

Climate sensitivity of global terrestrial ecosystems' subdaily carbon, water, and energy dynamics (an application of information flow process networks)

Workshop on Information Theory in the Geosciences
Schneefernerhaus, Zugspitze, Germany
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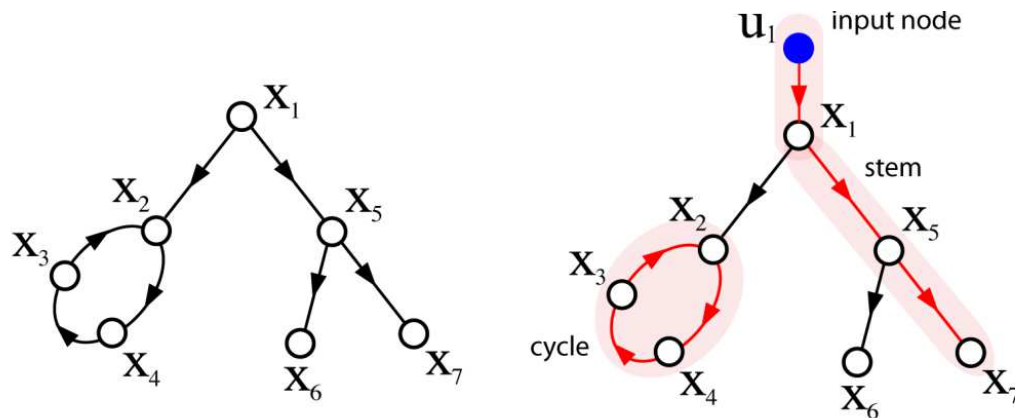


Macrosystems Biology Grant
2013-#1241960 (Conclusions
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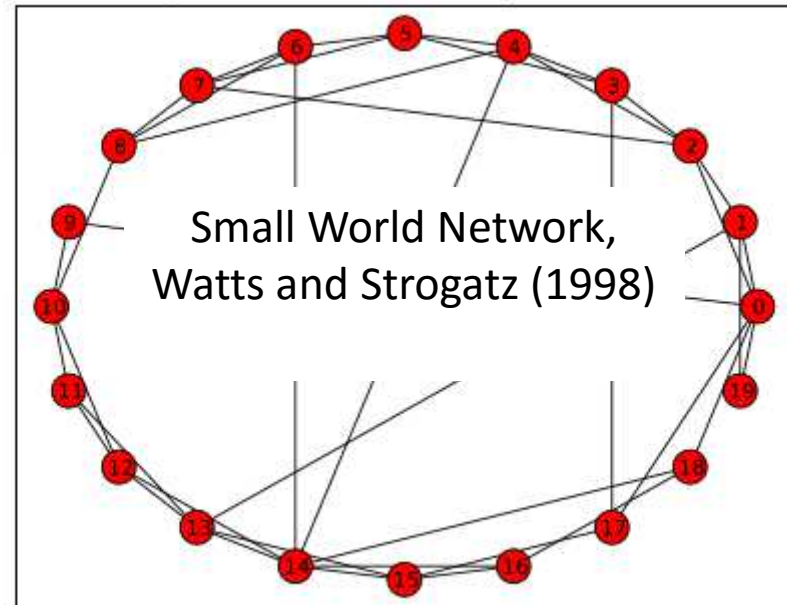
Benjamin L. Ruddell · bruddell@asu.edu
Dan Childers
Rong Yu
Minseok Kang
Joon Kim

Classic Network Theory Applications (are limited)

Control Centrality, Liu et al. (2012)

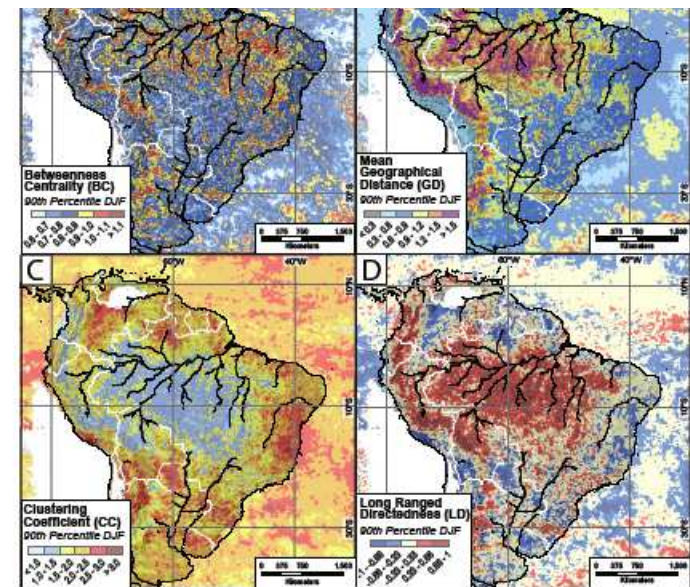
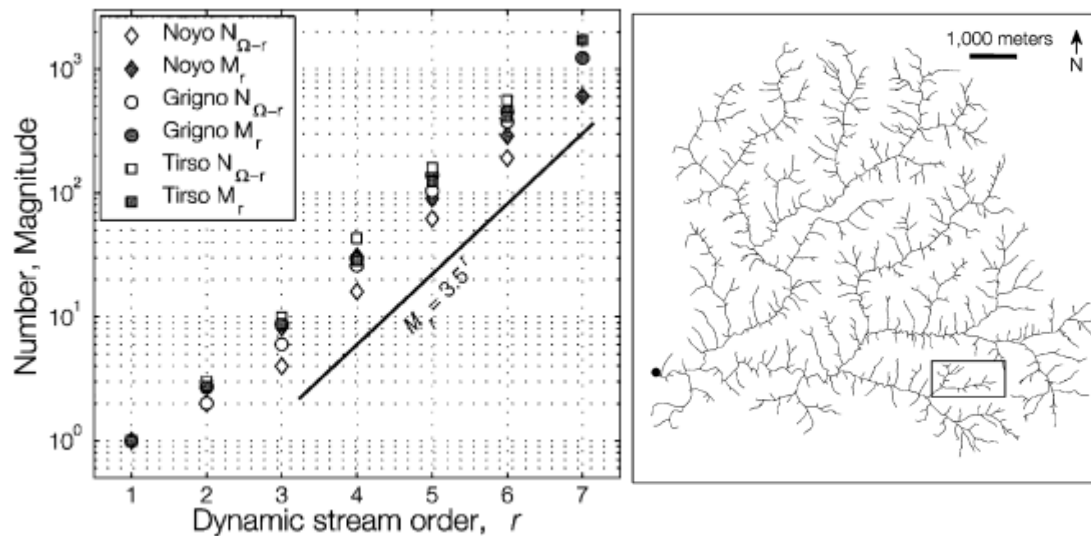


Watts-Strogatz model $N=20, K=4, \beta=0.2$



Betweenness Centrality for Amazon Rainfall, Boers et al. (2013)

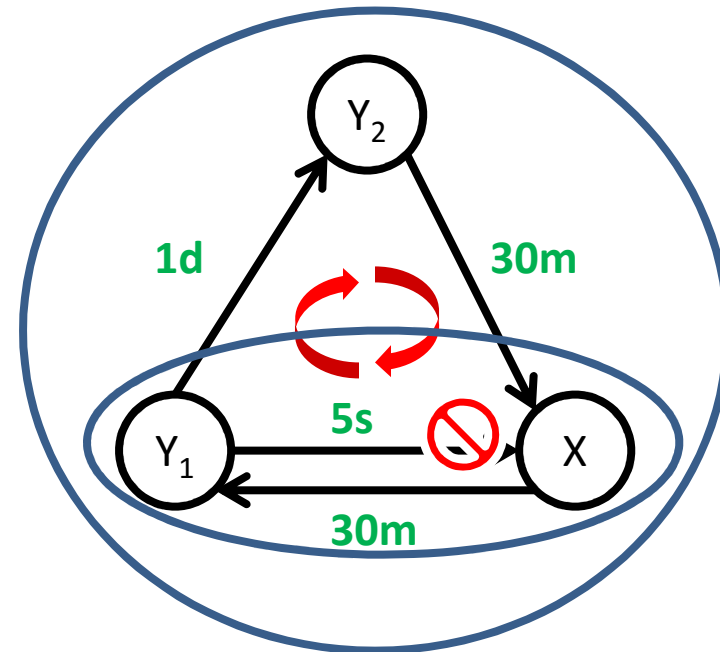
River Basin Power Law Scaling, Zaliapin et al. (2010)



Process Networks

Complex systems generally feature coupling and *feedback* between many nodes, producing self-organizing subsystem behavior, and/or *thresholds* where key couplings turn on and off and qualitatively different system states emerge (Kumar 2007, Liu et al. 2007).

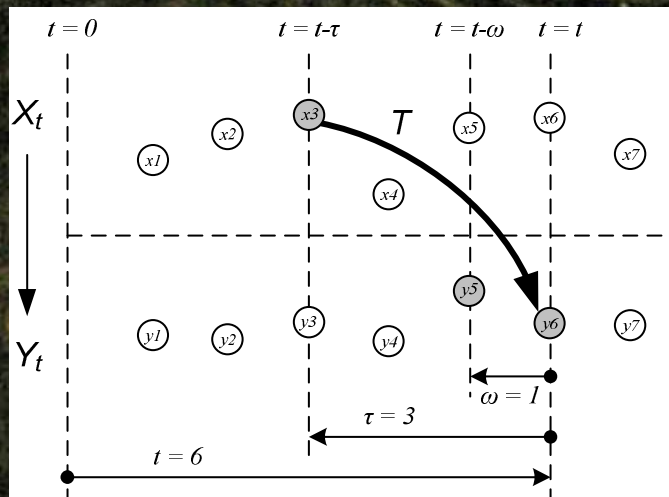
- Hierarchies of *self-organizing subsystems* can emerge via feedback.
- Connections have *characteristic timescales* at which processes operate.
- Connections have a *type, direction,* and *strength* (and possibly follow rules)
- In a *multitype network*, connections and nodes may be qualitatively different.



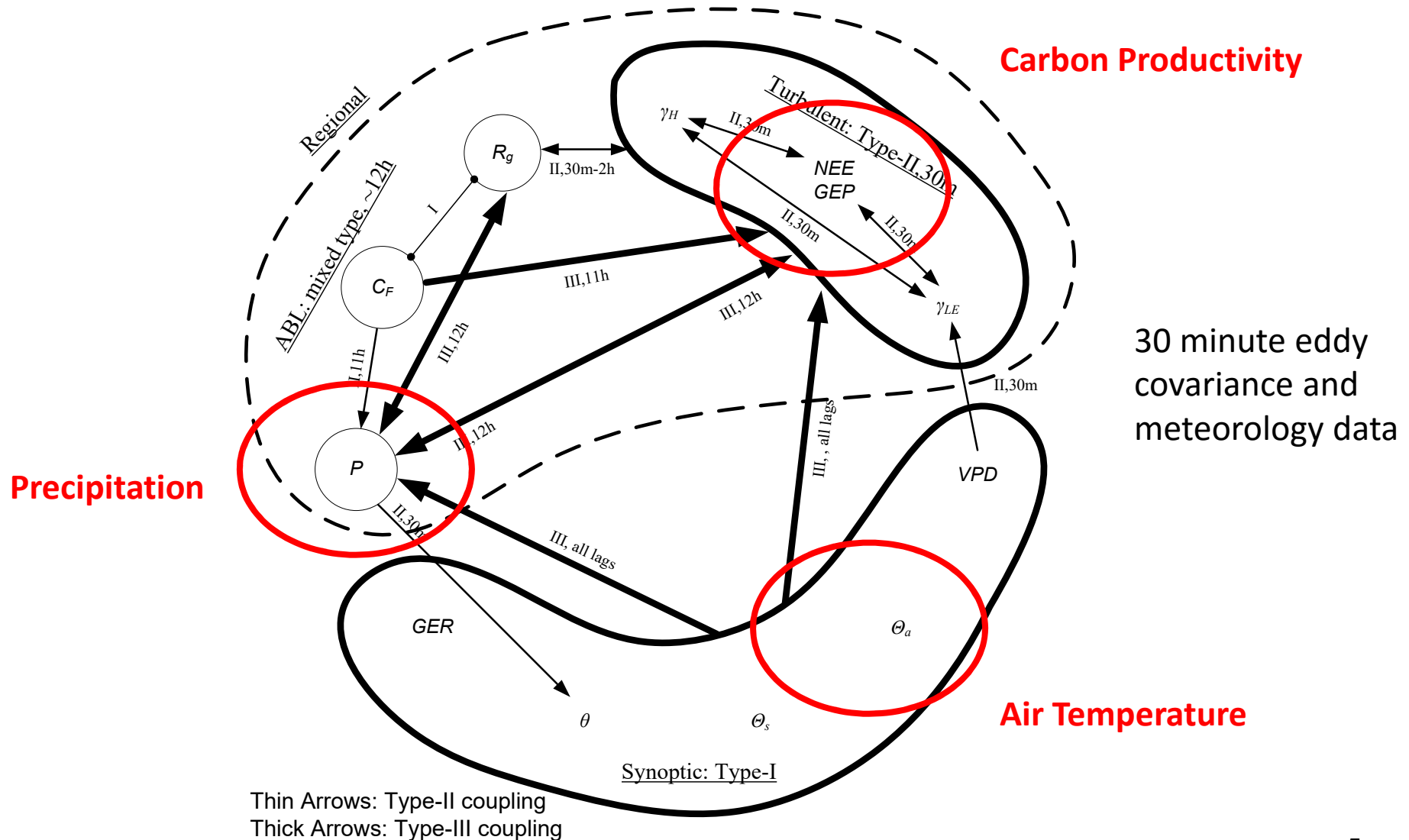
A **Process Network (PN)** is a network of feedback loops and the associated timescales that depicts the magnitude and direction of flow between the different subsystems. The PN graph itself defines the system state. (Ruddell and Kumar, 2009a)

We can robustly observe short-timescale dynamical sensitivity of ecosystems to climate forcings using advanced statistics to infer **DYNAMICAL PROCESS NETWORKS**. (Transfer Entropy, Schreiber 2000)

$$T(X_t \rightarrow Y_t, \tau) = \sum_{y_t, y_{t-1}, x_{t-\tau}} p(y_t, y_{t-1}, x_{t-\tau}) \log \frac{p(y_t | (y_{t-1}, x_{t-\tau}))}{p(y_t | y_{t-1})}$$

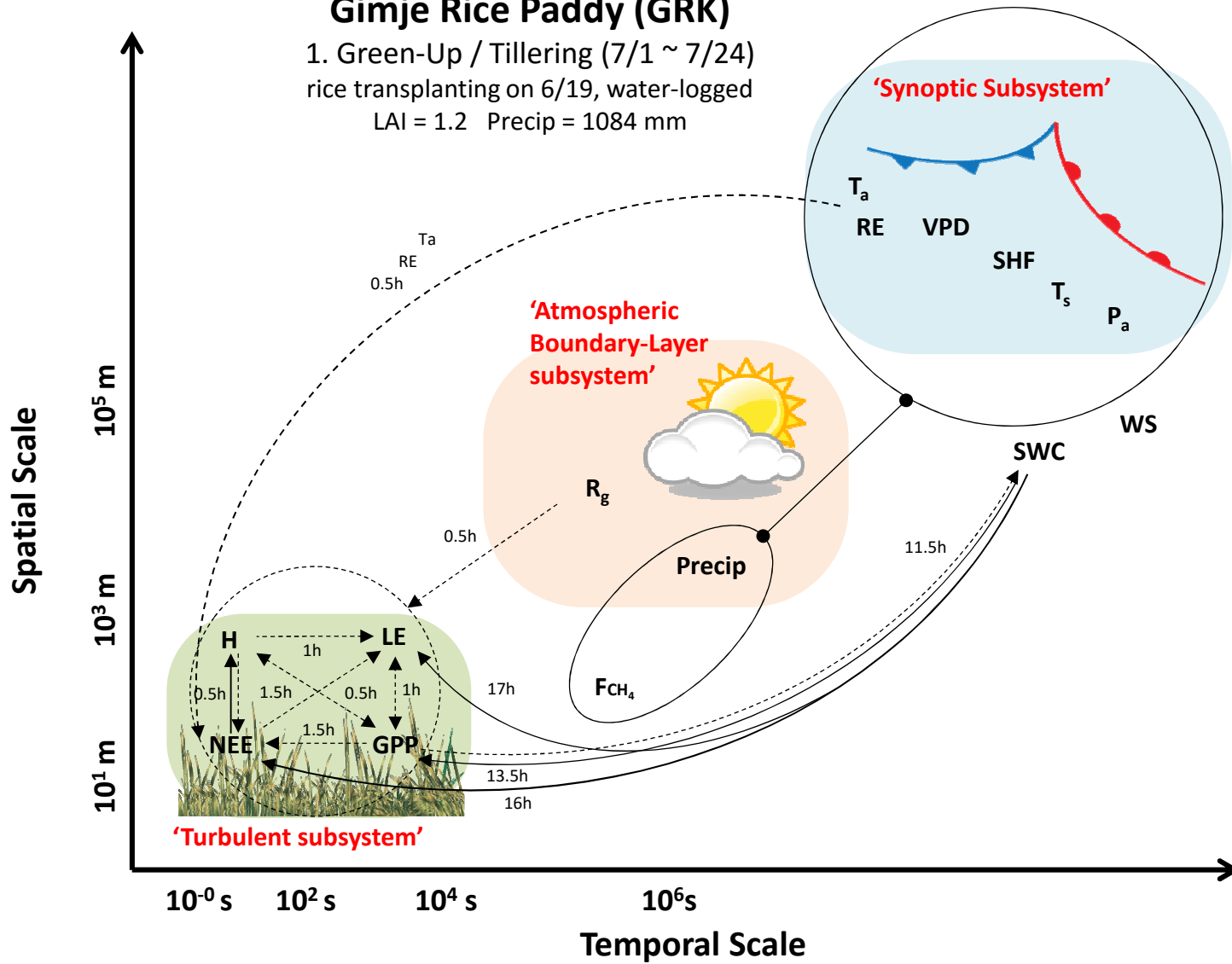


A Dynamical Process Network (DPN) for Illinois Corn+Climate (a graph-based state based on 'fast' functional dynamics)



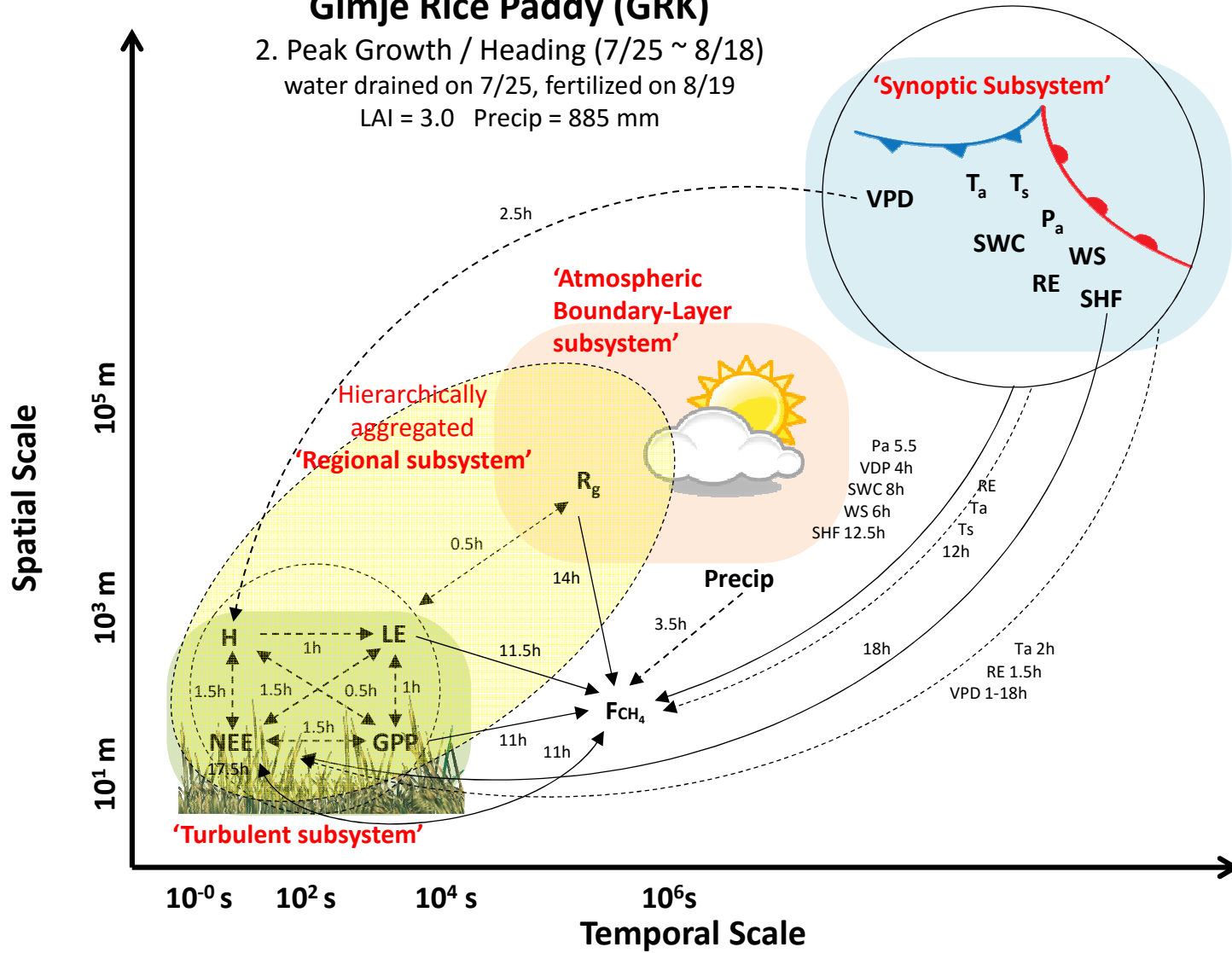
Gimje Rice Paddy (GRK)

1. Green-Up / Tillering (7/1 ~ 7/24)
 rice transplanting on 6/19, water-logged
 LAI = 1.2 Precip = 1084 mm



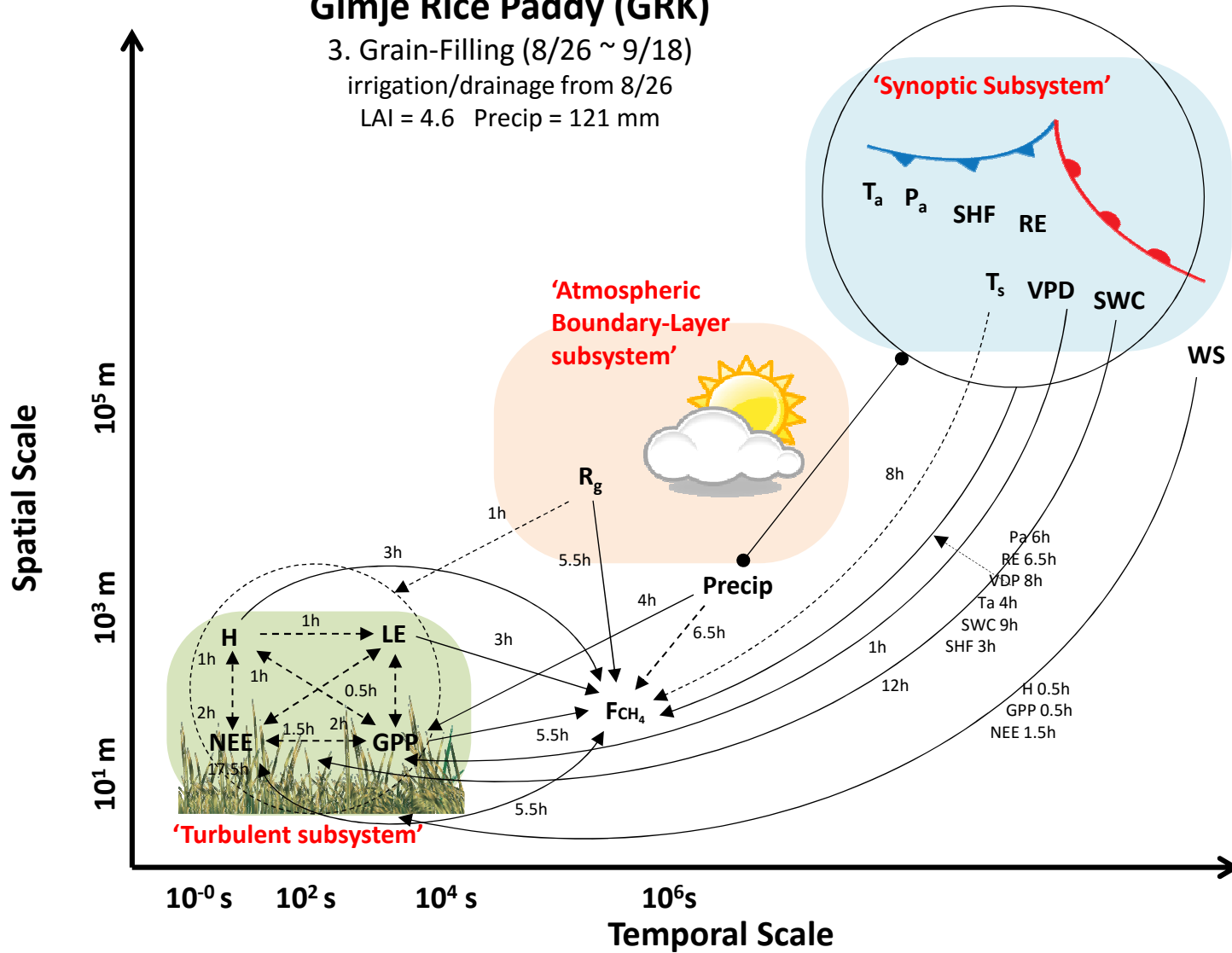
Gimje Rice Paddy (GRK)

2. Peak Growth / Heading (7/25 ~ 8/18)
 water drained on 7/25, fertilized on 8/19
 LAI = 3.0 Precip = 885 mm



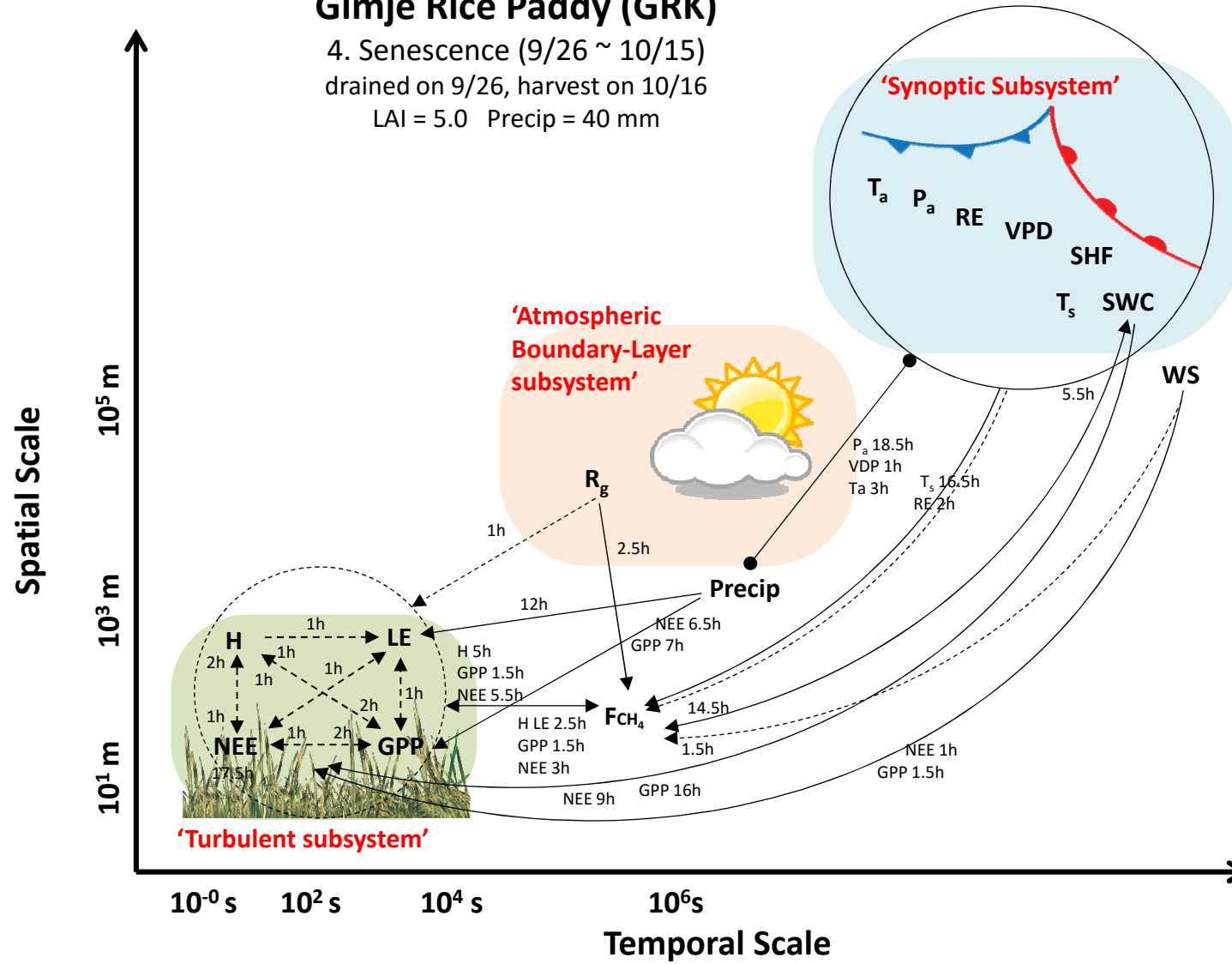
Gimje Rice Paddy (GRK)

3. Grain-Filling (8/26 ~ 9/18)
 irrigation/drainage from 8/26
 LAI = 4.6 Precip = 121 mm



Gimje Rice Paddy (GRK)

4. Senescence (9/26 ~ 10/15)
 drained on 9/26, harvest on 10/16
 LAI = 5.0 Precip = 40 mm

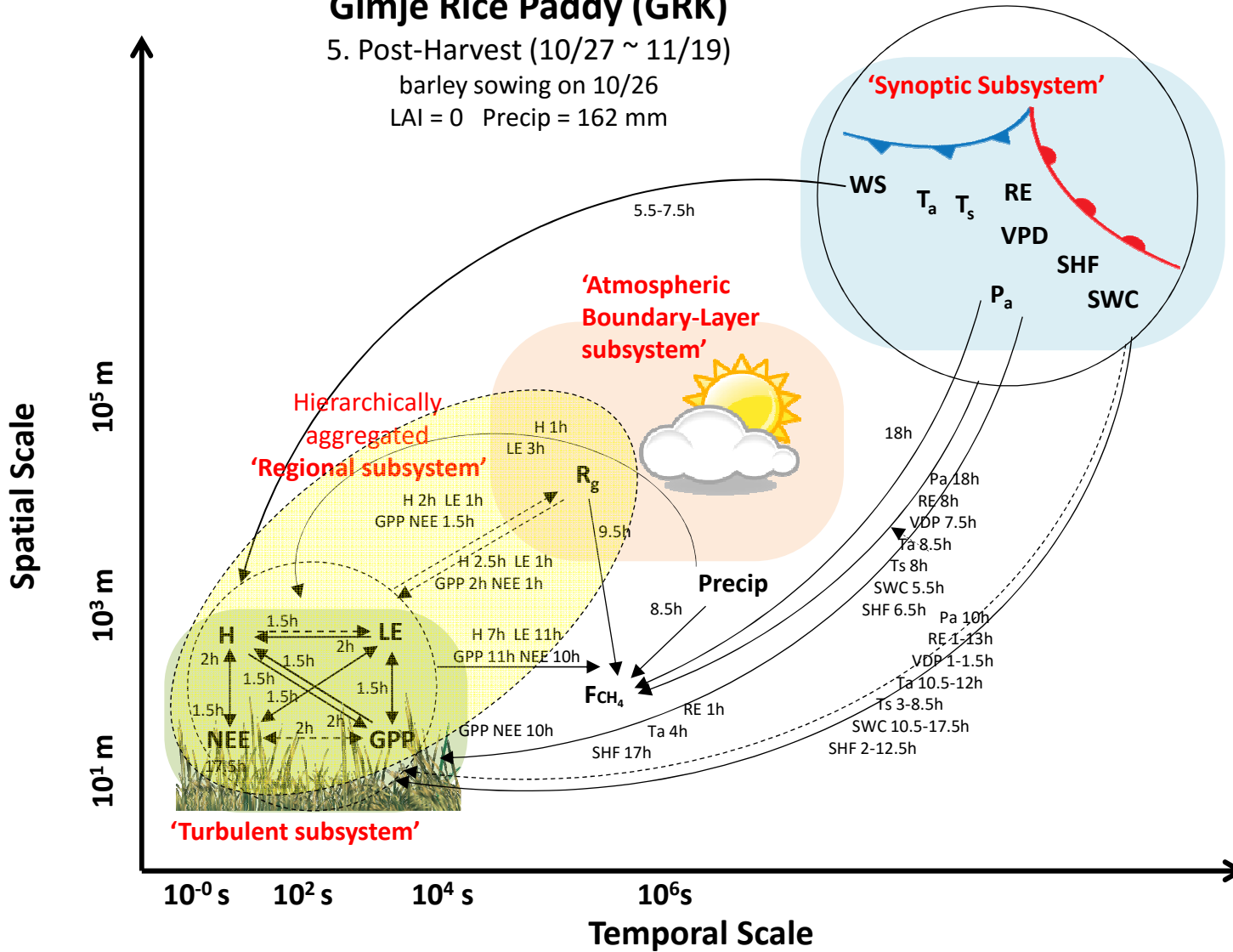


Gimje Rice Paddy (GRK)

5. Post-Harvest (10/27 ~ 11/19)

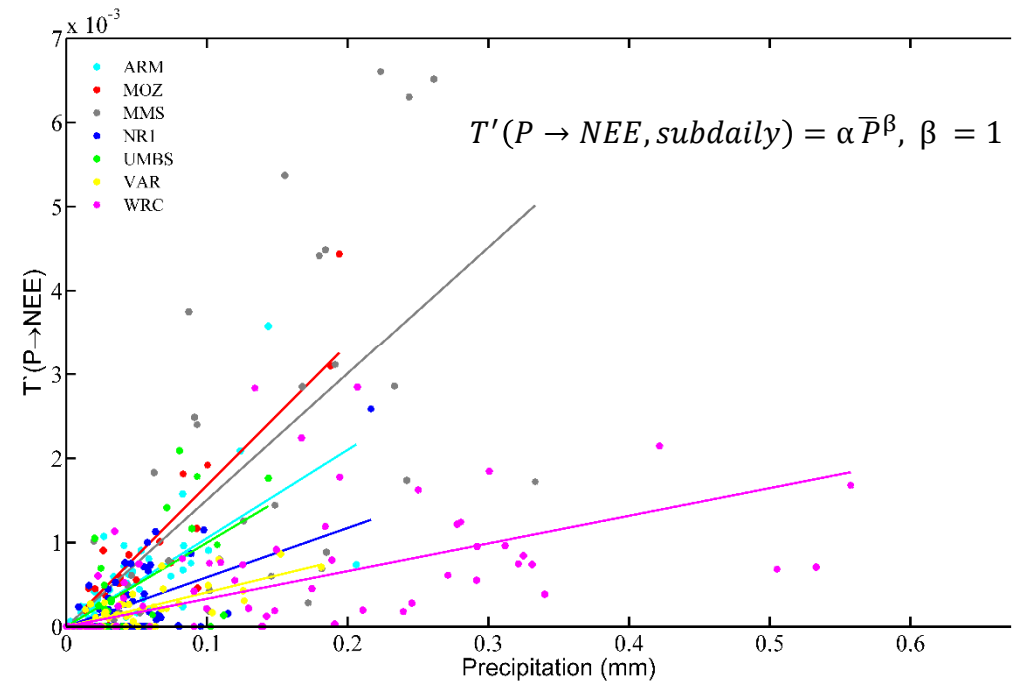
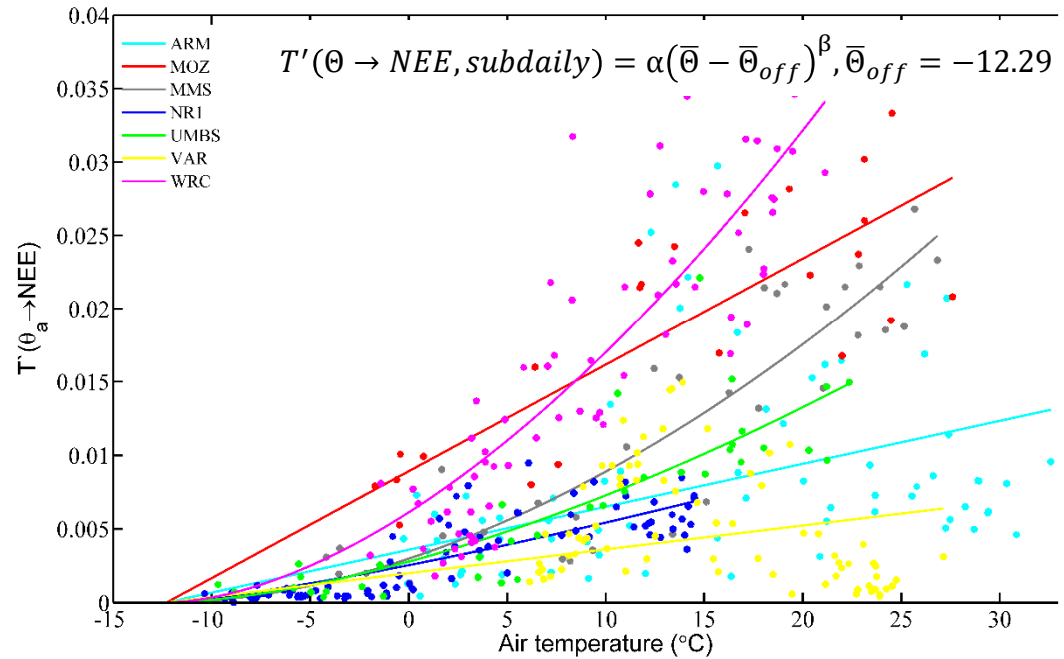
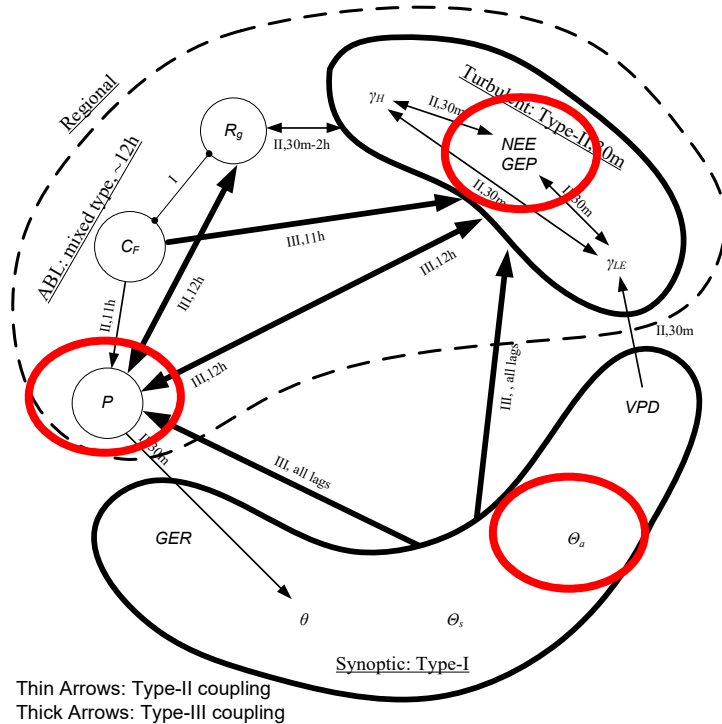
barley sowing on 10/26

LAI = 0 Precip = 162 mm



Elasticity of DPN Information Flows to “slow” forcing factors is often exponential

2003 July: Healthy System State



An Experimental Approach for Ecosystem Change

- ‘Slow’ Dynamics define **SYSTEM STATE** (or ‘macrostate’)
- To robustly predict long-term ecosystem change we need to observe **LONG TERM DYNAMICS OF STATE CHANGE**, but we don’t (yet) have data (paleo?)
- We can however observe the rich **SHORT TERM ‘FAST’ DYNAMICS** of function and structure of the subsystems within each macrostate (the Process Networks)
- By modeling (marginal) changes in those Process Networks as an **ELASTIC RESPONSE** to (marginal) changes in ‘slow’ forcing factors, we have the capability to estimate the response of ecosystem state (and process) to changes in forcings.

We find that Process Network Elasticity to ‘slow’ forcing factors can often be modeled using exponential functions derived from econometrics; these also separate combined and indirect effects.

Seasonal “slow” Forcing Factors are:

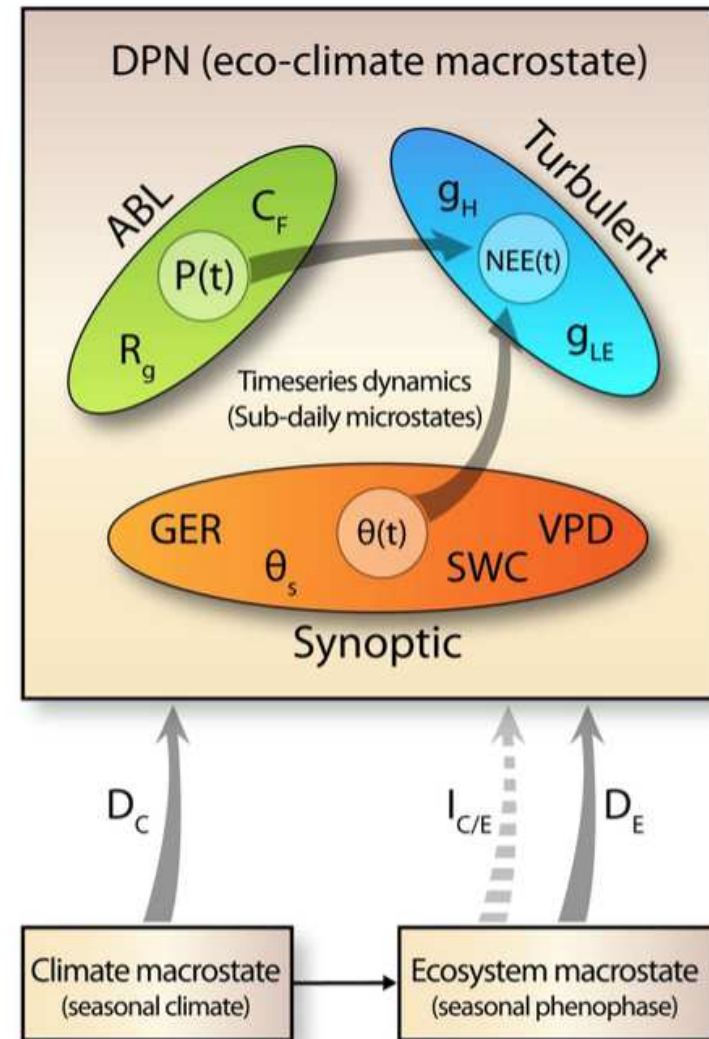
- Temperature
- Radiation
- Precipitation
- EVI/phenophase

Optimize for coefficients using OLS

Coefficients are the elasticities

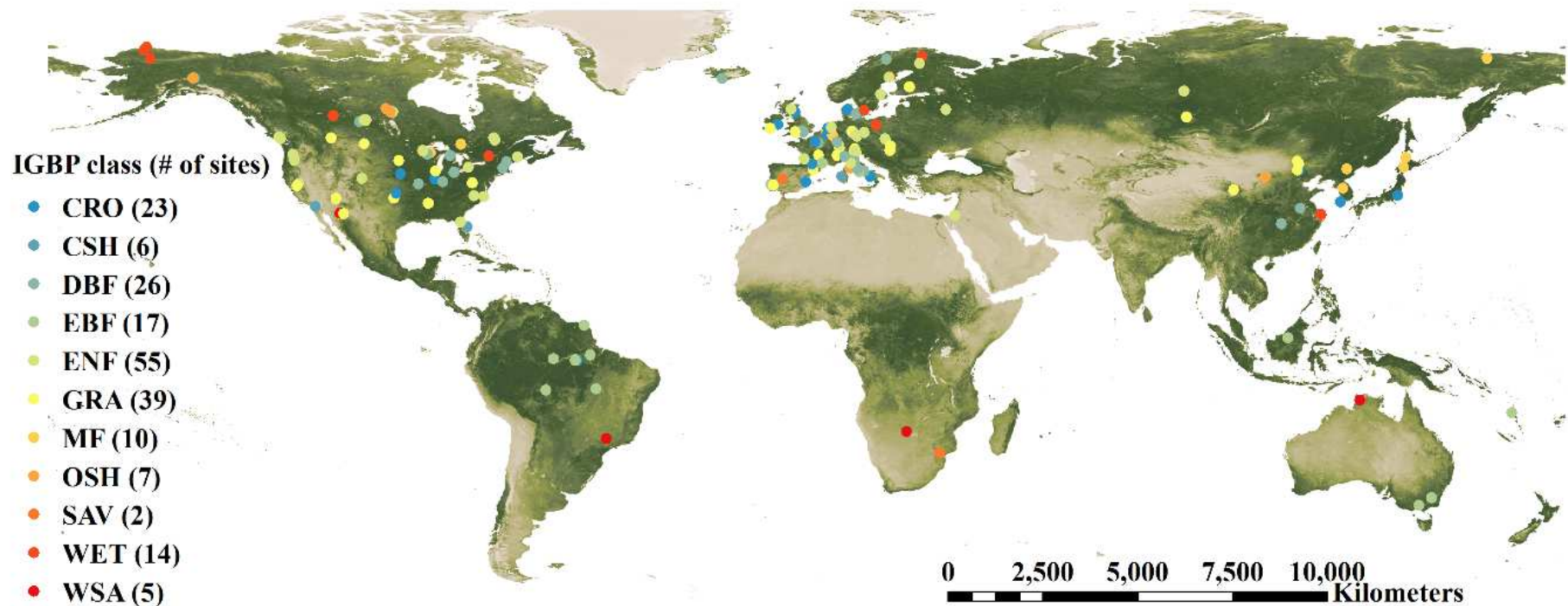
$$T^* = \alpha \cdot \bar{\theta}^{\beta_{\bar{\theta}}} \cdot \bar{P}^{\beta_{\bar{P}}} \cdot \bar{R}^{\beta_{\bar{R}}} \cdot \overline{EVI}^{C_{\bar{E}}}$$

$$C_{\bar{E}} = \beta_{\bar{E}} + \gamma_{\bar{\theta}} \ln \bar{\theta} + \gamma_{\bar{P}} \ln \bar{P}$$

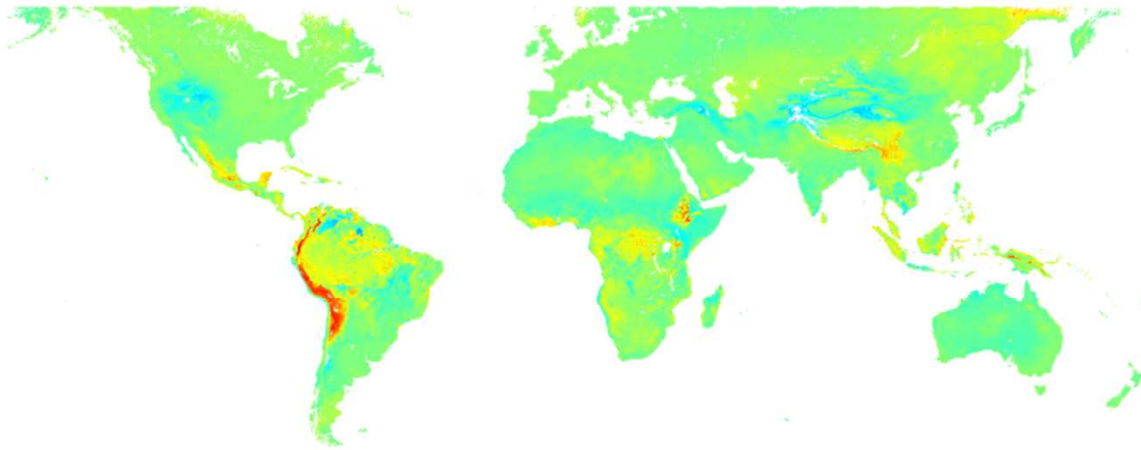


Big Ecological Data: FLUXNET

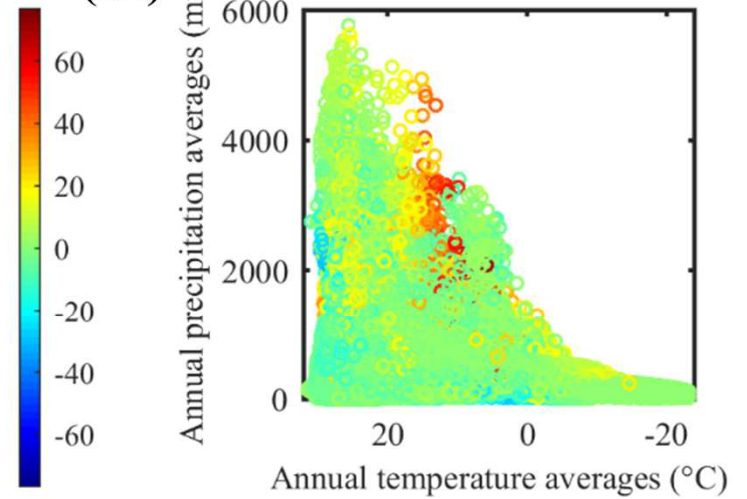
Thousands of site-years of 'fast' dynamics data for diverse ecosystems



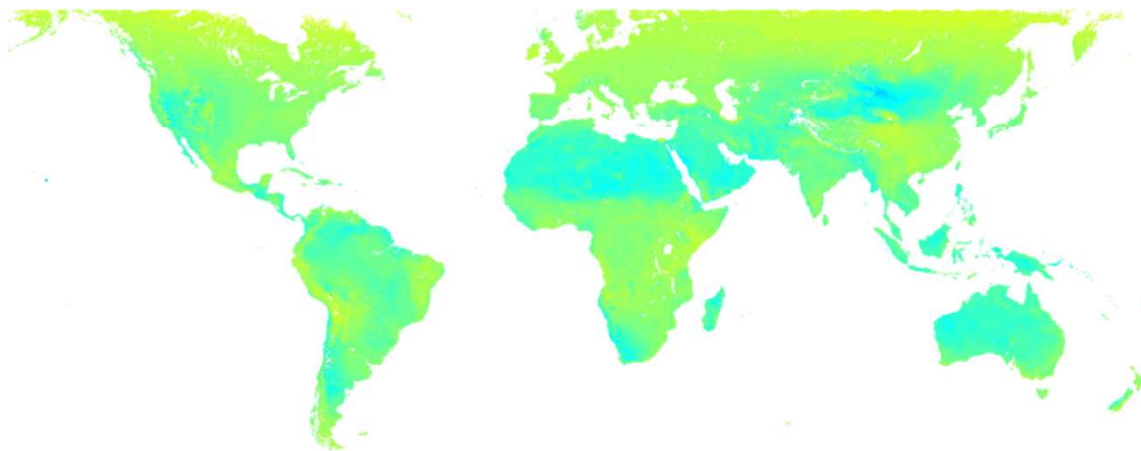
(a1) Elasticity of $T'(\theta \rightarrow NEE)$ to temperature ($e_{\bar{\theta}}$)



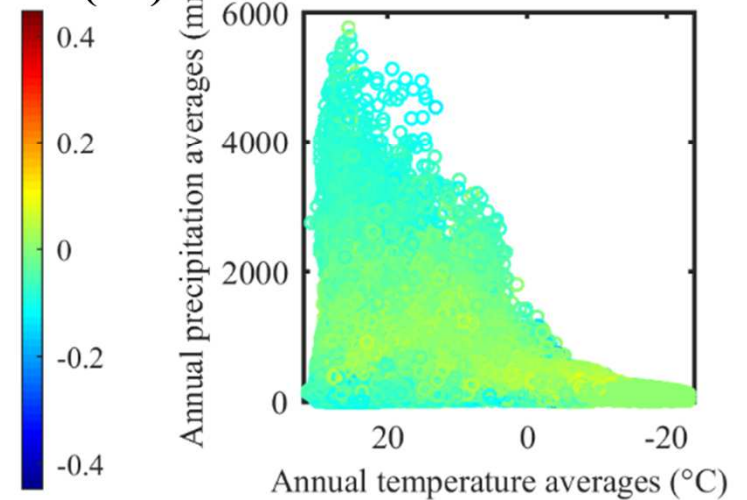
(a2)



(b1) Elasticity of $T'(\theta \rightarrow NEE)$ to precipitation ($e_{\bar{P}}$)

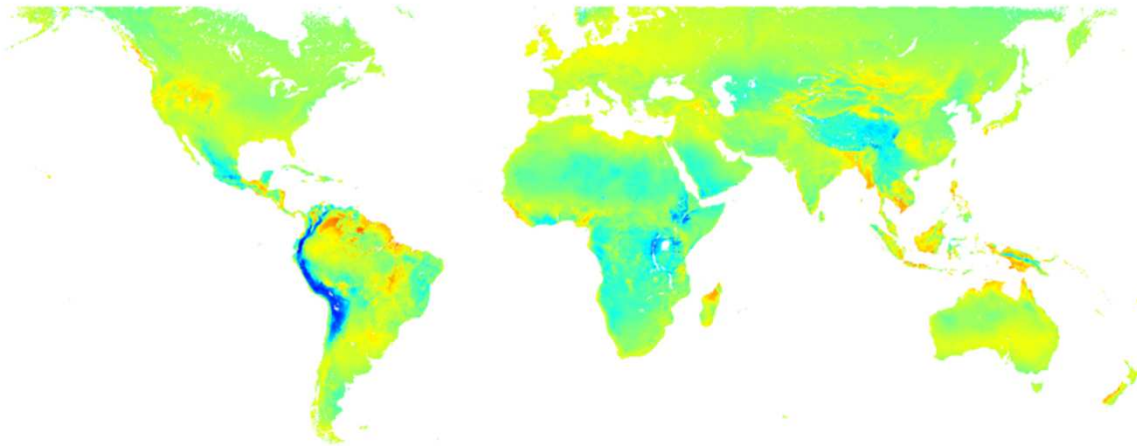


(b2)

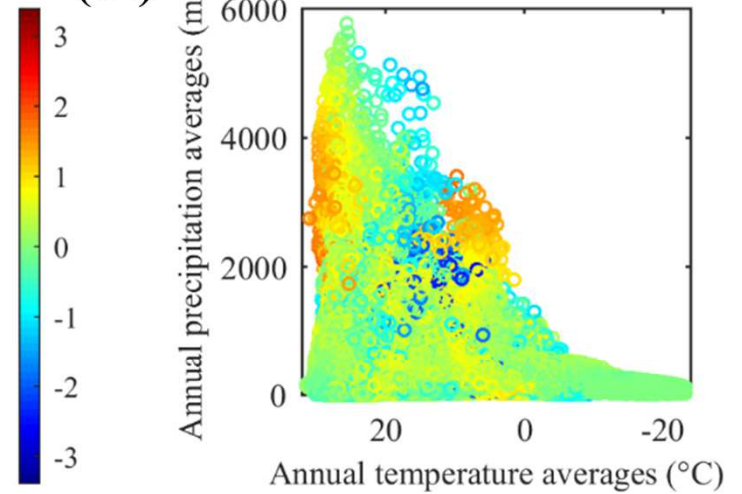


Elasticity coefficients are fitted based on OLS for all FLUXNET sites' observed monthly variations in DPN and 'slow' forcing factors, then extrapolated to the world using an ANN based on ecosystem type, geography, and climate

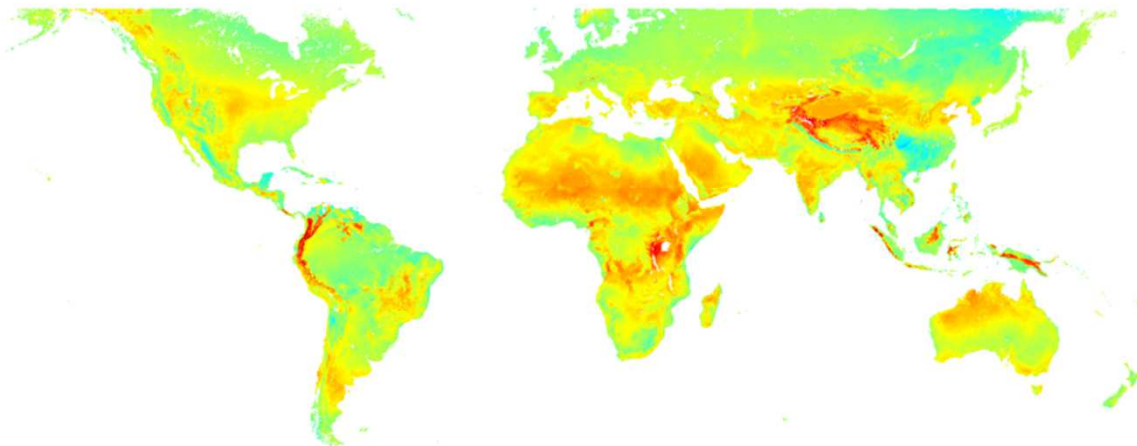
(c1) Elasticity of $T'(\theta \rightarrow NEE)$ to global radiation ($\beta_{\bar{R}}$)



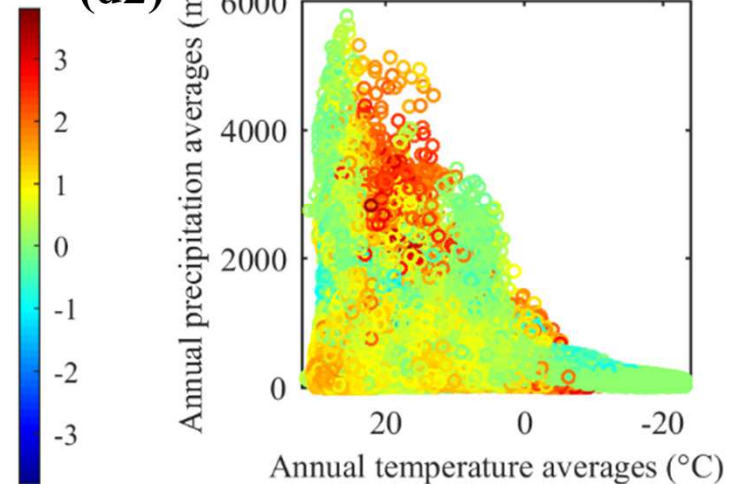
(c2)



(d1) Elasticity of $T'(\theta \rightarrow NEE)$ to phenology ($\beta_{\bar{E}}$)



(d2)



Elasticity coefficients are fitted based on OLS for all FLUXNET sites' observed monthly variations in DPN and 'slow' forcing factors, then extrapolated to the world using an ANN based on ecosystem type, geography, and climate

Model R² for Temperature > NEE coupling is better in some ecosystems than others, meaning we know more about the drivers of ecological state change for some ecosystems than others.

IGBP class	Mean	SD	Min	Max	N
OSH	0.875	0.040	0.832	0.912	3
DBF	0.793	0.117	0.471	0.987	23
SAV	0.724	0.169	0.604	0.843	2
EBF	0.716	0.138	0.363	0.935	15
MF	0.693	0.163	0.388	0.918	9
ENF	0.652	0.184	0.214	0.912	41
WET	0.631	0.241	0.282	0.879	7
CRO	0.621	0.207	0.206	0.941	19
GRA	0.599	0.225	0.114	0.910	31
CSH	0.554	0.185	0.390	0.855	6
WSA	0.513	0.262	0.160	0.778	4
Total	0.664	0.196	0.114	0.987	160

International Global Biosphere Programme (IGBP) vegetation class. CRO: croplands; CSH: closed shrublands; DBF: deciduous broadleaf forests; EBF: evergreen broadleaf forests; ENF: evergreen needleleaf forests; GRA: grasslands; MF: mixed forests; OSH: open shrublands; SAV: Savannas; WET: permanent wetlands; WSA: woody Savannas.

Conclusions

- Initial investigation succeeds in explaining changes in ecosystem functional state by studying a simplified subset of the DPN
- Elasticity to seasonal temperatures is by far the strongest, followed by phenophase and radiation, then precipitation
- Large differences in sign and magnitude of elasticity are present, meaning different ecosystems respond differently and are different distances from state transition thresholds
- There are large differences in state transition predictability based in part on ecosystem type. Poor predictability means the observed variation of DPN information content does not match the set of four 'slow' forcing factors we tested.
- This work can corroborate & critique ecological process models
- LaThuille-DPN v1.0 database contains complete monthly DPN's and climate & satellite subsets for all of FLUXNET (LaThuille)
- ProcessNetwork v1.5 is released, with wavelets etc.

Abstract

Under the context of global climate change, it is important to understand the direction and magnitude of different ecosystems respond to climate at the global level. In this study, we applied dynamical process network (DPN) approach combined with eco-climate system sensitivity model and used the global FLUXNET eddy covariance measurements (subdaily net ecosystem exchange of CO₂, air temperature, and precipitation) to access eco-climate system sensitivity to climate and biophysical factors at the flux site level. Eco-climate system sensitivity at flux timescales was estimated at the global flux sites and extrapolated to all possible land covers by employing artificial neural network approach. The extrapolation utilizes MODIS phenology and land cover products, the long-term climate GLDAS-2 product, and the GMTED2010 Global Grid elevation dataset. We found that the eco-climate system dynamical process structures are more sensitive to seasonal temperature, than to radiation, phenology, or (lowest sensitivity) precipitation. Interestingly, if global temperature continues rising, the temperature-to-NEE process coupling may increase in tropical rain forest areas while decreasing in tropical desert or Savanna areas. At the same time, phenology showed a positive effect on the temperature-to-NEE process coupling at all pixels, so increased greenness increases the importance of temperature to carbon dynamics and consequently carbon sequestration globally. This work is unique in that it provides a theoretically independent and complex system based means of assessing the sensitivity of global ecosystem processes to climate change, and it can therefore be used to critique or corroborate the findings of process based ecosystem models.