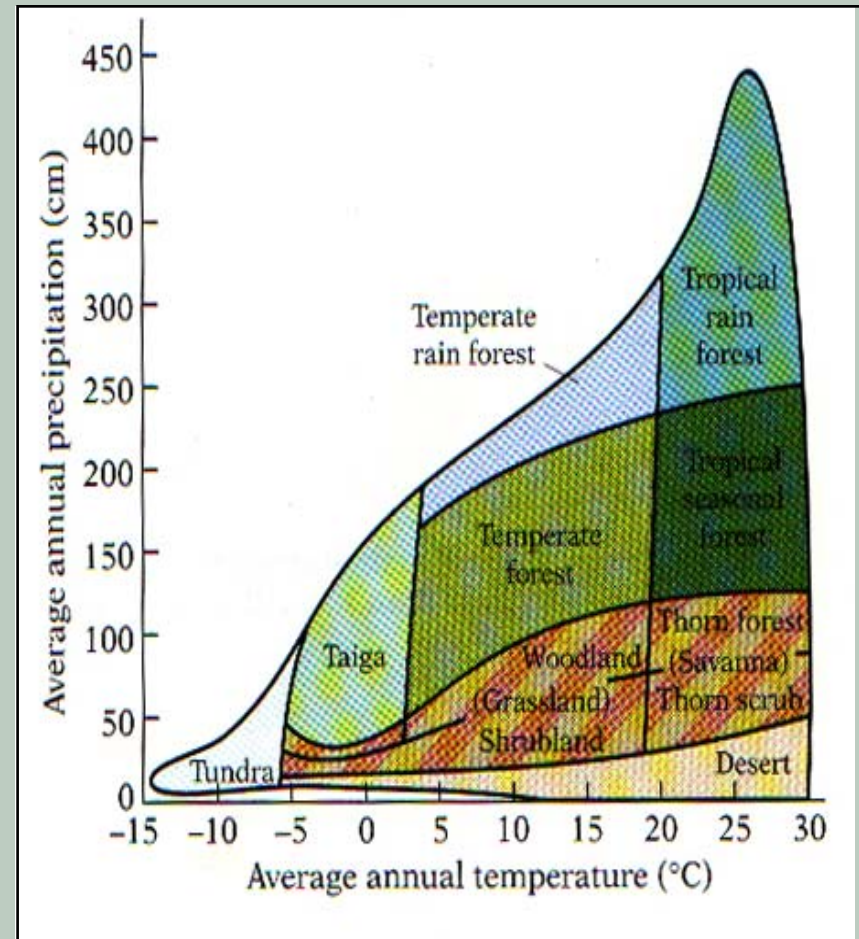
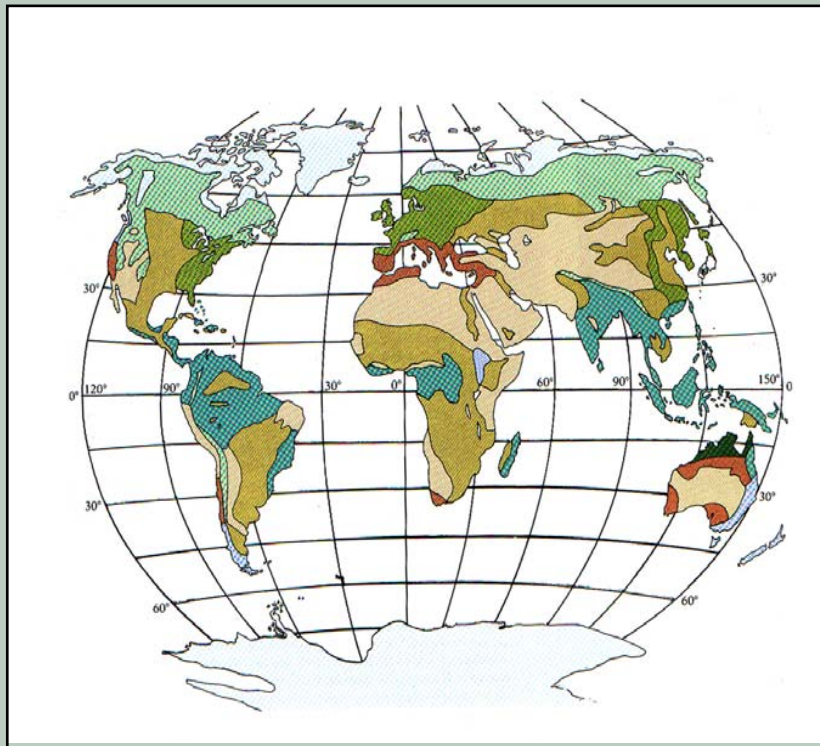
(Cox et al. 2000)

HadCM3LC: Predicted collapse of Amazon ecosystems in response to rising CO₂

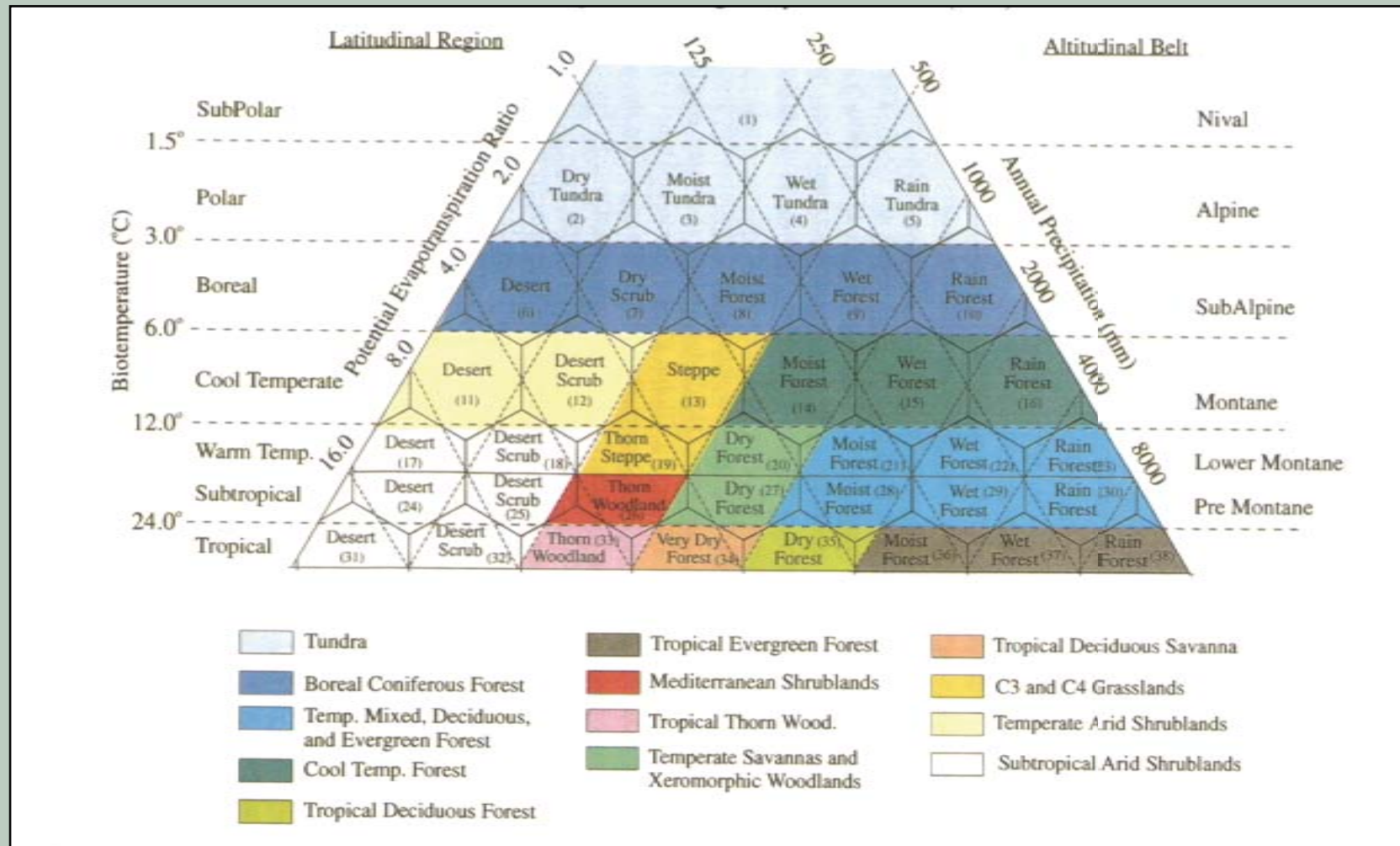


- Biogeographic models

Whittaker biogeographic scheme: biome type in relation to average annual temperature & precipitation



Holdridge classification scheme

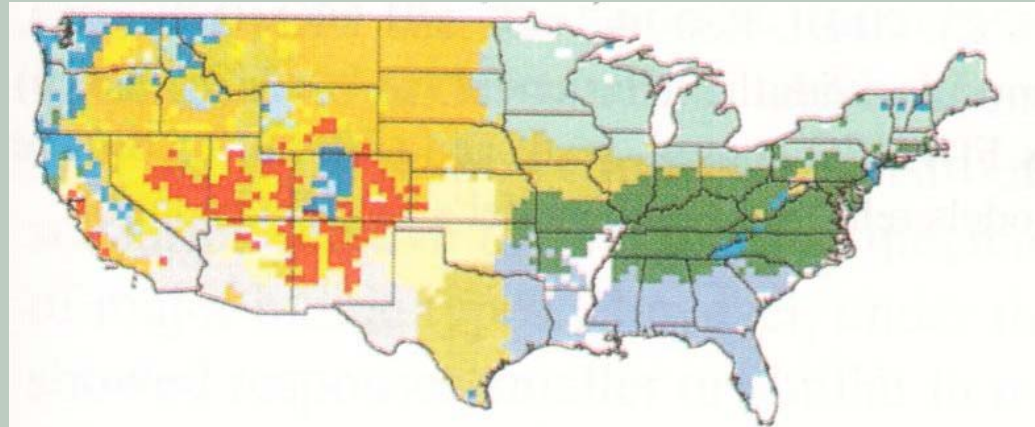


Potential Evapotranspiration Ratio = $\frac{\text{potential evapotranspiration}}{\text{precipitation}}$ Biotemperature = $\frac{\text{Average of temperatures} > 0^{\circ}\text{C}}$

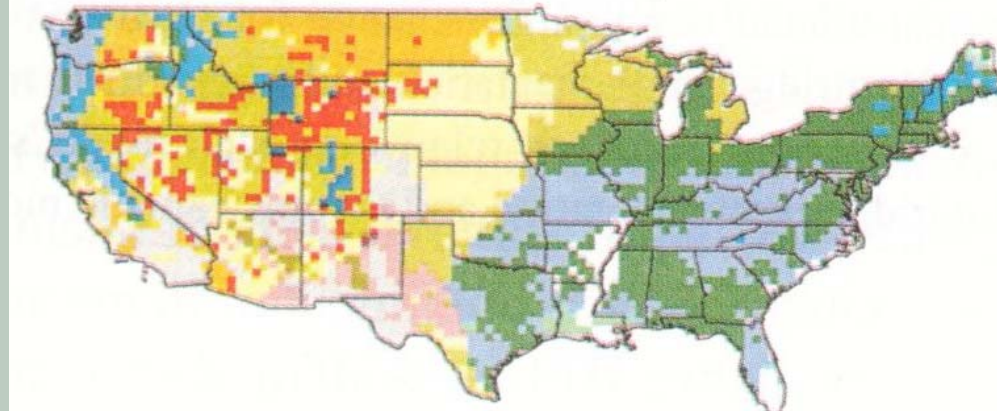
“Vegetation is crystallized, visible climate” (Koppen, 1936)

Using Biogeographic Vegetation Models to predict the effects of climate change

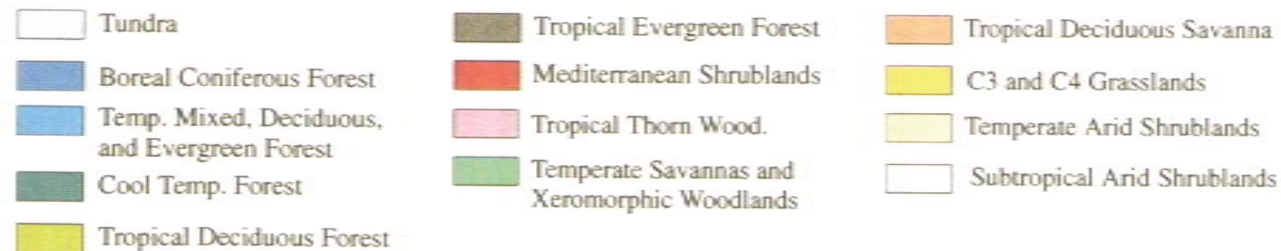
1 x CO₂



2 x CO₂

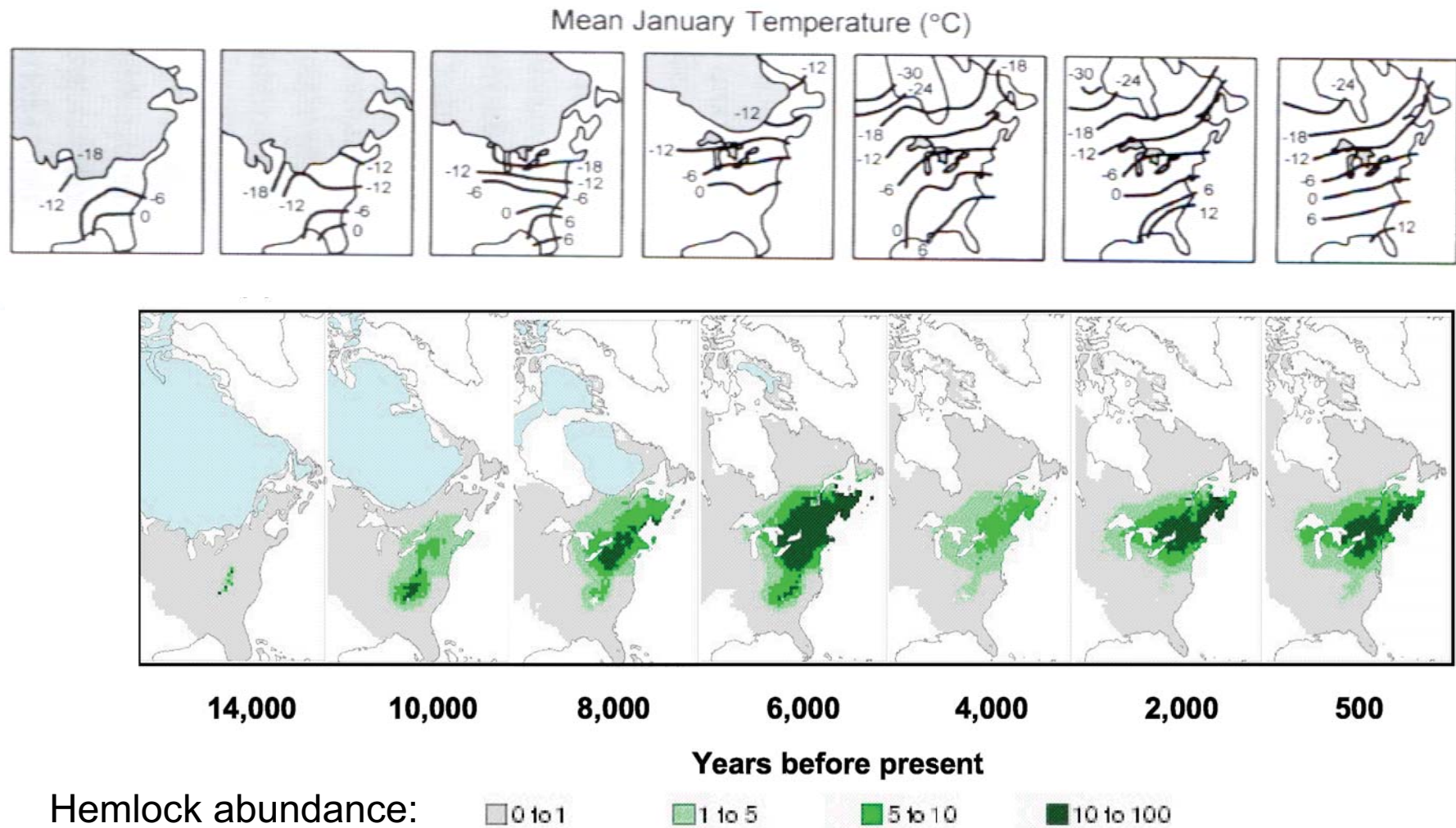


(Holdridge
classification
Scheme)



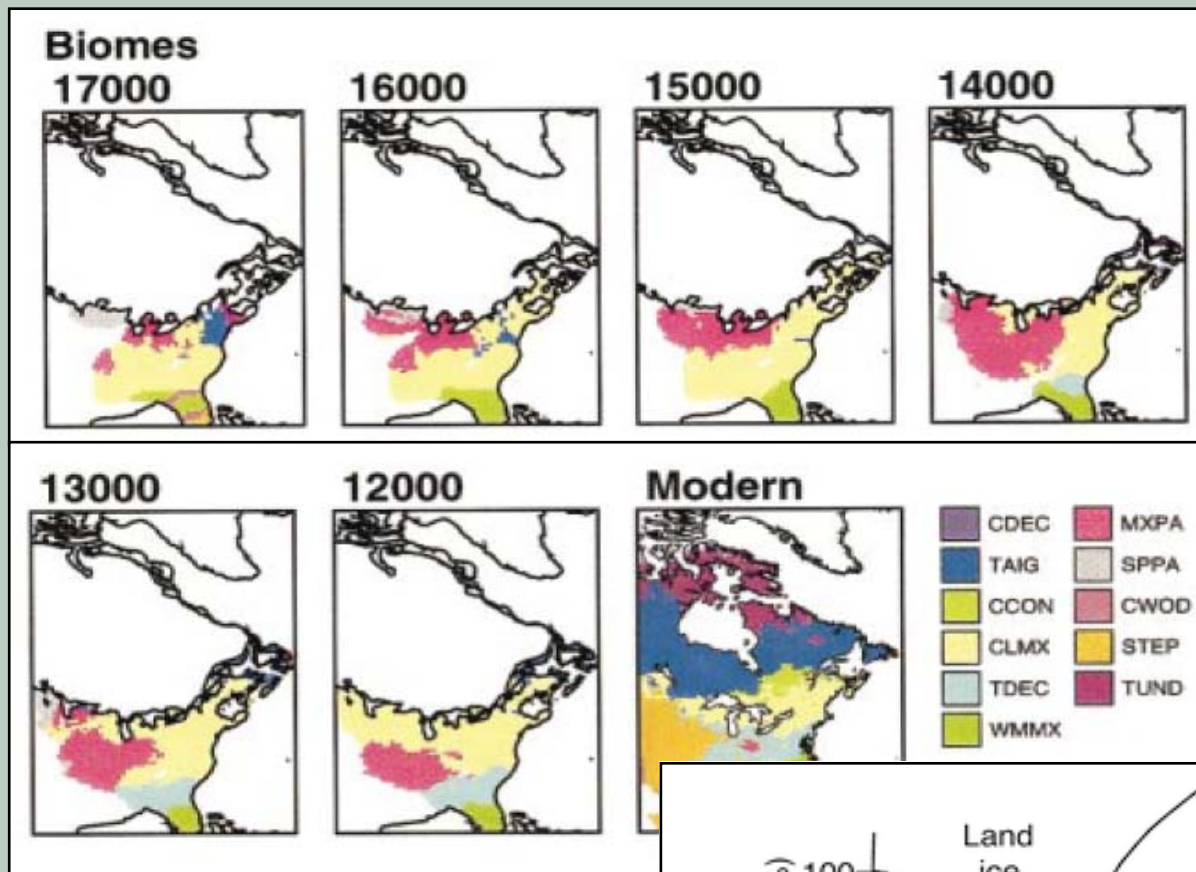
Problems with the Biogeographic approach

Issue #1: assumes ecosystems are in equilibrium with climate, both now & in the future.



Issue #2: “No-analog communities”

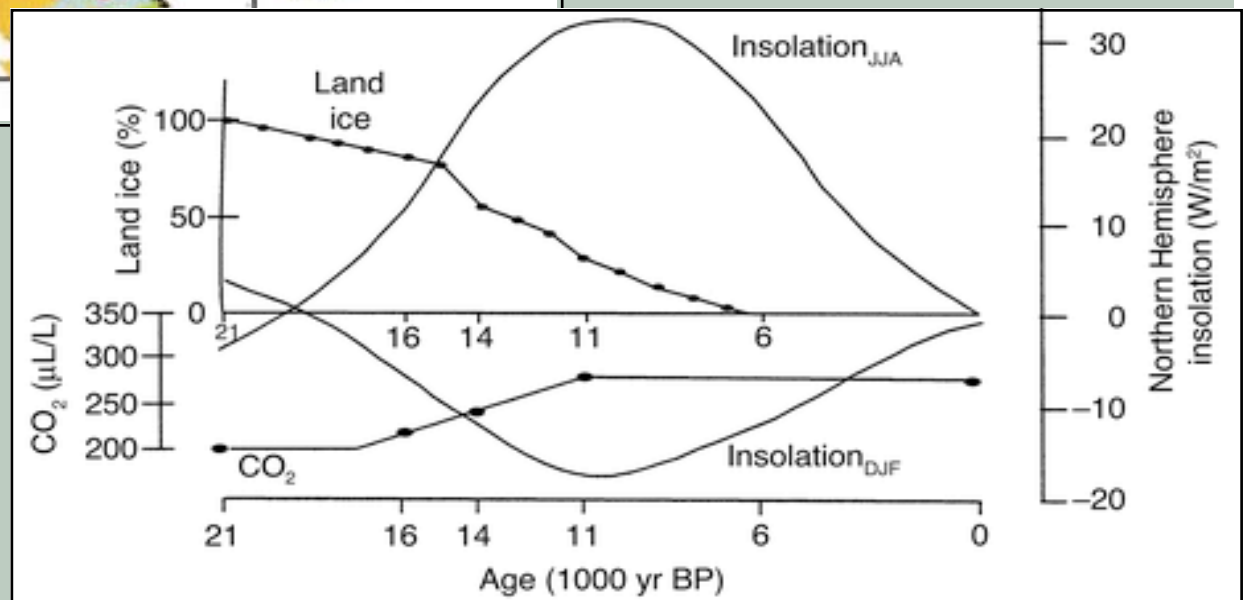
(Williams et al 2001)



MXPA= Mixed Parkland
[spruce, larch, ash,
hornbeam, poplar, willow,
sedge, and sage]

MXPA termed a a ‘no-analog community’ because these species are not found together in any present-day vegetation community.

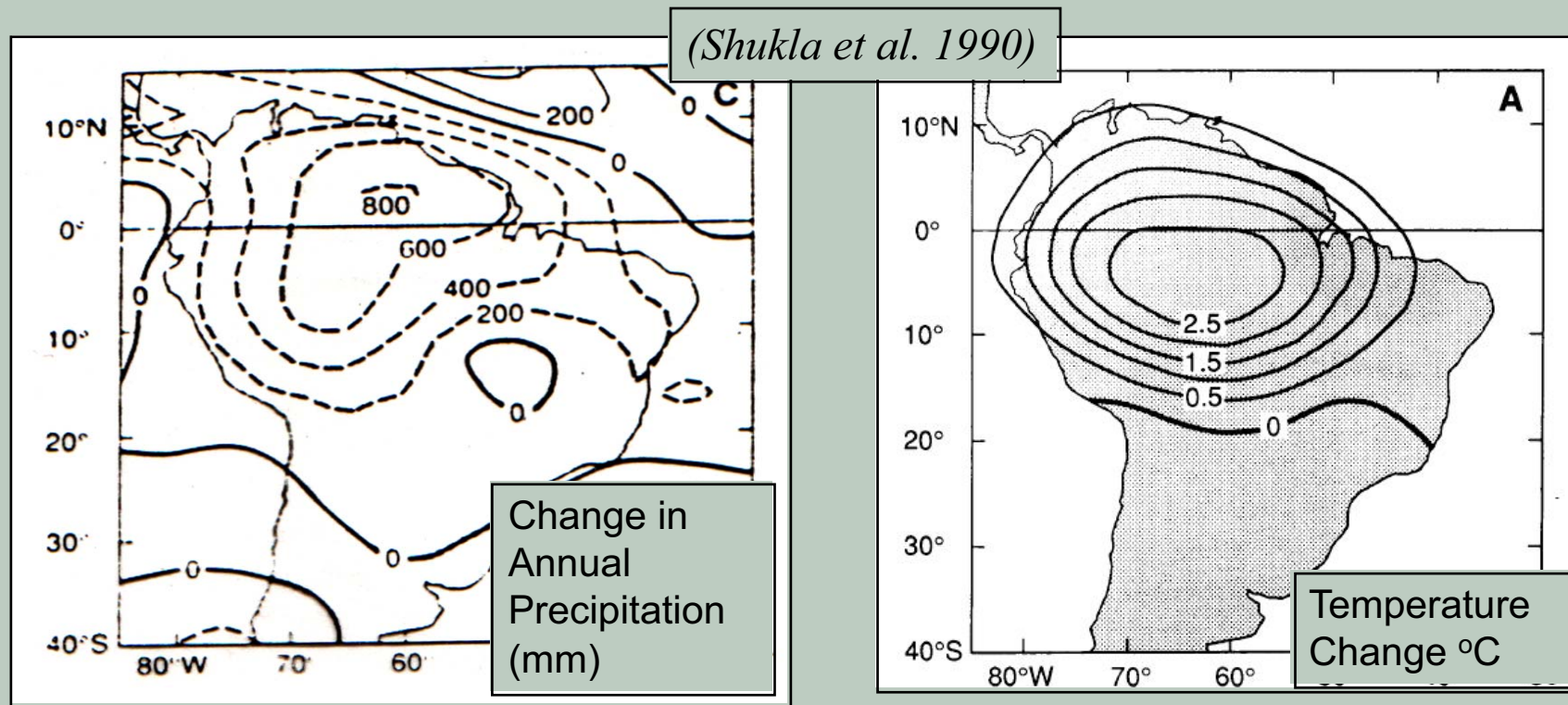
i.e. Distribution of biomes is a macroscopic pattern arising from the responses of the constituent species to climate forcing.



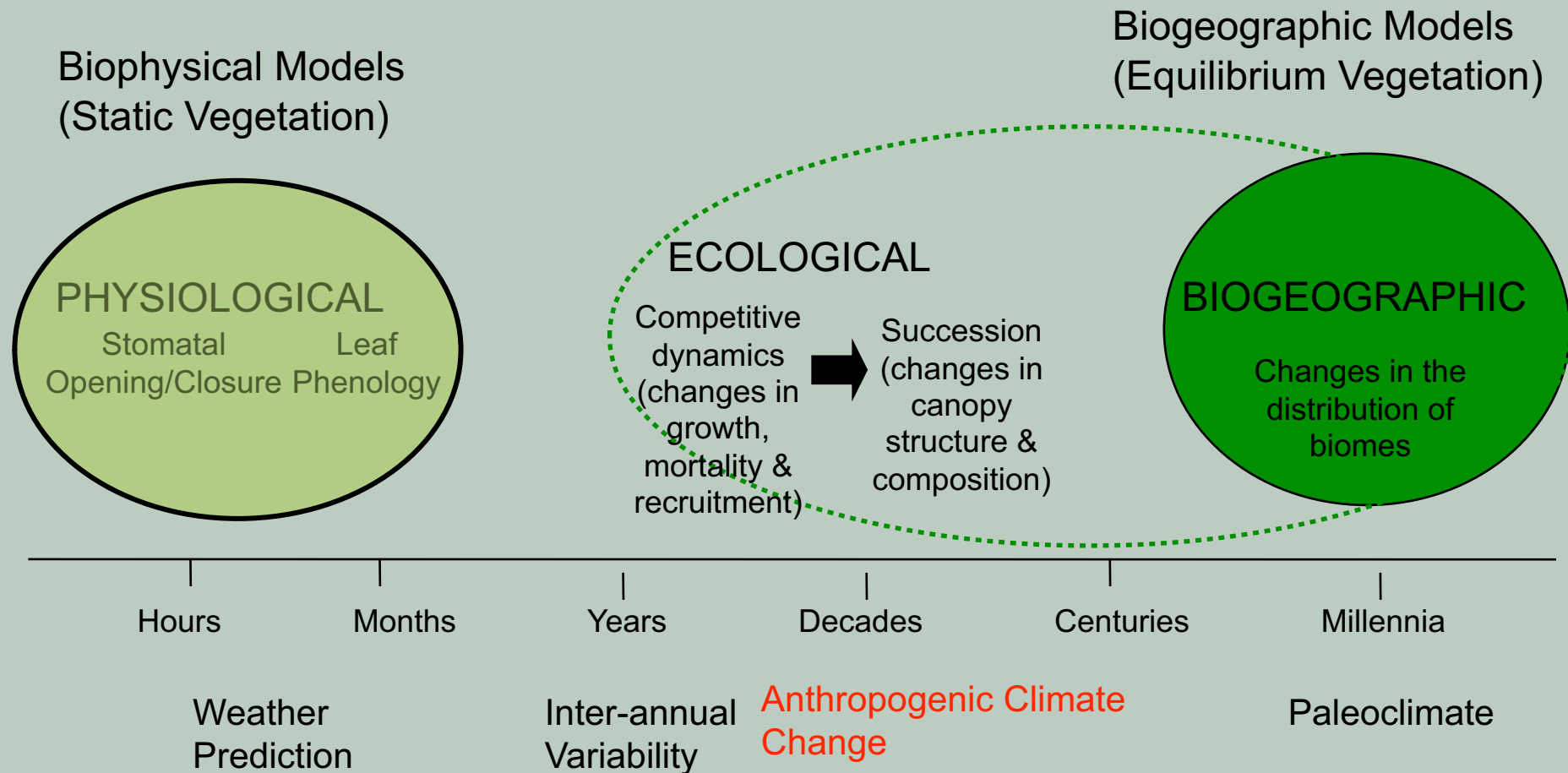
Issue #3: Biosphere-atmosphere feedbacks

- the biogeographic approach implicitly assumes a unidirectional relationship between ecosystems and climate:
- now know that this relationship is bi-directional.

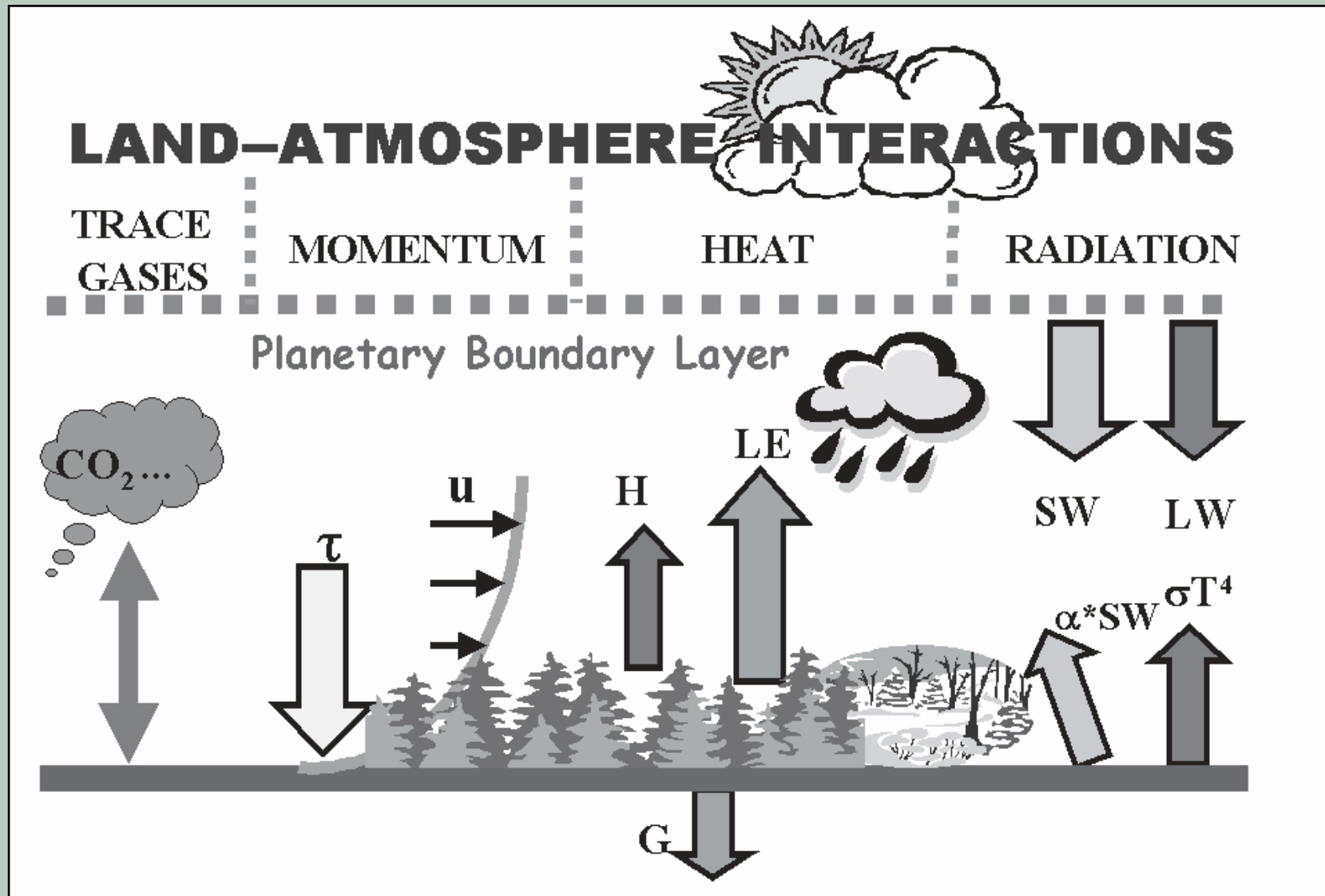
e.g: Amazonian deforestation predicted to change South American climate



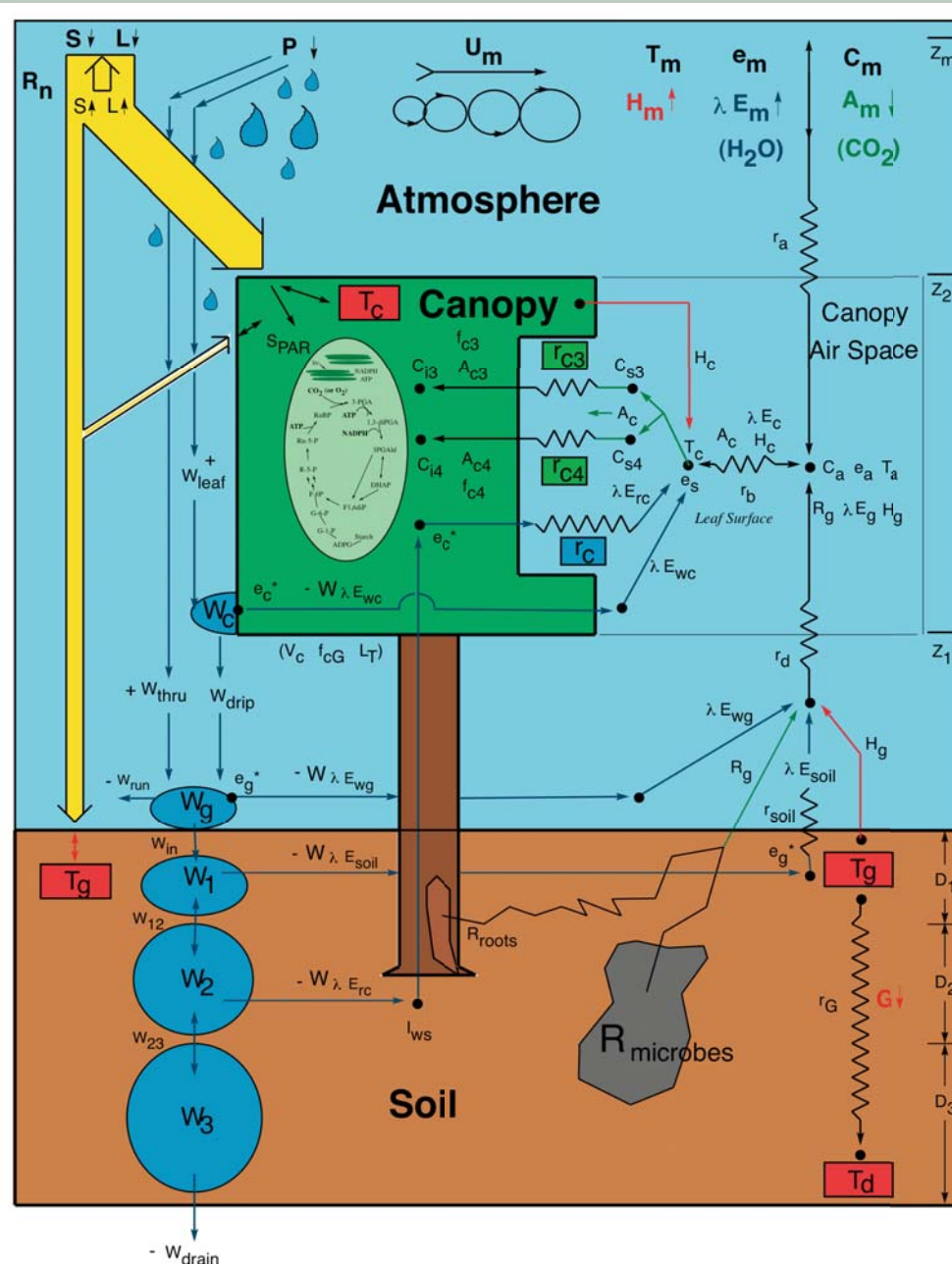
Timescales of Terrestrial Ecosystem Responses to the Atmosphere



Biosphere-Atmosphere Feedback Processes

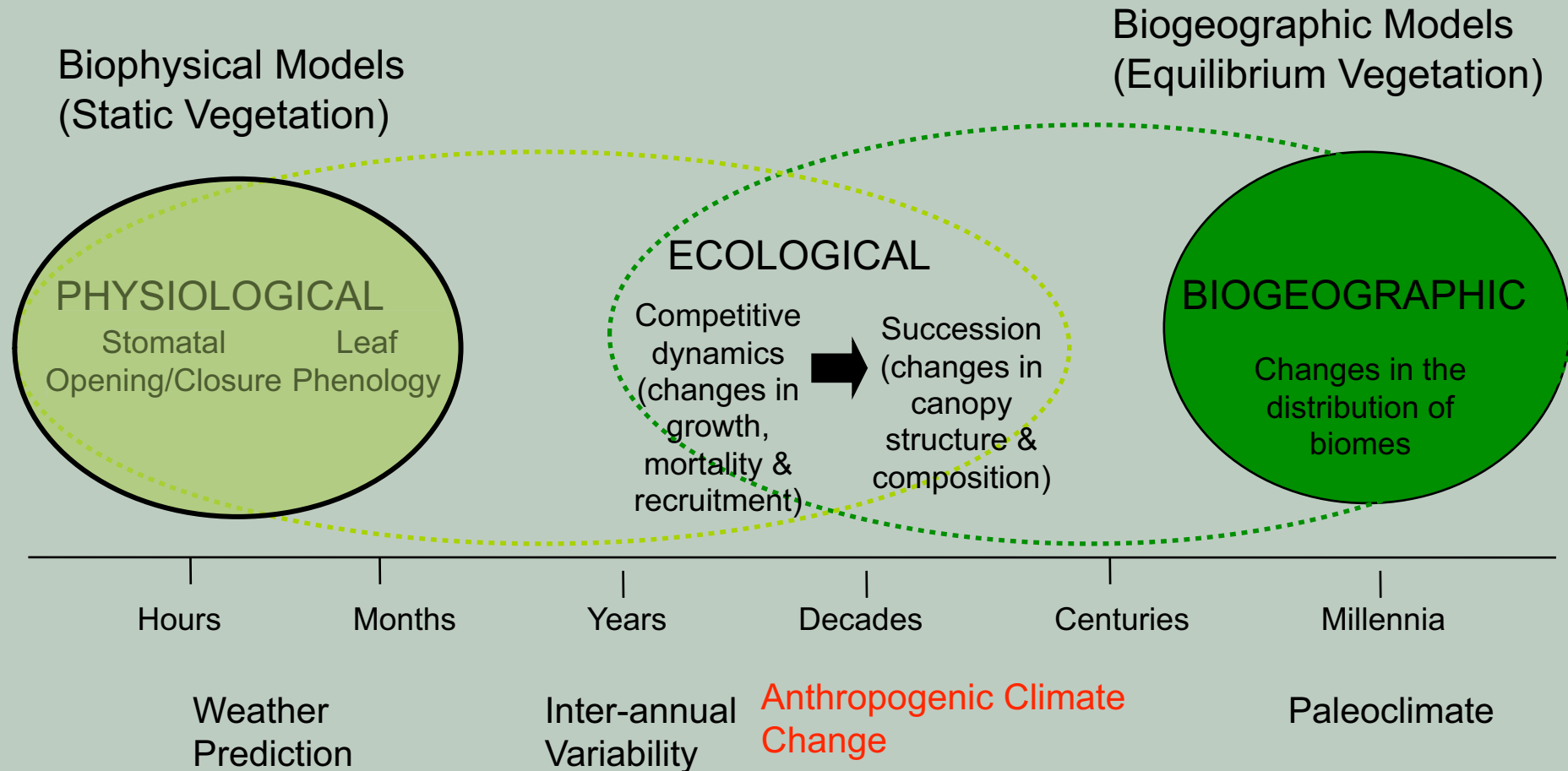


e.g. The Simple Biosphere Model, version 2 (SiB2) (Sellers et al. 1996)



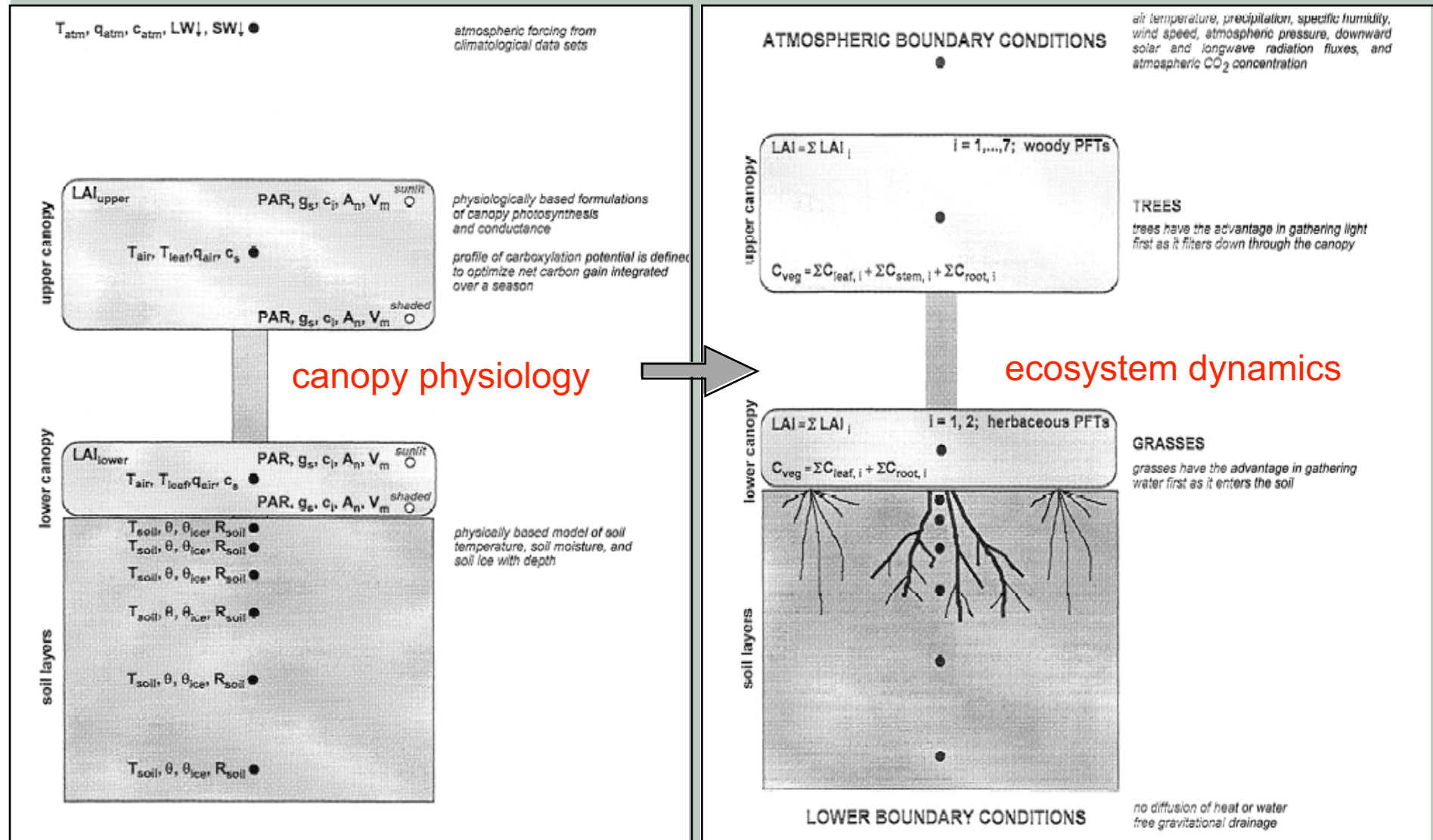
SiB2 was one of the first terrestrial biosphere models to explicitly link canopy biophysics with a biological of photosynthesis

Timescales of Terrestrial Ecosystem Responses to the Atmosphere



The Integrated Biosphere Simulator (IBiS) (Foley et al., 1996)

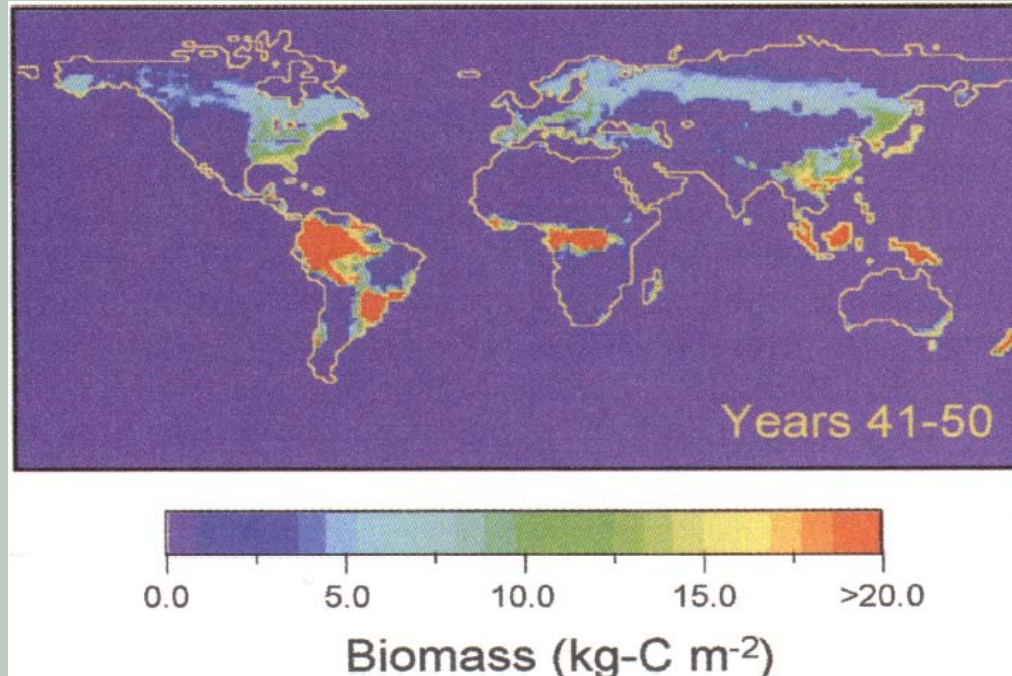
- IBiS was the first biosphere model to incorporate long-term ecosystem dynamics



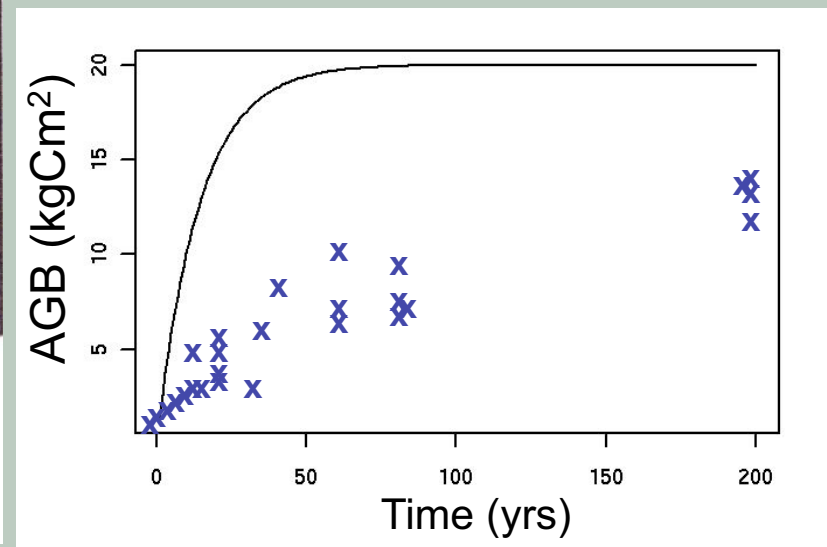
scale: $1^\circ \times 1^\circ$ ($\sim 10^4 \text{ km}^2$)

Long-term ecosystem dynamics in big leaf terrestrial biosphere models

Above-ground biomass dynamics of evergreen tree spp. in IBiS



Comparison of above-ground biomass dynamics to observations at San Carlos (tropical forest) 2°N,68°W

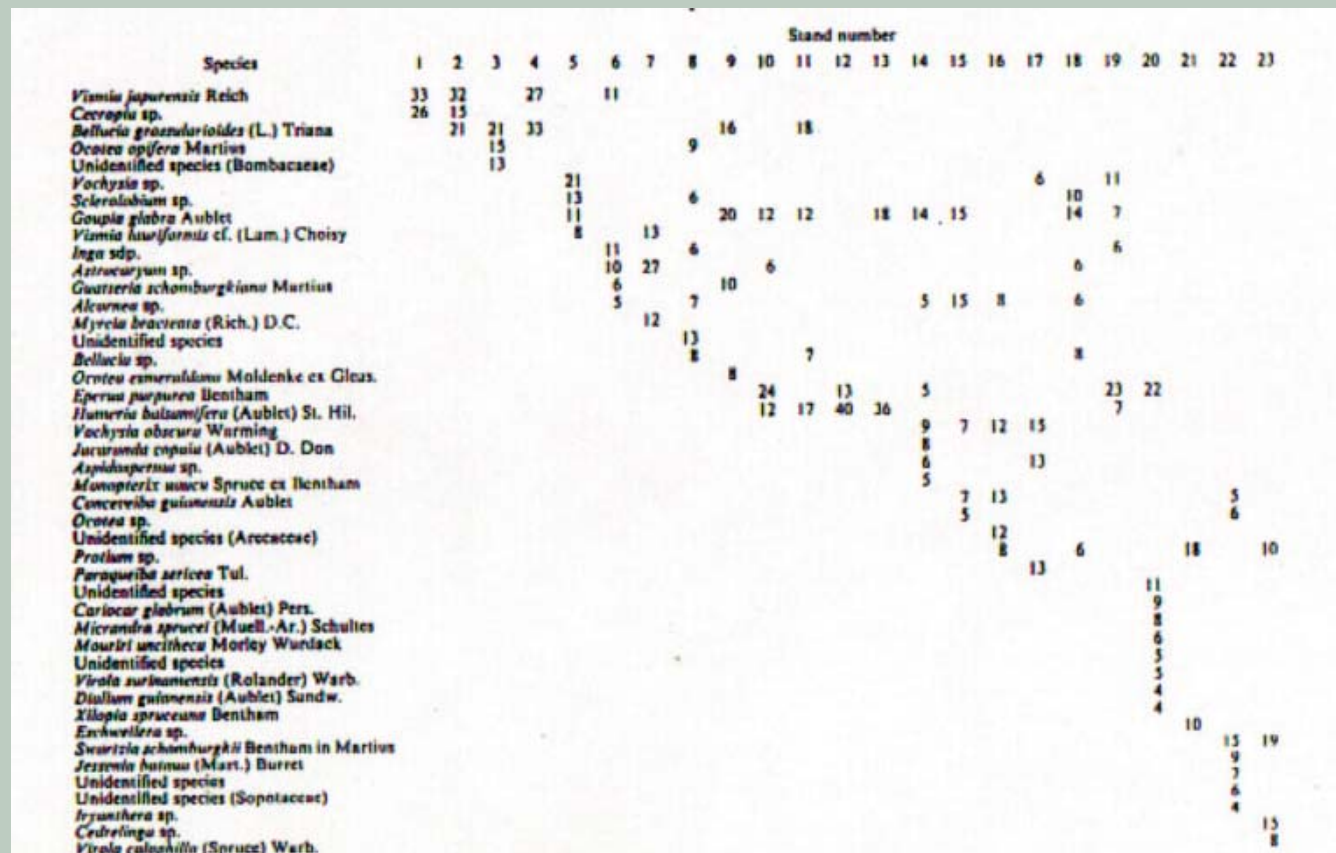


- Unrealistic timescales of response

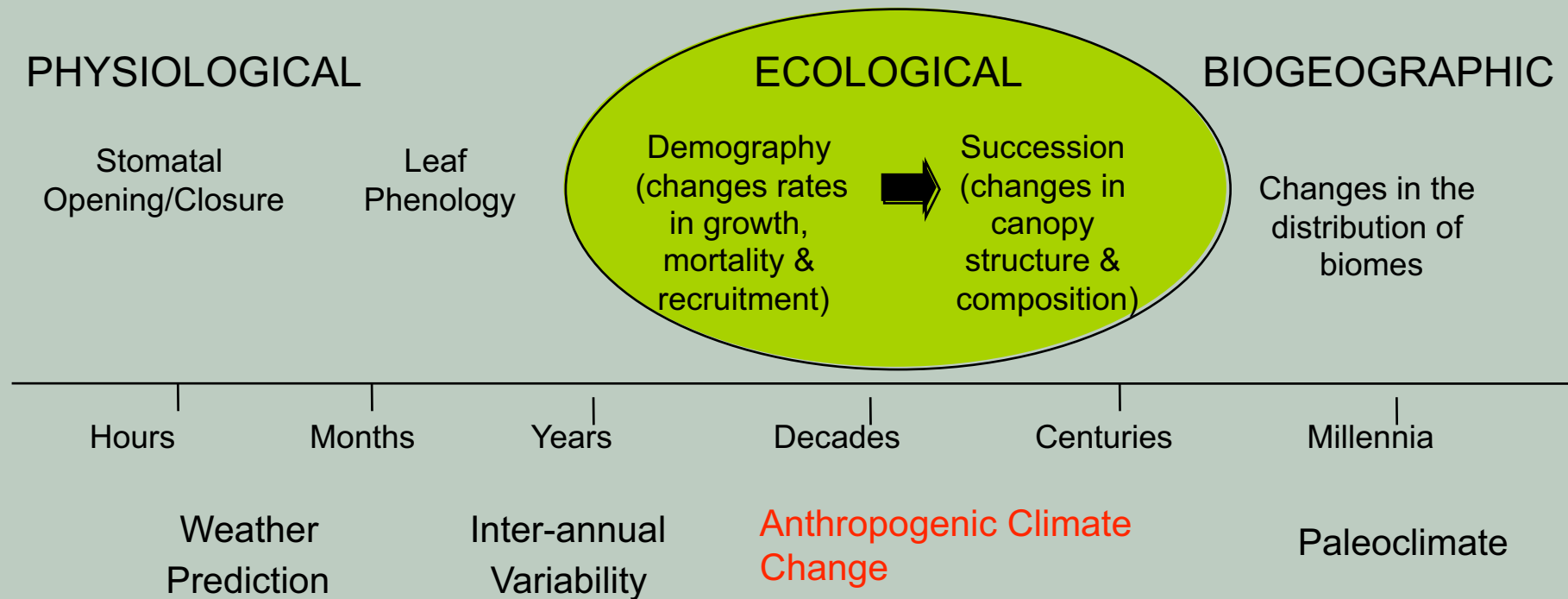
- Also the big-leaf assumption implies a single environmental niche within each grid cell. As a result, big-leaf models tend to predict homogeneous ecosystems (Gause competitive exclusion principle).

Species --->

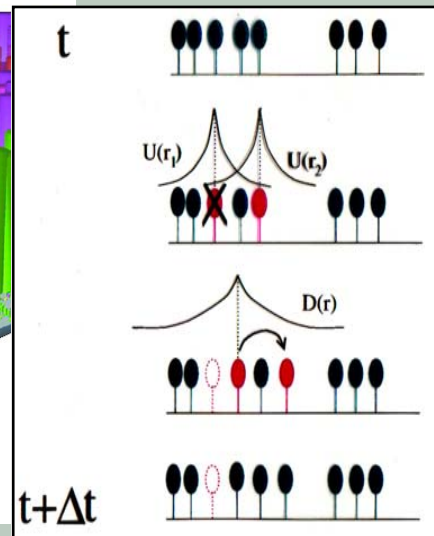
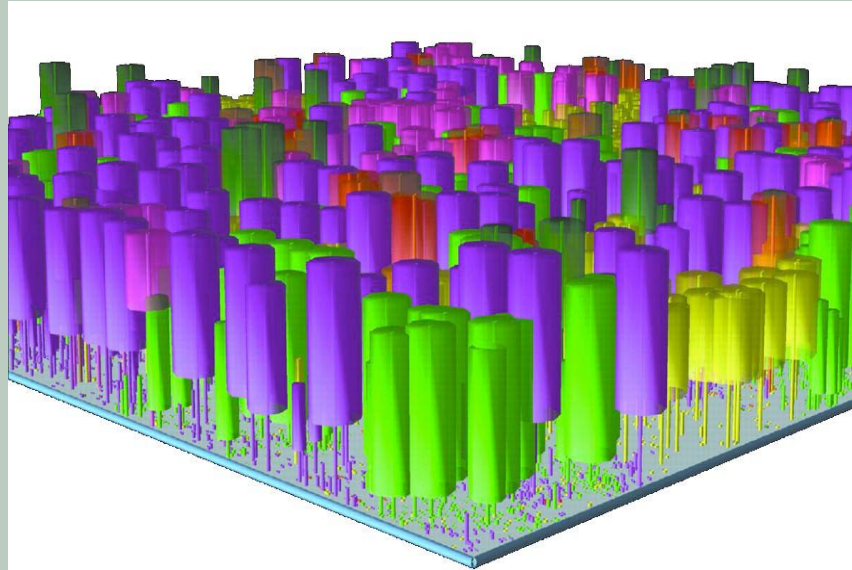
Stand Age (0-200 yrs) -->



Timescales of Terrestrial Ecosystem Processes



Individual-based vegetation models (gap models)

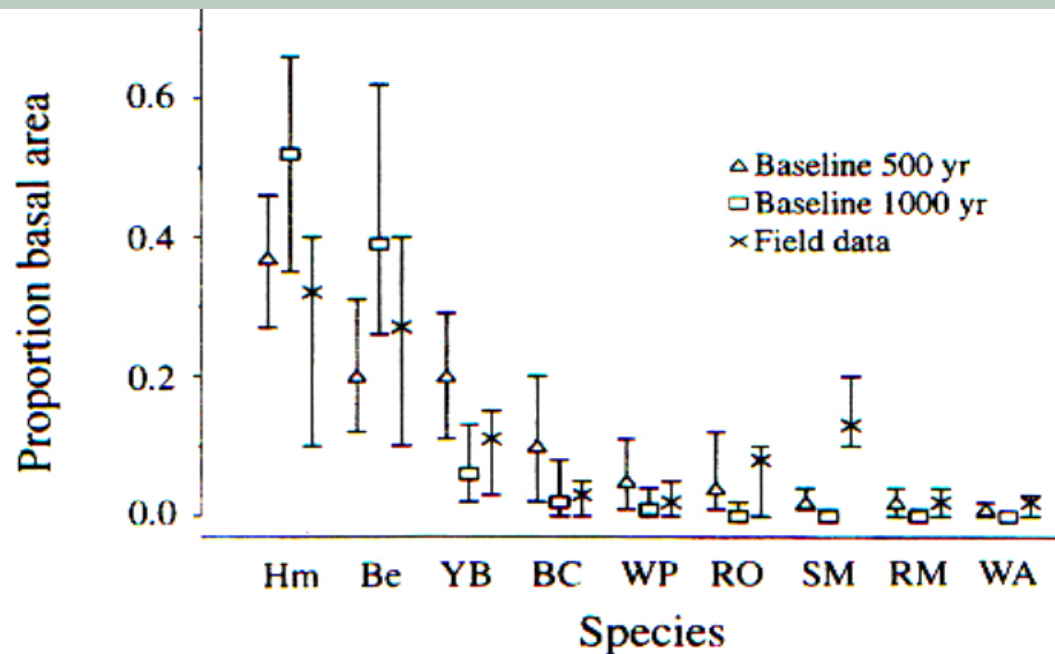


growth

height-structured
competition

mortality

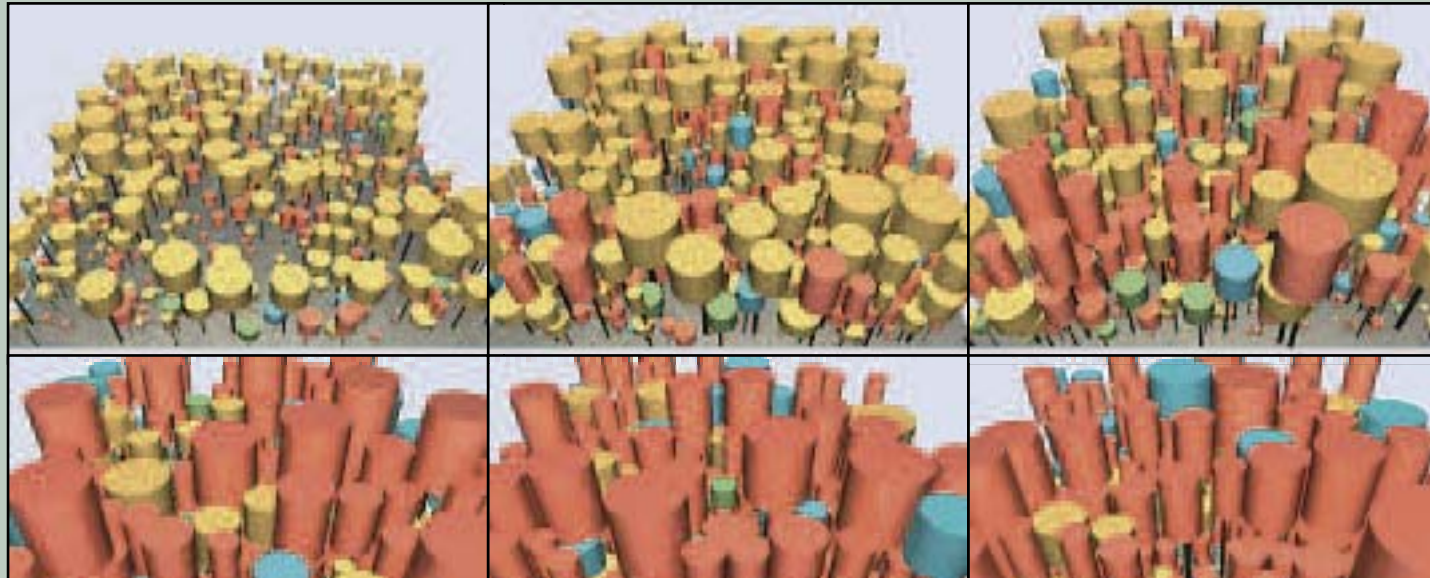
dispersal + recruitment



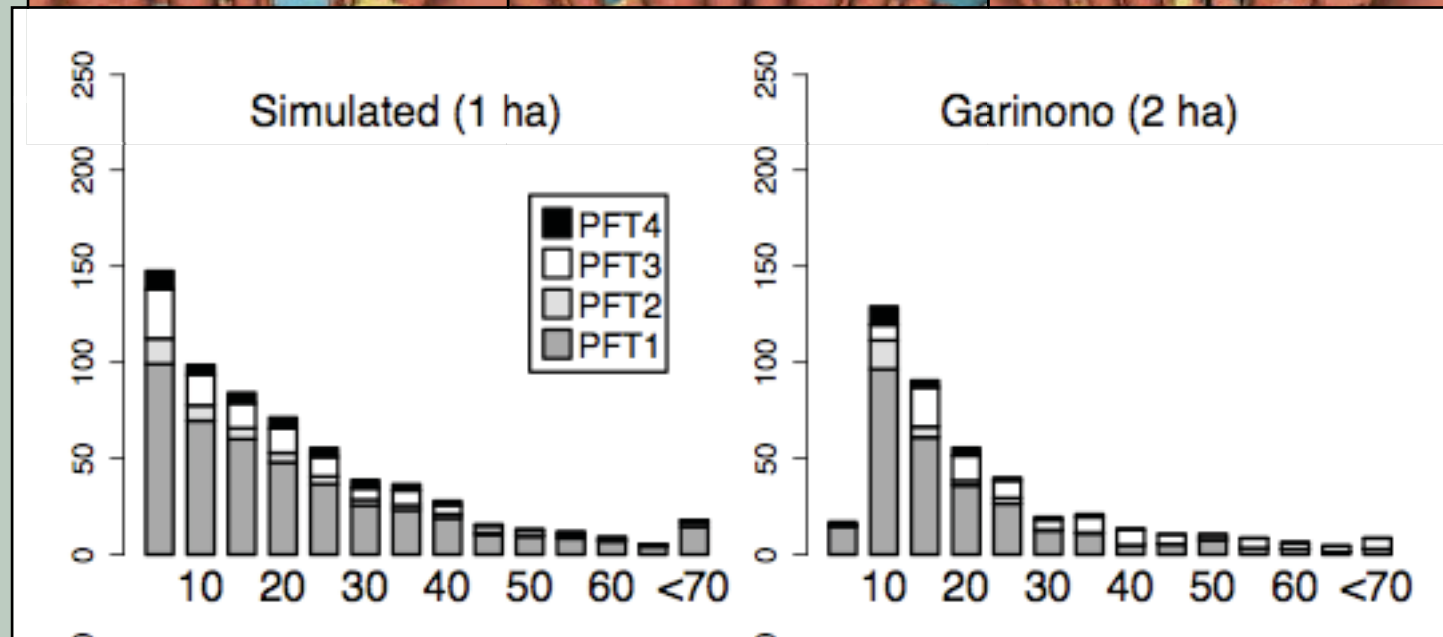
e.g.: SORTIE model of
New England forests

(Pacala & Canham et al. 1996)

Individual-based vegetation models (gap models)



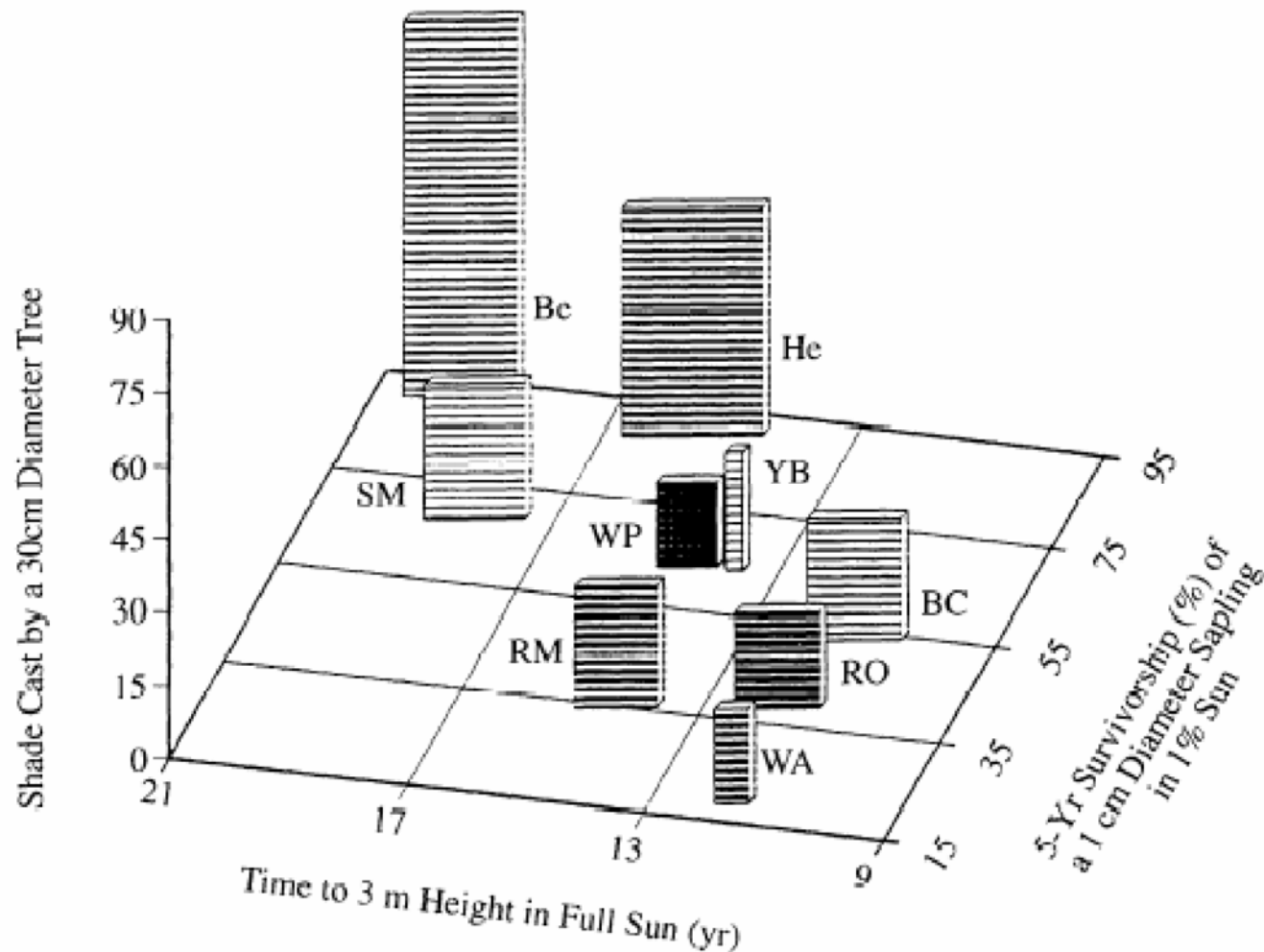
e.g.2: SEIB
model of
South-East
Asian Tropical
Forests



Sato et al. 2007

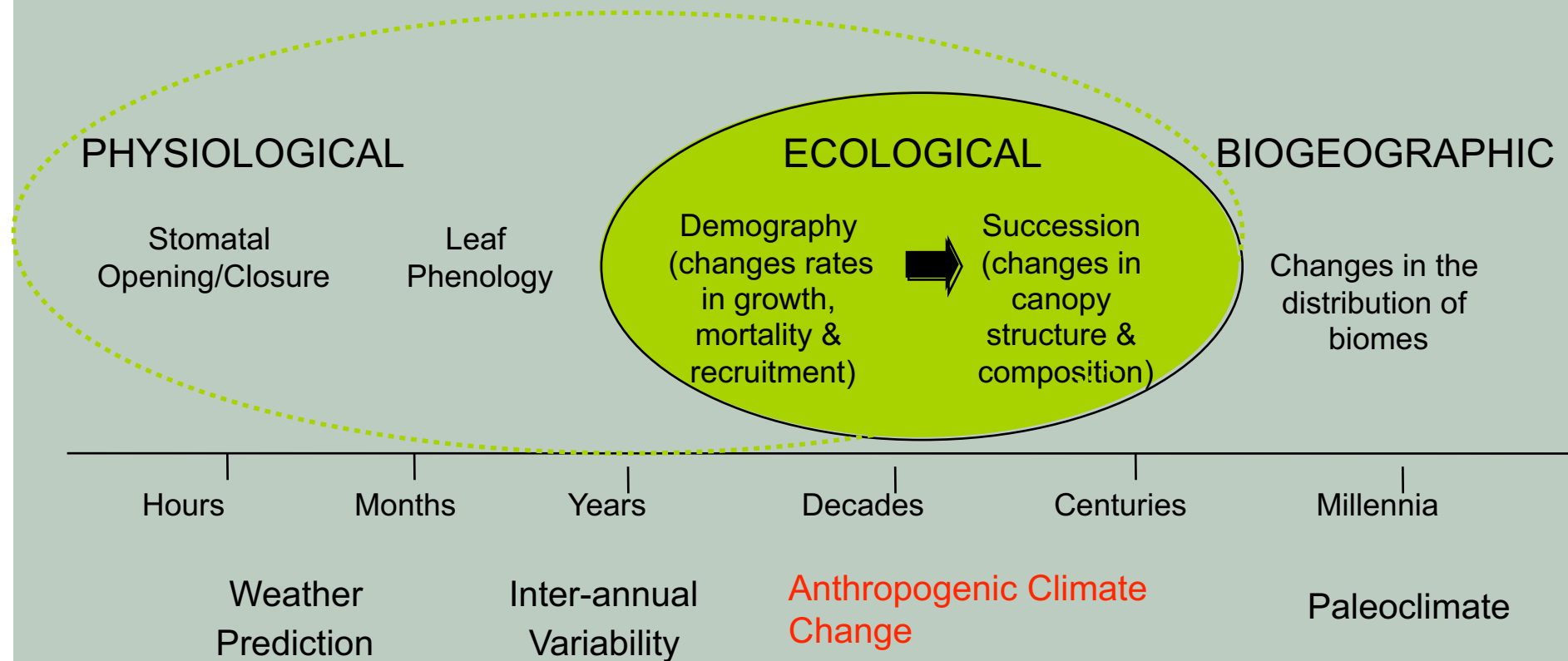
Traditionally gap-models were parameterized from field measurements of mortality, growth & recruitment in relation to resource availability.

e.g. Variation in growth, mortality and recruitment among New England tree species



*Pacala. Canham
et al (1996)*

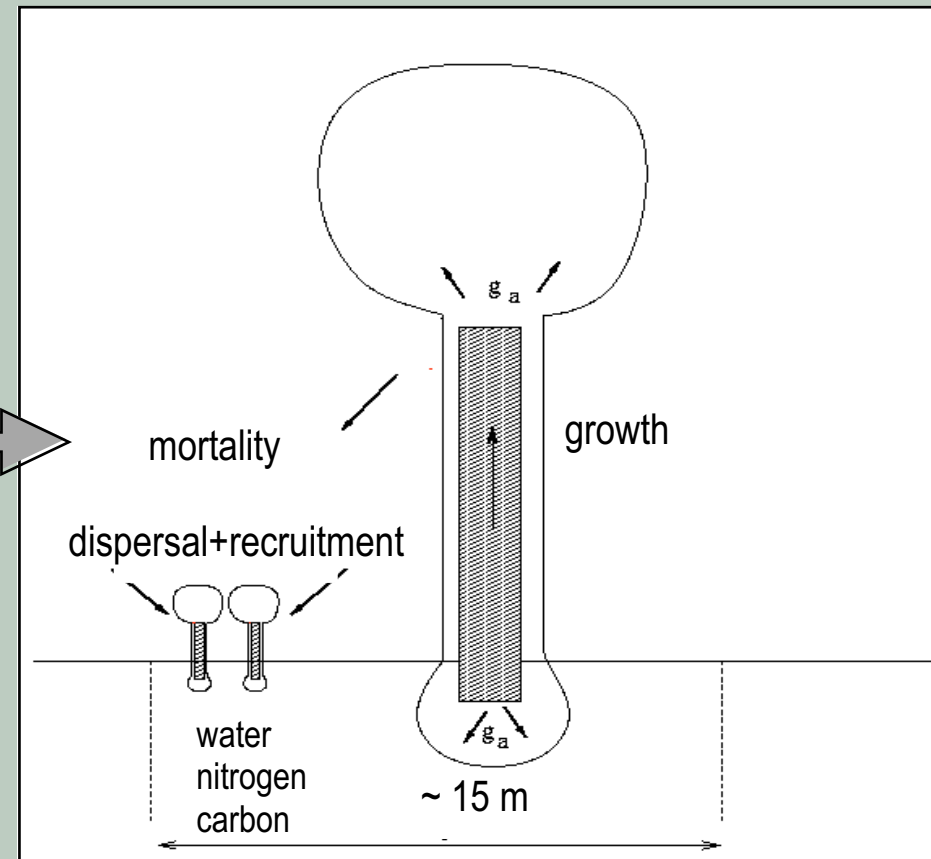
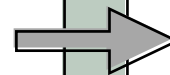
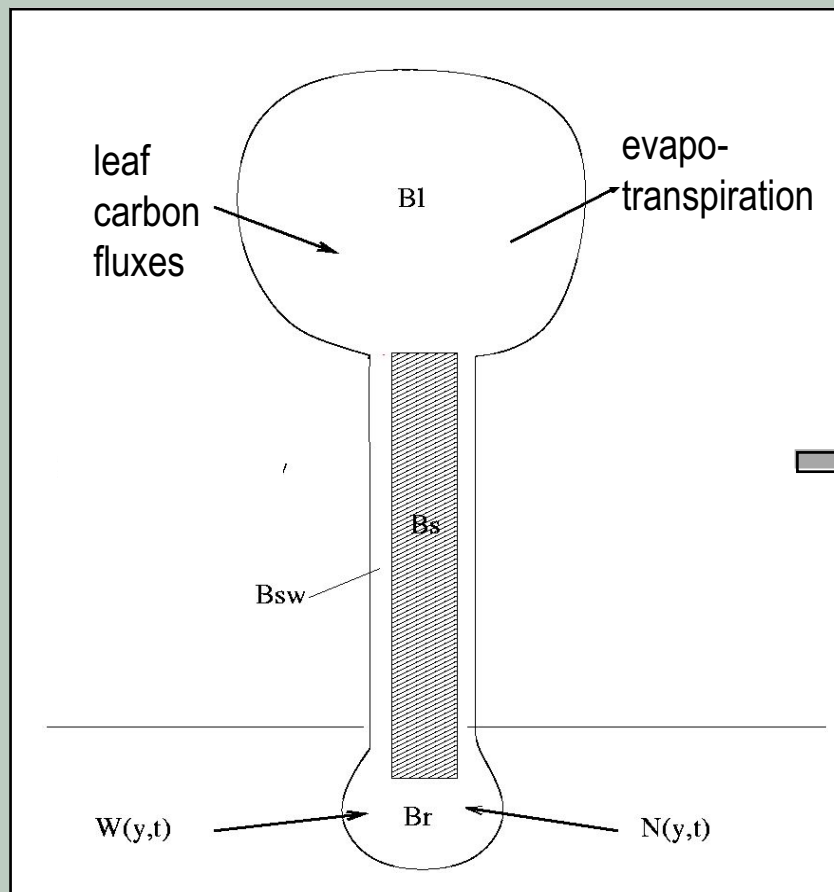
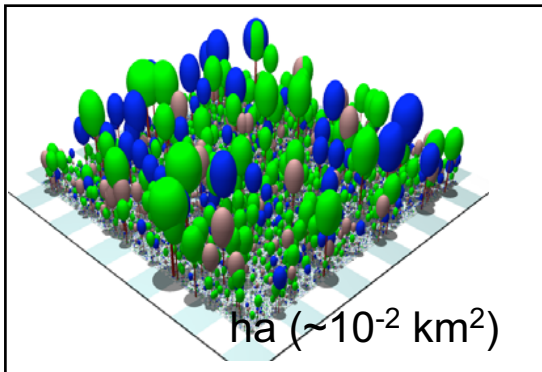
Timescales of Terrestrial Ecosystem Processes

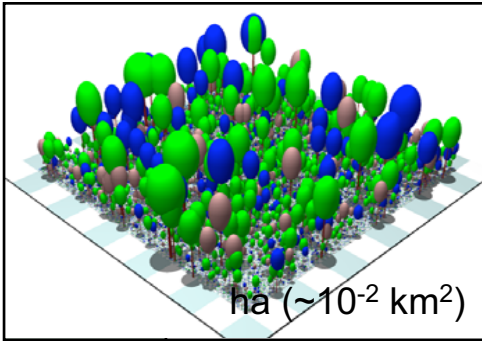


Physiologically-based gap models

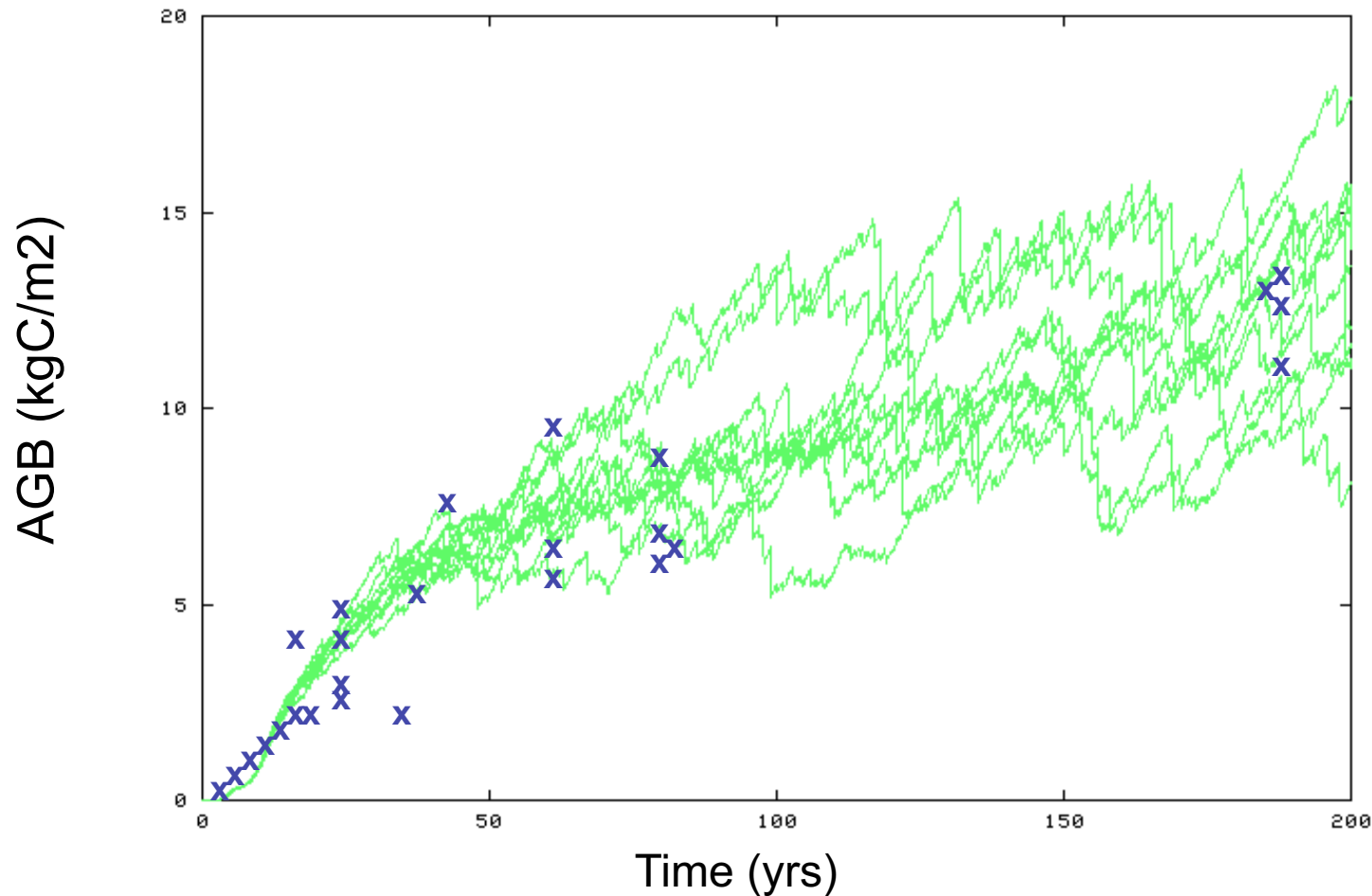
HYBRID (Friend et al. 1997)

ED (Moorcroft et al. 2001)

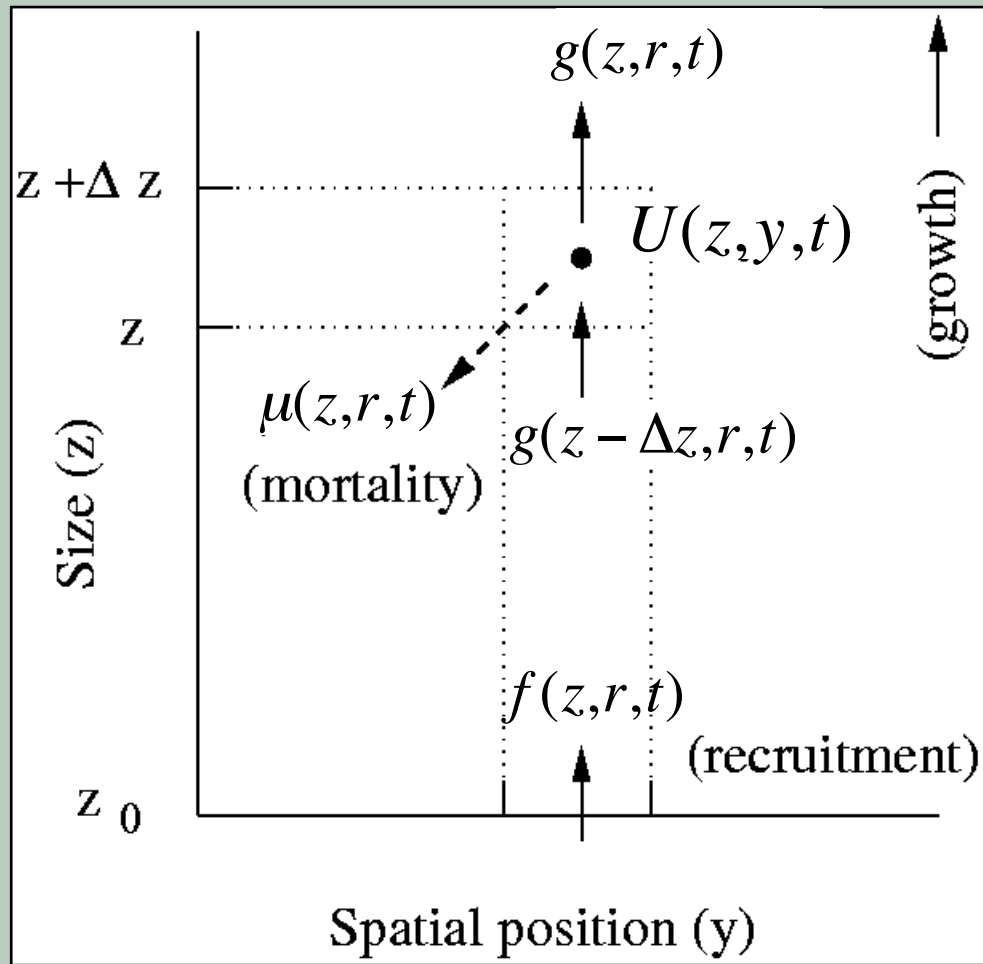




ED Model simulator dynamics at San Carlos (tropical forest) 2°N, 68°W: trajectory of above-ground biomass



Size-structured approximation



$r = r(z, y)$ = resource availability
at location y, z

$$u(z, t) = \langle U(z, y, t) \rangle$$

$$\Delta u(z, t) =$$

$$- \left\langle U(z, y, t) g(z, r, t) \frac{\Delta t}{\Delta z} \right\rangle$$

$$+ \left\langle U(z - \Delta z, y, t) g(z, r, t) \frac{\Delta t}{\Delta z} \right\rangle$$

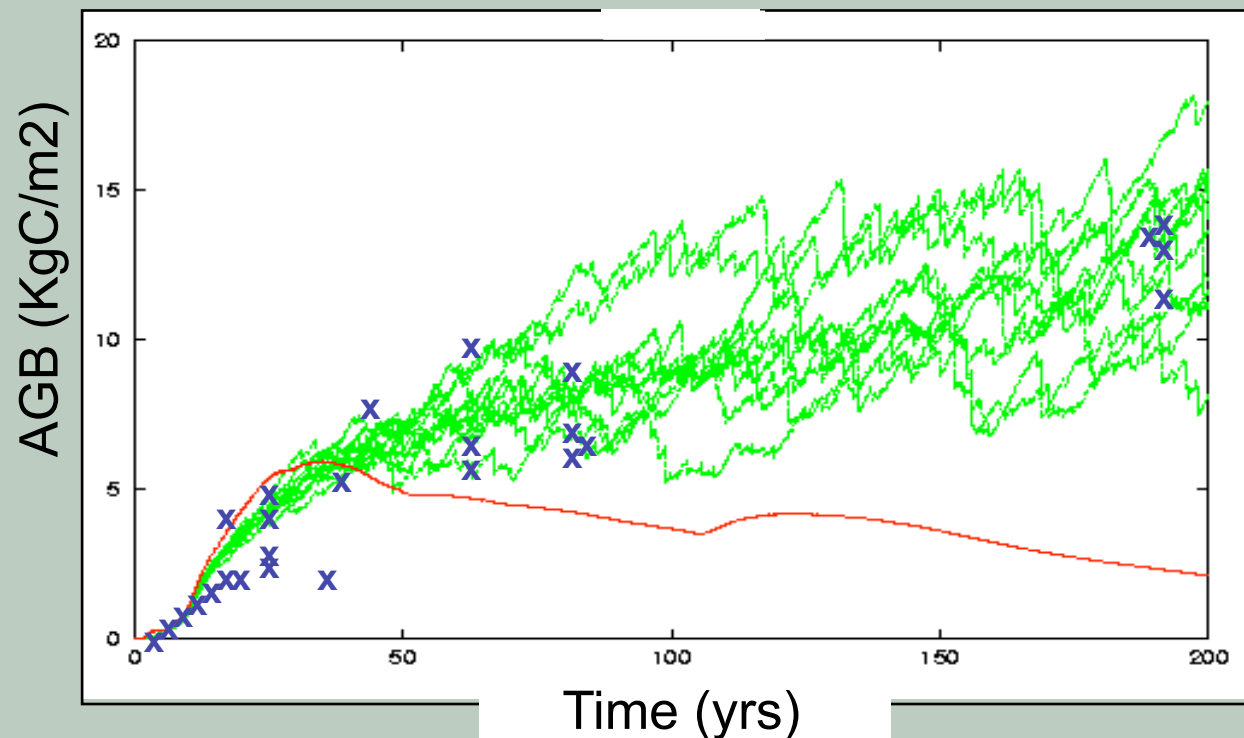
$$- \left\langle U(z, y, t) \mu(z, r, t) \frac{\Delta t}{\Delta z} \right\rangle$$

Size-structured approximation

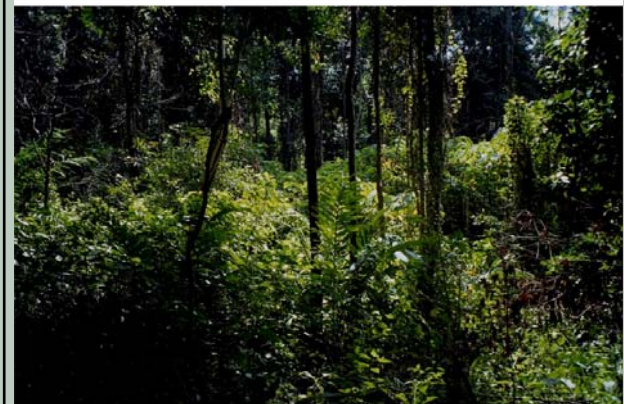
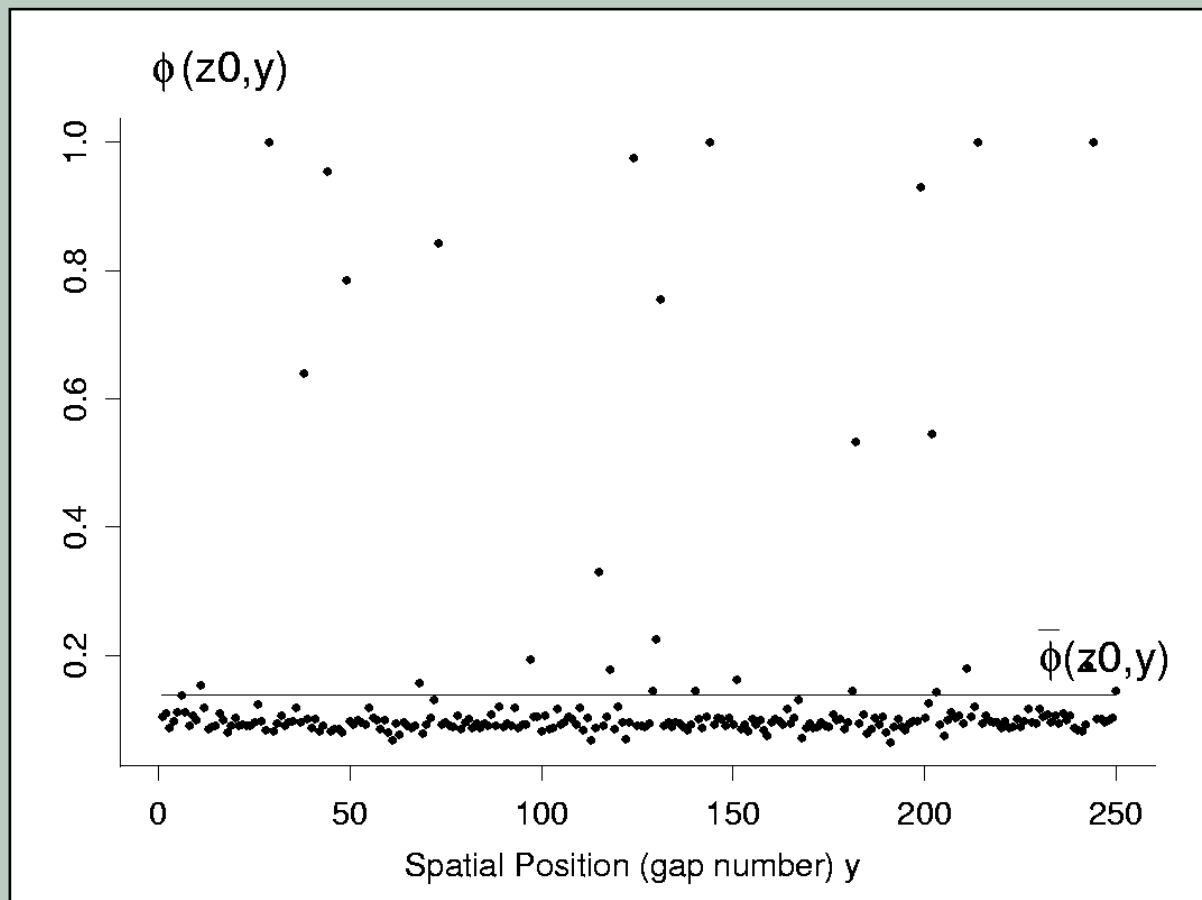
$$\text{PDE: } \underbrace{\frac{\partial n^{(i)}(z,t)}{\partial t}}_{\text{ch. in density of plant type } i} = \underbrace{-\frac{\partial}{\partial z} \left[g(z,t)^{(i)} n^{(i)}(z,t) \right]}_{\text{growth}} - \underbrace{\mu(z,t)^{(i)} n^{(i)}(z,t)}_{\text{mortality}}$$

$$\text{BC: } n^{(i)}(z_0,t) = \underbrace{\frac{1}{g^{(i)}(z_0,t)} \int_0^\infty f^{(i)}(z,t) n^{(i)}(z,t) dz}_{\text{reproduction}} \quad (i = 1 \dots n_{\text{plant types}})$$

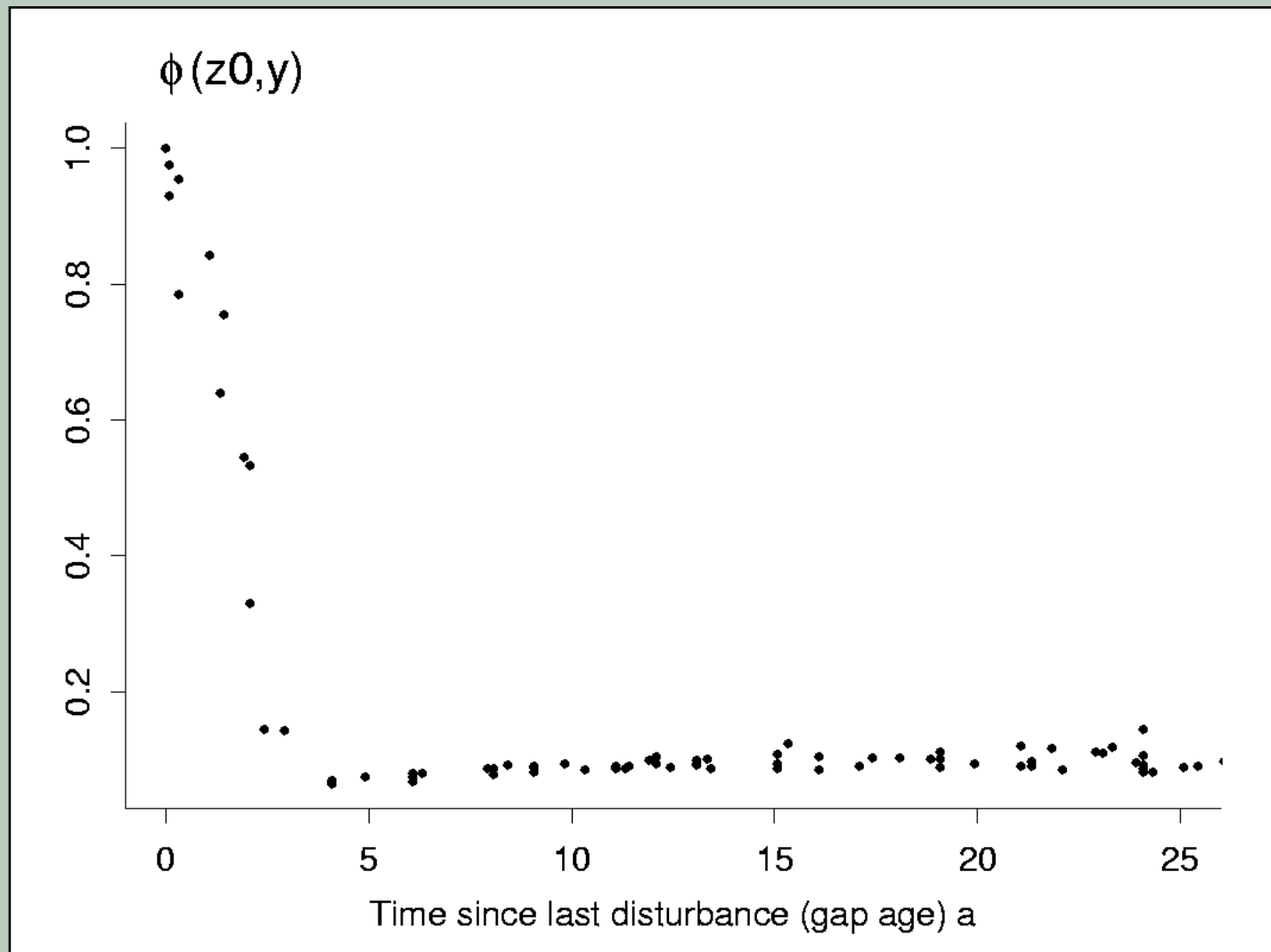
Size-structured approximation biomass dynamics at San Carlos:

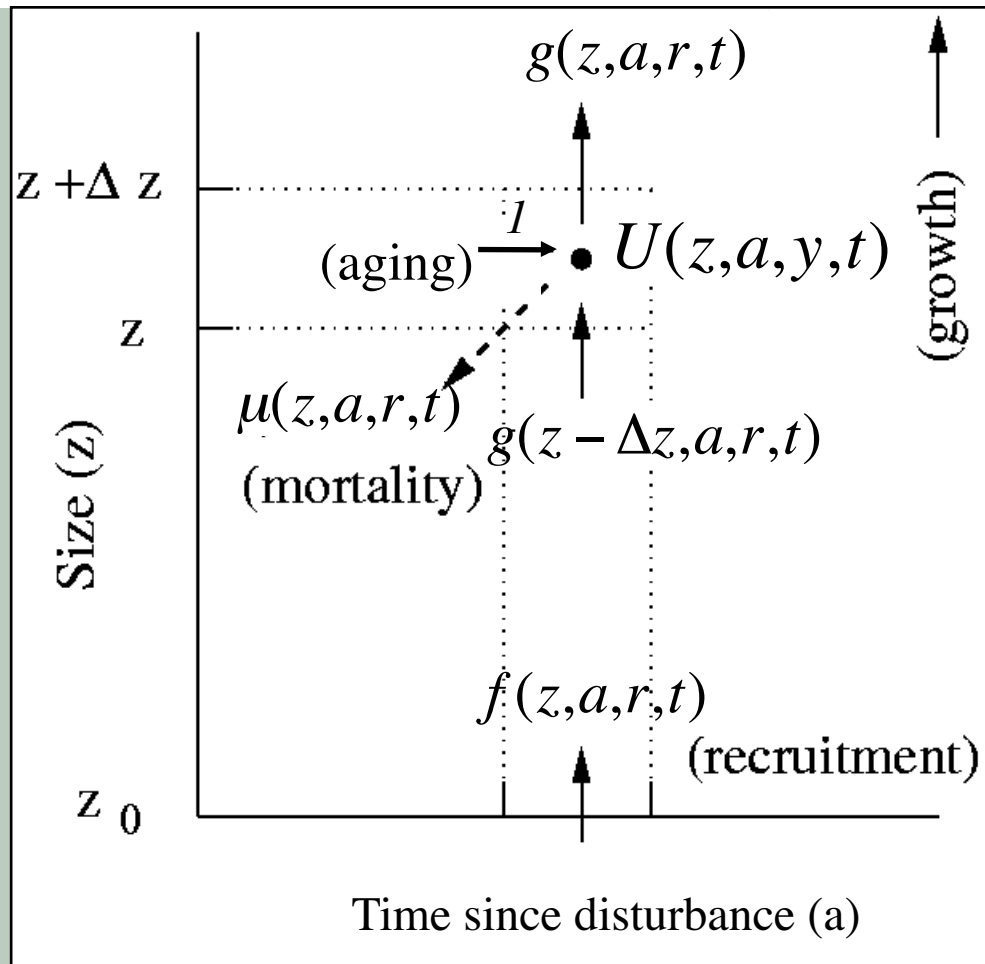


Light availability as a function of spatial position



Light availability as a function of time since disturbance





$$u(z,a,t) = \langle U(z,a,y,t) \rangle$$

Landscape age structure



Levin & Paine (1974)

Disturbance dynamics

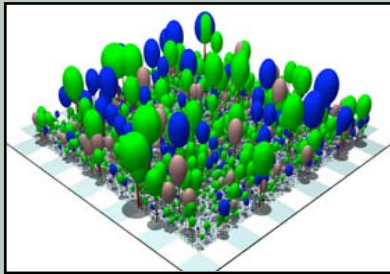
McKendrick Von-Forester Equation:

$$\frac{\partial p(a,t)}{\partial t} = -\frac{\partial}{\partial a} p(a,t) - \lambda(a,t)p(a,t) \quad p(0,t) = \int_0^{\infty} \lambda(a,t)p(a,t) da$$

Where:

$p(a)da$ = proportion of the grid-cell disturbed between a and $a + da$ years ago

$\lambda(a,t)$ = rate of disturbance



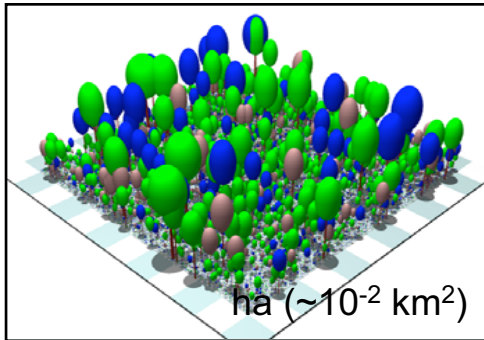
A size & age-structured terrestrial biosphere model

PDE:
$$\underbrace{\frac{\partial n^{(i)}(z,a,t)}{\partial t}}_{\text{ch. in density of plant type } i} = \underbrace{-\frac{\partial}{\partial z} \left[g(z,a,t) n^{(i)}(z,a,t) \right]}_{\text{growth}} - \underbrace{\frac{\partial}{\partial a} n^{(i)}(z,a,t)}_{\text{ageing}} - \underbrace{\mu(z,a,t) n^{(i)}(z,a,t)}_{\text{mortality}}$$

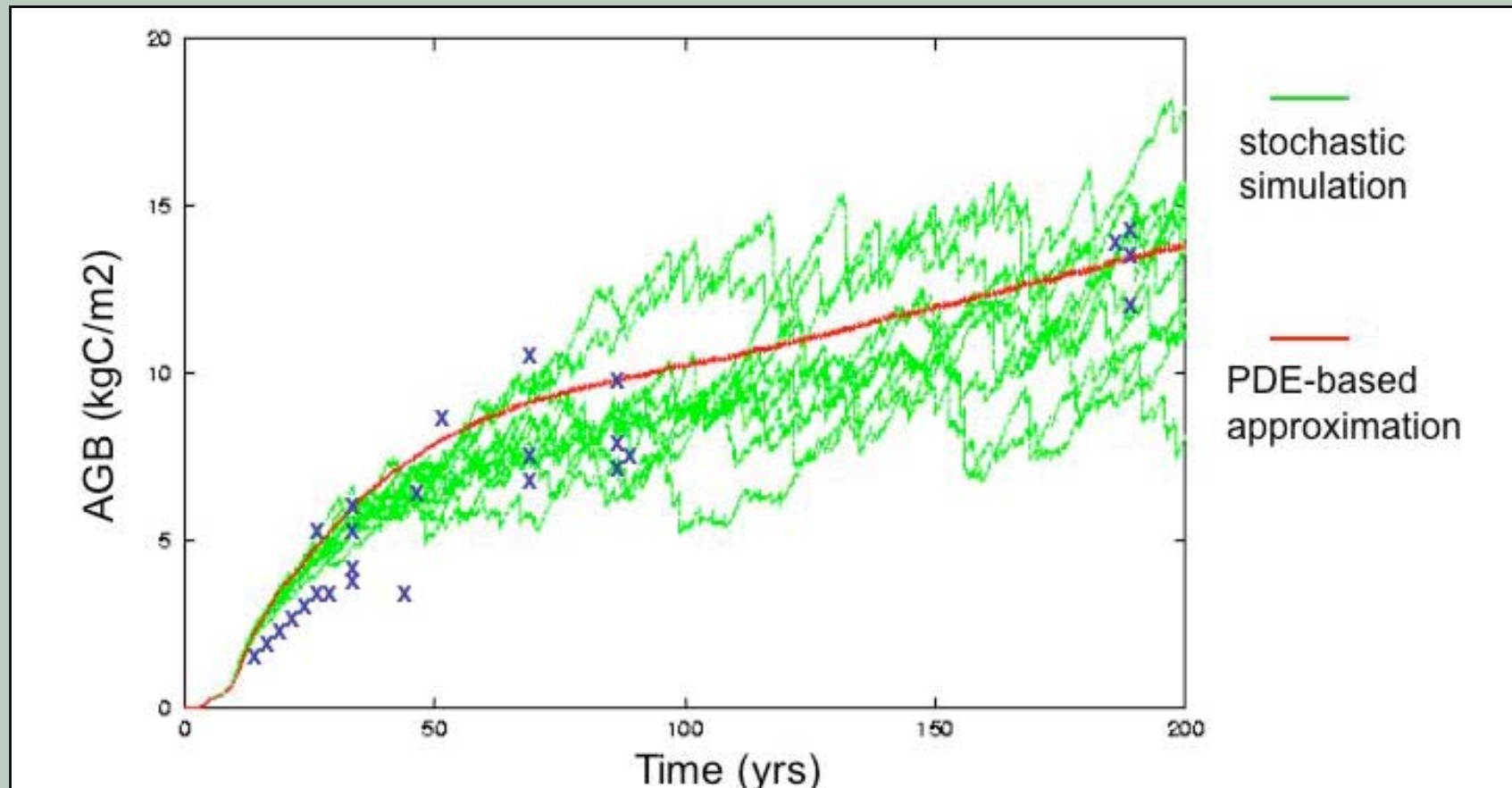
PDE:
$$\underbrace{\frac{\partial p(a,t)}{\partial t}}_{\text{ch. landscape age distribution}} = \underbrace{-\frac{\partial}{\partial a} p(a,t)}_{\text{ageing}} - \underbrace{\lambda(a,t) p(a,t)}_{\text{disturbance}}$$

BC:
$$n(z_0,t) = \frac{1}{g(z_0,t)} \int_0^\infty f(z,t) n(z,t) dz$$
 BC:
$$p(0,t) = \int_0^\infty \lambda(a,t) n(a,t) da$$

BC:
$$n^{(i)}(z,0,t) = s(z,a,t) \int_0^\infty s(z,a,t) n^{(i)}(z,a,t) da$$

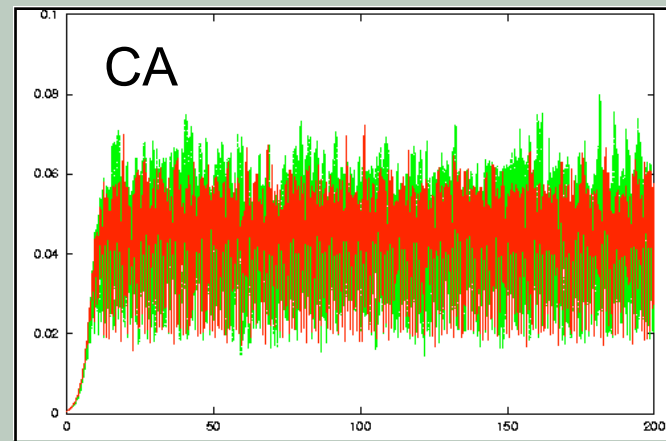
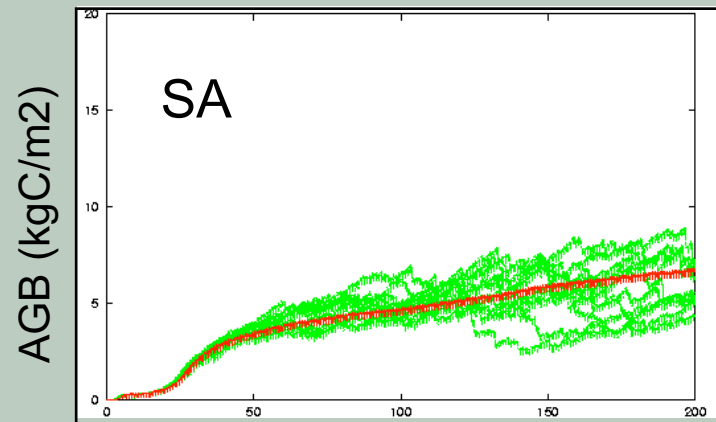
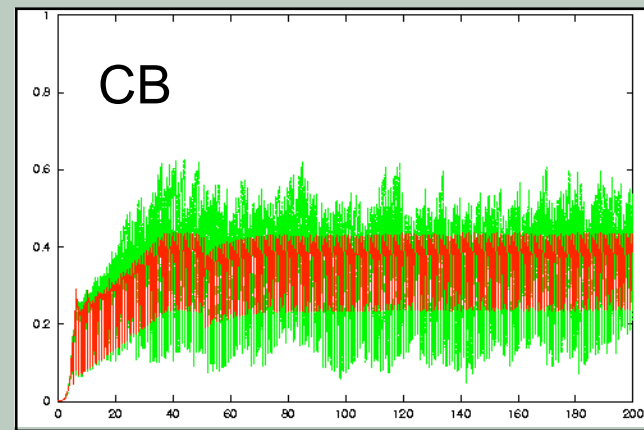
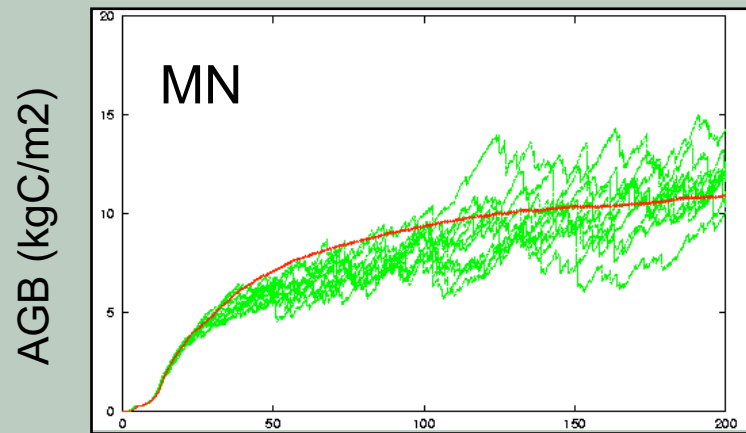
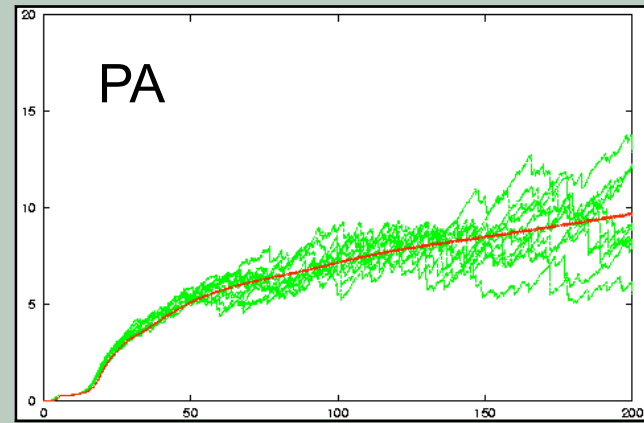
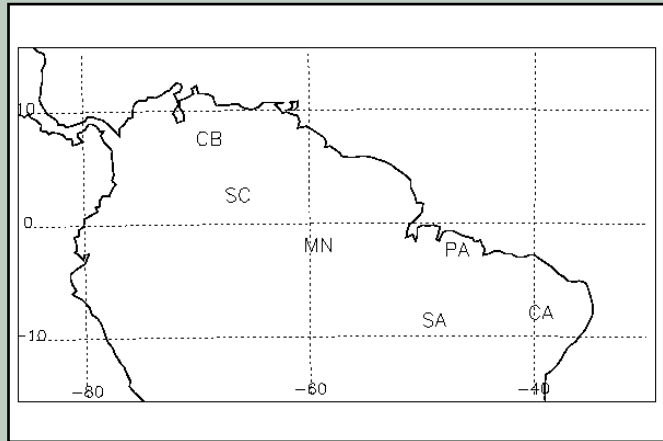


ED dynamics at San Carlos Tropical forest
(2°N, 68°W): trajectory of above-ground
biomass:



- accurately captures the behavior of corresponding individual-based model by tracking the dynamic horizontal & vertical sub-grid scale heterogeneity in canopy structure.

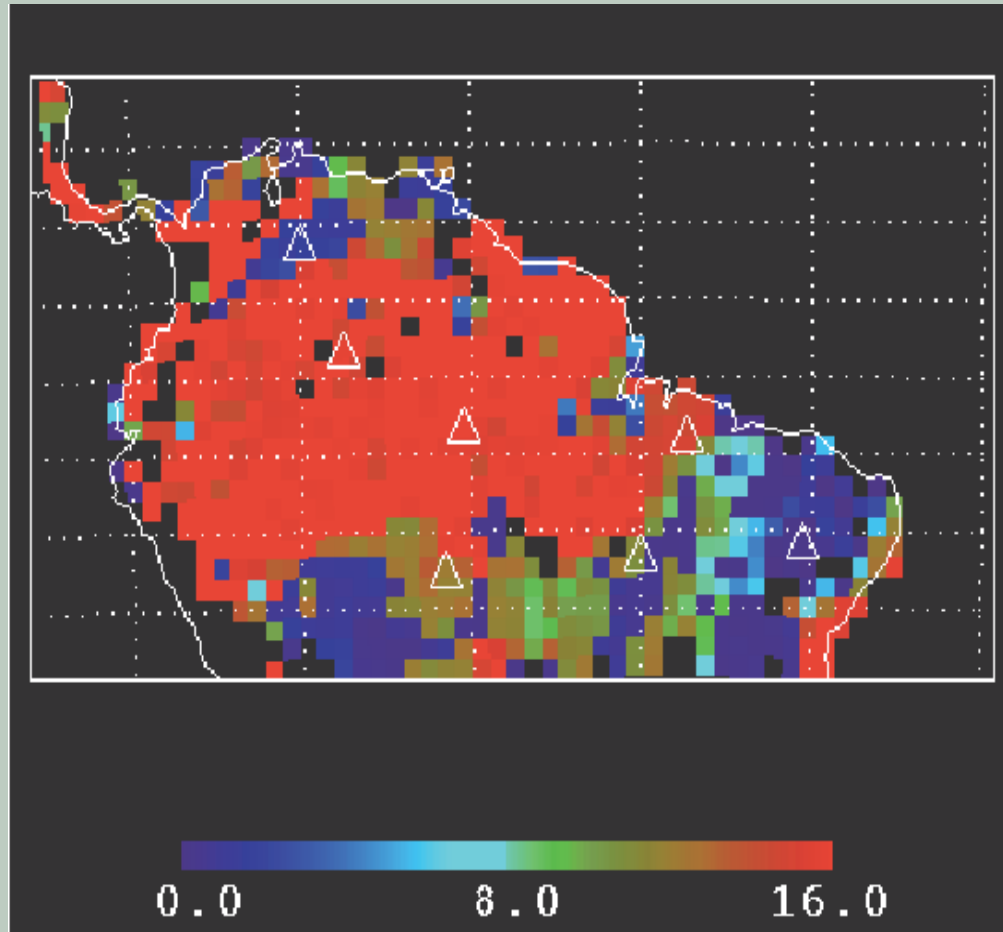
(Moorcroft et al 2001)



Time (yrs)

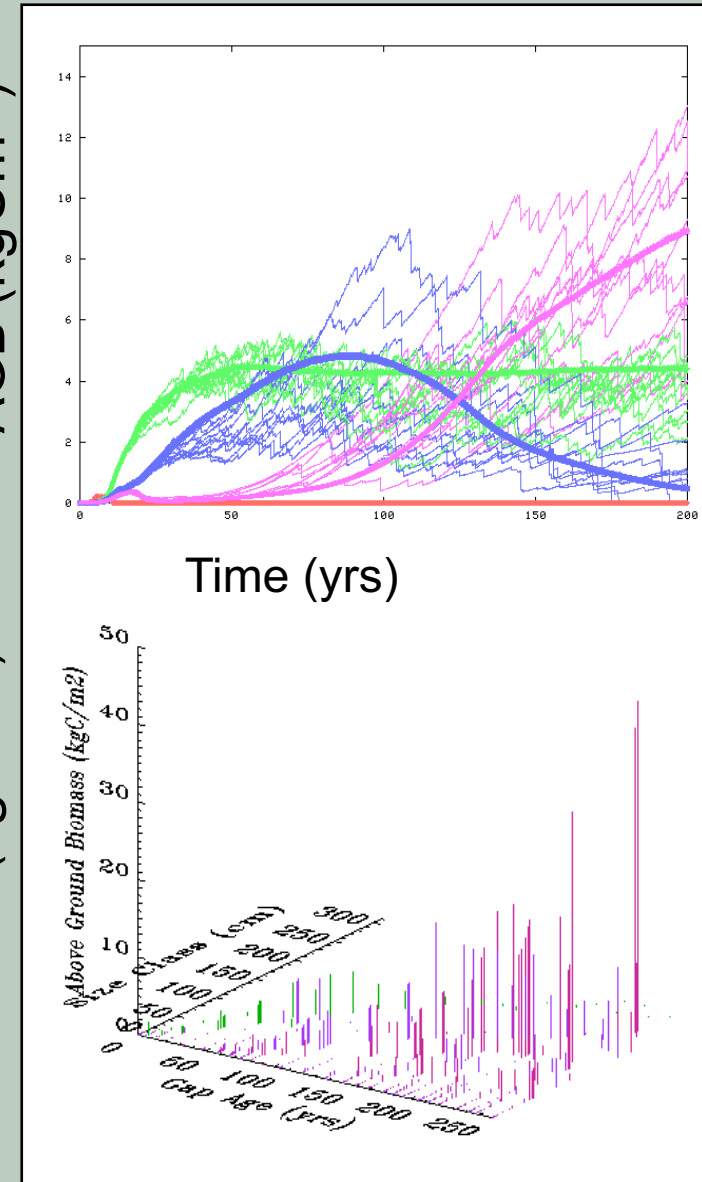
Time (yrs)

ED Model: Regional pattern of above-ground biomass (AGB) after 200 year simulation (kgCm^{-2})



AGB (kgCm^{-2})

AGB (kgCm^{-2})



Forms of disturbance

Treefall Gap Malaysia



Land-slip Malaysia



Fires in Amazonia



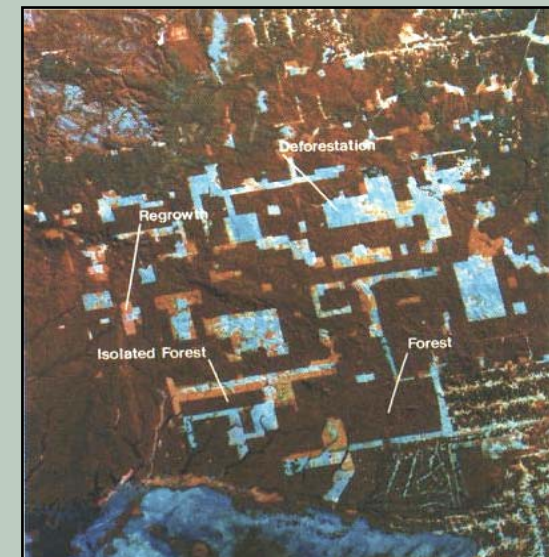
Forest Pathogens Alaska



Shifting Agriculture Venezuela

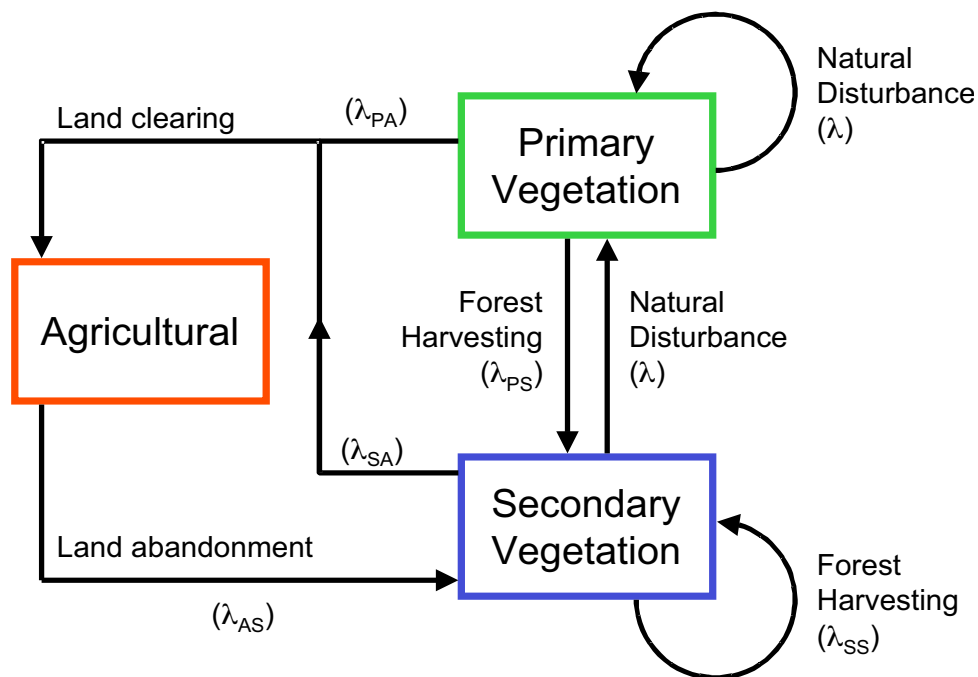


Land-Use Change Rondonia



Incorporating land-use change

$$\underbrace{\frac{\partial}{\partial t} \underline{p}(a, t)}_{\text{ch. in land class \& age distribution}} = \underbrace{-\frac{\partial}{\partial a} \underline{p}(a, t)}_{\text{aging}} - \underbrace{\Lambda(a, t) \underline{p}(a, t)}_{\text{disturbance}}$$



$$\underline{p}(a, t) = [p_1(a), p_j(a), \dots, p_m]]$$

$$\Lambda(a, t) = \begin{pmatrix} \lambda & \lambda & 0 \\ \lambda_{PS} & \lambda_{SS} & \lambda_{AS} \\ \lambda_{PA} & \lambda_{SA} & 0 \end{pmatrix}$$

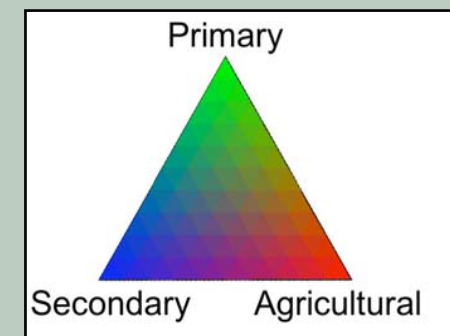
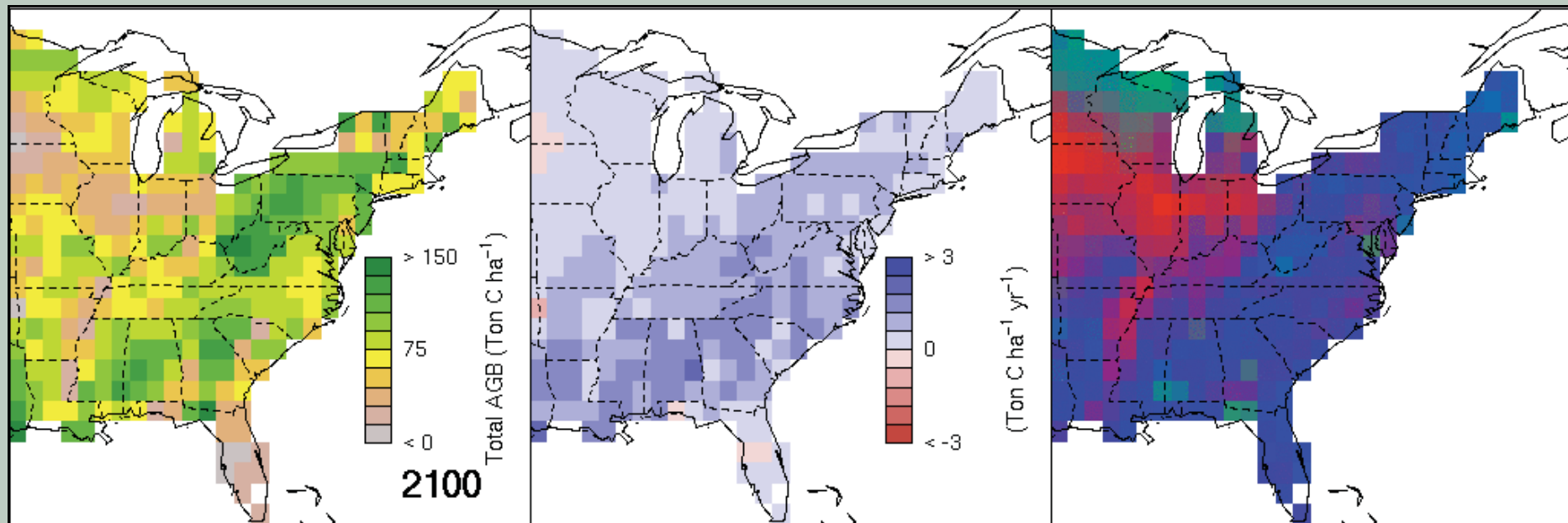
$$\underbrace{\underline{p}(0, t) = \sum_{j=1}^{n_{dist}} \int_0^{\infty} \lambda(a, t) p_j(a, t) da.}_{\text{formation of newly disturbed areas}}$$

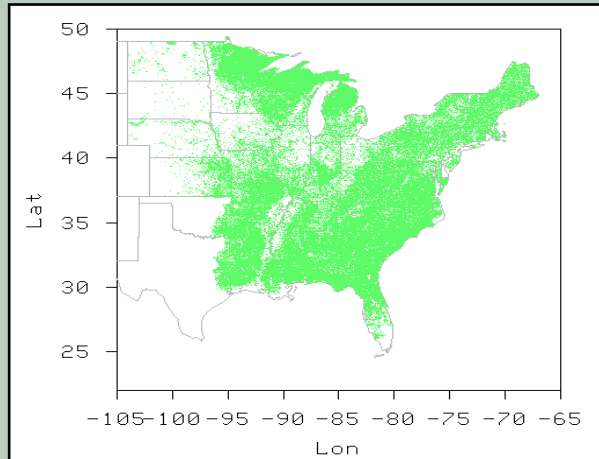
historical fraction of agricultural land in each county 1800-2100

regional historical patterns of forest harvesting (USFS)

ED model: predicted impacts of land-use history on the carbon dynamics of the Eastern US

above gnd. biomass (tC ha⁻¹) carbon uptake (NEP, tC ha⁻¹ y⁻¹) land use

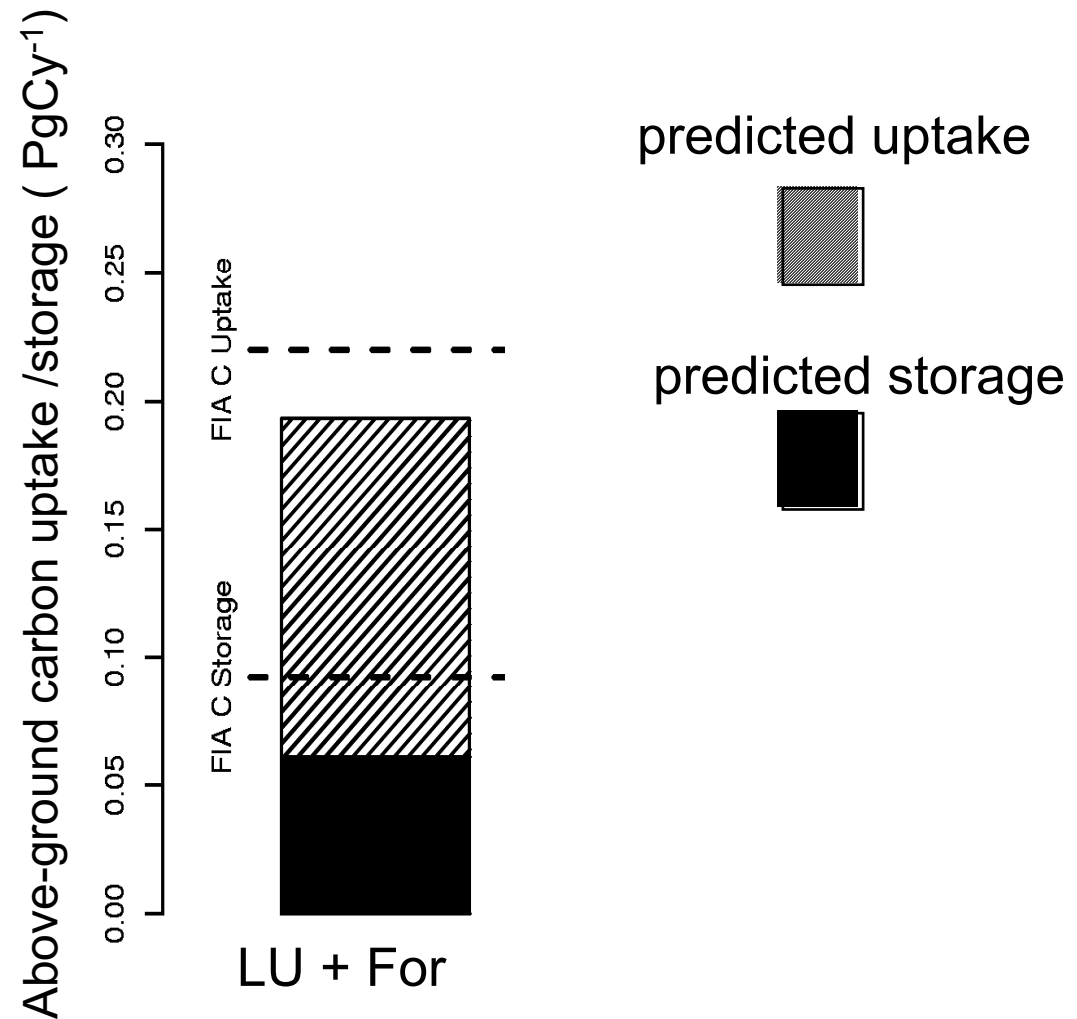




USFS FIA forest inventory measurements of current above-ground C uptake and storage

Results imply that:

- significant carbon uptake occurring as a result of land-use dynamics
- ~ 2/3 of the uptake is forest re-growth following harvesting, not carbon storage in forests.



Conclusions

In contrast to traditional 'big-leaf' biosphere models, structured biosphere models such as ED scale formally between fast timescale plant-level physiological responses to climate, and long-term large-scale ecosystem dynamics.

Enables them to:

- have both realistic short-term and long-term vegetation dynamics.
- incorporate the effects of natural and anthropogenic disturbances (wind-throw fire, land clearing, land abandonment, forest harvesting etc.) on ecosystem composition, structure & function.
- ability to connect to measurements