Constraining respiration flux and carbon pools in a simple ecosystem carbon model

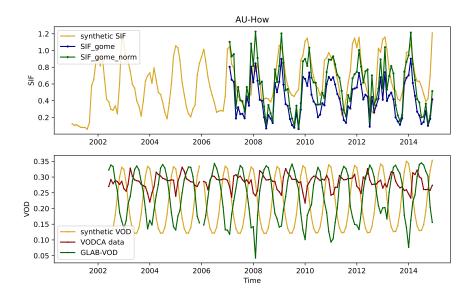
Olya Skulovich¹, Caroline Famiglietti², Alexandra Konings², and Pierre Gentine¹

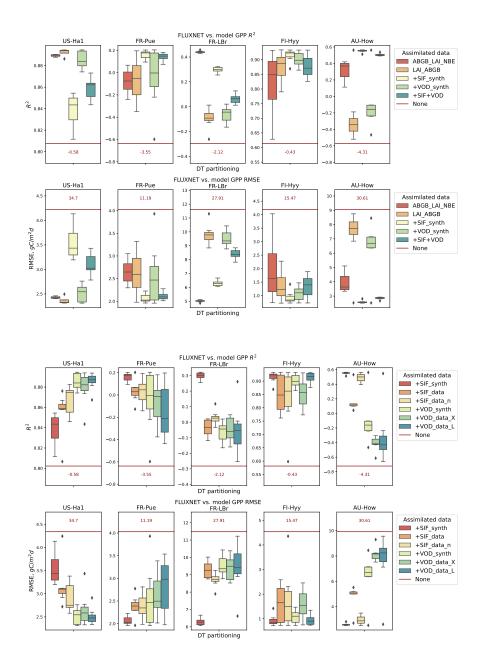
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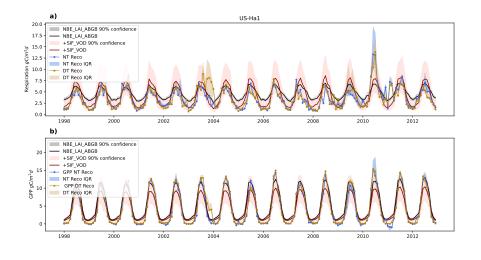
March 07, 2024

Abstract

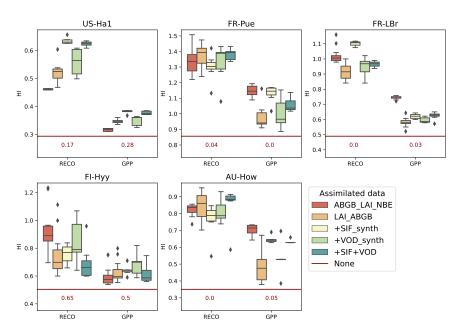
Incorporating observational data in carbon-cycle models provides a systematic framework for understanding complex ecosystem carbon dynamics, contributing essential insights for climate change mitigation and land ability to continue acting as a carbon sink. This study addresses the challenge of accurately quantifying carbon fluxes and pools, focusing on the information content of remote sensing observations. The research explores the impact of assimilating multiple observational datasets into the CARbon DAta MOdel fraMework (CARDAMOM). Satellite observations such as solar-induced fluorescence (SIF) and vegetation optical depth (VOD) are used as proxies for photosynthesis and aboveground biomass, respectively. The study aims to answer key questions about the reliability of remote sensing data in constraining the ecosystem respiration flux and sizes and dynamics of carbon pools and the relative usefulness of SIF and VOD across five FLUXNET sites. We conclude that assimilating remote SIF and VOD instead of site-based net ecosystem exchange did not deteriorate and even improved model predictions for all metrics except for interannual variability. Notably, the improved results correspond to a consistent shift in values for crucial model parameters across all five investigated sites.



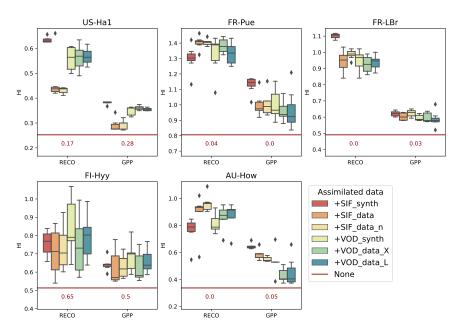




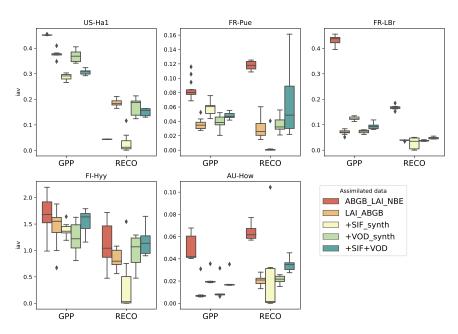
FLUXNET vs. model Histogram intersection



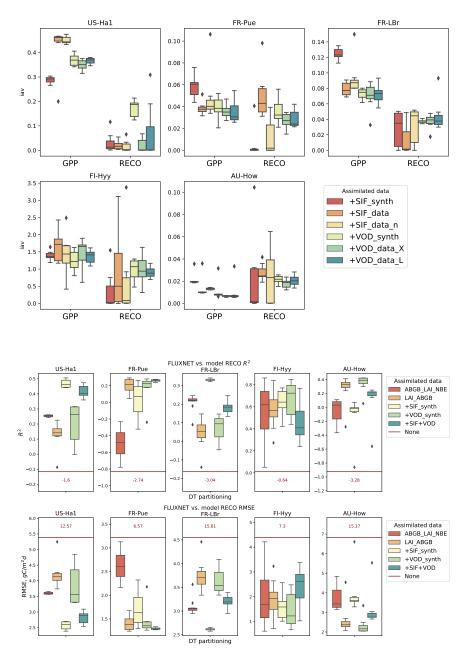
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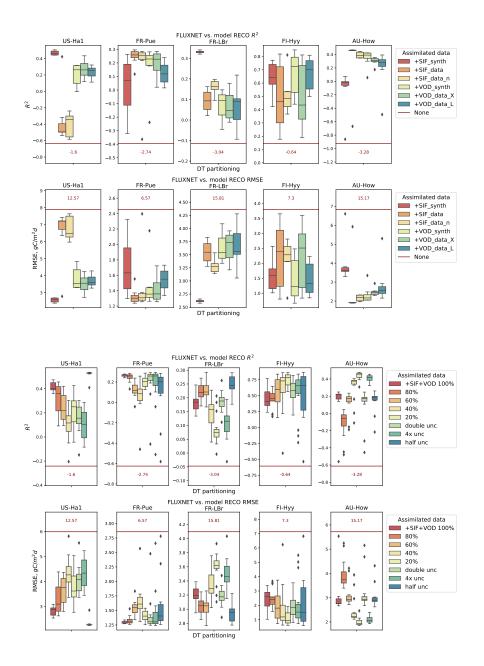


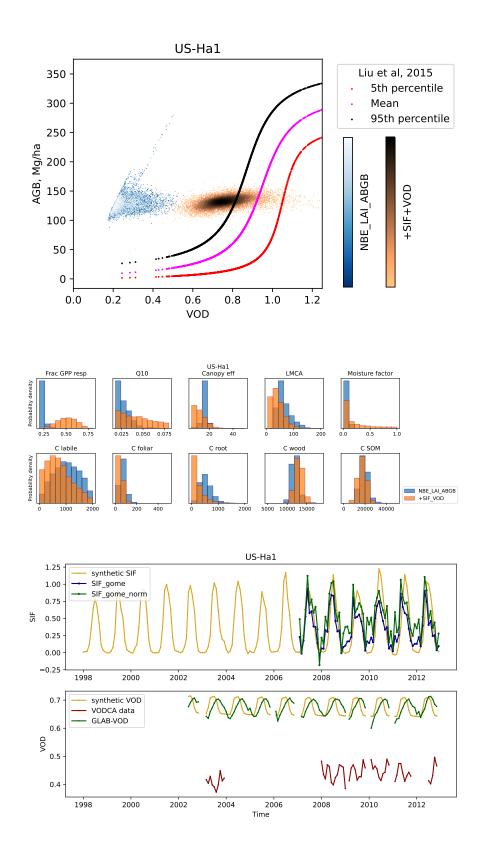
FLUXNET vs. model Interannual Anomaly Metric ratio

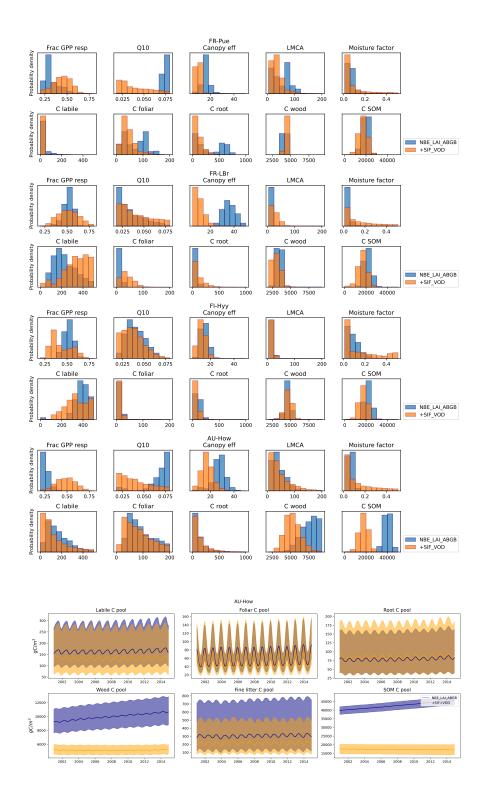


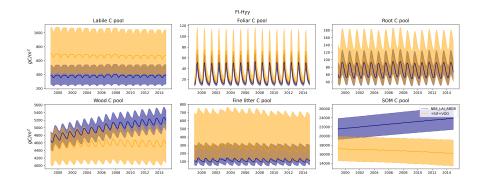
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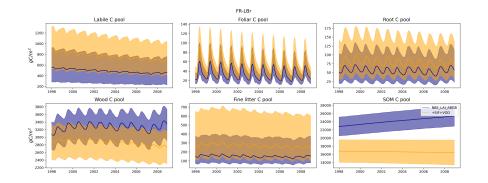


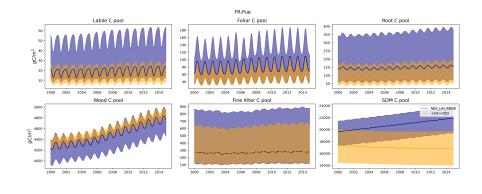


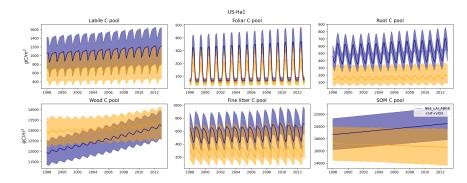












Constraining respiration flux and carbon pools in a simple ecosystem carbon model

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Key Points:

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8	•	Assimilation of globally available data like solar-induced fluorescence and vege-
9		tation optical depth improves model results.
10	•	The assimilated data helps constrain the fluxes and pools (e.g. soil) that are not
11		directly observed.
12	•	The effect of the additional data depends on the site-level conditions, data qual-

ity, and representation in the model.

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14 Abstract

Incorporating observational data in carbon-cycle models provides a systematic frame-15 work for understanding complex ecosystem carbon dynamics, contributing essential in-16 sights for climate change mitigation and land ability to continue acting as a carbon sink. 17 This study addresses the challenge of accurately quantifying carbon fluxes and pools, fo-18 cusing on the information content of remote sensing observations. The research explores 19 the impact of assimilating multiple observational datasets into the CARbon DAta MOdel 20 fraMework (CARDAMOM). Satellite observations such as solar-induced fluorescence (SIF) 21 and vegetation optical depth (VOD) are used as proxies for photosynthesis and above-22 ground biomass, respectively. The study aims to answer key questions about the reli-23 ability of remote sensing data in constraining the ecosystem respiration flux and sizes 24 and dynamics of carbon pools and the relative usefulness of SIF and VOD across five 25 FLUXNET sites. We conclude that assimilating remote SIF and VOD instead of site-26 based net ecosystem exchange did not deteriorate and even improved model predictions 27 for all metrics except for interannual variability. Notably, the improved results correspond 28 to a consistent shift in values for crucial model parameters across all five investigated 29 sites. 30

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Plain Language Summary

Carbon-cycle models allow us to study how terrestrial ecosystems absorb carbon from the atmosphere and release it back and what its afterlife is in plants and soil. Scientists use observed data to accurately quantify these processes and incorporate them into models to constrain model parameters. We study how two satellite-based measurements, one used to substitute for photosynthesis, which is not measurable directly, and one used to substitute for aboveground biomass, which is scarcely available, help to improve the model's performance.

³⁹ 1 Introduction

Accurately quantifying terrestrial carbon sinks/sources and carbon pools is critical for reliable projections of carbon emissions and climate change mitigation, yet large uncertainties still exist among the components of the global carbon budget (Arneth et al., 2017; Piao et al., 2018; Gasser et al., 2020; Friedlingstein et al., 2022). Carbon is taken up by the terrestrial biosphere through photosynthesis and released via autotrophic and

-2-

heterotrophic respiration as well as disturbances (wildfires, windthrows). Yet, how those 45 fluxes and their partitioning will change under elevated CO_2 remains a question of de-46 bate (e.g., Xu et al., 2015; Kirschbaum & McMillan, 2018; L. Liu et al., 2020). To tackle 47 this challenge, it is necessary to understand the potentially competing processes that af-48 fect carbon uptake under global change, such as plants' physiological response to elevated 49 CO₂, higher temperatures, increased aridity, increased extreme events frequency, and other 50 climatological shifts associated with climate change on plant-level and ecosystem-level 51 scales (Cox et al., 2000; Tharammal et al., 2019; Song et al., 2019; Denissen et al., 2022). 52 Nevertheless, large amounts of global and site-level observational data have become avail-53 able and can now be used to constrain many of those processes. Over the last couple of 54 decades, our understanding of photosynthesis and our capacity to constrain it at the global 55 scale has dramatically increased through the use, first, of vegetation indices, (e.g., Wu 56 et al., 2009) and then more recently of Solar Induced Fluorescence or SIF, (e.g., X. Yang 57 et al., 2015; Zhang et al., 2014), a proxy for gross primary productivity (GPP). How-58 ever, quantifying global respiration fluxes is currently impossible through direct measure-59 ments, and they can only be indirectly inferred, such as using statistical upscaling from 60 local measurements or using process-based models, (e.g., Jian et al., 2018; Jung et al., 61 2019; Nathaniel et al., 2023). In essence, from an observational standpoint, there are no 62 global constraints on the respiration part of the land carbon budget, leading to major 63 uncertainties in our capacity to understand and predict the terrestrial carbon cycle. 64

Process-based models allow us to combine knowledge of physical, chemical, and bi-65 ological processes with the collected data to achieve interpretable carbon cycle analy-66 sis. Process-based models depict ecological processes with models of varying complex-67 ity and different level of abstraction. Yet, while those models are good at capturing some 68 aspects of the system, they rely on several structural assumptions, and the model pa-69 rameters should be carefully calibrated to improve the model accuracy (Y.-P. Wang et 70 al., 2009). For example, Li et al. (2021) lists model structure and model assumptions un-71 certainties among the main processes contributing to the overall model uncertainties. These 72 uncertainties are due to our incomplete understanding of some ecological mechanisms 73 (for instance, belowground processes and microbial interactions, (e.g., Hartmann et al., 74 2020)), an abundance of empirical equations with parameters that are not necessarily 75 applicable globally, and model-specific simplifications. Model parameters can be calibrated 76 via data assimilation (also called "model-data fusion") to best match observational data, 77

-3-

including data uncertainty to quantify model parametric and prediction uncertainties, 78 (e.g., Li et al., 2020). Models can even constrain processes that are not directly observed 79 (Talagrand, 1997) because of the internal physical and biological constraints linking ob-80 served to unobserved variables (e.g., mass balance for carbon pools). In other words, since 81 these models quantify internal system interconnections and dynamics, constraints intro-82 duced via observational data are propagated through the model and can constrain the 83 rest of unobservable fluxes or pools. As such, unobserved fluxes and pools, such as res-84 piration and soil carbon, can potentially be constrained through the assimilation of ob-85 servable variables that are indirectly related to those processes. 86

A caveat of the data assimilation approach lies in the tradeoff between model com-87 plexity and the demand for data to be assimilated (e.g., Famiglietti et al., 2021). Indeed, 88 the more complex the models are, the more parameters are needed for their description. 89 Then, the more parameters the model comprises, the more likely it is that the model out-90 put will match equally well the observations with *different* combinations of the param-91 eters. This phenomenon is called equifinality (Beven, 1993; Beven & Freer, 2001). Sim-92 ply put, it means that the model can give the right answer (i.e., is optimized and cor-93 responds well with the observational data) but for the wrong reason (i.e., the resulting 94 combination of the model parameter is inconsistent with the *true* system dynamic). Gen-95 erally speaking, equifinality is reduced when more data is assimilated to constrain mul-96 tiple different components of the model. In practice, assimilating multiple datasets can 97 be limited by data availability and the technical complexity of the data assimilation pro-98 cess. 99

In this work, we study the effects of assimilating multiple observational datasets 100 in an ecosystem carbon model to quantify their impact on constraining unobserved res-101 piration flux and carbon pools. The physical models' equations constrain the relation-102 ship between non-observed variables and observed variables, which are assimilated. The 103 model used for this data assimilation framework is the CARbon DAta MOdel fraMework 104 model or CARDAMOM for short (A. Bloom & Williams, 2015; A. A. Bloom et al., 2016). 105 This model is actively used by the scientific community to model the terrestrial carbon 106 cycle for process understanding and has been proven to successfully capture spatial pat-107 terns and temporal trends, as well as inter-annual variability of the various variables of 108 interest (A. A. Bloom et al., 2018; López-Blanco et al., 2019; Quetin et al., 2019; Y. Yang 109 et al., 2019; Yin et al., 2020; A. Norton et al., 2021), including in a benchmarking effort 110

-4-

across more than 200 eddy-covariance sites (Y. Yang et al., 2022). The model has been 111 previously applied to study global and local mechanisms for diverse ecosystems, from Arc-112 tic (López-Blanco et al., 2019) to tropics (Yin et al., 2020). CARDAMOM was created 113 as a relatively simple model so that the low number of the model parameters could be 114 better constrained by observational data and thus reduce the risk of equifinality. Model 115 complexity and equifinality in the context of CARDAMOM's predictive skill were pre-116 viously examined by Famiglietti et al. (2021) and concluded that the model skill depends 117 on properly constraining the model parameters. Building on that, we look into the model 118 skill when the parameters and variables of interest are constrained by indirect observa-119 tions. In particular, we use SIF as a proxy for GPP, and vegetation optical depth (VOD) 120 as a proxy for above-ground biomass and use physical constraints to build additional con-121 straints on belowground carbon pools and ecosystem respiration. VOD is a variable de-122 rived from remote sensing observations as an attenuation of the surface reflectance by 123 the wet biomass and can serve as a measure of total biomass and water-related vegeta-124 tion stress (Konings et al., 2017). 125

When investigating the effects of assimilating additional datasets in CARDAMOM, 126 we employ a multi-objective approach and examine the impact of different model for-127 mulations, varying data quality (including information content and data uncertainty), 128 and implicit and explicit constraints introduced in the model. We investigate the follow-129 ing research questions: 1) Given the error-prone nature of NBE at the global scale (e.g., 130 Deng & Chen, 2011; Chevallier et al., 2019; Cui et al., 2022), how well can remote sens-131 ing data such as SIF and VOD help constrain the carbon cycle instead? 2) Which of the 132 SIF and VOD data is more useful in this task, and how does it depend on local condi-133 tions? 3) Can respiration flux and carbon pools be reliably inferred from data assimi-134 lation, given indirect observational constraints? 4) What is the role of data availability 135 and assumed uncertainty on the assimilation results? The analysis is conducted for five 136 FLUXNET sites across different biomes. 137

138 2 Methods

The CARDAMOM framework consists of two major parts: the carbon cycle model and the data assimilation infrastructure. Carbon cycle models have different "flavors" and are called DALEC (Data Assimilation Linked Ecosystem Carbon model) versions that vary depending on the physical processes and parameters included (e.g., with/without

-5-

fires, with/without water cycle, etc. A. Bloom and Williams (2015); Quetin et al. (2020)). 143 The general model structure is relatively simple and, in most of the DALECs, includes 144 six carbon pools: foliar, labile, wood, fine roots, litter, and soil organic matter (SOM). 145 An additional advantage of the CARDAMOM framework is the inclusion of a series of 146 ecological and dynamic constraints (EDC) on model parameters and initial conditions 147 (A. Bloom & Williams, 2015). These "common sense" constraints drive ecosystem vari-148 ables towards more consistent and realistic solutions, thus further helping to reduce equi-149 finality. The optimized model parameters and initial conditions are time-invariant and 150 site-specific. They are inferred using a Metropolis–Hastings Markov chain Monte Carlo 151 (MCMC) approach (Haario et al., 2001). The model is run at monthly resolution. 152

To test whether we can better constrain respiration fluxes and carbon pools by as-153 similating SIF and VOD, we first take a medium-complexity version of the DALEC model 154 (for a detailed description of the model, refer to A. Bloom and Williams (2015) and Famiglietti 155 et al. (2021)). In this configuration, 33 model parameters and initial conditions are be-156 ing optimized via data assimilation. Site-specific meteorological data (air temperature, 157 shortwave radiation, atmospheric CO_2 concentration, vapor pressure deficit, precipita-158 tion, and wind speed) drive the model dynamics, while observational data further con-159 strain model parameters. In particular, eddy covariance net ecosystem exchange (NEE) 160 measurements from FLUXNET (Pastorello et al., 2020), leaf area index (LAI) estimates 161 from the Copernicus Global Land Service (Fuster et al., 2020), and in situ biomass sur-162 veys are used for assimilation into the model. The same or similar combination of data 163 for data assimilation has previously been used in CARDAMOM (López-Blanco et al., 164 2019; Quetin et al., 2020; Famiglietti et al., 2021). In the previously reported configu-165 rations, NEE had the highest impact on the model performance (Famiglietti et al., 2021). 166 In CARDAMOM studies that have a regional or global focus, beyond FLUXNET sites, 167 atmospheric inversion of net biosphere exchange (NBE) is used in the assimilation with 168 the caveat that it can have large uncertainties (e.g., H. Wang et al., 2019; Cui et al., 2022). 169 Hence, we investigate how excluding it and including SIF and VOD datasets for data 170 assimilation, as well as other modeling choices and assumptions, affect the model per-171 formance. 172

The analysis is run over several eddy-covariance sites, including Harvard Forest EMS Tower, USA (US-Ha1), Puechabon, France (Fr-Pue), Le Bray, France (Fr-LBr), Hyytiala,

¹⁷⁵ Finland (Fi-Hyy), and Howard Springs, Australia (AU-How), see Table 1.

-6-

Table 1. Summary of eddy-covariance sites, showing their location, FLUXNET code, observational time period, mean climate information, and ecosystem type. Ecosystem type is denoted using the International Geosphere-Biosphere Programme (IGBP) classification. DBF: deciduous broadleaf forest; EBF: evergreen broadleaf forest; ENF: evergreen needleleaf forest; WSA: woody savanna. Simple aridity index is calculated as De Martonne aridity index (De Martonne, 1923) $AI = \frac{P}{T_a+10}.$

Site code	Lat	Lon	Elevation, m	IGBP	Data record	Mean annual temp, C	Mean annual precip- itation, mm/year	Simple aridity index	Reference
AU-How	-12.49	131.15	42	WSA	2001- 2014	27	1449	Moderate humid	(Beringer et al., 2007)
FI-Hyy	61.85	24.29	181	ENF	1999- 2014	3.8	709	Very humid	(Suni et al., 2003)
FR-LBr	44.72	-0.77	61	ENF	1998- 2008	13.6	900	Moderate humid	(Berbigier et al., 2001)
FR-Pue	43.74	3.59	270	EBF	2000- 2014	13.5	883	Moderate humid	(Rambal et al., 2004)
US-Ha1	42.54	-72.17	340	DBF	1998- 2012	6.2	1071	Excessive humid	(Munger & Wofsy, 2014)

2.1 Including new observational datasets

To include a new observational dataset in CARDAMOM, the following steps need to be performed: 1) define a functional form that represents the data as a function of existing and new model variables and parameters; 2) add data likelihood to the full model likelihood; 3) define a prior for any new model parameters. We discuss the assimilation of SIF and VOD in the following sections.

182 2.1.1 SIF model

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¹⁸³ SIF is assumed to be a linear function of GPP (Wood et al., 2017); hence, in the ¹⁸⁴ model, it is included as follows.

$$SIF = p_{sif}F_{GPP} \tag{1}$$

where F_{GPP} is the GPP flux and p_{sif} is a proportionality coefficient $[m^2 s^{-2} \mu m^{-1} s r^{-1}]$. Synthetic and observational SIF data are considered, with the synthetic SIF defined directly from FLUXNET GPP data for the corresponding FLUXNET site. This

allows assessing the effect of data quality (signal vs. noise) on the model performance. 189 Indeed, by construction, the synthetic SIF data contains the signal from the site-measured 190 GPP with no additional noise related to observational uncertainties, sensor uncertain-191 ties, SIF retrieval model uncertainties, as well as the scale mismatch between the site level 192 data and the grid level remote sensing data. The observational SIF, on the other hand, 193 contains the GPP-related signal along with the noise, with the unknown ratio of the two. 194 To which degree that affects the model performance defines the model sensitivity to the 195 data uncertainty. 196

¹⁹⁷ An average between daytime- GPP_{DT}^{FLUX} and nighttime- GPP_{NT}^{FLUX} derived GPP ¹⁹⁸ is considered the site GPP.

$$SIF_{synth} = p_{sif} \left(\frac{GPP_{DT}^{FLUX} + GPP_{NT}^{FLUX}}{2} \right)$$
(2)

The slope coefficient p_{sif} is individually fit to the FLUX site GPP data such that the final synthetic SIF data is in the same range as the observational SIF data. GOME-2 SIF data (Joiner et al., 2023) is used as SIF observational data. From GOME-2, both SIF and normalized SIF (normalized by photosynthetically active radiation – PAR) data are used. A similar amplitude of the synthetic and observational data is necessary to assess the effect of the data quality (noisiness) on the model performance and isolate it from other effects. An example data time series is shown in Fig. S5.

The SIF likelihood function is constructed similarly to the likelihood function for other assimilated data in CARDAMOM with the observation (O) probability given a set of model parameters \boldsymbol{x} is

$$P(\boldsymbol{O}|\boldsymbol{x}) = e^{-0.5\sum_{i=1}^{n} \frac{\left(M_{i}^{SIF} - O_{i}^{SIF}\right)^{2}}{\sigma_{i}^{2}}}$$
(3)

where O_i^{SIF} is the *i*th SIF observation, M_i^{SIF} is the corresponding model SIF, and σ_i is the *i*th error variance for each observation with no error covariance between observation errors assumed (A. Bloom & Williams, 2015).

2.1.2 VOD model

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Vegetation optical depth (VOD), based on microwave remote sensing, provides a 214 constraint on aboveground biomass. Unlike SIF, VOD representation in the model is not 215 as straightforward. VOD is measured as the attenuation of electromagnetic waves, which 216 is proportional to the mass of water in the vegetation (Jackson & Schmugge, 1991; Wigneron 217 et al., 2017; Konings et al., 2017). Hence, VOD reflects vegetation water content that 218 can be representative of both variability in total biomass and plant relative water con-219 tent that depends on meteorological conditions. There is no clear separation of these two 220 components even at larger-than-daily time scales (Konings, Holtzman, et al., 2021). Con-221 sidering the overall simplification of the given DALEC and aiming at keeping the num-222 ber of new parameters minimal to reduce equifinality, we assume VOD at the monthly 223 resolution to be a function of above-ground biomass only. That is, we assume that vari-224 ations in relative water content are small, which is a reasonable assumption. Relative 225 water content representation is missing in the carbon model, which is not uncommon for 226 a model of such simplicity but more complex models are starting to include plant hy-227 draulics (e.g., Kennedy et al., 2019) that could resolve water content. For simplicity, we 228 additionally assume that VOD is a linear combination of leaf and wood biomass, with 229 different learnable regression parameters of each biomass type, at each site: 230

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$$VOD = p_{fol}C_{fol} + p_{woo}C_{woo} \tag{4}$$

where C_{fol} is the foliar carbon pool, C_{woo} is the wood carbon pool $[gCm^{-2}]$ and p_{fol} and p_{woo} are the corresponding coefficients $[gC^{-1}m^2]$, since VOD is dimensionless.

In addition to simplifying the relationship between VOD and biomass, this repre-234 sentation poses a numerical issue. Indeed, for forests, in units of carbon mass, C_{woo} can 235 be several orders of magnitude larger than C_{fol} . For example, wood biomass averages 236 13,000 gCm^{-2} while foliar biomass reaches the maximum of 300 gCm^{-2} based on a CAR-237 DAMOM prediction for Harvard Forest. At the same time, the VOD dynamic is primar-238 ily due to C_{fol} variability because C_{fol} is much more variable than C_{woo} and because 239 VOD is more sensitive to upper canopy layers than to lower canopy layers (Konings, Saatchi, 240 et al., 2021). In CARDAMOM, both the coefficients p_{woo} and p_{fol} and carbon pool sizes 241 are inferred, so the contradicting forces of pools' dynamic and pool' sizes may result in 242 equifinality and a wide range of predicted pools sizes. This, in turn, would lead to no 243

or minimal additional information introduced by VOD in comparison to LAI (since LAI is defined in CARDAMOM as $LAI = p_{lai}C_{fol}$). There are several ways to tackle this issue, from changing VOD representation in the model to introducing new ecological and dynamic constraints (EDC). Here, we define narrow, not overlapping priors for p_{woo} and p_{fol} ranges that ensure the relationship between the sizes of the pools is preserved.

Similar to the SIF module, synthetic and observational VOD data are considered 249 to assess the effect of data quality (signal vs. noise) on the model performance. Since 250 the true foliar and wood carbon pool dynamic is unknown, we used the following pro-251 cedure to construct the synthetic VOD data. First, we run CARDAMOM assimilating 252 NBE, LAI, and ABGB and take model output median foliar and wood pools as true pools 253 for that site. With these pools, we fit p_{fol} and p_{woo} coefficients such that the constructed 254 VOD matches the observational data. Two observational datasets are considered – VODCA 255 (Moesinger et al., 2020) C-band VOD and GLAB-VOD (Skulovich et al., 2024) L-band 256 VOD (extracted for the grid cell closest to each site). VOD likelihood function is con-257 structed in the same way as Eq.3. 258

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2.2 Experimental setup

We compare the base case with no data assimilated ('none' case) to the previously reported combination that includes NBE, LAI, and ABGB ('NBE_LAI_ABGB') and then remove NBE and instead add either SIF, VOD, or both SIF and VOD synthetic and observational data (see Table 2). We examine to what extent remote sensing SIF and VOD can replace NBE and lead to similar or better results.

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2.3 Information content of observations

When assimilating observational data, the question of the information content of the available data is often reduced to the question of data uncertainty (e.g., Raupach et al., 2005) or data autocorrelation, as an indicator for data redundancy (e.g. Moore et al., 2011; Williams et al., 2009). Due to the overall data scarcity, it is often assumed that assimilating more data will improve model performance. In this series of experiments, aligned with the research question "What is the role of data availability and assumed uncertainty on the assimilation results?" we test this hypothesis by assimilating only a

Experiment	Case name	Assimilated data		
1	'none'	-		
2	'ABGB_LAI_NBE'	ABGB, LAI, NBE		
3	'LAI_ABGB'	ABGB, LAI		
4	'+SIF+synth'	ABGB, LAI, synthetic SIF		
5	'+SIF+data'	ABGB, LAI, GOME2 SIF		
6	+SIF+data_n'	ABGB, LAI, GOME2 SIF normalized		
7	'+VOD_synth'	ABGB, LAI, synthetic VOD		
8	'+VOD_data_C'	ABGB, LAI, VODCA VOD		
9	+VOD_data_L'	ABGB, LAI, GLAB-VOD VOD		
10	+SIF+VOD'	ABGB, LAI, synthetic SIF, GLAB-VOD		

Table 2. Data assimilation cases

²⁷³ portion of the available data and compare the results with assimilating data with reduced²⁷⁴ or increased uncertainty.

In particular, we consider the following cases:

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- Comparing assimilating all available LAI, ABGB, SIF, and VOD observations with 276 assimilating just a part of all available observations, namely 20, 40, 60, and 80% 277 of the data points. In each run, the corresponding number of indices are selected 278 at random, each run is repeated 25 times so that each time it is a different 20%279 of the data that is used in assimilation (for the 20% case, the principle is the same 280 for all percentage values). 281 • Assimilating 100% of the available LAI, ABGB, SIF, and VOD, and increasing 282 or reducing the data uncertainty for SIF and VOD. We consider doubling, qua-283
- drupling, and reducing the uncertainty in half.

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Note that 100% of the available data do not necessarily cover 100% of the time steps

defined by the forcing meteorological data. For the sites and time period of this study,

NBE and LAI data have more coverage than SIF and VOD data (see Table 3).

 Table 3.
 Number of data points and percentage of the available forcing data for each of the assimilated variables per site

	US-Ha1	FR-Pue	FR-LBr	AU-How	FI-Hyy
Forcing time series	180 (100%)	180 (100%)	132 (100%)	168(100%)	192(100%)
LAI	151 (84%)	180 (100%)	120 (91%)	150 (89%)	150 (78%)
NBE	156 (87%)	170 (94%)	108 (82%)	134 (80%)	182 (95%)
ABGB	15 (8%)	13 (7%)	8 (6%)	3 (2%)	6 (3%)
VOD (data)	113 (63%)	44 (24%)	41 (31%)	150 (89%)	105 (55%)
SIF (data)	68 (38%)	93 (52%)	22 (17%)	95 (57%)	58 (30%)

When considering the information content coming from the observations, we can hypothesize the following relationships:

• More data is better than less data. However, to what degree additional data improves the model results depends on the local conditions. For instance, some sites can exhibit minimal stress and can be almost fully described by the drivers only, so additional observational data might not improve the results as much as for sites with more complex vegetation feedback.

Sometimes, the true uncertainty of the observations is unknown. In that case, there
 should exist an optimal uncertainty that can be used as an empirical parameter
 in the model. Indeed, if the uncertainty is too high, the observations do not add
 much information, and if the uncertainty is too low, the model tries to match noise
 along with the actual signal from observations.

We examine how these hypotheses hold for assimilating satellite observations in CAR-

301 DAMOM.

2.4 Performance metrics

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For every assessed combination of CARDAMOM inputs, model formulations, and parameters, the model runs at least ten times. Within these runs, the convergence is assessed using Gelman-Rubin diagnostics. Next, the model output is compared to FLUXNET eddy-covariance site data, and different model outputs are compared to each other.

• Flux comparison: R^2 , RMSE, HI. The flux comparison is conducted for GPP 307 and respiration based on eddy-covariance partitioning. Median model output is 308 compared with FLUXNET GPP and RECO derived using friction velocity thresh-309 old (VUT, Barr et al. (2013)) method and daytime and nighttime partitioning sep-310 arately. To estimate uncertainties, the FLUXNET data spread is based on the 25th 311 and 75th percentile of the corresponding parameter. The model and site data are 312 compared using the coefficient of determination R^2 and root mean square error 313 (RMSE). 314

These metrics allow us to assess the model accuracy; however, in the Bayesian framework, distribution comparison is more meaningful. To achieve this, a histogram intersection (HI, Famiglietti et al. (2021)) is used as an additional skill metric. HI measures the similarity of two (discretized) distributions, with larger HI corresponding to more similar distributions. HI is calculated for GPP and RECO.

• Interannual variability. Through the data assimilation framework, CARDAMOM can capture interannual variability in the fluxes. However, the tightness of the fit to the interannual peaks might not necessarily be clear from metrics like R^2 , RMSE, or HI. To assess how different combinations of model parameters affect the interannual variability, we calculate the following metric (IAM - interannual anomaly metric):

$$IAM = \sum_{i=1}^{n} (V_i - V_i^{seas})^2 \tag{5}$$

where V_i is the *i*th observation or model output of a parameter V (for example, GPP), V_i^{seas} is the *i*th value of the seasonal cycle calculated for this parameter, and *i* can vary from 1 to *n*, where *n* is the total number of observations or model outputs. The seasonal cycle in this approach is identical from year to year. It is calculated by matching a periodic signal, a sine wave with a period of 365.25 days, to the observations, with other parameters of the sine wave fitted using *curve_fit()*

function in Python. Comparison to a sine-wave for a signal with a strong seasonal 333 component is defined to emphasize year-to-year anomalies. For a signal with a weaker 334 seasonality, the fitted sine wave will have a small amplitude and be close to the 335 long-term average. We calculate this metric for the observational data and then 336 compare the value to the metric calculated for different model configurations. With 337 similar R^2 and RMSE, IAM helps to assess another aspect of the model perfor-338 mance. For consistent comparison, we take the ratio IAM^{model_output}/IAM^{data}. 339 If this value is close to 1, the model and the data have very similar interannual 340 variability, and if the value is close to 0, the model significantly underestimates 341 interannual variability present in the data. 342

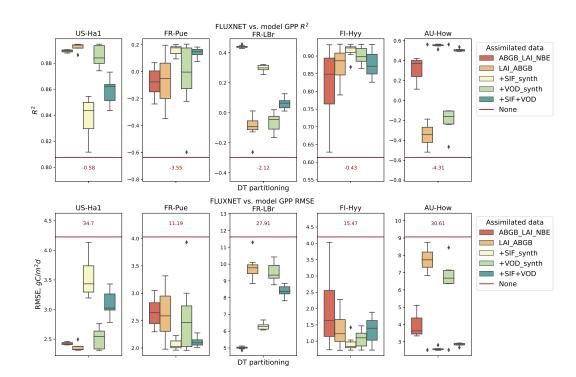
• Pool constraints: relative change and biomass-VOD relationship. Biomass 343 data is mainly constrained through indirect proxies (LAI, VOD, NDVI, etc.) apart 344 from temporarily and spatially sparse surveys. For example, for the sites discussed 345 in this study, ABGB measurements are available only for 2-8% of the total length 346 of the meteorological observations (See Table 3). In addition, ABGB comprises 347 the sum of labile, foliar, root, and wood carbon pools in CARDAMOM without 348 any partitioning information. Hence, we assume the comparison of the model pool 349 dynamic to the ABGB observational data is insufficient. Instead, we compare the 350 relative model performance for different combinations of the assimilated data, an-351 swering the question, "Can carbon pools be reliably inferred from data assimila-352 tion, given indirect observational constraints?" comparing the output carbon pools' 353 mean and distribution. 354

Additionally, the modeled biomass is compared to empirical above-ground biomass 355 for a given VOD using the relationship from Y. Y. Liu et al. (2015) (Supplemen-356 tary Eq. 2 and Supplementary Figure 4 of the original article). The authors used 357 reliable VOD observations and benchmark biomass maps to obtain the VOD-biomass 358 relationship. While this relationship does not necessarily represent the ground truth 359 biomass for a given VOD and depends on the choice of a reference for VOD in the 360 original paper (since VOD derived from different frequency bands can be mutu-361 ally biased), the closeness of the model output to the paper-derived limits shows 362 us the ability of CARDAMOM model in a particular configuration to constrain 363 carbon pools. 364

365 **3 Results**

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Overall, more than 300 different scenarios were considered, and more than 5 000 366 CARDAMOM runs were performed (with 10 to 25 repetitions per scenario). Gelman-367 Rubin metric indicates convergence of most of the runs and most of the scenarios, with 368 the exception of stochastic information content-related runs. In some cases, for exam-369 ple, for the FR-LBr site, when SIF (both synthetic and observational data) was assim-370 ilated, it resulted in divergence in some model parameters (e.g., Decomposition rate, Frac-371 tion of GPP respired, Leaf Lifespan, Canopy efficiency), however, when both SIF and 372 VOD data were assimilated, all runs converged. 373



3.1 Can SIF and VOD observations substitute NBE?

Figure 1. Effect of assimilating different data combinations on determination coefficient R^2 and RMSE between median CARDAMOM results and FLUXNET GPP data. The 'none' case (no data assimilated) is denoted as a red line not in scale with the corresponding value of the parameter denoted.

375 376 We start by comparing the model performance in matching FLUXNET daytime GPP for all five sites. Figure 1 illustrates the base case 'ABGB_LAI_NBE' in compar-

ison to the case without NBE data directly assimilated ('LALABGB') and with SIF, VOD, 377 and both SIF and VOD consequently added to the set of the assimilated observations. 378 Comparing the first two cases, we notice that NBE data is only crucial for FR-LBr and 379 AU-How sites. For the rest of the sites, without NBE, CARDAMOM can converge to 380 the same or even better solution. When SIF and VOD are added, it further improves 381 the results for FR-Pue, FR-LBr, and AU-How sites. For US-Ha1, the '+SIF+VOD' case 382 results are worse than the 'ABGB_LAI_NBE' case, however, the fit is still tight - R^2 = 383 0.86 in comparison to the initial $R^2 = 0.89$. For FI-Hyy site, there was no apparent ef-384 fect related to the SIF and VOD inclusion, which is likely due to this Finnish evergreen 385 needle-leaved forest site with cool summer and no dry season experiencing little to no 386 water stress. In all cases, significant improvements in the model performance are observed 387 with any data assimilated in comparison to no data assimilated (the 'none' case). In all 388 cases except for the US-Ha1 site, adding SIF and VOD, especially together, can success-389 fully substitute NBE data. 390

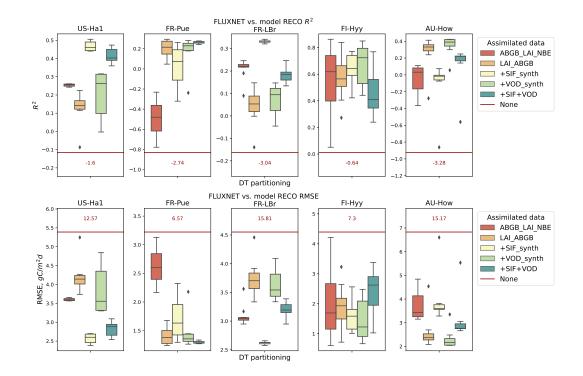


Figure 2. Effect of assimilating different combinations of data on determination coefficient R^2 and RMSE between median CARDAMOM results and FLUXNET RECO data. The 'none' case (no data assimilated) is denoted as a red line not in scale with the corresponding value of the parameter denoted.

Next, we compare the model performance in the same setup to match the FLUXNET 391 estimated ecosystem respiration. Figure 2 illustrates the change in R^2 and RMSE for 392 ecosystem respiration when assimilating different combinations of observational datasets 393 for all five sites. Here, NBE was improving the model results only for US-Ha1 and FR-394 LBr sites. From Table 3, we notice that there are less available data points for SIF and 395 VOD observational data in comparison to NBE and LAI. However, even with that, as-396 similating SIF and VOD results in higher R^2 and lower RMSE than for 'ABGB_LAI_NBE' 397 and/or 'LALABGB' cases for all sites except for FI-Hyy. For example, for respiration 308 at US-Ha1, R^2 increases from 0.25 for 'ABGB_LAI_NBE' and 0.16 for 'LAI_ABGB' to 399 0.42 with inclusion of SIF and VOD. Correspondingly, respiration RMSE reduces from 400 3.6 and 4.2 to 2.8 gC/m^2d . For the FR-Pue site, assimilating LAI and NBE results in 401 the worst performance among all cases, whereas the '+SIF+VOD' shows the best and 402 most consistent results with the narrowest interquartile interval for both R^2 and RMSE. 403 indicating model convergence across optimized parameters. This evergreen oak forest is 404 located in a Mediterranean climate with long dry summers and has large interannual vari-405 ation (Rambal et al., 2004), which can explain why additional observational data can 406 improve the model results. For FR-LBr, '+SIF+VOD' cannot achieve the results obtained 407 for the 'ABGB_LAI_NBE' case, however, the SIF and VOD addition significantly improves 408 the results in comparison to the 'LALABGB' case. In other words, SIF and VOD can-409 not fully substitute NBE for this site but still bring improvement in contrast to the case 410 without either. The Finish site again does not show any improvements with respect to 411 different combinations of data assimilated. Finally, the AU-How results show that the 412 assimilation of SIF and VOD improves the results compared to the 'ABGB_LAI_NBE' 413 case but not the 'LALABGB' case. That might be due to the peculiarities of this site 414 and LAI and VOD formulations in the model that will be discussed separately in Sec-415 tion 3.1.1. Peculiar case of Australian site. 416

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For all sites, SIF and VOD data can either substitute or improve the results obtained with NBE, however, the degree of improvement and the effect of a particular combination of the assimilated data varies from site to site, depending on the local conditions and, potentially, data availability and quality.

⁴²¹ Overall, the results measured with R^2 and RMSE are similar to the result based ⁴²² on histogram intersection (Fig. 3) and interannual anomaly metric ratio (Fig. 4). For ⁴²³ the histogram intersection, the '+SIF+VOD' case improves the results in comparison

to the 'ABGB_LAI_NBE' case for US-Ha1 both respiration and GPP, FR-Pue and AU-424 How respiration, and for the rest of the cases, improves the results in comparison to 'LALABGB'. 425 The only exception is the FI-Hyy site, where the addition of SIF and VOD degrades the 426 respiration histogram intersection. The fact that SIF and VOD assimilation can improve 427 both the metrics related to the model's median output (like R^2 and RMSE) and the model's 428 distribution output is an important result. Improving on just the median metrics can 429 indicate overfitting of the model, whereas improvement on histogram intersection alone 430 can signal about the model underfitting. The improvement on the two types of metrics 431 indicates the model indeed matched the observational distributions better. However, the 432 '+SIF+VOD' case never reached the interannual variability, measured as the IAM ra-433 tio, achieved for the 'ABGB_LAI_NBE' case except for Harvard Forest respiration and 434 FI-Hyy site. At the same time, assimilating SIF and VOD consistently improves the in-435 terannual variability compared to the 'LAI_ABGB'. In other words, while SIF and VOD 436 cannot fully substitute the degree of variability introduced by NBE, they still improve 437 the results in comparison to the case when NBE is simply not used. 438

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3.1.1 Peculiar case of Australian site

Another reason why VOD was less effective for the Australian (AU-How) site might 440 lie in the peculiarity of the VOD and LAI dynamic for this site. As shown in Fig. S7, 441 in this particular case, the synthetic and observational VOD data seem decoupled. That 442 is because we defined synthetic VOD as being proportional to above ground biomass (Eq. 443 4). Considering that LAI is also linearly proportional to the leaf mass, by definition, VOD 444 and LAI will always be coupled in the model. Yet, Tian et al. (2018) showed that VOD 445 and LAI can be decoupled for some regions. In particular, this pattern was found for African 446 tropical woodlands (centered at 11.5°S, 18.5°E). While the Australian site is classified 447 as a woody savanna, it is located at the same latitude as the African study region and 448 may exhibit similar dynamics. In addition, grass and trees in woody savannas have dif-449 ferent seasonal dynamics, with LAI dynamics driven mainly by the strong seasonal cy-450 cle of grass, whereas the dynamics of VOD are driven by trees. Due to the way LAI and 451 VOD are represented in the model, CARDAMOM is unable to reproduce such a decou-452 pling pattern. The tradeoff between model complexity (and, correspondingly, its abil-453 ity to replicate a wide variety of natural phenomena and, here, multiple plant functional 454

types within a single pixel) and model fidelity should be considered when carbon mod-els like CARDAMOM are used.

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3.2 Which of the SIF and VOD is more useful in this task?

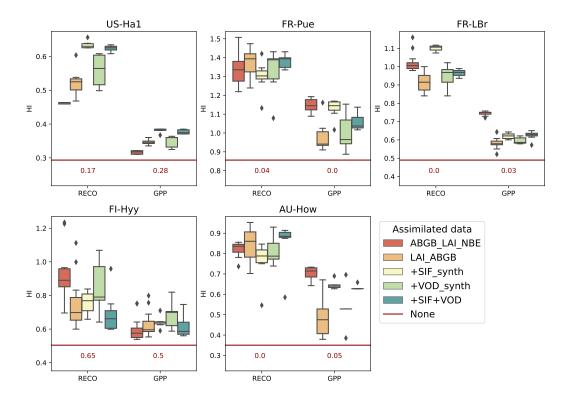
Further, different configurations of the assimilated data can help answer the research 458 question: Which of the SIF and VOD is more useful in this task, and how does it de-459 pend on the local conditions? Supplementary figures S1–S4 illustrate the same metric 460 as discussed above for assimilating either synthetic or observational SIF and VOD. For 461 all sites, including SIF in data assimilation immediately improves the model results for 462 GPP and respiration. For US-Ha1, FR-LBr, and FI-Hyy, the best results were achieved 463 with the assimilation of synthetic SIF data, which were outperformed by the observa-464 tional SIF only for FR-Pue and AU-How. Note that from Table 3, these are the two sites 465 with the most SIF data available, suggesting that more data can help further improve 466 the model skill in constraining respiration. The difference between the two versions of 467 the observational SIF data (GOME SIF and normalized GOME SIF) is minimal and was 468 only evident for the FR-LBr site. Synthetic and observational VOD performed surpris-469 ingly similarly when assessed by respiration R^2 and RMSE, except FR-Pue and AU-How 470 sites that had slightly better performance with the synthetic VOD. GLAB-VOD data 471 and VODCA VOD data are very different in the mean and the amplitude of year-to-year 472 variation (for illustration, see Fig. S5 and S7). Despite that, there is an apparent sim-473 ilarity of assimilating either of the VOD datasets on the model performance. We can as-474 sume some level of flexibility in this model configuration, and note that the effect of as-475 similating VOD should be more pronounced when looking at the simulated pools, rather 476 than fluxes (like GPP and respiration). At the same time, assimilating both SIF and VOD 477 together leads to improved results across sites and on an aggregate basis across metrics. 478

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3.3 Can respiration flux and carbon pools be reliably inferred from data assimilation, given indirect observational constraints?

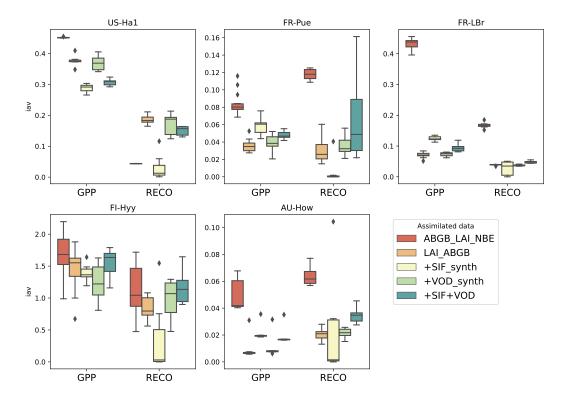
Figure 5 illustrates the effect of assimilating LAI, ABGB, SIF, and VOD ('+SIF+VOD' case) in comparison to 'NBE_LAI_ABGB' case and FLUXNET data on GPP and respiration. For all time series, the uncertainty is presented. The median output for GPP in the '+SIF+VOD' case slightly underestimates the summer peak. Still, the model output has a more prominent uncertainty range than the 'NBE_LAI_ABGB' case that in-



FLUXNET vs. model Histogram intersection

Figure 3. Effect of assimilating different combinations of data on Histogram intersection between CARDAMOM results distribution and FLUXNET RECO and GPP data. The 'none' case (no data assimilated) is denoted as a red line not in scale with the corresponding value of the parameter denoted.

cludes the observational data. In turn, the FLUXNET respiration time series are bet-486 ter matched with the SIF and VOD assimilation. Figure 6 sheds some light on the po-487 tential reasons behind the shift from the 'NBE_LAI_ABGB' case to the '+SIF+VOD' 488 case. It shows the posterior distribution for 10 sample model parameters, reflecting the 489 changes in model dynamics (parameters Fraction of GPP respired - "Frac GPP resp"; 490 temperature sensitivity Q10 - "q10"; canopy efficiency - "Canopy eff"; leaf mass carbon 491 per area, gC/m^2 - "LMCA," moisture factor) and initial values for five carbon pools ("C 492 labile" to "C SOM," gC/m^2). Most parameters converge equally well for both 'NBE_LAI_ABGB' 493 and '+SIF+VOD' cases with the posterior distributions clearly defined. While the frac-494 tion of GPP respired varies across a wider interval for the '+SIF+VOD' case, it takes 495 more realistic values around 50% (Van Oijen et al., 2010), than less than 25% obtained 496 in 'NBE_LAI_ABGB' case. At the same time, the posterior distributions for Q10 and 497



FLUXNET vs. model Interannual Anomaly Metric ratio

Figure 4. Effect of assimilating different combinations of data on relative interannual anomaly metric between median CARDAMOM results and FLUXNET GPP data. The 'none' case (no data assimilated) is denoted as a red line not in scale with the corresponding value of the parameter denoted.

canopy efficiency (that is used to calculate GPP using aggregated canopy model (ACM)) 498 have broader ranges for the '+SIF+VOD' case, which is expected since NBE is removed 499 from the assimilated data, and hence, GPP-related parameters are less constrained. There 500 is also a shift in LMCA. For deciduous forests (e.g., at US-Ha1), the typical value of leaf 501 mass per area is 75 q/m^2 (Poorter et al., 2009), which translates into a mean leaf car-502 bon mass per area 37.5 gC/m^2 , that corresponds better with the '+SIF+VOD' case re-503 sults. A slight shift in moisture factor – a parameter used in CARDAMOM to scale de-504 composition rate based on water availability – in the '+SIF+VOD' case shows that pre-505 cipitation influences the decomposition rate more than in the 'NBE_LAI_ABGB' case. 506 Another apparent shift is in initial carbon pool partitioning – the initial leaf carbon pool 507 is smaller, with the initial value for the wood carbon pool being larger for the '+SIF+VOD' 508 case in comparison to the previous case. At the same time, the '+SIF+VOD' case sug-509

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gests very low initial values for the root carbon pool, which might be infeasible. Fig. S6 shows the same 10 parameters for the remaining four sites. For all sites, there is a significant difference in distribution for at least some of the parameters, especially for the fraction of GPP respired, Q10, canopy efficiency, and wood carbon pool. This suggests that assimilating SIF and VOD positively affects the model and, indeed, favors better constraining respiration and carbon pools.

Overall, assimilating SIF and VOD instead of NBE does not deteriorate the model results and even seems to improve them at some sites, leading to a reduction in respiration RMSE for US-Ha1, FR-Pue, and AU-How. Furthermore, the changes in the model results are likely due to improvement of the model parameters, which shift towards more realistic values. Note that good metrics and a decent fit for the 'NBE_LAI_ABGB' case were achieved with a very different combination of CARDAMOM parameters than in the '+SIF+VOD' case as shown in Fig. 6.

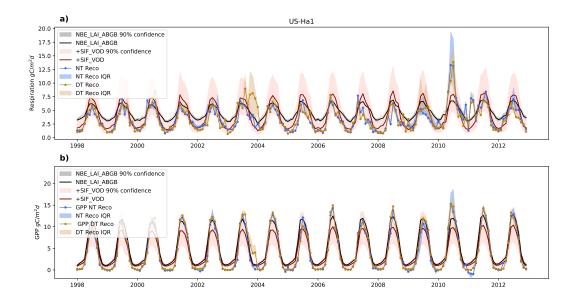


Figure 5. GPP and Respiration time series comparing FLUXNET data and median model outputs for 'NBE_LAI_ABGB' and '+SIF+VOD' cases with inter-quantile range for all variables

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3.4 Constraining carbon pools

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Assimilating SIF and VOD allows for constraining the carbon pools. Fig. 7 represents the temporal dynamics of the Harvard Forest carbon pools for the 'NBE_LAI_ABGB' and '+SIF+VOD' cases. The median and the interquantile range correspond to the full

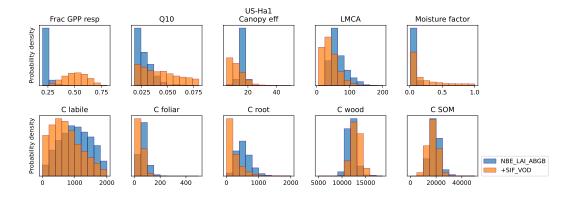


Figure 6. The effect of assimilating SIF and VOD in comparison to the 'NBE_LAI_ABGB' case on a selection of CARDAMOM parameters for the Harvard Forest site. All parameters are given in the ranges of their prior, except for C foliar pool (the original prior is $1 - 2000 \ gC/m^2$), C wood (the original prior is $1 - 100,000 \ gC/m^2$), and C SOM (the original prior is $1 - 200,000 \ gC/m^2$). The ranges were changed for these three parameters to highlight the visual differences between the two cases.

model outputs from all runs for a given scenario combined. Essentially, the difference 527 in the carbon pool partitioning represents the effect of substituting NBE with SIF and 528 VOD in data assimilation. This shift corresponds to the shift in the initial values of car-529 bon pools discussed above. In addition to it, the '+SIF+VOD' case modifies the dynam-530 ics of all pools, effectively propagating the constraints through the model. The same ef-531 fect persists for all sites examined in this study (See Supplementary Fig. S8 - S11). For 532 the Harvard Forest site, SIF and VOD reduced the size of all pools except for the wood 533 pool compared to the 'NBE_LAI_ABGB' case. For example, in the 'NBE_LAI_ABGB' 534 case, the mean values for the root carbon pool are close to $500 \ gC/m^2$, while in the '+SIF+VOD' 535 case, the mean value is reduced to 150 qC/m^2 . Additionally, for this pool, the seasonal 536 amplitude is reduced from 260 gC/m^2 to 60 gC/m^2 . Another interesting note is related 537 to the change in the wood carbon pool dynamics. In the 'NBE-LAI-ABGB' case, it has 538 a significant trend growing from 11,900 qC/m^2 on average in 1998 to 13,100 qC/m^2 by 539 the end of 2012, whereas, for the '+SIF+VOD' case, the overall growth over these years 540 is about 300 qC/m^2 . The more stable size of the wood biomass and moderate variabil-541 ity of the root biomass are expected for a stable ecosystem of the US-Ha1 site (Finzi et 542 al., 2020). 543

Another interesting result is revealed when above-ground biomass (ABG, calcu-544 lated as the sum of foliar and wood biomass) is compared to VOD and the empirical range 545 defined in Y. Y. Liu et al. (2015) as shown in Fig. 8. While the estimated biomass in 546 the '+SIF+VOD' case still mainly lies outside the range defined in Y. Y. Liu et al. (2015), 547 the VOD-AGB relationship is closer to the empirically determined bounds than the one 548 obtained from the 'NBE_LAI_ABGB' case. By definition, in-model VOD is linear with 549 respect to above-ground biomass and hence, will not follow the empirical relationship 550 exactly. Yet, the shift in carbon pools distribution and size brings the system towards 551 a different equilibrium than the 'NBE_LAI_ABGB' case. This new state is more aligned 552 with the Y. Y. Liu et al. (2015) findings. Hence, substituting NBE with SIF and VOD 553 allows constraining carbon pools more effectively. 554

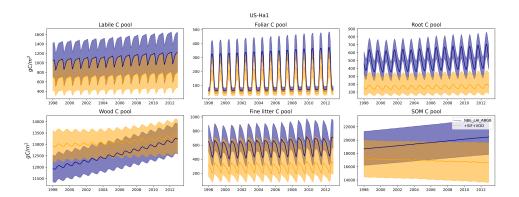


Figure 7. Carbon pools' temporal dynamic for 'NBE_LAI_ABGB' and '+SIF+VOD' cases for Harvard Forest. Median and interquantile range over the full output of all runs for a given scenario.

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3.5 Information content

The real-world observational data can be sparse and uncertain. We noted this effect already when examining the VOD data availability for different sites. We further explore the effect of assimilating more or less data and data uncertainty more systematically here.

Fig. 9 shows how varying the number of assimilated data points and data uncertainty affect ecosystem respiration estimates in terms of R^2 and RMSE for our five FLUXNET sites. The behavior at the Harvard Forest site follows our expectations when reducing

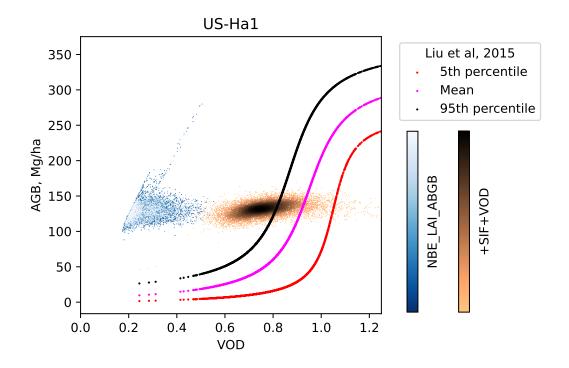


Figure 8. VOD vs. biomass relationship for 'NBE_LAI_ABGB' and '+SIF+VOD' cases in comparison to the empirical range defined in (Y. Y. Liu et al., 2015)

the amount of assimilated data deteriorated the model performance. Reducing the data 563 to 40 and 20% of the initially available data is equivalent to doubling and quadrupling 564 data uncertainty. Interestingly, reducing the data uncertainty for US-Ha1 improves the 565 results for both metrics (R^2 0.52 vs. 0.42, RMSE 2.29 vs 2.82 gC/m^2d for the initial un-566 certainty), suggesting that the used uncertainty is not optimal and the model can ex-567 tract more signal information from the assimilated data. For the FR-Pue site, the re-568 sults changing the data amount follow a similar pattern; however, reducing and increas-569 ing data uncertainty deteriorates the results. Based on that, we can conclude that the 570 initially used data uncertainty for this site is close to the optimal values. For the sec-571 ond French site, FR-LBr, the pattern of the results changes – using 80 and 60% of the 572 initially available data improves the respiration metrics (followed by further deteriora-573 tion of the results for even smaller amount of the used data that is also equivalent to qua-574 drupling the data uncertainty). Given the overall scarcity of SIF and VOD data for this 575 site and an apparent improvement of the results with the reduced data uncertainty (R^2 576 0.25 vs 0.18, RMSE 2.93 vs. 3.19 gC/m^2d for the initial uncertainty), we can assume that 577 the model struggles to fully differentiate between the signal and the noise in the initial 578

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configuration. Based on the results for the Finnish site, the model might be overfitted 579 when all available data is used. Here, the best results (yet with quite a large uncertainty) 580 are achieved for the smallest portion of the data -20%. For this site, it was already noted 581 that assimilating SIF and VOD does not improve the model results. Here, it is further 582 confirmed – an attempt to impose more constraints for this evergreen needle-leaved for-583 est site with little year-to-year variability deteriorates CARDAMOM predictions. Inter-584 estingly, the results for the AU-How site do not follow any pattern, most likely due to 585 the previously mentioned issue of LAI and VOD varying in anti-phase. Since the data 586 points for the assimilation are selected at random (Section 2.3 Information content 587 of observations), this subset can either improve or deteriorate the model output de-588 pending on the selected points. 589

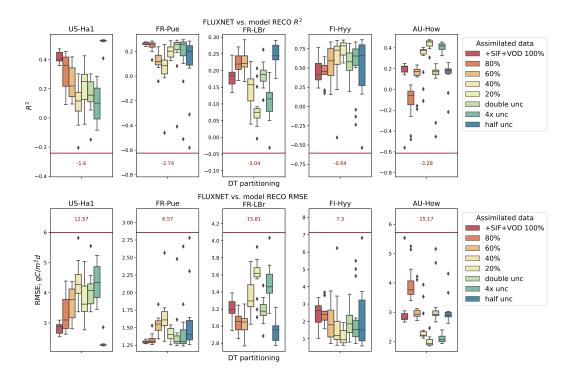


Figure 9. The effect of information content on CARDAMOM results: respiration R^2 and RMSE for all five sites

590 4 Conclusions

Process-based carbon cycle models like CARDAMOM can effectively model the car bon cycle and reconstruct carbon fluxes so that they match well with observations. The
 quality of the skill of the model in adequately simulating carbon pools and predicting

meaningful system behavior in contrafactual scenarios is based on the model structure 594 and how well it can reproduce natural phenomena with model equations – and model 595 parameters that are tuned through assimilation to observational data. Given the absence 596 of direct observations for respiration, photosynthesis, and net primary productivity and 597 the scarcity of biomass data on the global scale, this study investigates CARDAMOM's 598 ability to effectively utilize physical model constraints and indirect observations connected 599 to those processes, namely Solar-Induced Fluorescence (SIF) (as a proxy for photosyn-600 thesis) and Vegetation optical Depth (VOD) (used as a proxy for biomass). The results 601 indicate that these indirect observations can lead to improvements in the estimates of 602 carbon pools and respiration flux, often better than the ones obtained using net ecosys-603 tem exchange observations. This effect is even more witnessable when we consider the 604 scales of the observations since these results were achieved when comparing and assim-605 ilating FLUXNET site level net ecosystem exchange with a 25 km grid cell remote sensing-606 based SIF and VOD. The discrepancy between the site-level and remote sensing level 607 scales as a potential source of inconsistencies between CARDAMOM results and data 608 was also previously noted by A. J. Norton et al. (2023). 609

Moreover, the better performance achieved when assimilating net ecosystem ex-610 change (e.g., higher R^2 for US-Ha1 GPP) can be due to overfitting the model to follow 611 net ecosystem exchange variability. Bacour et al. (2023) used a different carbon cycle 612 model with a different combination of observational data assimilated, yet came to sim-613 ilar conclusions. Assimilation of more data, in our case, VOD, and SIF, leads to more 614 consistent results. Yet, similarly to Bacour et al. (2023), observational errors of multi-615 ple observations should be addressed to ensure an adequate signal-to-noise ratio and avoid 616 under- and overfitting. 617

The effect to which SIF and VOD observations improve the model estimates of car-618 bon pools and respiration flux depends on various factors. Among them are data avail-619 ability, data uncertainty and local conditions (interannual variability, stresses). Infor-620 mational content analysis reveals that for the locations that experience seasonal stresses. 621 more data helps better constrain the model. However, even for the sites with more sta-622 ble climate conditions, assimilating any data was beneficial compared to not assimilat-623 ing any SIF and VOD. The potential of SIF for carbon cycle modeling was noted ear-624 lier (MacBean et al., 2018; Bacour et al., 2019; MacBean et al., 2022; A. J. Norton et 625 al., 2023). Indeed, for FR-LBr, FI-Hyy, and AU-How, SIF not only successfully substi-626

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tuted but outperformed NBE (the performance measures as R^2 and RMSE for GPP). 627 VOD, on the other hand, while was proposed for data assimilation in carbon cycle mod-628 els (Scholze et al., 2017), and was assimilated in other instances (Kaminski et al., 2018; 629 Smith et al., 2020), was not previously used in models like CARDAMOM. We show that 630 assimilating VOD improves model performance in matching respiration flux and constrain-631 ing carbon pools. Indeed, respiration and carbon pool sizes are related (Ma et al., 2022), 632 and connected through the model parameters like respiration rate and canopy efficiency, 633 that were better constrained with the new assimilated data. 634

Additionally, we show that SIF and VOD assimilation improves metrics like R^2 and 635 RMSE, without deteriorating metrics like histogram intersection – a metric that assesses 636 the output distribution. Nevertheless, SIF and VOD could not achieve the same level 637 of interannual variability in the results that were previously achieved with net ecosys-638 tem exchange. This is expected since the site-level net ecosystem exchange comprises 639 more signal than averaged over the grid cell observational SIF and VOD. However, given 640 less reliable global net ecosystem exchange assessments (e.g., Peylin et al., 2013; Cui et 641 al., 2022), this finding can be wavered on a global scale. Satellite-based observational 642 SIF and VOD may be expected to further outperform the globally available version of 643 net ecosystem exchange obtained from CO_2 inversion. 644

Future work should focus on extending the study's geographical area to include other 645 climate zones. Observational data can have varying quality depending on the ecosystem 646 specifics (e.g., dense or sparse vegetation, frozen ground, and complex topography can 647 pose technical challenges for space-born sensors) or even human activity (e.g., radio-frequency 648 interference contaminating the signal). Further, model parameters' sensitivity to the as-649 similated data and its uncertainty can be spatially variable (Ma et al., 2022). As we have 650 already shown here, for the sites not subjected to stresses, the model can be overfitted 651 to data so that assimilating less data leads to better model performance. Finding more 652 regions prone to this behavior is essential to properly utilizing carbon cycle model-data 653 assimilation on the global scale. In addition to this, more attention should be dedicated 654 to the process representation in the model. As it was shown for LAI in A. J. Norton et 655 al. (2023), process representation plays a crucial role in the model's ability to effectively 656 extract information from the assimilated data to constrain model parameters. In this 657 study, we chose a simple VOD representation with minimal new parameters introduced 658 into the model. Even in this configuration, VOD successfully constrained carbon pools, 659

- including aboveground biomass, and improved model performance in comparison to as-
- similating just SIF. Future work should include VOD and SIF within a broad range of
- datastreams to best constrain the different carbon fluxes and pools.
- 663 Open Research Section

The data and scripts are publicly available at https://github.com/os2328/CARDAMOM SIF_VOD

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671 References

Arneth, A., Sitch, S., Pongratz, J., Stocker, B. D., Ciais, P., Poulter, B., ... others
(2017). Historical carbon dioxide emissions caused by land-use changes are
possibly larger than assumed. *Nature Geoscience*, 10(2), 79–84.

Bacour, C., MacBean, N., Chevallier, F., Léonard, S., Koffi, E. N., & Peylin, P.
(2023). Assimilation of multiple datasets results in large differences in regional-

- to global-scale nee and gpp budgets simulated by a terrestrial biosphere model. *Biogeosciences*, 20(6), 1089–1111.
- Bacour, C., Maignan, F., MacBean, N., Porcar-Castell, A., Flexas, J., Frankenberg,
 C., ... Bastrikov, V. (2019). Improving estimates of gross primary productivity by assimilating solar-induced fluorescence satellite retrievals in a terrestrial
 biosphere model using a process-based sif model. Journal of Geophysical Research: Biogeosciences, 124 (11), 3281–3306.
- Barr, A., Richardson, A., Hollinger, D., Papale, D., Arain, M., Black, T., ... others (2013). Use of change-point detection for friction-velocity threshold evaluation in eddy-covariance studies. *Agricultural and forest meteorology*, 171, 31–45.
- Berbigier, P., Bonnefond, J.-M., & Mellmann, P. (2001). Co2 and water vapour
 fluxes for 2 years above euroflux forest site. Agricultural and Forest Meteorology, 108(3), 183–197.

690	Beringer, J., Hutley, L. B., Tapper, N. J., & Cernusak, L. A. (2007). Savanna fires
691	and their impact on net ecosystem productivity in north australia. Global
692	Change Biology, 13(5), 990–1004.
693	Beven, K. (1993). Prophecy, reality and uncertainty in distributed hydrological mod-
694	elling. Advances in water resources, $16(1)$, $41-51$.
695	Beven, K., & Freer, J. (2001). Equifinality, data assimilation, and uncertainty esti-
696	mation in mechanistic modelling of complex environmental systems using the
697	glue methodology. Journal of hydrology, $249(1-4)$, 11–29.
698	Bloom, A., & Williams, M. (2015). Constraining ecosystem carbon dynamics in a
699	data-limited world: integrating ecological" common sense" in a model–data
700	fusion framework. Biogeosciences, $12(5)$, $1299-1315$.
701	Bloom, A. A., Exbrayat, JF., Van Der Velde, I. R., Feng, L., & Williams, M.
702	(2016). The decadal state of the terrestrial carbon cycle: Global retrievals
703	of terrestrial carbon allocation, pools, and residence times. Proceedings of the
704	National Academy of Sciences, 113(5), 1285–1290.
705	Bloom, A. A., Liu, J., Bowman, K. W., Konings, A., Saatchi, S., Worden, J. R.,
706	\dots Schimel, D. (2018). Observing the tropical carbon balance sensitivity to
707	memory and climate extremes. In Agu fall meeting abstracts (Vol. 2018, pp.
708	B51G–2016).
709	Chevallier, F., Remaud, M., O'Dell, C. W., Baker, D., Peylin, P., & Cozic, A.
710	(2019). Objective evaluation of surface-and satellite-driven carbon dioxide
711	atmospheric inversions. Atmospheric Chemistry and Physics, 19(22), 14233–
712	14251.
713	Cox, P. M., Betts, R. A., Jones, C. D., Spall, S. A., & Totterdell, I. J. (2000). Accel-
714	eration of global warming due to carbon-cycle feedbacks in a coupled climate
715	model. Nature, 408(6809), 184–187.
716	Cui, Y. Y., Zhang, L., Jacobson, A. R., Johnson, M. S., Philip, S., Baker, D.,
717	others (2022). Evaluating global atmospheric inversions of terrestrial net
718	ecosystem exchange co2 over north america on seasonal and sub-continental $% \mathcal{A}$
719	scales. Geophysical Research Letters, $49(18)$, e2022GL100147.
720	De Martonne, E. (1923). Aridité et indices d'aridité. Académie des Sciences.
721	Comptes Rendus, $182(23)$, $1935-1938$.
722	Deng, F., & Chen, J. (2011). Recent global co 2 flux inferred from atmospheric co 2

-30-

723	observations and its regional analyses. Biogeosciences, $8(11)$, $3263-3281$.
724	Denissen, J. M., Teuling, A. J., Pitman, A. J., Koirala, S., Migliavacca, M., Li, W.,
725	\ldots Orth, R. (2022). Widespread shift from ecosystem energy to water limita-
726	tion with climate change. Nature Climate Change, $12(7)$, 677–684.
727	Famiglietti, C. A., Smallman, T. L., Levine, P. A., Flack-Prain, S., Quetin, G. R.,
728	Meyer, V., \ldots others (2021). Optimal model complexity for terrestrial carbon
729	cycle prediction. <i>Biogeosciences</i> , 18(8), 2727–2754.
730	Finzi, A. C., Giasson, MA., Barker Plotkin, A. A., Aber, J. D., Boose, E. R.,
731	Davidson, E. A., others (2020). Carbon budget of the harvard forest long-
732	term ecological research site: Pattern, process, and response to global change.
733	Ecological Monographs, $90(4)$, e01423.
734	Friedlingstein, P., Jones, M. W., O'Sullivan, M., Andrew, R. M., Bakker, D. C.,
735	Hauck, J., others (2022). Global carbon budget 2021. Earth System
736	Science Data, 14(4), 1917–2005.
737	Fuster, B., Sánchez-Zapero, J., Camacho, F., García-Santos, V., Verger, A., Lacaze,
738	R., Smets, B. (2020). Quality assessment of proba-v lai, fapar and fcover
739	collection 300 m products of copernicus global land service. Remote Sensing,
740	12(6), 1017.
741	Gasser, T., Crepin, L., Quilcaille, Y., Houghton, R. A., Ciais, P., & Obersteiner, M.
742	(2020). Historical co 2 emissions from land use and land cover change and
743	their uncertainty. Biogeosciences, $17(15)$, $4075-4101$.
744	Haario, H., Saksman, E., & Tamminen, J. (2001). An adaptive metropolis algorithm.
745	Bernoulli, 223-242.
746	Hartmann, H., Bahn, M., Carbone, M., & Richardson, A. D. (2020). Plant car-
747	bon allocation in a changing world–challenges and progress: introduction to a
748	virtual issue on carbon allocation. New Phytologist, $227(4)$, 981–988.
749	Jackson, T., & Schmugge, T. (1991). Vegetation effects on the microwave emission of
750	soils. Remote Sensing of Environment, 36(3), 203–212.
751	Jian, J., Steele, M. K., Thomas, R. Q., Day, S. D., & Hodges, S. C. (2018). Con-
752	straining estimates of global soil respiration by quantifying sources of variabil-
753	ity. Global Change Biology, 24(9), 4143–4159.
754	Joiner, J., Yoshida, Y., Koehler, P., Frankenberg, C., & Parazoo, N. (2023). L2 daily

rss solar-induced fluorescence (sif) from metop-a gome-2, 2007-2018. ornl daac,

oak ridge, tennessee, usa.

756

757	Jung, M., Koirala, S., Weber, U., Ichii, K., Gans, F., Camps-Valls, G., Reich-
758	stein, M. (2019). The fluxcom ensemble of global land-atmosphere energy
759	fluxes. Scientific data, $6(1)$, 74.
760	Kaminski, T., Scholze, M., Knorr, W., Vossbeck, M., Wu, M., Ferrazzoli, P., oth-
761	ers (2018) . Constraining terrestrial carbon fluxes through assimilation of smos
762	products. In Igarss 2018-2018 ieee international geoscience and remote sensing
763	symposium (pp. 1455–1458).
764	Kennedy, D., Swenson, S., Oleson, K. W., Lawrence, D. M., Fisher, R., Lola da
765	Costa, A. C., & Gentine, P. (2019). Implementing plant hydraulics in the com-
766	munity land model, version 5. Journal of Advances in Modeling Earth Systems,
767	11(2), 485-513.
768	Kirschbaum, M. U., & McMillan, A. (2018). Warming and elevated co2 have oppos-
769	ing influences on transpiration. which is more important? Current forestry re-
770	ports, 4(2), 51-71.
771	Konings, A. G., Holtzman, N. M., Rao, K., Xu, L., & Saatchi, S. S. (2021). Inter-
772	annual variations of vegetation optical depth are due to both water stress and
773	biomass changes. Geophysical Research Letters, $48(16)$, e2021GL095267.
774	Konings, A. G., Piles, M., Das, N., & Entekhabi, D. (2017). L-band vegetation opti-
775	cal depth and effective scattering albedo estimation from SMAP. Remote Sens-
776	ing of $Environment, 198, 460-470.$
777	Konings, A. G., Saatchi, S. S., Frankenberg, C., Keller, M., Leshyk, V., Anderegg,
778	W. R., others (2021). Detecting forest response to droughts with global
779	observations of vegetation water content. Global change biology, $27(23)$, 6005–
780	6024.
781	Li, X., Liu, F., & Fang, M. (2020). Harmonizing models and observations: Data
782	assimilation in earth system science. Science China Earth Sciences, 63, 1059–
783	1068.
784	Li, X., Ma, H., Ran, Y., Wang, X., Zhu, G., Liu, F., \dots Huang, C. (2021). Terres-
785	trial carbon cycle model-data fusion: Progress and challenges. Science China
786	Earth Sciences, 64(10), 1645–1657.

Liu, L., Gudmundsson, L., Hauser, M., Qin, D., Li, S., & Seneviratne, S. I. (2020).
 Soil moisture dominates dryness stress on ecosystem production globally. Na-

789	$ture\ communications,\ 11(1),\ 1-9.$
790	Liu, Y. Y., Van Dijk, A. I., De Jeu, R. A., Canadell, J. G., McCabe, M. F., Evans,
791	J. P., & Wang, G. (2015). Recent reversal in loss of global terrestrial biomass.
792	Nature Climate Change, 5(5), 470–474.
793	López-Blanco, E., Exbrayat, JF., Lund, M., Christensen, T. R., Tamstorf, M. P.,
794	Slevin, D., Williams, M. (2019). Evaluation of terrestrial pan-arctic car-
795	bon cycling using a data-assimilation system. Earth System Dynamics, $10(2)$,
796	233–255.
797	Ma, R., Xiao, J., Liang, S., Ma, H., He, T., Guo, D., Lu, H. (2022). Pixel-
798	level parameter optimization of a terrestrial biosphere model for improving
799	estimation of carbon fluxes with an efficient model–data fusion method and
800	satellite-derived lai and gpp data. $Geoscientific Model Development, 15(17),$
801	6637 - 6657.
802	MacBean, N., Bacour, C., Raoult, N., Bastrikov, V., Koffi, E., Kuppel, S., others
803	(2022). Quantifying and reducing uncertainty in global carbon cycle predic-
804	tions: Lessons and perspectives from 15 years of data assimilation studies with
805	the orchidee terrestrial biosphere model. Global Biogeochemical Cycles, $36(7)$,
806	e2021GB007177.
807	MacBean, N., Maignan, F., Bacour, C., Lewis, P., Peylin, P., Guanter, L., Dis-
808	ney, M. (2018). Strong constraint on modelled global carbon uptake using
809	solar-induced chlorophyll fluorescence data. Scientific reports, $8(1)$, 1973.
810	Moesinger, L., Dorigo, W., de Jeu, R., van der Schalie, R., Scanlon, T., Teubner, I.,
811	& Forkel, M. (2020). The global long-term microwave vegetation optical depth
812	climate archive (VODCA). Earth System Science Data, 12(1), 177–196.
813	Moore, A. M., Arango, H. G., Broquet, G., Edwards, C., Veneziani, M., Powell,
814	B., Robinson, P. (2011) . The regional ocean modeling system (roms)
815	4-dimensional variational data assimilation systems: part ii–performance and
816	application to the california current system. Progress in Oceanography, $91(1)$,
817	50-73.
818	Munger, W., & Wofsy, S. (2014). Biomass inventories at harvard forest ems tower
819	since 1993. Harvard Forest Data Archive: HF069.
820	Nathaniel, J., Liu, J., & Gentine, P. (2023). Metaflux: Meta-learning global carbon

fluxes from sparse spatiotemporal observations. Scientific Data, 10(1), 440.

821

822	Norton, A., Bloom, A. A., Parazoo, N., Levine, P. A., Ma, S., & Braghiere, R. K.
823	(2021). The impacts of a climate-sensitive leaf phenology model on predict-
824	ing the terrestrial carbon cycle using the cardamom framework. In $Agu \ fall$
825	meeting 2021.
826	Norton, A. J., Bloom, A. A., Parazoo, N. C., Levine, P. A., Ma, S., Braghiere,
827	R. K., & Smallman, T. L. (2023). Improved process representation of leaf
828	phenology significantly shifts climate sensitivity of ecosystem carbon balance.
829	Biogeosciences, 20(12), 2455-2484.
830	Pastorello, G., Trotta, C., Canfora, E., Chu, H., Christianson, D., Cheah, YW.,
831	others (2020) . The flux net2015 dataset and the oneflux processing pipeline for
832	eddy covariance data. Scientific data, $7(1)$, 1–27.
833	Peylin, P., Law, R., Gurney, K., Chevallier, F., Jacobson, A., Maki, T., oth-
834	ers (2013). Global atmospheric carbon budget: results from an ensemble of
835	atmospheric co 2 inversions. Biogeosciences, $10(10)$, 6699–6720.
836	Piao, S., Huang, M., Liu, Z., Wang, X., Ciais, P., Canadell, J. G., others (2018).
837	Lower land-use emissions responsible for increased net land carbon sink during
838	the slow warming period. Nature Geoscience, $11(10)$, 739–743.
839	Poorter, H., Niinemets, Ü., Poorter, L., Wright, I. J., & Villar, R. (2009). Causes
840	and consequences of variation in leaf mass per area (lma): a meta-analysis.
841	New phytologist, $182(3)$, $565-588$.
842	Quetin, G. R., Bloom, A. A., Bowman, K. W., Diffenbaugh, N. S., Liu, J., & Kon-
843	ings, A. G. (2019). Attribution of historical terrestrial carbon uptake due to
844	climate change. In Agu fall meeting abstracts (Vol. 2019, pp. B11B–03).
845	Quetin, G. R., Bloom, A. A., Bowman, K. W., & Konings, A. G. (2020). Carbon
846	flux variability from a relatively simple ecosystem model with assimilated data
847	is consistent with terrestrial biosphere model estimates. Journal of Advances in
848	Modeling Earth Systems, $12(3)$, e2019MS001889.
849	Rambal, S., Joffre, R., Ourcival, J., Cavender-Bares, J., & Rocheteau, A. (2004).
850	The growth respiration component in eddy $co2$ flux from a quercus ilex
851	mediterranean forest. Global Change Biology, $10(9)$, $1460-1469$.
852	Raupach, M. R., Rayner, P. J., Barrett, D. J., DeFries, R. S., Heimann, M., Ojima,
853	D. S., Schmullius, C. C. (2005). Model–data synthesis in terrestrial carbon
854	observation: methods, data requirements and data uncertainty specifications.

855	Global Change Biology, 11(3), 378–397.
856	Scholze, M., Buchwitz, M., Dorigo, W., Guanter, L., & Quegan, S. (2017). Reviews
857	and syntheses: Systematic earth observations for use in terrestrial carbon cycle
858	data assimilation systems. $Biogeosciences$, $14(14)$, $3401-3429$.
859	Skulovich, O., Li, X., Wigneron, JP., & Gentine, P. (2024). GLAB-VOD:
860	Global L-band AI-BasedVegetation Optical Depth dataset based on machine
861	learning and remote sensing. Scientific Data (submitted). Retrieved from
862	https://zenodo.org/doi/10.5281/zenodo.10306094
863	Smith, W. K., Fox, A. M., MacBean, N., Moore, D. J., & Parazoo, N. C. (2020).
864	Constraining estimates of terrestrial carbon uptake: New opportunities us-
865	ing long-term satellite observations and data assimilation. $New Phytologist$,
866	225(1), 105-112.
867	Song, J., Wan, S., Piao, S., Knapp, A. K., Classen, A. T., Vicca, S., others
868	(2019). A meta-analysis of 1,119 manipulative experiments on terrestrial
869	carbon-cycling responses to global change. Nature ecology & evolution, $\mathcal{I}(9)$,
870	1309-1320.
871	Suni, T., Rinne, J., Reissell, A., Altimir, N., Keronen, P., Rannik, U., Vesala, T.
872	(2003). Long-term measurements of surface fluxes above a scots pine forest
873	in hyptiala, southern finland, 1996-2001. Boreal Environment Research, $\mathcal{S}(4)$,
874	287–302.
875	Talagrand, O. (1997) . Assimilation of observations, an introduction (gtspecial is-
876	sueltdata assimilation in meteology and oceanography: Theory and practice).
877	Journal of the Meteorological Society of Japan. Ser. II, 75(1B), 191–209.
878	Tharammal, T., Bala, G., Narayanappa, D., & Nemani, R. (2019). Potential roles
879	of co 2 fertilization, nitrogen deposition, climate change, and land use and land
880	cover change on the global terrestrial carbon uptake in the twenty-first century.
881	Climate Dynamics, 52, 4393–4406.
882	Tian, F., Wigneron, JP., Ciais, P., Chave, J., Ogée, J., Peñuelas, J., others
883	(2018). Coupling of ecosystem-scale plant water storage and leaf phenology
884	observed by satellite. Nature ecology & evolution, $2(9)$, 1428–1435.
885	Van Oijen, M., Schapendonk, A., & Höglind, M. (2010). On the relative magnitudes
886	of photosynthesis, respiration, growth and carbon storage in vegetation. An -
887	nals of Botany, 105(5), 793–797.

888	Wang, H., Jiang, F., Wang, J., Ju, W., & Chen, J. M. (2019). Terrestrial ecosys-
889	tem carbon flux estimated using gos at and oco-2 xco 2 retrievals. $Atmospheric$
890	Chemistry and Physics, 19(18), 12067–12082.
891	Wang, YP., Trudinger, C. M., & Enting, I. G. (2009). A review of applications
892	of model–data fusion to studies of terrestrial carbon fluxes at different scales.
893	Agricultural and forest meteorology, $149(11)$, $1829-1842$.
894	Wigneron, JP., Jackson, T., O'neill, P., De Lannoy, G., de Rosnay, P., Walker, J.,
895	\dots others (2017). Modelling the passive microwave signature from land sur-
896	faces: A review of recent results and application to the l-band smos & smap
897	soil moisture retrieval algorithms. Remote Sensing of Environment, 192,
898	238–262.
899	Williams, M., Richardson, A. D., Reichstein, M., Stoy, P. C., Peylin, P., Verbeeck,
900	H., \dots others (2009). Improving land surface models with fluxnet data.
901	$Biogeosciences, \ 6(7), \ 1341-1359.$
902	Wood, J. D., Griffis, T. J., Baker, J. M., Frankenberg, C., Verma, M., & Yuen, K.
903	(2017). Multiscale analyses of solar-induced florescence and gross primary
904	production. Geophysical Research Letters, 44(1), 533–541.
905	Wu, C., Niu, Z., Tang, Q., Huang, W., Rivard, B., & Feng, J. (2009). Remote
906	estimation of gross primary production in wheat using chlorophyll-related
907	vegetation indices. Agricultural and Forest Meteorology, 149(6-7), 1015–1021.
908	Xu, Z., Jiang, Y., & Zhou, G. (2015). Response and adaptation of photosynthesis,
909	respiration, and antioxidant systems to elevated $co2$ with environmental stress
910	in plants. Frontiers in plant science, 6, 701.
911	Yang, X., Tang, J., Mustard, J. F., Lee, JE., Rossini, M., Joiner, J., Richard-
912	son, A. D. (2015). Solar-induced chlorophyll fluorescence that correlates with
913	canopy photosynthesis on diurnal and seasonal scales in a temperate deciduous
914	forest. Geophysical Research Letters, 42(8), 2977–2987.
915	Yang, Y., Bloom, A. A., Ma, S., Levine, P., Norton, A., Parazoo, N. C., others
916	(2022). Cardamom-fluxval version 1.0: a fluxnet-based validation system for
917	cardamom carbon and water flux estimates. Geoscientific Model Development,
918	15(4), 1789-1802.
919	Yang, Y., Saatchi, S., Bloom, A. A., & Yu, Y. (2019). Validation of cms carbon flux

In

and stock datasets across the fluxnet eddy covariance flux tower network.

920

Agu fall meeting abstracts (Vol. 2019, pp. B13F-2435). 921 Yin, Y., Bloom, A. A., Worden, J., Saatchi, S., Yang, Y., Williams, M., ... others 922 (2020).Fire decline in dry tropical ecosystems enhances decadal land carbon 923 sink. Nature communications, 11(1), 1–7. 924 Zhang, Y., Guanter, L., Berry, J. A., Joiner, J., van der Tol, C., Huete, A., ... 925 Köhler, P. (2014).Estimation of vegetation photosynthetic capacity from 926 space-based measurements of chlorophyll fluorescence for terrestrial biosphere 927 models. Global change biology, 20(12), 3727-3742. 928

929 5 Supplemental

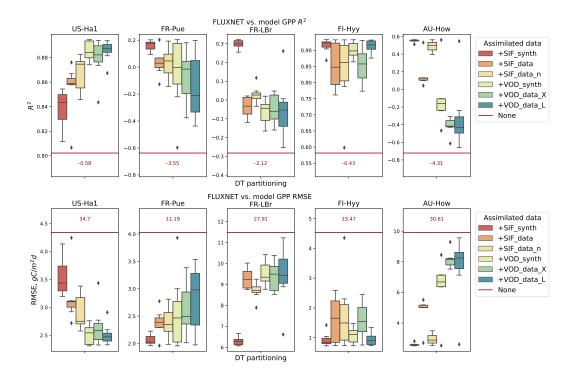


Figure S1. Effect of assimilating different data combinations on determination coefficient R^2 and RMSE between median CARDAMOM results and FLUXNET GPP data. The 'none' case (no data assimilated) is denoted as a red line not in scale with the corresponding value of the parameter denoted.

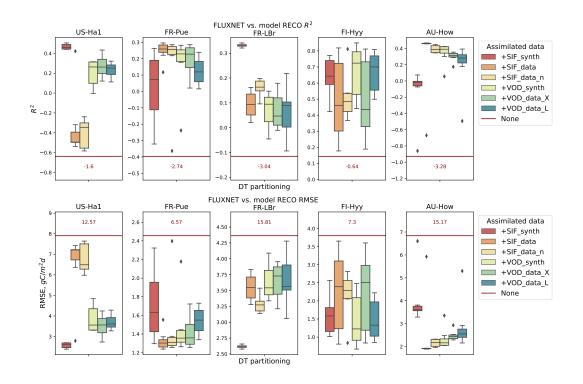
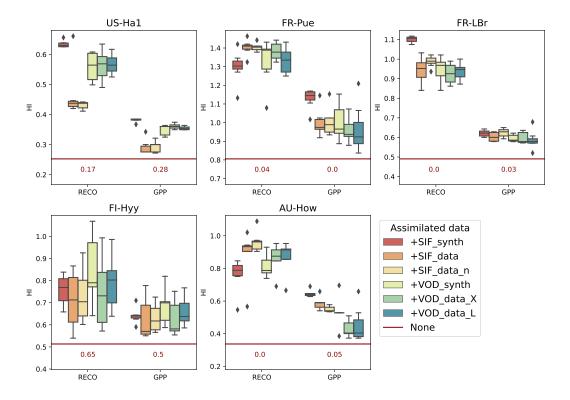
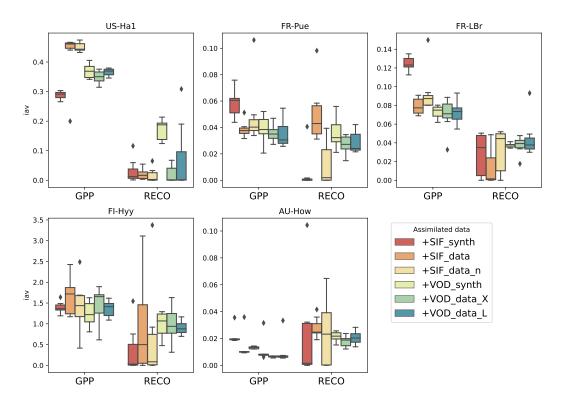


Figure S2. Effect of assimilating different combinations of data on determination coefficient R^2 and RMSE between median CARDAMOM results and FLUXNET RECO data. The 'none' case (no data assimilated) is denoted as a red line not in scale with the corresponding value of the parameter denoted.



FLUXNET vs. model Histogram intersection

Figure S3. Effect of assimilating different combinations of data on Histogram intersection between CARDAMOM results distribution and FLUXNET RECO and GPP data. The 'none' case (no data assimilated) is denoted as a red line not in scale with the corresponding value of the parameter denoted.



FLUXNET vs. model Interannual Anomaly Metric ratio

Figure S4. Effect of assimilating different combinations of data on relative interannual anomaly metric between median CARDAMOM results and FLUXNET GPP data. The 'none' case (no data assimilated) is denoted as a red line not in scale with the corresponding value of the parameter denoted.

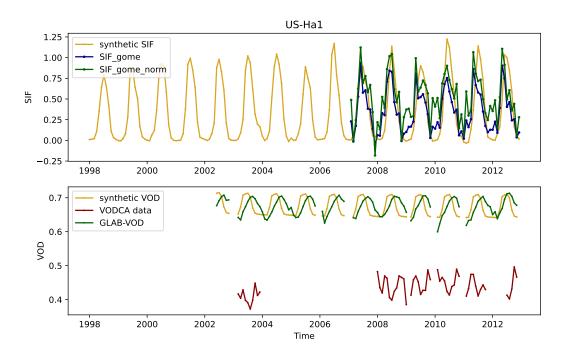


Figure S5. Synthetic and observational SIF and VOD data for US-Ha1 site

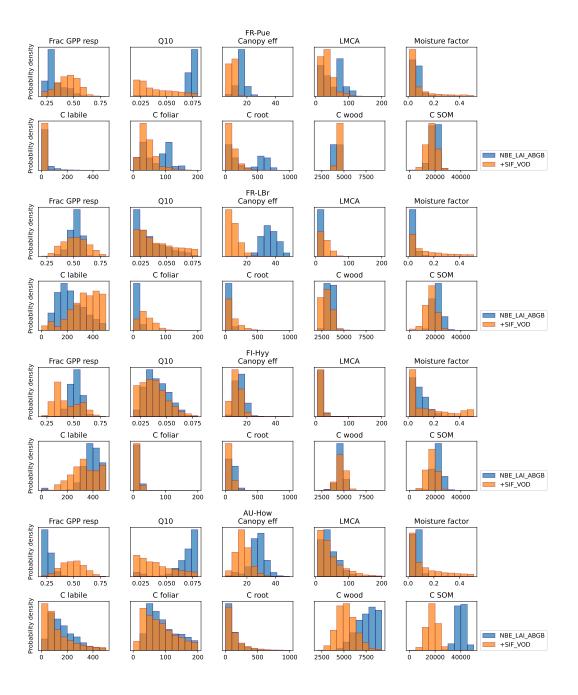


Figure S6. The effect of assimilating SIF and VOD in comparison to the "NBE_LAI_ABGB" case on a selection of CARDAMOM parameters for FR-Pue, FR-LBr, FI-Hyy, and AU-How sites.

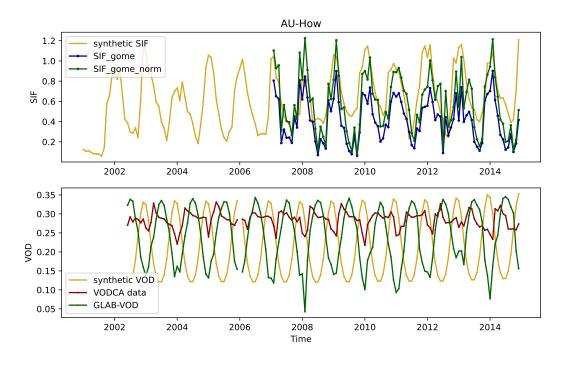


Figure S7. Synthetic and observational SIF and VOD data for AU-How site

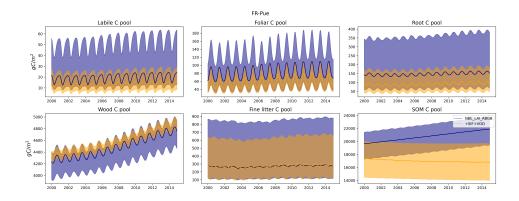


Figure S8. Carbon pools temporal dynamic for 'NBE_LAI_ABGB' and '+SIF+VOD' cases for the FR-Pue site. Median and interquantile range over the full output of all runs for a given scenario.

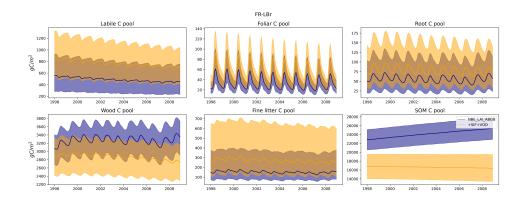


Figure S9. Carbon pools temporal dynamic for 'NBE_LAI_ABGB' and '+SIF+VOD' cases for the FR-LBr site. Median and interquantile range over the full output of all runs for a given scenario.

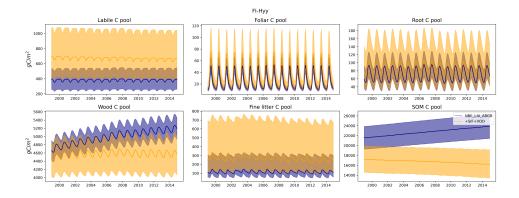


Figure S10. Carbon pools temporal dynamic for 'NBE_LAI_ABGB' and '+SIF+VOD' cases for the FI-Hyy site. Median and interquantile range over the full output of all runs for a given scenario.

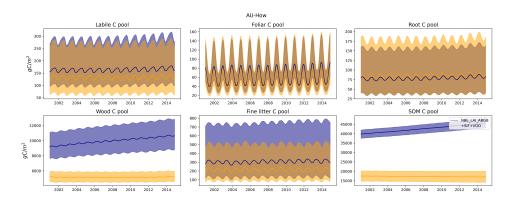


Figure S11. Carbon pools temporal dynamic for 'NBE_LAI_ABGB' and '+SIF+VOD' cases for the AU-How site. Median and interquantile range over the full output of all runs for a given scenario.

Figure 1.

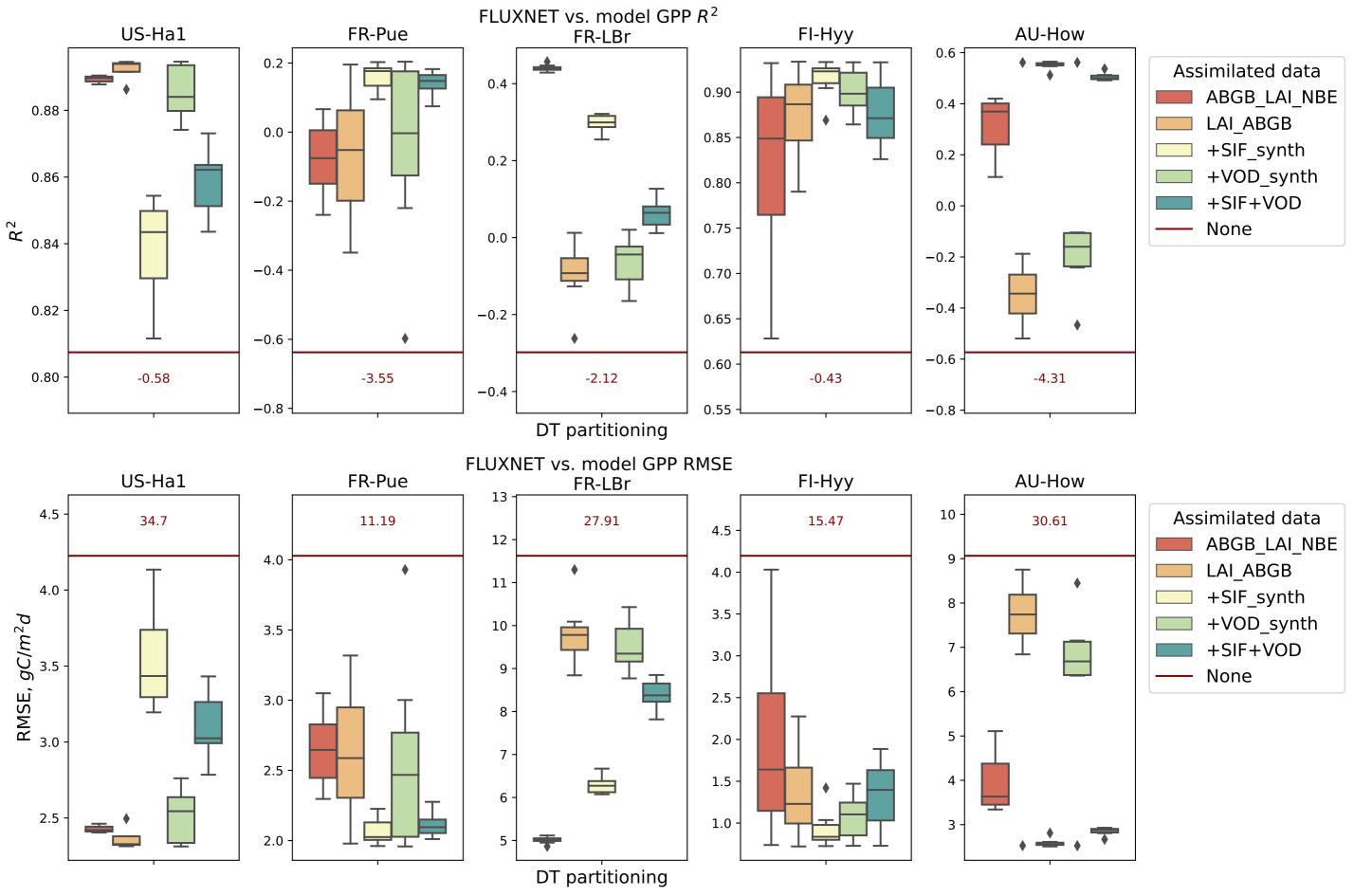


Figure 2.

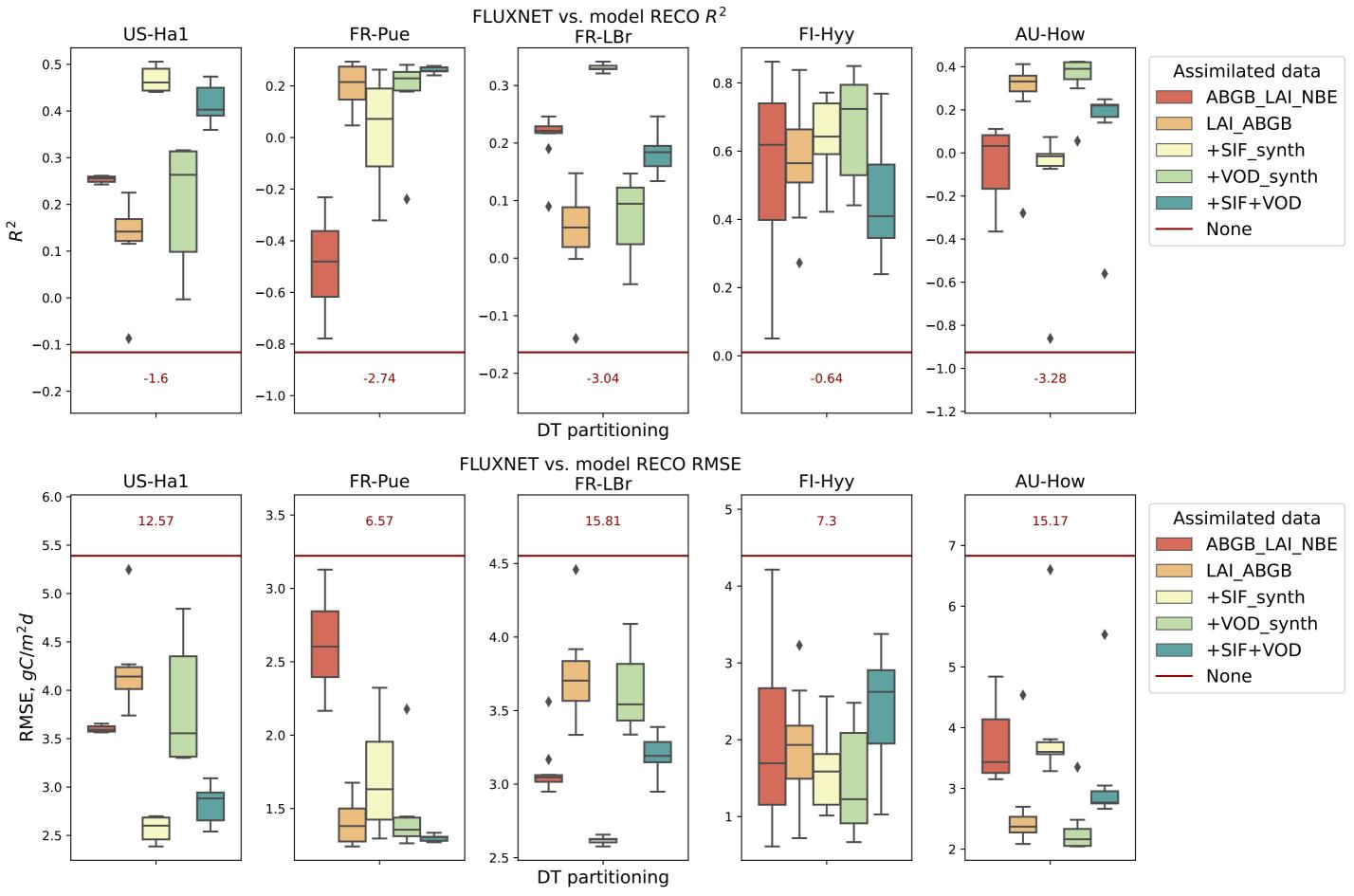


Figure 3.

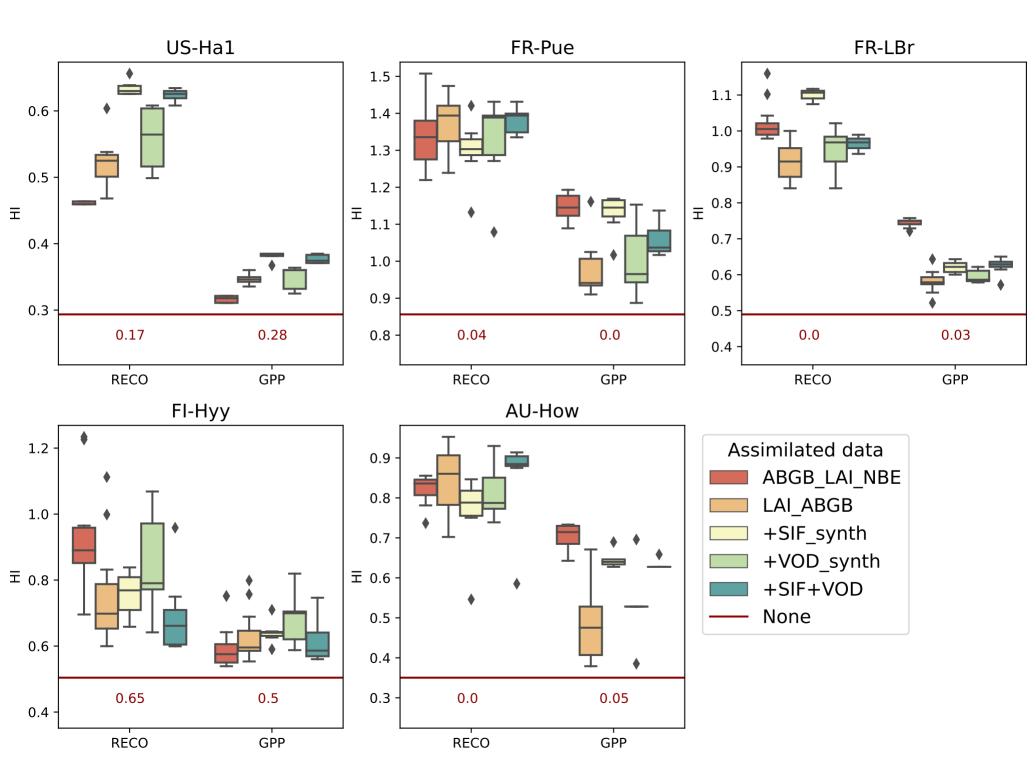


Figure 4.

FLUXNET vs. model Interannual Anomaly Metric ratio

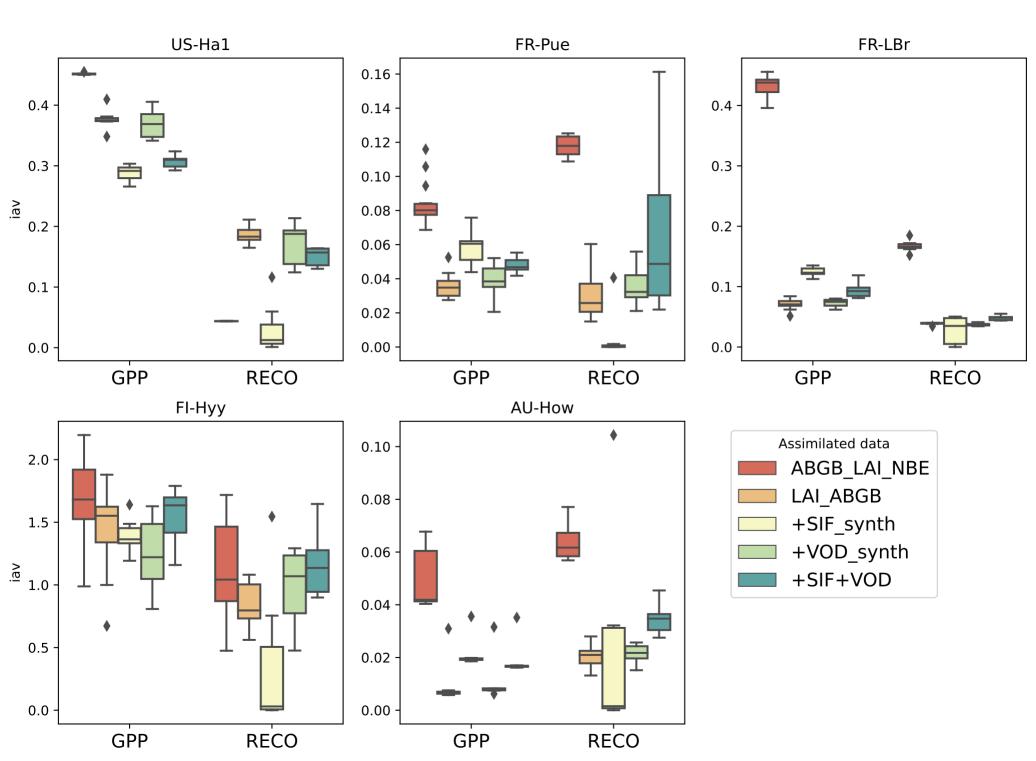


Figure 5.

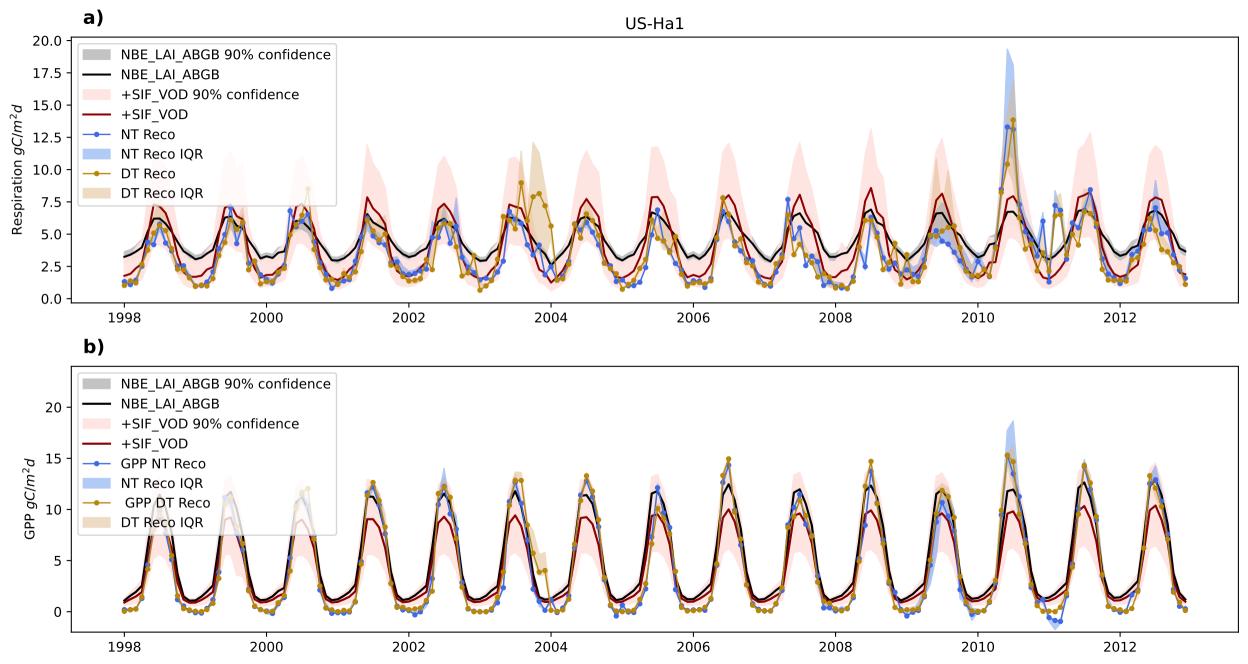


Figure 6.

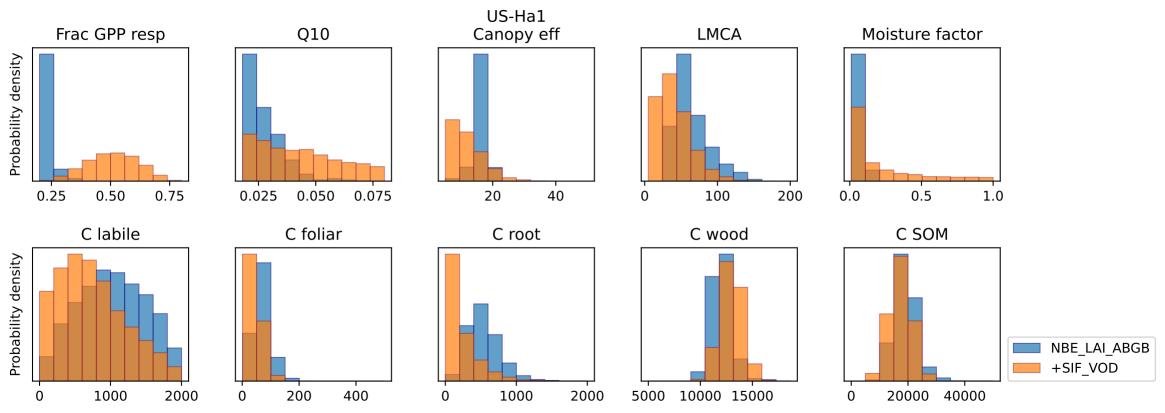


Figure 7.

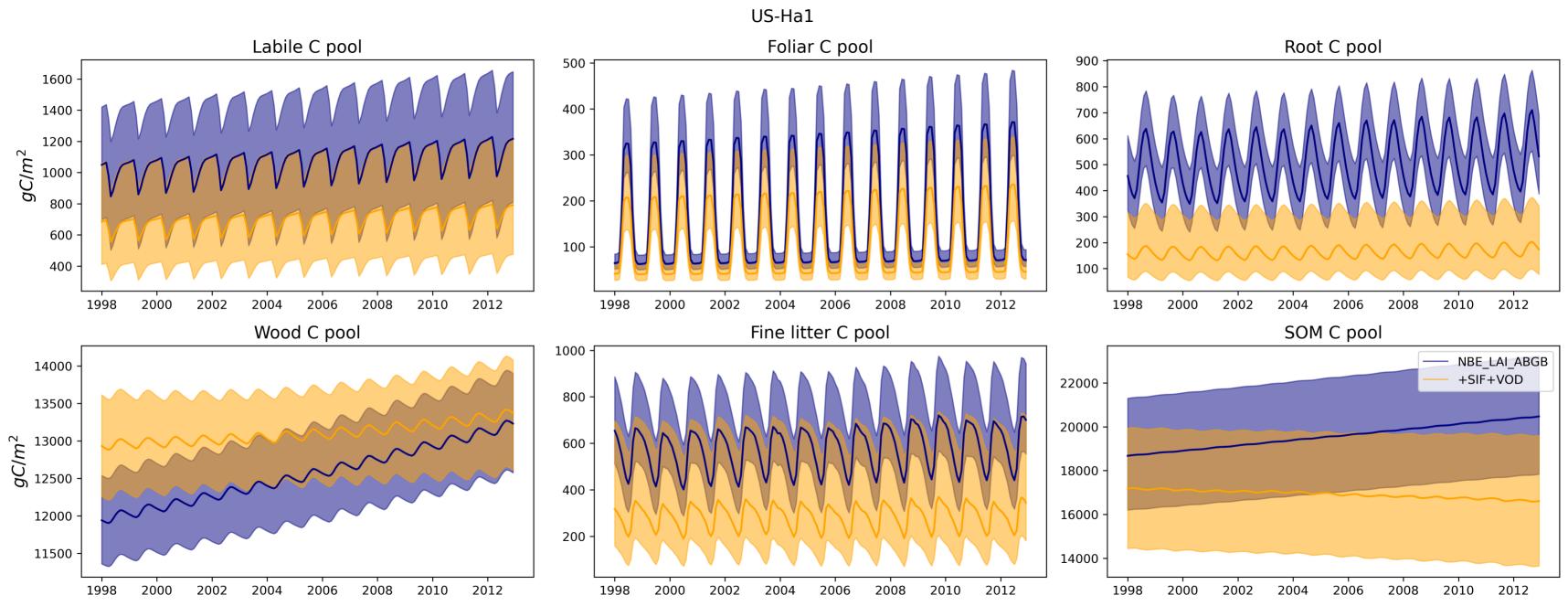


Figure 8.

US-Ha1

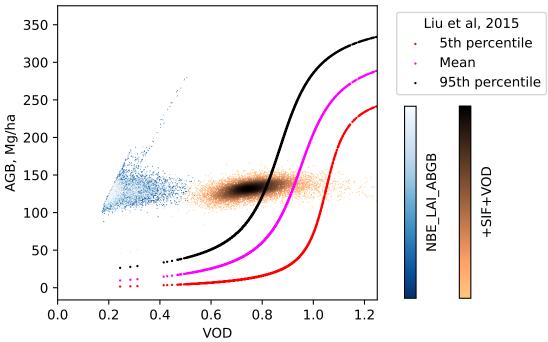
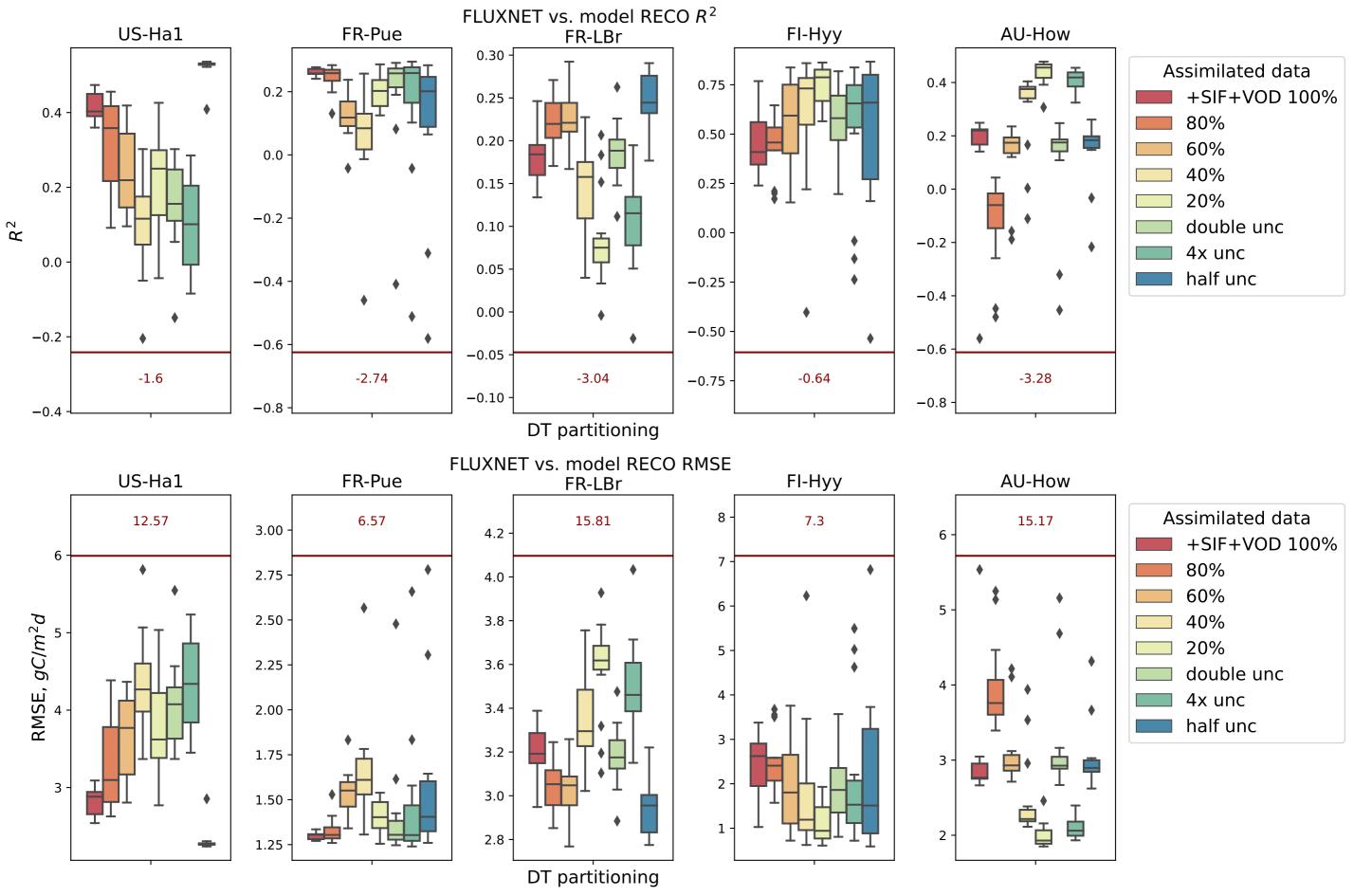
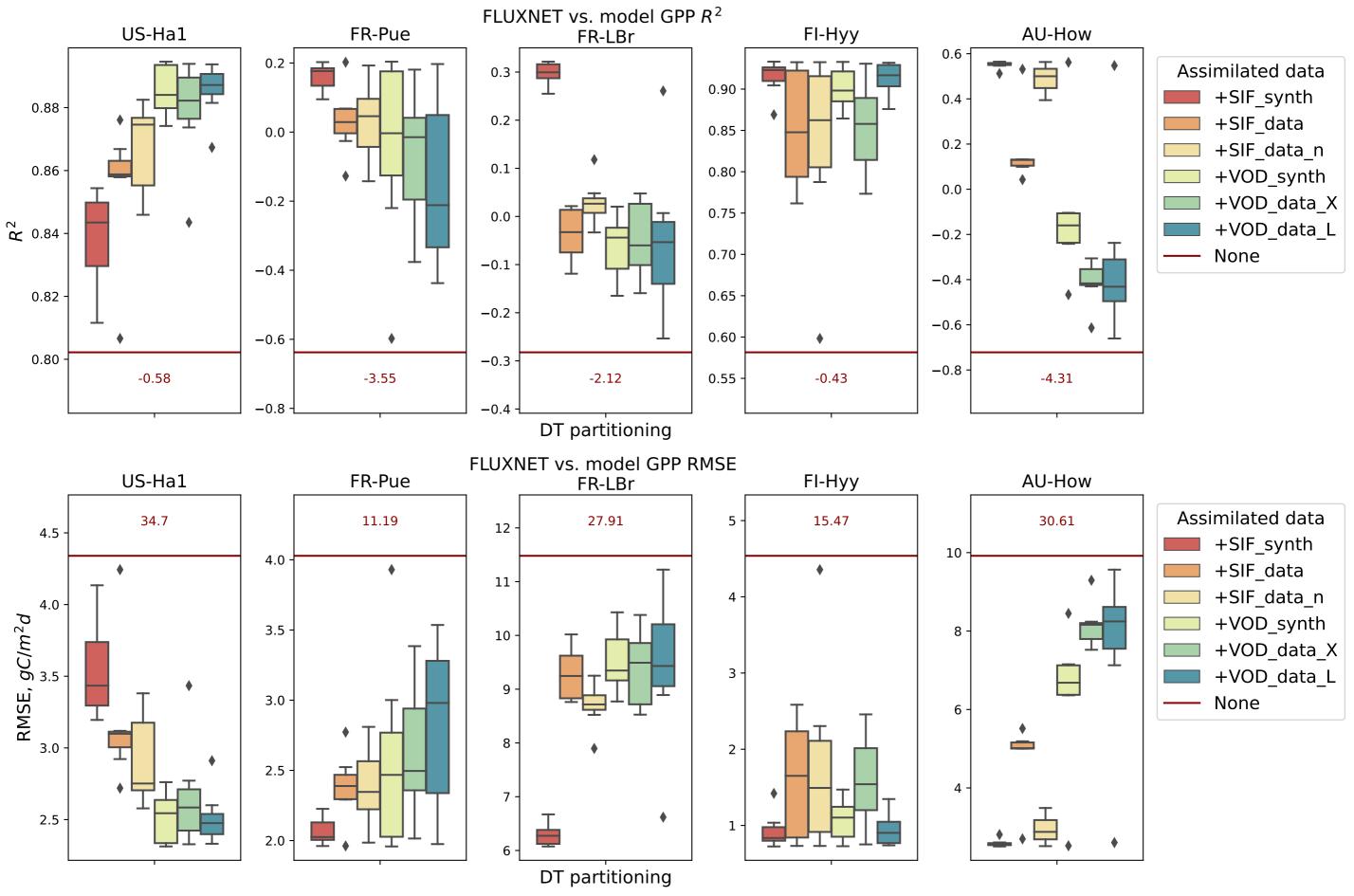


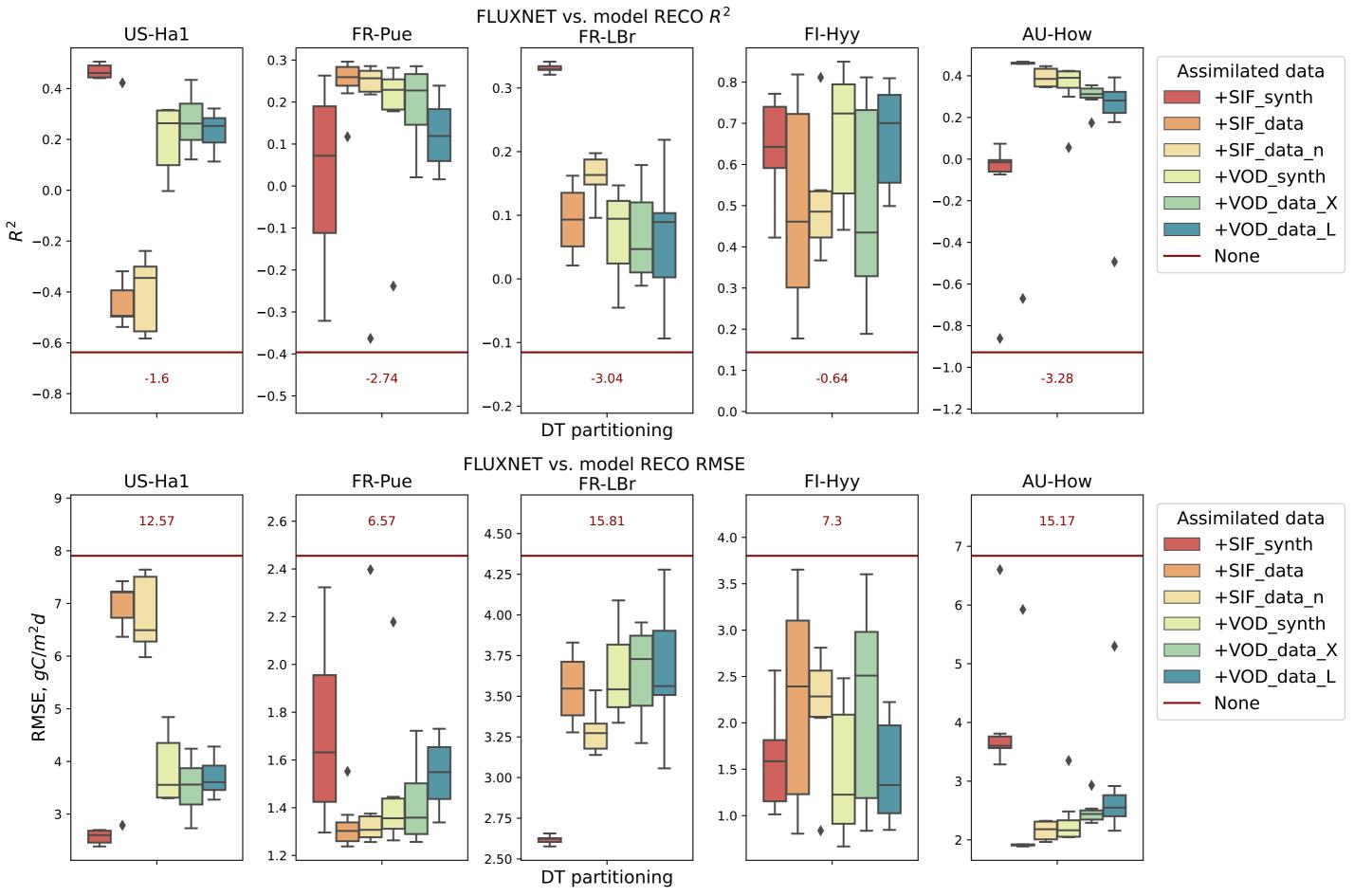
Figure 9.



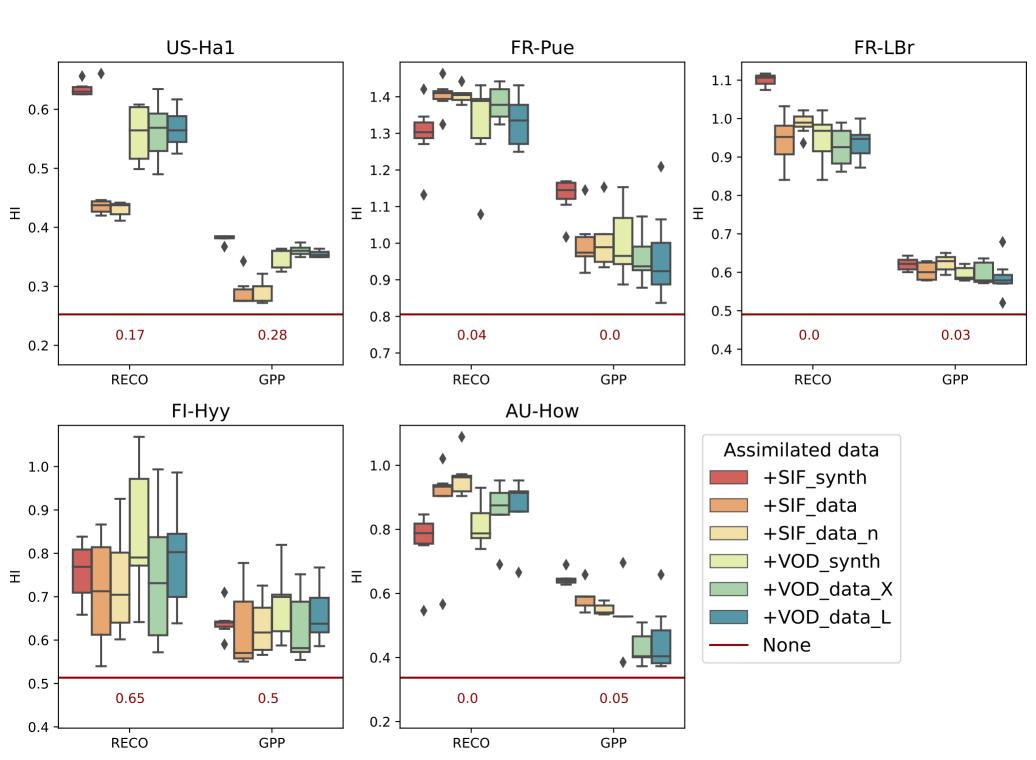
Supplementary Figure 1.



Supplementary Figure 2.

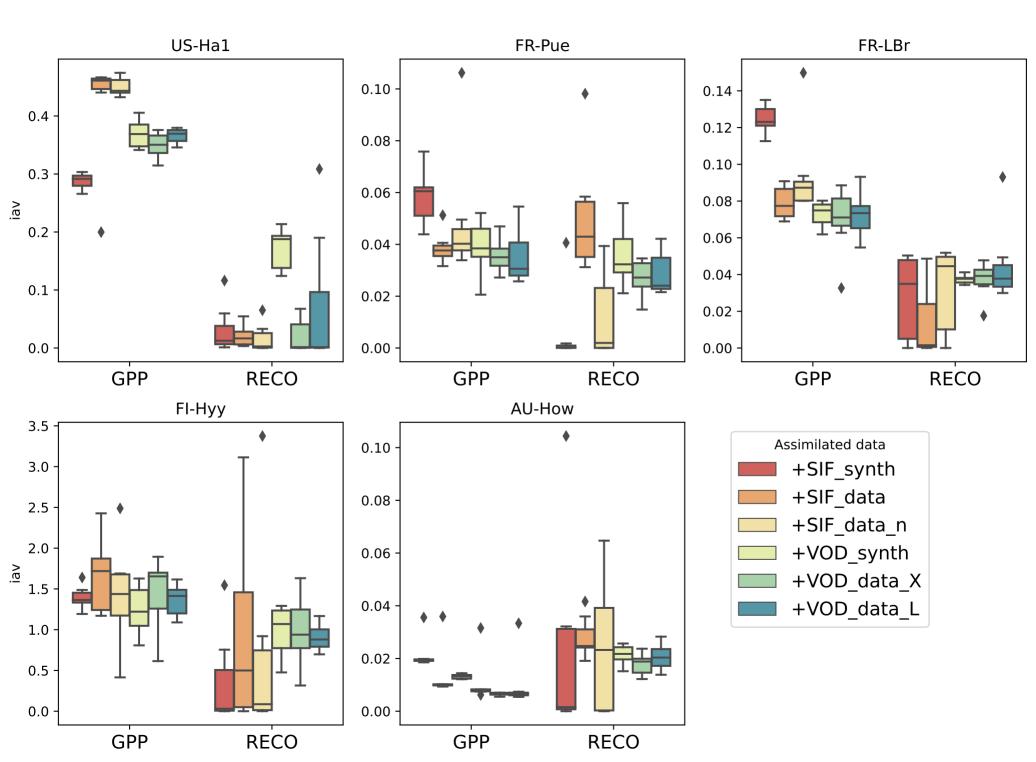


Supplementary Figure 3.

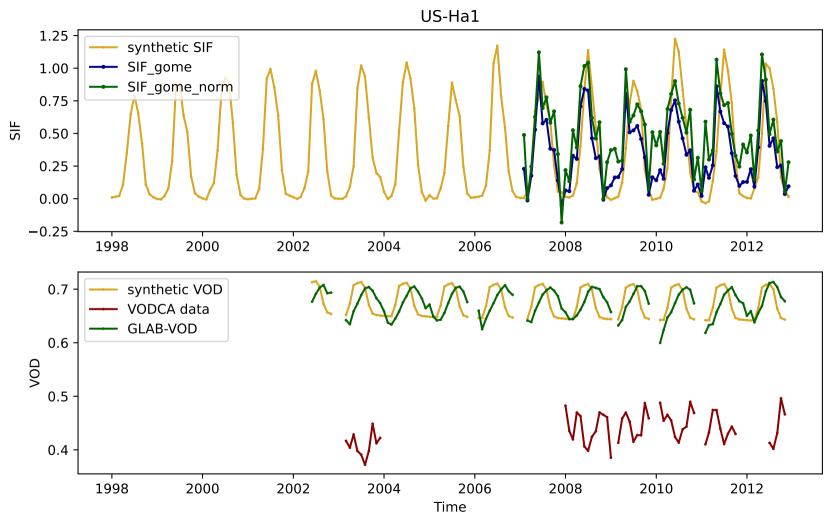


Supplementary Figure 4.

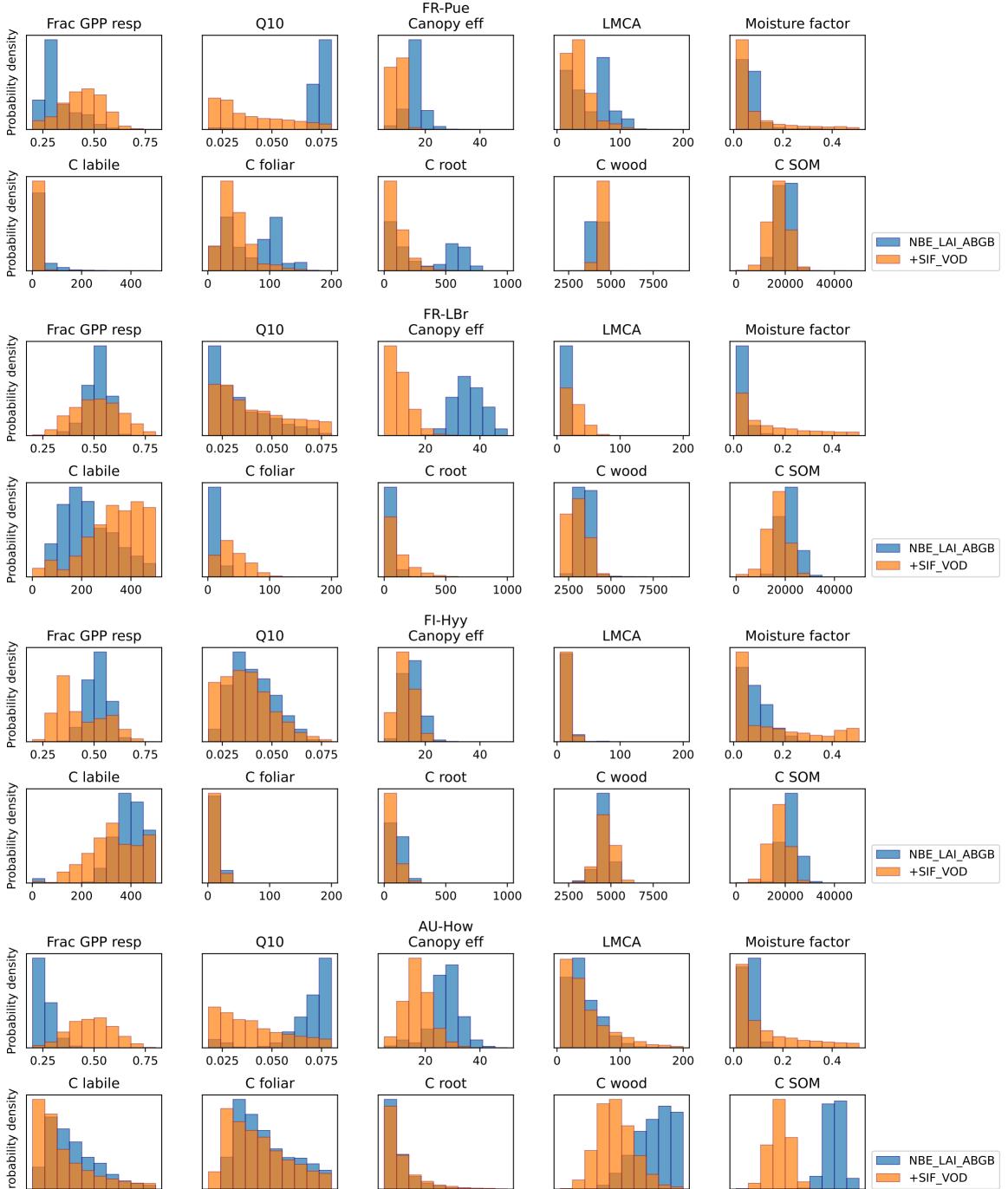
FLUXNET vs. model Interannual Anomaly Metric ratio



Supplementary Figure 5.



Supplementary Figure 6.



200 400

Probability density

0

100 200

0

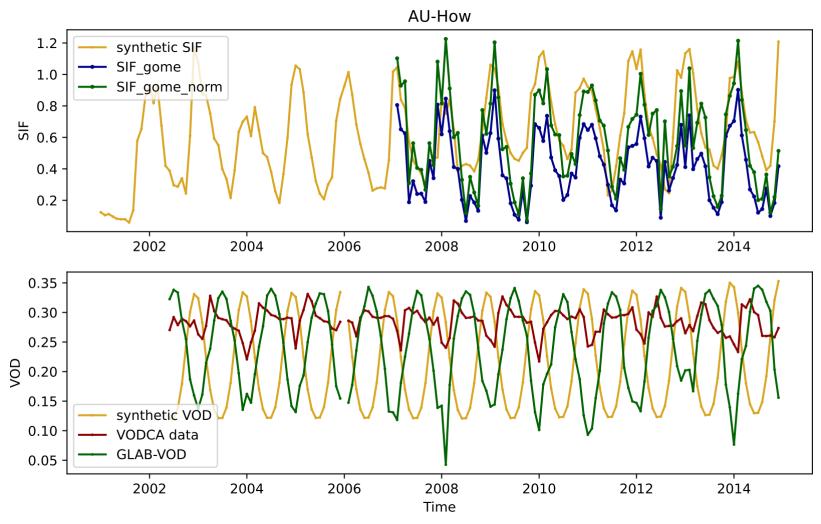
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0

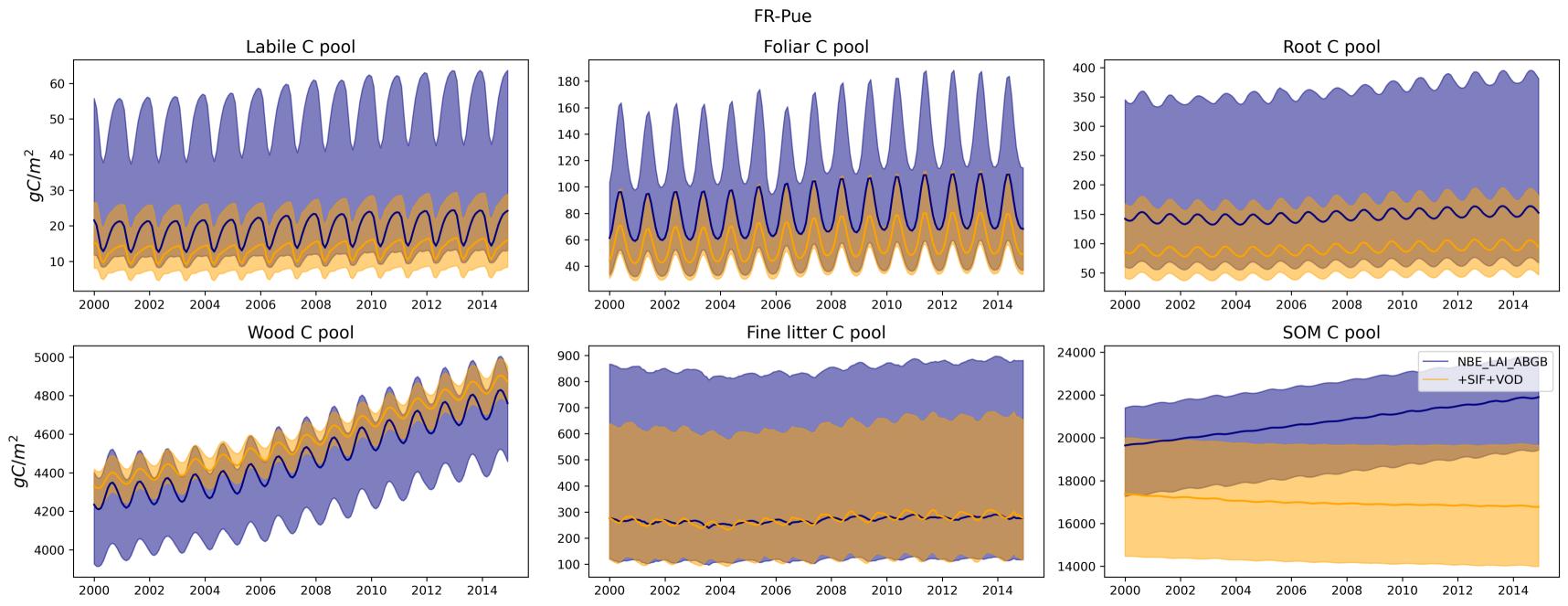
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20000 40000 0

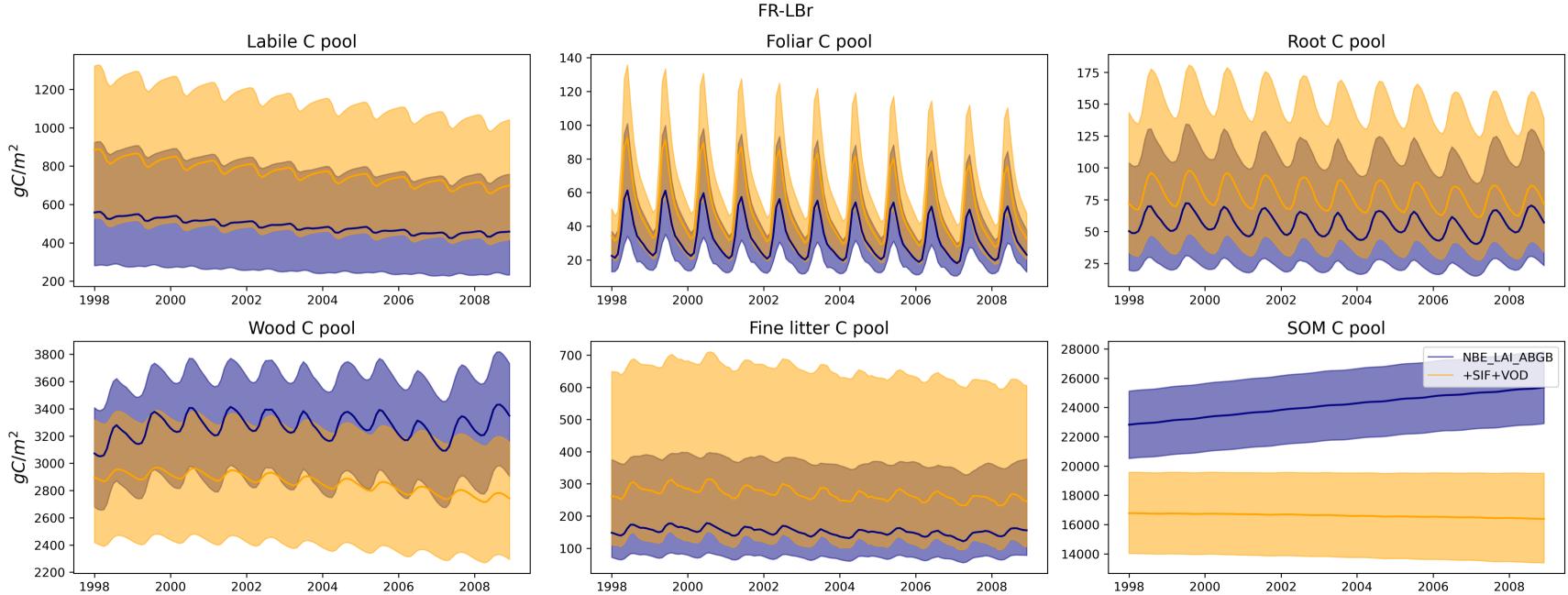
Supplementary Figure 7.



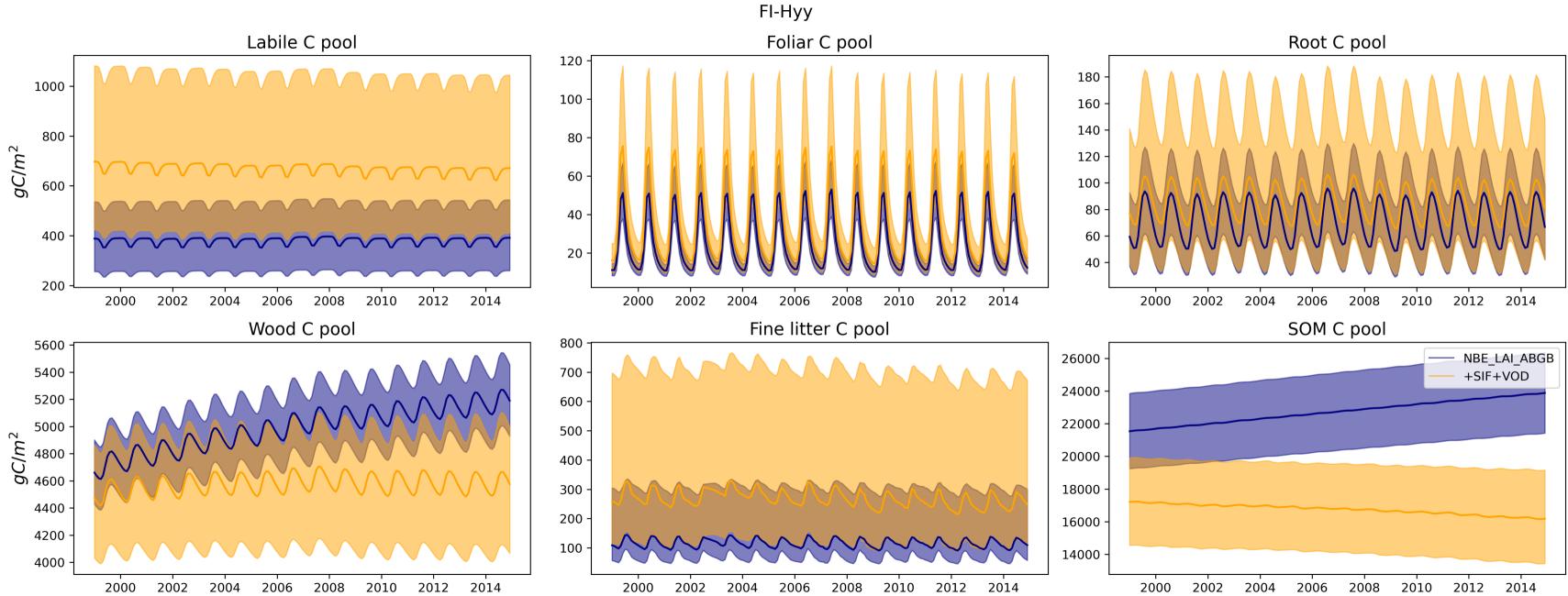
Supplementary Figure 8.



Supplementary Figure 9.



Supplementary Figure 10.



Supplementary Figure 11.

