

**Title: FORCCHN V2.0: An individual-based model for predicting multiscale forest carbon
dynamics**

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1 **Abstract**

2 Process-based ecological models are essential tools to quantify and predict forest growth and carbon cycle
3 under the background of climate change. The accurate description of phenology and tree growth processes
4 enables an improved understanding and predictive modeling of forest dynamics. An individual tree-based
5 carbon model, FORCCHN2 (FORest ecosystem Carbon budget model for CHiNa Version 2.0), used the
6 non-structural carbohydrates (NSC) pools to couple tree growth and phenology. This model performed
7 well in reducing uncertainty in predicting forest carbon fluxes. Here, we describe the framework in detail
8 and provide the source code of FORCCHN2. We also present a Dynamic Link Library (DLL) package
9 containing the latest version of the FORCCHN2 model. This package has the advantage of using Fortran
10 as an interface to make the model runs fast on a daily step, the package also allows the users to call it with
11 their preferred computer tools (e.g., Matlab, R, Python, *etc.*). FORCCHN2 model can be used directly to
12 predict the [spring and autumn phenological dates](#) as well as the daily carbon fluxes (including
13 photosynthesis, above- and belowground autotrophic respiration, and soil heterotrophic respiration) and
14 biomass on plot, regional, and hemispheric scales. As case studies, we provide an example of the
15 FORCCHN2 running, model validations in 78 forest sites, and an example model application for the
16 carbon dynamics of Northern Hemisphere forests. We demonstrate the FORCCHN2 model can produce
17 a reasonable agreement with flux observations. Given the potential importance of the application of this
18 ecological model in many studies, there is substantial scope for using the FORCCHN2 model in fields as

19 diverse as forest ecology, climate change, and carbon estimations.

20

21 **Keywords:** ecological models, forest ecosystems, carbon cycle, non-structural carbohydrates, individual

22 tree, leaf phenology

23 **1. Introduction**

24 Forests contribute an enormous carbon flux to terrestrial ecosystems (Pan et al. 2011, Keenan and
25 Williams 2018). Thus, accurate estimation and prediction of forest dynamics play an important role in
26 understanding the carbon cycle in the background of global change (Beer et al. 2010, Harris et al. 2021).
27 Over past decades, process-based ecological models have often been considered as effective tools for
28 evaluating forest dynamics at multiple scales (Friedlingstein et al. 2020).

29 Even though the ecological models are widely used in the prediction of forest dynamics, large
30 uncertainties remain (Huntzinger et al. 2012, Friedlingstein et al. 2020). Some of these uncertainties can
31 be attributed to the lack of effective phenological parameterization in the models and the neglect of
32 autumn phenology modeling (Raczka et al. 2013), both of which need to be based on an improved
33 understanding and coupling of mechanisms regulating forest phenology (Piao et al. 2019). Furthermore,
34 the previous models assumed that the reserve carbon of trees acts merely as a carbon buffer pool between
35 sink and source (Schiestl-Aalto et al. 2015). Recent studies considered the stored carbon as the non-
36 structural carbohydrates (NSC), which may have an active role on tree growth and carbon dynamics
37 (Martínez-Vilalta et al. 2016, Piper 2020). For example, trees rely on NSC to resume growth after the
38 non-growing season (Furze et al. 2019). The individual tree-based model, FORCCHN version 2.0
39 (FORCCHN2), has been developed to treat these considerations by integrating two NSC pools (NSC
40 active and slow pool) and optimizing phenological parameters (Fang et al. 2020a, Fang et al. 2021).

41 FORCCHN2 has improved performance for predicting forest carbon sinks compared to other models in
42 North American forests (Fang et al. 2020b).

43 This model provides the temporal predictions of individual tree growth processes as well as the
44 spatially explicit estimations of carbon dynamics on biomass, photosynthesis, autotrophic respiration, and
45 heterotrophic respiration (Fang et al. 2020b). The latest version can capture forest carbon dynamics, but
46 current runs of FORCCHN2 have limitations that prevent a seamless integration of the model into the
47 data-oriented software environment (e.g., Matlab, R, Python, etc.). FORCCHN2 and its previous versions
48 are designed originally for the daily calculation of individual trees in a given plot and implemented in
49 Fortran (Ma et al. 2017, Zhao et al. 2019, Fang et al. 2020a). Fortran ensures the calculation efficiency
50 and shortens the model runtime, but the model code and the implementation are not designed for the end-
51 users with appropriate help and instruction files. Moreover, until now, the FORCCHN2 model has only
52 been validated and applied in North America, and there has been no comprehensive publication describing
53 the model itself and no hemispheric-scale validation using this model.

54 Here, we present a DLL package aimed to provide a flexible and user-friendly interface for
55 implementing the newest FORCCHN2 model. Meanwhile, we provide the source code and the detailed
56 description of this model and demonstrates that FORCCHN2 model can predict a realistic and stable
57 carbon dynamics in the hemispheric-scale forests. With the package, users can conveniently run model
58 predictions on the individual, plot, regional, continental, and hemispheric scales according to their

59 computer tools. This package is compiled by Fortran 95 and thus can keep the high calculation efficiency.
60 We also demonstrate the functionality and example of FORCCHN2 model, perform the model validation
61 at the carbon flux sites, apply the model on a hemispheric scale (i.e. Northern Hemisphere), and provide
62 an open-access dataset of carbon outputs across the Northern Hemisphere.

63

64 **2. FORCCHN2 description**

65 FORCCHN2, an individual tree-based carbon dynamic model, predicts the daily processes of NSC,
66 photosynthesis, growth, [phenophase](#), vegetation (autotrophic) respiration, and soil dynamics in forests
67 (**Fig. 1** and **Method S1-S2**). This model is driven by the daily climate data and uses the leaf area index
68 (LAI) to initialize the vegetation information (i.e., trees' number, DBH, height, and biomass) on a fixed
69 area (**Method S3**).

70 For an individual tree, the NSC produced by photosynthesis is considered as the substrate supply for
71 vital activities, such as participating [in](#) the autotrophic respiration and forming the structural carbon pools
72 (i.e., leaves, wood, and fine roots) through growth (Sala et al. 2012, Richardson et al. 2013). The NSC
73 production is limited by external environmental factors (e.g., water, temperature, CO₂, etc.), and the NSC
74 consumption for the growth of each structural carbon pool (i.e., leaves, wood, and fine roots) is regulated
75 by phenology factors and daily climate (Schiestl-Aalto et al. 2015, Delpierre et al. 2019). [The phenophase](#)
76 [of spring and autumn in FORCCHN2 is controlled by heat and chilling requirements, respectively](#) (Fang

et al. 2022). The spring phenophase is decided by the effective temperature with the Thermal Time model (Eqn 39-40), and the autumn phenophase is decided by the effective temperature and photoperiod with the Cold Degree-Day model (Eqn 41-42). The model divides NSC into an active NSC pool and a slow NSC pool. The active pool provides the essential NSC consumption for daily activities; the slow pool is an NSC storage pool providing the necessary NSC for requirements when the contemporaneous active pool is insufficient, such as maintaining vegetation respiration during the non- and early-growing seasons. These NSC pools allow trees to be dead if the NSC storage drops below zero.

Dynamic changes of NSC production, allocation and consumption drive change in the NSC active pool (NSC_{active} , kg C) at a daily time step. The NSC slow pool (NSC_{slow} , kg C) is defined as the NSC storage pool. The changes in the daily active pool and yearly slow pool are:

$$\frac{dNSC_{active}}{dt} = \frac{dGPP}{dt} - \sum \frac{dR_j}{dt} - \sum \frac{dR_j^G}{dt} - \sum \frac{dG_j}{dt} \quad (1)$$

$$NSC_{slow}(y) = NSC_{active,y} \quad (2)$$

Where t is the day of the year; y is the y th year; j is each part of the tree (i.e. leaf, fine roots, and wood); GPP is gross primary productivity (kg C); R is the maintenance respiration (kg C); R^G is the growth respiration (kg C); G is the carbon demand of growth (kg C); $NSC_{active,y}$ is the size of NSC active pool at the end of y th year (kg C). The NSC active pool is initialized to zero on the first day of the next year. The calculation of GPP, maintenance respiration, growth respiration, and growth processes can be found in Method S1 and S2.

For the relationship between an individual tree with its neighbors, the model uses a distance-independent gap model to describe the light competition. To simplify the physiological and ecological parameters, each individual tree is assumed to belong to the plant functional types (PFTs) instead of specific tree species (**Table S2**). The PFT of one tree is decided by tree species when using the inventory data or it is estimated by forest types and random function when using the satellite data. The phenological parameters are parameterized by the local climate and observed phenological time in the first year (**Eqn S43-S45**). A part of structural carbon pools is then transferred into the soil pools by litter-fall. The main soil processes in the FORCCHN2 model are soil organic matter (SOM) decomposition, N mineralization, and water dynamics. According to the attribute, soil pools include above- and belowground metabolic and structural pools; fine and coarse woody litter pools; active, slow, and resistant SOM pools (**Table S4**). Except for these pools, the soil nitrogen pool also includes the inorganic nitrogen pool.

After each time step, the predicted vegetation and soil statements are converted into output variables such as biomass and carbon fluxes. The carbon fluxes of plot scale include GPP (kg C m⁻²), net primary productivity (NPP, kg C m⁻²), and net ecosystem productivity (NEP, kg C m⁻²). The NPP of a given plot at the daily step is determined by the GPP, R (kg C m⁻²), R^G (kg C m⁻²). The NEP of a given plot at the daily step is determined by the GPP, R, R^G, and soil respiration (R_S, kg C m⁻²):

$$\frac{dNPP}{dt} = \sum \frac{dGPP_n}{dt} - \sum \frac{dR_n}{dt} - \sum \frac{dR_n^G}{dt} \quad (3)$$

$$\frac{dNEP}{dt} = \frac{dNPP}{dt} - \frac{dR_S}{dt} \quad (4)$$

109 Where n is the n th tree of the plot.

110 The more detailed description, including inputs, outputs, calculation processes, and parameter sets
111 of FORCCHN2, can be found in **Table 1**, **Method S1-S3**, and **Table S2-S5**.

112

113 **3. Example runs**

114 Here, we provide an integrated DLL package ('FORCCHN2.dll') to simplify the usage of the
115 FORCCHN2 model. This file is highly flexible and it allows users to adapt model runs to their own
116 computer language (e.g., Matlab, R, Fortran, Python, etc.). Except for the model inputs, using only one
117 command can call the calculation of the model. We provide users with 32- and 64-bit DLL packages to
118 choose the most suitable version.

119 We take the Harvard Forest (a deciduous broadleaf forest in the eastern United States) and use Matlab
120 as an example run to demonstrate the functionality of the FORCCHN2 model (the code of example also
121 can be accessed via https://github.com/JingF1/FORCCHN2_model.git). First, we install and load the
122 package:

```
123 >>name1=('XXX');%load path of the FORCCHN2 DLL package
```

```
124 >>name2=[name1,'FORCCHN2_64.dll']; %input 64-bit or 32-bit DLL file
```

```
125 >>name3=[name1,'FORCCHN2.h'];%input header file
```

```
126 >>loadlibrary(name2,name3);%load the DLL package
```

Then, we input the data of Harvard Forest during 1991-2012. The inputs include the year information, the initialization data (i.e., geography, vegetation, and soil data), and the driven data (i.e., climate data). The more detailed information and format of these input data can be found in the example code ('FORCCHN2_run_example.m').

After inputting all data, we predict the dynamics of this forest for a period of 22 years. We can choose four output results of the FORCCHN2:

```
>>[fj,ycx,dayout,yearout]=calllib('FORCCHN2_64','forcchn2',fj,ycx,dayout,yearout,ntrees,ny0,ny,nday  
s,lat,lon,ele,tmax,tmin,tmean,pho,prec,ra,rh,wind,sfc,pwp,vw,sc0,sn0,silt,sand,class1,evergr0,deci0,lai0,  
co2);% run model with DLL file  
  
>>unloadlibrary FORCCHN2; %unload the DLL package
```

Where the **four** outputs include: 'fj' is the phenology dates, which included the start time of leaf growth (SOS) and the end time of leaf growth (EOS); 'ycx' is the allocation parameter of each soil pool, which can be used as input instead of the initial soil allocation parameters; 'dayout' is the daily carbon dynamics, which included above- and belowground biomass, gross primary productivity (GPP), above- and belowground respiration, soil heterotrophic respiration, litter-fall biomass, and soil carbon; 'yearout' is the yearly carbon dynamics.

4. External validation

145 The comparison between model simulations and external observations is considered as the rigorous model
146 test (Houlahan et al. 2017). Among the various observation methods, the eddy-covariance (EC) technique
147 can provide high-frequency and accurate measurements of relevant data (Keenan and Williams 2018).
148 The FLUXNET2015 dataset (Pastorello et al. 2020, <https://fluxnet.org/>) from the EC tower is an ideal
149 dataset to validate the FORCCHN2 model in predicting carbon flux dynamics. This dataset is developed
150 by using the EC (Eddy Correlation) technique to measure the net ecosystem CO₂ exchange (NEE, which
151 equaled to the negative of NEP) directly in the footprint of the EC tower. [The Variable Ustar Threshold](#)
152 [\(VUT\)](#) Mean values of FLUXNET2015 are used in this work. We extracted the flux data from the mean
153 value of the nighttime and the daytime method. The nighttime method uses nighttime NEE data to
154 parameterize a respiration-temperature model that is then applied to the whole dataset to estimate
155 Ecosystem Respiration (ER). The vegetation GPP is then calculated as the difference between ER and
156 NEE (Lasslop et al. 2010). The daytime method uses daytime and nighttime NEE data to parameterize a
157 model with one component based on a light-response curve and vapor pressure deficit for GPP, and a
158 second component using a respiration-temperature relationship similar to the nighttime method
159 [\(Pastorello et al. 2020\)](#). Due to the different phenological phasing in the Northern and Southern
160 Hemisphere, our predictions focus on the Northern Hemisphere. We chose the 78 active forest sites with
161 continuous daily observations in the Northern Hemisphere (i.e., a total of 232664 observations). These
162 sites cover the most forest types, including the evergreen broadleaf forest (EBF), evergreen needleleaf

163 forest (ENF), deciduous broadleaf forest (DBF), and mixed forest (MF). The distribution and information
164 of all sites are shown in **Fig. S1** and **Table S1**. We also extract the climate data from the FLUXNET2015
165 dataset to drive the model. Soil data are taken from the Harmonized World Soil Database (HWSD) V1.2
166 (<http://www.fao.org/soils-portal/soil-survey/soil-maps-and-databases/>).

167 We predict the daily carbon flux in the 78 forest sites and then validate the predictions with the
168 observations. As the overall performance, **Fig. 2** shows the direct daily comparison between predictions
169 and observations. Overall, the model had the best performance in capturing GPP dynamics, followed by
170 ER and NEP (i.e. the predicted GPP has the highest R). In the FORCCHN2 model, we use the phenology
171 model and the optimized phenological parameters to predict the leaf growth, which could improve the
172 predicted performance of GPP (Fang et al. 2020b). We did the statistics for the results in all sites. The
173 validation statistics include the correlation coefficient (R), model efficiency (E , calculated by **Eqn S60**),
174 root mean square error ($RMSE$), mean absolute error (MAE), and bias ($Bias$, calculated by **Eqn S61**). The
175 calculation of each statistic can be found in **Methods S4**. Each site had one group of statistics. **Fig. 3**
176 shows that the FORCCHN2 model could reproduce the daily dynamics of the carbon flux in all sites,
177 particularly for predicting daily GPP (median of all sites: $R=0.86$, $E=0.62$, $RMSE=2.29\text{ g C m}^{-2}\text{ d}^{-1}$,
178 $MAE=1.61\text{ g C m}^{-2}\text{ d}^{-1}$). The predicted ER performs lower than GPP (i.e. the median of R and E from the
179 predicted ER is less than GPP) but shows a high correlation with the observed ER (median: $R=0.83$,
180 $E=0.25$, $RMSE=1.46\text{ g C m}^{-2}\text{ d}^{-1}$, $MAE=1.04\text{ g C m}^{-2}\text{ d}^{-1}$). NEP results had the lowest performance in all

181 flux variables (median: $R=0.61$, $E=-0.16$, $RMSE=1.91$ g C m⁻² d⁻¹, $MAE=1.43$ g C m⁻² d⁻¹). The highest
182 uncertainty in predicting NEP maybe because of the compounding effect of GPP and ER errors (Balzarolo
183 et al. 2014). In terms of bias, FORCCHN2 overestimates the GPP and ER (median: $Bias=0.49$ and 0.56
184 g C m⁻² d⁻¹, respectively) but slightly underestimates the NEP (median: $Bias=-0.14$ g C m⁻² d⁻¹). For the
185 different forest types, the predictions present well in DBF and MF ($R=0.84$ and 0.57 , $E=0.53$ and 0.64 ,
186 respectively), whereas the lowest performance is found in EBF ($R=0.61$, $E=0.31$). These results are
187 consistent with the previous studies: EBF reveals subtle changes in the leaf phenology and thus increases
188 the difficulty in modeling photosynthesis (i.e., GPP) (Raczka et al. 2013, Yuan et al. 2014, Piao et al.
189 2019).

190

191 **5. Applications in the Northern Hemisphere**

192 As a case application on large scale, we predict the carbon dynamics in the Northern Hemisphere forests
193 during 1980-2016 (spatial resolution: 0.5×0.5 degree). For the Hemisphere, we use the Simple Biosphere
194 (SiB) model of the International Satellite Land Surface Climatology Project (ISLSCP II) to represent
195 forest types (**Fig. S1**, https://daac.ornl.gov/ISLSCP_II) (Friedl et al. 2010). The LAI data are extracted
196 from the Global Land Surface Satellite (GLASS) Product (<http://www.glass.umd.edu/Download.html>).
197 The climate data are from the daily analysis of ERA-Interim in the European Centre for Medium-range
198 Weather Forecasts (ECMWF) dataset (Hersbach et al. 2020). Soil data are taken from the HWSD V1.2.

Fig. 4 reported the spatial distribution of 37-year averaged GPP, above- and belowground autotrophic

respiration, soil heterotrophic respiration, net primary productivity (NPP), and net ecosystem productivity

(NEP) for forest area. All results show a similar spatial pattern with the largest fluxes occurring around

the equator, such as the northern part of the Amazon and Central African tropical rainforests; secondly,

the monsoonal subtropical regions such as South Asia and eastern North America show the large fluxes;

the northern forests near the Arctic Circle had the smallest fluxes. Overall, our predictions demonstrate

that the forests in Northern Hemisphere had a huge carbon sink potential by the vegetation (i.e.,

$\text{NPP}=16.76 \text{ Pg C year}^{-1}$ or $61.45 \text{ Gt CO}_2 \text{ year}^{-1}$) and the total ecosystem ($\text{NEP}=3.19 \text{ Pg C year}^{-1}$ or 11.70

$\text{Gt CO}_2 \text{ year}^{-1}$) during 1980-2016, which is within the range of the newest estimation of forest carbon

sinks (Harris et al. 2021). As the comparisons, we use the aboveground biomass (AGB) from the GLASS

product (a satellite-derived product, <http://www.glass.umd.edu/Download.html>) and the carbon fluxes

from the FluxCom dataset (<https://www.bgc-jena.mpg.de/geodb/projects/Data.php>) to test our predictions

(**Fig. S2** and **Fig. S3**). Both predictions and GLASS observations present the tropical forests own the

highest AGB and the boreal forests had the smallest AGB (**Fig. S2**). In terms of carbon fluxes (i.e. GPP,

ER, and NEP), the resulting spatial pattern is consistent with the FluxCom dataset (**Fig. S3**). However,

the GPP and ER derived from FORCCHN2 for some boreal forests are approximately $0.5 \text{ kg C m}^{-2} \text{ year}^{-1}$

¹ smaller and for parts of eastern North America are approximately $0.5 \text{ kg C m}^{-2} \text{ year}^{-1}$ larger than those

of FluxCom GPP and ER, respectively. Compared to the FluxCom NEP, the model overestimates NEP in

217 some tropical forests and underestimates NEP in some boreal forests.

218 The predicted carbon results including the variables of ‘dayout’ and ‘yearout’ in this case (i.e.,
219 Northern Hemisphere forests) are deposited at an open-access repository (Fang 2022:
220 <https://doi.org/10.6084/m9.figshare.18318722.v1>).

221

222 **6. Conclusions**

223 We develop the FORCCHN2 model and design the corresponding DLL package with the intention to
224 simplify the input and processing of the model and make it more accessible to ecologists interested in the
225 forest ecosystem, climate change, carbon cycle, and modeling. This package provides convenient access
226 and allows high computational efficiency with the Fortran-language-based model predicting the daily
227 dynamics of individual trees. With this new package, we have demonstrated the workflow, functions, and
228 applications of the FORCCHN2 model.

229 In addition, the FORCCHN2 model is tested at 78 flux sites, and then it is applied in predicting the
230 carbon dynamics in the whole Northern Hemisphere forests (1980-2016). Our assessment indicated that
231 FORCCHN2 is able to predict the satisfactory carbon dynamics. While we provided publicly available
232 data in the Northern Hemisphere with 0.5 degrees, our hope is that end-users can offer a wide range of
233 applications and analyses of the FORCCHN2 model, such as providing the new dataset with finer
234 resolution and estimating future changes of forest carbon fluxes. We are also open to further suggestions

235 on enhanced functions that ecologists may find helpful in the subsequent model versions.

236

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242

243 **Authors' contributions**

244 JF planned the project. XY and JF conducted the modeling. XY, JF, YS, and FL contributed to data
245 collection. HHS and JF contributed to data analysis and interpretation of the results. HHS, FL and JF took
246 the lead in writing the manuscript. JF feedback and approval from co-authors. The authors have no
247 conflicts of interest to report.

248

249 **Data availability statement**

250 The source code, instructions, and example run, together with FORCCHN2 DLL package are publicly
251 available via <https://doi.org/10.5281/zenodo.6351153> (Fang et al. 2022). The datasets predicted by
252 FORCCHN2 model include the 37-year (1980-2016) GPP, above- and belowground autotrophic

253 respiration, soil heterotrophic respiration for Northern Hemisphere forests ($0.5^{\circ} \times 0.5^{\circ}$) are publicly
254 available via <https://doi.org/10.6084/m9.figshare.18318722.v1> (Fang 2022).

255

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Table 1. Description of functions and variables in the FORCCHN2 model. A detailed explanation of functions and variables can be found in the FORCCHN2 DLL package documentation. SOS: the start time of leaf growth; EOS: the end time of leaf growth; DOY: day of the year; GPP: gross primary productivity.

Model functions and variables	Description
Time step	Daily and yearly
Initialization data (inputs)	Vegetation: maximum LAI (m ² m ⁻²), forest types, SOS dates (DOY), EOS dates (DOY)
	Soil: field capacity (cm), permanent wilting point (cm), soil volume weight (kg m ⁻³), total organic carbon (kg C m ⁻²), total nitrogen (kg C m ⁻²), silt percent (%), sand percent (%)
	Geography: latitude (°), longitude (°), elevation (m)
Driven data (inputs)	Daily climate: Mean temperature (°C), maximum temperature (°C), minimum temperature (°C), air pressure (hPa), wind (m s ⁻¹), relative humidity (%), precipitation (mm), shortwave radiation (W m ⁻²), CO ₂ concentration (ppm)

Outputs

Daily: aboveground vegetation biomass (kg C m^{-2}),
belowground vegetation biomass (kg C m^{-2}), GPP (kg C m^{-2}), aboveground autotrophic respiration (kg C m^{-2}),
belowground autotrophic respiration (kg C m^{-2}), soil
heterotrophic respiration (kg C m^{-2}), litter-fall (kg C m^{-2}),
soil total organic carbon (kg C m^{-2})

Yearly: same as the daily outputs, with the SOS dates
(DOY) and EOS dates (DOY)

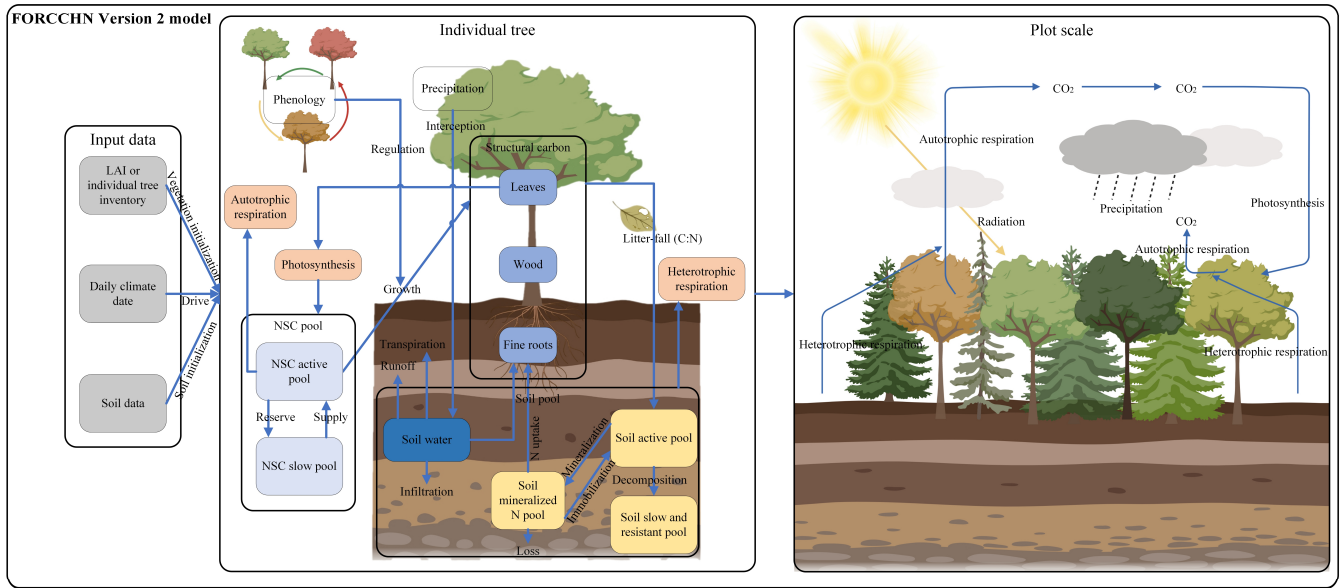


Fig. 1. Schematic representation of the FORCCHN2 model. LAI: leaf area index; NSC: non-structural carbohydrates; C: carbon; N: nitrogen.

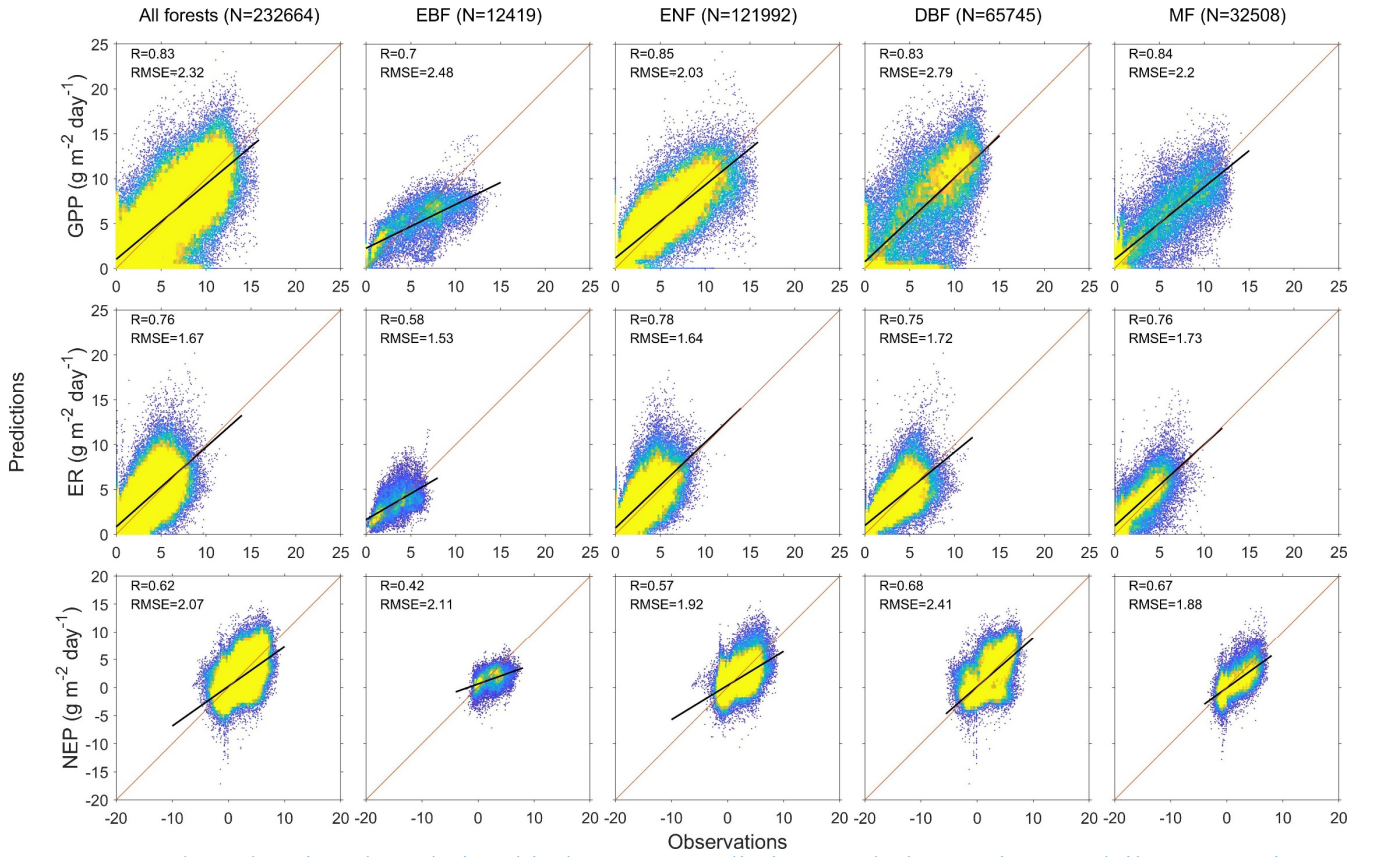


Fig.2. Heat plots showing the relationship between predictions and observations of daily gross primary productivity (GPP), ecosystem respiration (ER), and net ecosystem productivity (NEP) of the studied EC sites. N: the total days of all sites; R: correlation coefficient; RMSE: root mean square error (unit: $\text{g C m}^{-2} \text{ day}^{-1}$). EBF: evergreen broadleaf forest; ENF: evergreen needleleaf forest; DBF: deciduous broadleaf forest; MF: mixed forest. Diagonal lines are 1:1 lines, indicating perfect agreement between predicted and observed fluxes. Black lines represent the linear regression. Colors indicate the percentage of pixels in each bin area (yellow is the densest).

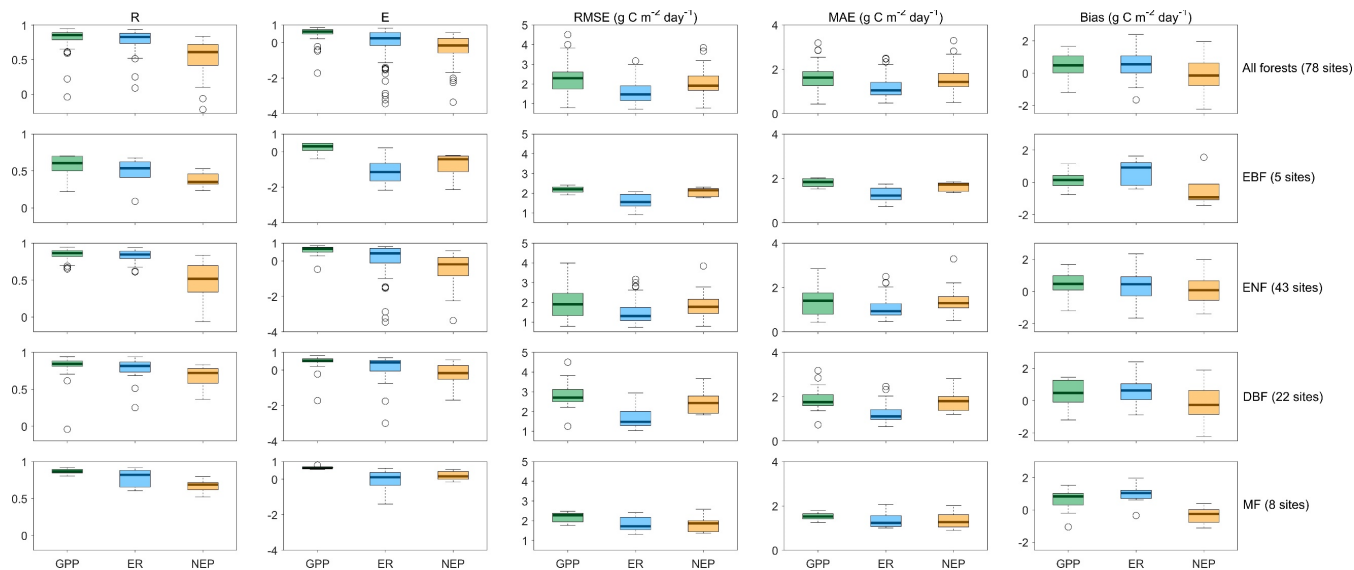


Fig. 3. The statistical results of daily gross primary productivity (GPP, green), ecosystem respiration (ER, blue), and net ecosystem productivity (NEP, tan) observations versus predictions in the studied EC sites. R: correlation coefficient; E: model efficiency; RMSE: root mean square error; MAE: mean absolute error; Bias: bias. EBF: evergreen broadleaf forest; ENF: evergreen needleleaf forest; DBF: deciduous broadleaf forest; MF: mixed forest.

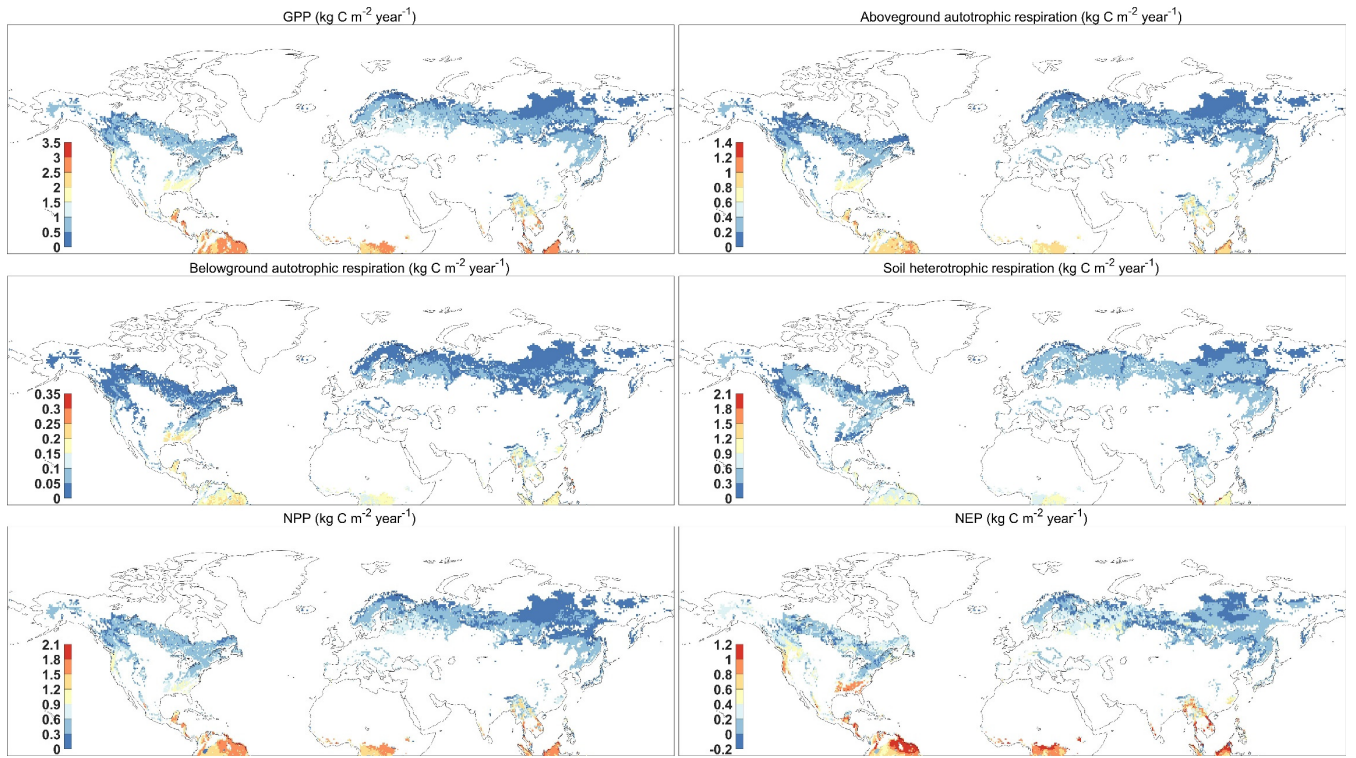


Fig. 4. The spatial distribution of mean GPP (Gross Primary Productivity), above- and belowground autotrophic respiration, soil heterotrophic respiration, NPP (Net Primary Productivity), and NEP (Net Ecosystem Productivity) predicted by the FORCCHN2 model for forest ecosystems of the Northern Hemisphere during 1980–2016. The spatial resolution is $0.5^\circ \times 0.5^\circ$.