

Point-to-point response to reviewers' comments

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1. *I have a concern about the advantage of the new model. Several models have been presented to simulate cropland processes (e.g., DNDC by Li et al., 1992; LPJmL by Bondeau et al., 2007; AgroIBIS by Kucharik et al., 2007; JULES-SUCROP by Van den Hoof et al., 2011). What are the characteristics and advantages of the model presented in this study? Please clarify the point in discussion.*

Response: Thank you for your comments. Indeed, there are a lot of models are developed to simulate the croplands processes. All those models can be grouped into two categories: crop specific models (such as Crop-DNDC, STICS, etc.) and Agro-land surface models (such as, LPJmL, JULES-SUCROP, and ORCHIDEE-crop). ORCHIDEE-crop is designed as an Agro-LSM, so here we discussed the progress of different Agro-LSMs, which can simulate both managed and natural vegetation dynamics interacting with climate change. Some of the existed DGVM crop modes are suffered from any/some of the following short-comings: a). adopted a simplification strategy for representing the growth dynamics of varied kinds of crops in some Agro-LSM (e.g., the CFTs in LPJmL); b). model performance is limited to specific crops (e.g., Agro-IBIS for corn, soybean and wheat specifically) and/or regions (e.g., Agro-IBIS for Northern America specifically); and c). processes of crop growth and carbon allocation is not explicitly simulated but rather with some artificial approximations (e.g., the emergence start of crops and the initial carbon allocations in JULES-SUCROP is artificially fixed and the carbon allocations into of dry matter into different organs are also parameterized with fixed factors within different growth stages). Rather, the ORCHIDEE-crop is developed as an Agro-LSM by adopting a generic framework through integrating the crop processes of STICS into a DGVM model ORCHIDEE. Thus, managed and natural vegetation share the same fundamental biophysical and biochemical functions within the DGVM framework. ORCHIDEE-crop can simulate many different kinds of crops over the global with a generic crop development structure. The crop phenology, developments and carbon allocations during crop growth are with a relatively complete scope. The emergence, growth, maturity and senescence of crops are determined by climate conditions and some kinds of limitations (e.g., nitrogen limitation). The carbon allocations of dry matter into different organs are determined by specific component growth rates, which is dynamically determined by climate conditions and limiting factors. Additionally, crop processes were modelled differently within different DGVM crop models and it is the purpose of some MIP projects to compare model outputs and model skills.

2. *Page 4655 Abstract In abstract, you mentioned about the comparison with ORCHIDEEv196 but not with STICS. Please add some statements.*

Response: In the revision, we add some statements for the comparison between the ORCHIDEE-crop and STICS.

3. *Page 4656 Line 17 “ressources” should be replaced by “resources”.*

Response: Done.

4. *Page 4657 Line 21 “Incomplete” should be replaced by “incomplete”.*

Response: Done.

5. *Page 4662 Line 19 I could not understand the statement “If the NPP available after the grain demand is satisfied is not sufficient to meet the allocation to grain, ...”.*

Response: We checked this sentence and found that the latter “*grain*” should be leaf. Therefore, the sentence has been revised into “*If the NPP available after the grain demand is satisfied is not sufficient to meet the allocation to leaf, ...*”.

6. *Page 4665 Line 5 “long-wave incoming radiation” appears twice.*

Response: we deleted the redundant one.

7. *Page 4668 Line 7 Results section should be, in general, described with the past tense.*

Response: Done.

8. *Page 4672 Line 1 The simplified root distribution could also account, at least partly, for the discrepancy in LE. Is it correct?*

Response: Yes, we agree with you completely. We include your suggestion into our discussion part for explaining the discrepancy in LE simulation.

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1. *The biggest deficiency in the approach is clearly the failure to include a mechanism for irrigation. This is mentioned in the process description only briefly on 4663. It would benefit the paper if this could be expanded somewhat to describe why irrigation was not included. Irrigation is not generally considered to be a difficult thing to include in a process model, with most models using simple rules where additional water is applied if topsoil moisture falls below some threshold. Given that water cycles and especially large-scale irrigation patterns have been shown to have significant impacts on regional climate in several recent studies, this seems like a surprising oversight in what is otherwise a very comprehensive treatment of crops in a land-surface scheme.*

Response: Thank you Joshua for your valuable comments. ORCHIDEE-crop is built by integrating the detailed crop development and carbon allocation processes of STICS, a generic crop model, into the DGVM ORCHIDEE. There are two modules in the standard ORCHIDEE (version Tag196): SECHIBA (simulating energy and water dynamics within the SPA continuum and plant photosynthesis, etc) and STOMATE (simulating carbon allocation, decomposition, vegetation dynamics, etc.). In ORCHIDEE-crop (V0 version), the crop development is driven by the thermal-hydro variables (e.g., soil temperature, water infiltration, soil moisture conditions, etc.) simulated in SECHIBA module. Both managed and natural vegetation share the same thermal-hydro dynamics within the same mosaic (something like a pixel with different vegetation types mixed together). Therefore, if we want to be rigorous we need to change the water and energy processes only on cropland, which is more complicated. It has been done now in a new version but the detailed estimation for that process was out of the main scope of this paper.

2. *I also have some further questions about the root distribution. Root biomass is accrued dynamically but as stated it seems that root depth and distribution is static, with 65% in the top 20cm. Is that correct?*

Response: I agree with you that root dynamic is a key process for better representing the crop development and its interaction with water conditions. The water limitation effects on crop growth is estimated by the mean water condition within the root zone. In the current version, we did not consider the dynamic of root growth, but with a static distribution of root during the whole growing season, with 65% in the top 20 cm.

3. *Since the model considers winter wheat, it would be good to comment on whether/how cold temperature effects (leaf kill and full plant kill) are considered and (in the case where leaves are killed but the plant survives) whether/how this affects allocation during subsequent growth.*

Response: Yes, you are right. Cold temperature during growing season (from

planting to harvest) has important impacts on winter wheat growth. In ORCHIDEE-crop, the effects of cold temperature on winter wheat growth are different in different development stages. From planting to emergence, the cold temperature can impact the crop seeding emergence by both vernalization and thermal accumulation effect. Since emergence, the cold temperature can reduce or even stop (the cold temperature can lead to a no increment in daily LAI) the increment of LAI on a daily step and thus lead to changes in daily leaf growth (leaf growth is determined by both leaf growth increment and daily senescence dynamic) and hence other processes (e.g., photosynthesis, carbon allocations, etc.).

4. *It may not be in the scope of this paper, but I'm certainly very curious to know also how different the new scheme is in terms of large-scale factors that could affect climate feedbacks. Some integrated measure of surface albedo, total carbon budgets, etc. if possible, it would be good to add to the conclusion some statement about this. Does it greatly increase or decrease the land-surface climate feedback in any way that you expect to have implications for future IPSL coupled runs?*

Response: This is really an important issue within the terrestrial-atmosphere interactions. As illustrated in previous literatures that the standard ORCHIDEE simulated crops (C3 and C4) as “super grass”, with same phenology routines as grasses but different photosynthesis abilities. However, in ORCHIDEE-crop we integrated a generic crop module into ORCHIDEE and it simulated the crop phenology, development and carbon allocations based on explicit processes of crop development. The ORCHIDEE-crop can at least improve both the seasonality and magnitudes of LAI for different crops (as illustrated in our manuscript). Thus, ORCHIDEE-crop can improve the simulations of surface roughness, surface albedo, water, energy and carbon budgets for land surface (mixed with both natural and managed vegetation types). Therefore, an improvement in the land-surface climate feedbacks is reasonably expected for the future IPSL coupling.

5. *Small things. Some editorial work is needed to improve readability. For example, page 4659 line 10 “in the crop module same to STICS”.*

Response: We went through the manuscript carefully and made our presentation much clearer.

1

2 **ORCHIDEE-CROP (v0), a new process based Agro-Land Surface**
3 **Model: model description and evaluation over Europe**

4

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43

44 Abstract:

45 The responses of crops to changing climate and atmospheric CO₂ concentration ([CO₂])
46 could have large effects on food production, and impact carbon, water, and energy
47 fluxes, causing feedbacks to the climate. To simulate the responses of temperate crops
48 to changing climate and [CO₂], ~~that~~ which ~~account~~ sing for the specific phenology of
49 crops mediated by management practice, we ~~present~~ describe here the development of
50 a process-oriented terrestrial biogeochemical model named ORCHIDEE-CROP (v0),
51 which integrates a generic crop phenology and harvest module, and a very simple
52 parameterization of nitrogen fertilization, into the land surface model (LSM)
53 ORCHIDEEv196, in order to simulate biophysical and biochemical interactions in
54 croplands, as well as plant productivity and harvested yield. The model is applicable
55 for a range of temperate crops, but ~~it~~ is tested here using ~~for~~ maize and winter wheat,
56 with the phenological parameterizations of two European varieties originating from the
57 STICS agronomical model. We evaluate the ORCHIDEE-CROP (v0) model against
58 eddy covariance and biometric measurements at seven ~~7~~ winter wheat and maize sites
59 in Europe. The specific ecosystem variables used in the evaluation are CO₂ fluxes (net
60 ecosystem exchange (NEE)), latent heat, and sensible heat fluxes. Additional
61 measurements of leaf area index (LAI), aboveground biomass and yield are used as
62 well. Evaluation results revealed that ORCHIDEE-CROP (v0) reproduced ~~s~~ the
63 observed timing of crop development stages and the amplitude of the ~~pertaining~~ LAI
64 changes. This is in contrast to ORCHIDEEv196 in which ~~where~~, by default, crops have

65 the same phenology ~~than as~~ grass. A ~~near~~-halving of the root mean square error ~~for~~
66 LAI from $2.38 \pm 0.77 \text{ m}^2 \text{ m}^{-2}$ to $1.08 \pm 0.34 \text{ m}^2 \text{ m}^{-2}$ ~~was~~ obtained ~~between when~~
67 ORCHIDEEv196 and ORCHIDEE-CROP (v0) ~~were compared~~ across the ~~seven~~7 study
68 sites. Improved crop phenology and carbon allocation ~~lead~~ to a ~~general~~-good match
69 between modelled and observed aboveground biomass ~~{(with a normalized root mean~~
70 squared error (NRMSE) of ~~11.0%–54.2%}, %),~~ crop yield, ~~as well as of the~~ daily
71 carbon and energy fluxes ~~(with a~~ NRMSE of ~~~9.0%–20.1%~~ and ~~~9.4%–22.3%~~ for
72 NEE), and sensible and latent heat fluxes, ~~respectively~~. ~~The simulated yields for~~
73 ~~winter wheat and maize from ORCHIDEE-CROP (v0) showed a good match with the~~
74 ~~simulated results from STICS for three sites with available crop yield observations,~~
75 ~~where the~~ ~~with an average NRMSE was~~ ~~of~~ ~~~8.8%~~. The model data ~~mis~~fit for energy
76 fluxes ~~were~~ ~~are~~ within ~~the~~ uncertainties of the measurements, which themselves showed
77 an incomplete energy balance closure within the range ~~80.6%–86.3%~~. The remaining
78 discrepancies between ~~the~~ modelled and observed LAI and other variables at specific
79 sites ~~were~~ ~~are~~ partly attributable to ~~unrealistic impossible-unrealistic~~ representations
80 of management events ~~from by the model~~. ~~In addition,~~ ORCHIDEE-CROP (v0) ~~is shown~~
81 ~~to have~~ ~~has~~ the ability to capture the spatial gradients of carbon and energy-related
82 variables, such as gross primary productivity, NEE, ~~and~~ sensible ~~heat fluxes~~ and latent
83 heat fluxes, across the sites in Europe, ~~which is~~ an important requirement for future
84 spatially explicit simulations. Further improvement of the model, with an explicit
85 parameterization of nutritional dynamics and ~~of~~ management, is expected to improve
86 its predictive ability to simulate croplands in an Earth System Model.

87

88 Keywords: Crop model, ORCHIDEE-CROP, STICS, Crop yield, Energy balance~~s~~, Net

89 ecosystem exchange~~s~~

90

91 **Introduction**

92

93 Croplands cover ~~–about~~ ~12% of the world land surface (Ramankutty and Foley, 1998),
94 with temporal and spatial variations being subject to population increase, changes in
95 diet, market prices, and other socio-economic factors (IPCC, 2014; Ramankutty et al.,
96 2002; Vuichard et al., 2008). The responses of croplands to climate change ~~are~~ is
97 expected to have significant, but uncertain, consequences for 1) global food production
98 and 2) land surface water, carbon, and energy fluxes, which affects food security as well
99 as regional climate and water resources (Bonan, 2008, 2001; Loarie et al., 2011;
100 Rosenzweig et al., 2014).

101 Along with improving understanding of crop physiology to ~~improve~~ increase
102 production and yield quality, research has focused on investigating the climate impacts
103 on crop functioning by combining historical observations with statistical models
104 (Lobell and Field, 2007; Lobell et al., 2011; Rosenzweig and Parry, 1994) or by running
105 crop models from site to global scales. Impact studies have always pointed ~~to~~ out to
106 ~~at~~ the significant ~~contribution of effect of~~ climate on crop yield variability (Lobell and
107 Field, 2007; Parry et al., 2005; Rosenzweig et al., 2013). However, discrepancies in the
108 response to climate change between different crop models have highlighted the
109 uncertainties that are related to model structure, parameterization, and external drivers
110 (Asseng et al., 2013; Müller, 2011; Rosenzweig et al., 2013).

111 ~~Besides, there~~ There is an increasing need ~~of better~~ to improve understanding of the

112 environmental and climate consequences of changes in cropland area and in
113 management practices, via modification of biophysical and biogeochemical land-
114 atmosphere fluxes (Foley et al., 2011; Lobell et al., 2006; Osborne et al., 2009; Tubiello
115 et al., 2007). ~~Multiple-Many~~ lines of evidence show that changes of cropland plant
116 properties can strongly modify ~~strongly enough~~ the biophysical characteristics (albedo,
117 roughness, turbulent fluxes) of the land surface, ~~which affecting in order to have an~~
118 ~~effect on~~ local and regional climates (Davin et al., 2014; Foley et al., 2011; Georgescu
119 et al., 2009; Loarie et al., 2011; Osborne et al., 2009).

120 Investigation of cropland-climate interactions has led to new model developments ~~to~~
121 that improve Land Surface Models (LSMs) ~~for including also that they give a~~ more
122 realistic representation of crop processes (Bondeau et al., 2007; Gervois et al., 2004;
123 Kucharik, 2003). ~~The,~~ aim ising to simulate the spatial distribution and variability of
124 crop production ~~as well as and their-its~~ water, energy, and carbon fluxes, all of which
125 affect climate. These efforts have improved the seasonal dynamics of modeled foliar
126 and biomass developments (Bondeau et al., 2007; Gervois et al., 2008; Gervois et al.,
127 2004; Kucharik, 2003; Valade et al., 2014; Van den Hoof et al., 2011) and long-term
128 soil carbon changes (Ciais et al., 2011). Despite progress, these “Agro-LSM” models
129 have ~~shown~~ some limitations, such as 1) ~~static~~ or crop/region specific
130 parameterizations (Berg et al., 2011; Kucharik, 2003); 2) ~~idealized~~ representation of
131 different crop types and cultivation practices (Bondeau et al., 2007); and 3) ~~Incomplete~~
132 incomplete coupling between crop growth parameterizations and LSM processes (de
133 Noblet-Ducoudré et al., 2004; Gervois et al., 2004; Valade et al., 2014).

134 In this study, we integrate a generic crop phenology and allocation module ~~based~~
135 ~~on~~from the STICS agronomical model—STICS, which has been extensively validated
136 ~~as a generic crop model and can~~to simulate ~~various kinds of different~~
137 maize, soybean, bananas) (Brisson et al., 1998; Brisson et al., 2002)— into the carbon-
138 water-energy LSM ORCHIDEE model (Krinner et al., 2005), resulting ~~into~~ a new
139 Agro-Land Surface Model, ORCHIDEE-CROP (at version v0, hereafter referred to as
140 ORCHIDEE-CROP, <https://forge.ipsl.jussieu.fr/orchidee/wiki/DevelopmentActivities>).

141 ORCHIDEE-CROP is developed for two applications: offline and online.
142 Offline applications (presented here) are useful to improve understanding of the
143 mechanisms controlling yield, given climate and management forcing. Online
144 simulations require the crop model to be coupled with an atmospheric model (GCM)
145 when studying feedbacks of crop vegetation on climate. There exist
146 Several crop models have been developed for offline applications and impact studies,
147 but very few of these models can be coupled with GCMs, e.g. because they do not
148 represent albedo, roughness, and sensible and latent heat fluxes on the typical time
149 step of ≈ 30 min, which are needed when being required to couple with a GCM.

150 Our efforts have focused on improving the representation of phenology, ~~and~~
151 simulation of biophysical and biogeochemical fluxes, ~~as well as~~ and on biomass and
152 grain yields. ORCHIDEE-CROP can solve the incomplete coupling problems in the
153 existing ORCHIDEE-STICS model (Gervois et al., 2004).

154 ~~In the following, we~~We first describe the structure of ORCHIDEE-CROP (section 2) and
155 evaluate the new model for phenology, ~~and~~ CO_2 , and energy fluxes over winter wheat

156 and maize sites across a large climate gradient in Europe, using observations of
157 biophysical and carbon variables (~~leaf area index~~ [LAI], biomass, latent (LE) and
158 sensible heat (H) fluxes, and net ecosystem exchanges, NEE) from seven eddy
159 covariance sites (section 3). Finally, we discuss the general performance of
160 ORCHIDEE-CROP, ~~and~~ its limitations ~~as well as~~ and the future research that is
161 needed (section 4).

162 **2. Materials and methods**

163 **2.1 Model description**

164 Two key processes of crop plants were introduced into a ~~crop specific~~ module integrated
165 in ORCHIDEEv196 (version Tag196,
166 <http://forge.ipsl.jussieu.fr/orchidee/wiki/Tags/196>, ~~we mention it as~~ called ORCHIDEE
167 hereafter). This module simulates crop phenology and the specific carbon allocation
168 ~~of carbon~~ to grain filling prior to harvest (Fig. 1~~Fig. 1~~Fig. 1). This crop module is used
169 to calculate 1) the seasonal dynamics of LAI, a key variable that impacts surface
170 biophysical properties (albedo, roughness) and water, energy and carbon fluxes, and 2)
171 the timing and amount of grain filling that determines yield.

172 In ORCHIDEE, the vegetation is ~~discretized~~ divided into 13 plant function types -
173 (PFTs), including bare soil, 10 natural PFTs (e.g., evergreen and deciduous trees, C3,
174 and C4 grass) and two crop PFTs (C3 and C4 crops) that are assumed to have the
175 same phenology as ~~than~~ natural grasslands, but with higher carboxylation rates
176 (Krinner et al., 2005). More vegetation types can be simulated using a new PFT

177 external definition module (<http://labex.ipsl.fr/orchidee/index.php/about-orchidee>).

178 Several PFTs can coexist within the same grid cell (also referred to as mosaic

179 vegetation) which can have any size, generally given by the spatial resolution of

180 climate forcing data. All PFTs that co-exist within a grid cell share the same climate

181 forcing but different carbon, energy and water dynamics, due to their specific

182 parameterizations. The sum of fluxes from the different PFT tiles is averaged before

183 being ~~given to~~entered into the atmospheric model, in order to avoid~~ease of~~ coupled

184 simulations.

185

186 **2.1.1 Crop development stages and phenology in ORCHIDEE-CROP**

187 A thermal index (degree-day) adjusted for photoperiodic and vernalization effects

188 according to crop types, controls the developments of temperate crops, such as winter

189 wheat and maize considered here. Seven development stages are sequentially simulated

190 for crop growth and grain filling in the crop module, which is the same as ~~the~~ the

191 processes in STICS ~~{(in detail see~~ Fig. 1 in Brisson et al., 1998). The timing and

192 duration of each stage is calculated based on development units, which describe ~~the~~ ing

193 physiological requirements of crops. These development units are calculated, ~~d~~ just as

194 in STICS, as growing degree days weighted by limiting functions to account for

195 photoperiodism (e.g., winter wheat and soybean) and vernalization (e.g., winter wheat).

196 Vernalization requirement is defined as a given number of vernalizing days (JVC) since

197 the crop germination, and requires a minimum of 7 vernalizing days. The vernalizing

198 value of a given day (JVI) is a function of air temperature. The vernalization status

199 (RFVI) ~~for of~~ the vernalization sensitive crop increases gradually to ~~reach~~ one when
200 the vernalization requirement is met (Supplementary Eqn. 1). The photoperiodic
201 slowing effect, RFPI, is determined by two photoperiod thresholds, PHOBASE and
202 PHOSAT, for photoperiodic crops. In the case of short-day crops, the PHOBASE is
203 higher than PHOSAT, whereas in the case of long-day crops, the PHOBASE is lower
204 than PHOSAT. The current photoperiod PHOI is calculated on the basis of calendar
205 days and latitude (Sellers, 1965) (Supplementary Eqn. 2). Transition between stages
206 occurs when the threshold values of development units are reached, which are specific
207 to different crops or cultivars, but also depend upon management intensity and local
208 climate. Using generic terms for the various ~~stages of~~ plant development stages makes
209 it possible to simulate different kinds of crops if crop-specific parameter values are
210 provided (Bassu et al., 2014; Brisson et al., 2002; Valade et al., 2014).

211 Crop emergence occurs during the sowing-emergence stage, and is divided into ~~a phase~~
212 ~~of~~ seed germination and ~~a phase of~~ epicotyl extension. Germination occurs when the
213 sum of degree-days, using the soil temperature (TSOL) at the sowing depth
214 (PROFSEM), reaches a given threshold (STPLTGER) ~~with a condition on and is~~
215 dependent on soil dryness (Supplementary Eqn. 3). The growth rate of the epicotyl is
216 assumed to be a logistic function that depends ~~ing~~ on soil temperature and water status
217 at the sowing depth (Supplementary Eqn. 4). Crop emergence occurs when the
218 ~~elongation of~~ epicotyl elongates and is dependent on planting depth (PROFSEM).
219 The actual density of emerged plants is calculated from the initial sowing density, a
220 fixed parameter, considering the which takes into account some lack of germination and

221 the death of a fraction of young plants due to unsuitable soil moisture (humectation or
222 drought) and/or to thermal time deficit (Brisson et al., 2008). ~~At~~During this stage, the
223 extremely cold temperatures can reduce the seedling density through its effects on both
224 vernalization and thermal limits for cold-sensitive crops (e.g., winter wheat). From
225 emergence to physiological maturity, the temporal evolution of LAI is calculated in the
226 crop module as the net balance between leaf growth and senescence. The daily growth
227 rate of LAI (ΔLAI) is calculated based on a logistic function of development units
228 (ΔLAI_{dev} , related to different development stages) multiplied by an effective crop
229 temperature, an effective plant density, which takes the inter-plant competition into
230 account, and stress functions (ΔLAI_{stress}) related to water and nitrogen limitations
231 (Supplementary Eqn. 5) (Brisson et al., 1998). The leaf senescence ~~of LAI~~ depends
232 upon the evolution of temperature and leaf lifespan as a function of leaf development
233 and stresses (e.g., water stress). Consequently, leaf senescence ~~of LAI~~ is updated each
234 day (Brisson et al., 2008). Extremely hot and/or cold temperatures during from crop
235 emergence to maturity can affect leaf dynamics through its effects on both the daily leaf
236 growth increment and leaf senescence of crops, and thus exert great large significantly
237 affects on the consequent photosynthesis and carbon allocations.

238

239 **2.1.2 Photosynthesis, carbon allocation and yield**

240 In ORCHIDEE-CROP, photosynthesis is calculated ~~with the equations of~~ using
241 ORCHIDEE (Krinner et al., 2005), which is based on the Farquhar leaf photosynthesis
242 model for C3 crops (Farquhar et al., 1980) and on the model developed by Collatz *et*

243 *al.* for C4 crops (Collatz et al., 1992). In both cases, photosynthetic rate is the minimum
244 of the Rubisco-limited rate ~~for~~ CO₂ assimilation and the electron transport-limited
245 rate ~~for~~ CO₂ assimilation, whose maximal values are the model parameters (V_{cmax}
246 and V_{jmax} , respectively). These two parameters can be calibrated using, for instance,
247 the leaf-level measurements for different kinds of crops and varieties.

248 In ORCHIDEE, the carbon allocation model common to all PFTs is adapted from
249 Friedlingstein *et al.* (Friedlingstein et al., 1999) and accounts for ~~eight~~ biomass
250 compartments (leaves, roots, fruits/harvested organs, reserves, aboveground sapwood,
251 belowground sapwood, aboveground heartwood, and belowground heartwood) for
252 ~~natural~~ trees, and considers ~~five~~ carbon pools for grass and crop PFTs (leaves, roots,
253 fruits/harvested organs, reserves, and aboveground sapwood). The fractions of newly
254 formed assimilates or reserves allocated to these pools are parameterized as a function
255 of soil water content, temperature, light, and soil nitrogen availability.

256 In ORCHIDEE-CROP, we modified the carbon allocation scheme of the two crop PFTs
257 to reconcile the calculations ~~for~~ leaf and root biomass and grain yield (fruits/harvested
258 organs), which are described—driven by the phenology and LAI development
259 parameterizations described in section 2.1.1. Specifically, the daily increment of leaf
260 biomass for crops, Δ_{leaf_m} , is calculated by dividing the daily change ~~of in~~ LAI, Δ_{LAI} ,
261 by specific leaf area (sla), which is weighted by the water and nitrogen stress factors—
262 ~~weighted specific leaf area (sla)~~ (Brisson et al., 2008) as given by:

$$263 \quad \Delta_{leaf_m} = \Delta_{LAI} / sla \quad (1)$$

264 The daily increment ~~for~~ root biomass is determined by the daily total biomass

265 increment and a daily dynamic belowground-to-total biomass partition coefficient,
266 which depends on root development through a normalized root development unit. After
267 the start of the grain filling stage, the ~~quantity of~~ dry matter accumulation in grains is
268 calculated ~~by~~ using a ~~variable~~ “*harvest index*” function that determines the daily
269 fraction of the ~~daily~~ increment ~~for the of~~ total biomass ~~progressively~~ allocated to grain
270 filling. This “*harvest index*” function increases linearly with time from the start of grain
271 filling to the physiological maturity of the crop (when crop is harvested), and is
272 restricted by an upper limit. The effects of extreme temperature ~~impact on~~ the grain
273 filling process ~~and can stop carbon filling of harvested organs as are~~ described ~~by in the~~
274 (Supplementary Eqn. 6) (Brisson et al., 2008). The remaining daily net primary
275 production ~~from ORCHIDEE (NPP)~~, once allocation ~~into~~ leaf, root, and grain biomass
276 is performed (the latter occurring only after the start of the grain filling phase), and is
277 allocated to the stem compartment to conserve mass. ~~In this case,~~ This “residual” stem
278 “residual” compartment denotes ~~in fact~~ both the actual stem biomass and additional
279 reserves. At harvest, a small part of the carbon (with the same amount allotted to planted
280 seeds) is moved from harvested organs to the reserves pool. This mimics the amount of
281 carbon ~~that for~~ seeds needed for the next crop season.

282 In ORCHIDEE-CROP, the ~~priority of~~ carbon allocation priority to different
283 compartments was changed ~~to be so that it was~~ consistent with the growth development
284 phases derived from STICS. In the vegetative stages, the leaf and root have the highest
285 priority. ~~In cases~~ If the ~~net primary product (NPP)~~ supply cannot satisfy the leaf and root
286 biomass demand, no carbon is allocated to stems and the required amount of carbon

287 demanded for leaf and root growth is removed from the reserves. If the extreme case
288 occurs, in which the reserves are not sufficient, the amount of NPP allocated to leaf and
289 root is reduced in ~~the~~ proportion ~~to~~ of the shoot/root ratio (~~yet~~ no carbon being allocated
290 to the stem). However, in such extreme cases, the consistency between LAI and leaf
291 biomass is lost. Conversely, during the reproductive stage, carbon allocation is
292 prioritized to grain filling and leaf biomass, followed by stem and root allocation ~~in~~
293 ~~ease~~ of the remaining NPP. If the NPP available after satisfying the grain demand ~~is~~
294 ~~satisfied~~ is not sufficient to meet support the allocation to the grain ~~leaf~~, then carbon is
295 remobilized from stem and root according to a fixed shoot/root ratio. ~~-(the reserve pool~~
296 ~~was used out before reproductive stages) according to a fixed shoot/root ratio.~~

297

298 2.1.3 Soil moisture limitation effect on plant growth

299 Water limitation for crop development and biomass production is accounted for through
300 a water stress index calculated from ORCHIDEE, and ranges from ~~in the interval~~ ~~[0~~
301 ~~to -1], 1,~~ ~~which is calculated from ORCHIDEE and applied to reduce~~ It allows for
302 ~~reduced~~ ing leaf growth and accelerated ~~ing~~ leaf senescence rates. The root water
303 uptake function in ORCHIDEE is based on the assumption that the vertical root density
304 ~~profile distribution is~~ exponentially decreasing with depth (Krinner et al., 2005) and
305 that water uptake is a function of root zone extractible water weighted by ~~this~~ the root
306 profile. Relative water content in the root zone is an index defined by the difference
307 between actual water content and the wilting point, divided by the difference between
308 field capacity and the wilting point. This index always varies between 0 and 1. Below

309 a fixed relative root zone ~~integrated relative~~ water content threshold of 0.5, the
310 ORCHIDEE stress index value decreases from 1 (no stress) to zero (wilting point). T;
311 ~~and this~~ stress index is used as a multiplier ~~for~~ both V_{cmax} and stomatal
312 conductance, ~~and acting to leads to a~~ decrease in both gross primary productivity and
313 transpiration.

314 Two different soil hydrological schemes, namely a ~~{(the two} 2 layer soil scheme,~~
315 referred to as 2LAY ~~hereafter~~, and ~~the an~~ 11 layer soil diffusion scheme, referred to as
316 11LAY ~~(hereafter, in detail see (Guimberteau et al., 2014))~~ (Guimberteau et al., 2014)
317 ~~can be used alternatively in ORCHIDEE~~ were used in this study to calculate soil
318 moisture, and all dependent ecosystem state variables. In ORCHIDEE-CROP (V0), soil
319 water dynamics between soil hydrology is simulated for three separate soil tiles in each
320 grid cell. These three tiles are covered by bare soil, short vegetation (including crops),
321 and by forest vegetation, respectively. Here, for site-scale simulations, we assumed a
322 grid cell with single tile entirely covered by crops.

323 ~~different soil layers for all PFTs within a vegetation mosaic share the same hydrological~~
324 ~~framework in SECHIBA module but with different parameterizations for different PFTs.~~

325 Relative root extractible soil moisture in the different soil layers ~~is was~~ computed by in
326 each hydrological scheme, as the mean relative soil moisture over the different soil
327 layers, weighted by the fraction of roots within each layer (Krinner et al., 2005). The
328 stress index defined ~~as above~~ was then calculated based on relative root extractible
329 water, which differs between the 2LAY and the 11LAY versions. Application of water
330 irrigation was not taken into account in the current version of ORCHIDEE-CROP

331 ~~and. w~~We keep the default hydrological framework for both natural and crop PFTs as
332 ~~developed in ORCHIDEE (also see the discussion section).~~The typical exponential
333 (~~and~~-static) root profile assumed for grass and crop PFT in ORCHIDEE ~~assumes~~
334 ~~that~~locates ~65% of the roots ~~are above~~in the upper 20 cm ~~of the soil~~. This root
335 distribution profile ~~was~~is different from the one that was used in STICS₂ where ~~only~~
336 fewer roots were assumed ~~to be~~ in the upper 20 cm of soil and more below (Brisson et
337 al., 2008; Gervois et al., 2004). ~~But i~~n ORCHIDEE-CROP we kept~~ep~~ the root profile
338 as parameterized in ORCHIDEE.

339

340 2.1.4 Simplified nitrogen limitation and fertilization effects

341 Nitrogen fertilization ~~allows to~~increases₂ crop productivity and ~~the~~ LAI, which
342 consequently impacts ~~on~~ crop phenology, carbon allocation₂ and turbulent fluxes
343 exchanged with the atmosphere (Mueller et al., 2012). ORCHIDEE-CROP is currently
344 unable to account for dynamic nitrogen stress within the crop growing season due to
345 the lack of an explicit parameterization of nitrogen processes and nitrogen-carbon
346 interactions. We thus defined a simple nitrogen limitation index (*innlai*) ~~and~~ expressed
347 ~~it~~ as a parameter ranging from 0 ~~{(the maximum limitation of nitrogen)}~~ to 1 ~~{(without~~
348 ~~nitrogen limitation)}~~. To account₂ in a very simple manner₂ for the effects of nitrogen
349 fertilization on plant productivity, we introduced_d an additive nitrogen response
350 ~~parameter~~, N_{add} , ~~which is of~~linked to photosynthetic parameters, V_{cmax_opt} and
351 J_{max_opt} , using the following equation:

$$352 N_{add} = 1 + N_{max} - N_{max} \times 0.75^{(N_{fert}/30)} \quad (2)$$

353 ~~w~~Where N_{max} is the maximum additive effects of nitrogen fertilization during the
354 growing season, N_{fert} , on the photosynthetic parameters (~~for~~ details see Chang et
355 al., 2015). The N_{max} is a PFT-specific parameter that can be calibrated by the
356 observed additive nitrogen fertilization effects on plant productivity (e.g., using field
357 trials). This simple function ~~allow~~ed us to estimate the impacts of different levels of
358 nitrogen fertilization on crop productivity (Chang et al., 2015).

359

360 2.2 Simulation set-up

361 2.2.1 Site description

362 We tested ORCHIDEE-CROP ~~using~~ for winter wheat and maize at ~~seven~~ eddy-
363 covariance sites, which are part of the CarboEurope-IP project
364 (<http://www.carboeurope.org/>). ~~The~~se sites span different climatic conditions (Table 1
365 and Fig S1). All the sites recorded the meteorological half-hourly variables necessary
366 to run ORCHIDEE-CROP as well as CO₂ fluxes (NEE), ~~and~~ latent and sensible
367 heat fluxes. The NEE half-hourly data were gap-filled and partitioned into gross
368 primary productivity (GPP) and total ecosystem respiration (TER) using the online
369 eddy covariance processing tool (Moffat et al., 2007; Papale, 2006; Reichstein et al.,
370 2005). Management information (e.g., sowing and harvest date, irrigation and
371 fertilization) and crop development monitoring data ~~{~~(e.g., LAI, aboveground biomass
372 (AGB) and crop yield~~}~~) were available ~~for~~ each site and were used either for
373 parametrization (sowing date, fertilization) or evaluation purposes. The geographic
374 locations, climate regimes, and management informations are provided in Table 1,

375 Table 2, and Fig. S1. More details about the ~~seven~~⁷ sites can be found in (Kutsch et al.,
376 2010; Vitale et al., 2007).

377

378 2.2.2 Climate forcing data and atmospheric CO₂

379

380 At each site, meteorological forcing ~~measured on the top of each flux tower~~ on a half-
381 hour time step, was ~~directly~~ used as ~~a~~ model input. ~~This included~~ air temperature,
382 precipitation, wind speed, atmospheric water vapor pressure, shortwave and longwave
383 incoming radiation, ~~and long-wave incoming radiation~~, mean near-surface atmospheric
384 pressure. Annual CO₂ atmospheric concentration ~~is~~ ~~was~~ ~~prescribed~~ ~~derived~~ from
385 background atmospheric measurements. ~~Because of~~ ~~There were~~ gaps in ~~the~~
386 meteorological data, ~~mainly~~ caused ~~mainly~~ by instrumentation malfunction. ~~Therefore~~,
387 we reprocessed ~~them~~ ~~the data~~ using standardized procedures ~~for~~ gap-filling and
388 quality control (Moffat et al., 2007; Papale, 2006). A significant source of systematic
389 errors ~~when in~~ ~~comparisons~~ ~~between~~ modeled and ~~eddy covariance~~ ~~observed~~ fluxes
390 ~~is~~ ~~were attributed to~~ the lack of energy balance closure ~~in the eddy covariance using~~
391 ~~eddy covariance data~~ ~~measurements~~ (Foken, 2008). Our evaluation revealed ~~an~~ obvious
392 problem regarding the energy balance closure in the eddy covariance observations on
393 these crop sites, ~~where~~ ~~the~~ the energy closure rate rang~~ed~~ ~~ing~~ ~~from~~ ~~~~~ ~~-80.6%~~ ~~to~~ ~~-86.3%~~
394 (e.g., Fig. S2). We ~~thus performed corrections of~~ ~~corrected the~~ daily LE and H
395 measurements ~~in a~~ similar ~~way~~ to Twine *et al.* (Twine et al., 2000) and Jung *et al.* (Jung
396 et al., 2011), which preserved the Bowen ratio:

397 $E_{corr} = \alpha \times E_{uncorr} = (R_n - G)/(H_{uncorr} + LE_{uncorr}) \times E_{uncorr}$ (3)

398 where, E is either the LE or H flux, α is a daily correction factor, and R_n and G ~~is-are~~
399 the net radiation and ~~ground-soil~~ heat ~~storage-flux~~, respectively. In our correction, we do
400 not consider the ~~ground-soil~~ heat ~~storage-flux~~ due to the lack of observations. Although
401 the magnitude and causes of energy ~~balance-budget~~ imbalance ~~likely-probably~~ vary
402 among sites and across time scales (~~Barr et al., 2006; Franssen et al., 2010~~), (Barr et al.,
403 2006; Franssen et al., 2010), ~~this~~ This simplified approach can correct the energy
404 balance closure gap and yields consistent energy fluxes with other independent
405 estimates (Jung et al., 2011).

406

407 **2.2.3 Simulation experiments**

408 A set of simulations were performed for each crop-site (~~in detail see~~ Table 1), using
409 STICS (JavaStics-v11, <http://www6.paca.inra.fr/stics/>), ORCHIDEE, and
410 ORCHIDEE-CROP to evaluate the performance of ORCHIDEE-CROP and the
411 impacts of the parameterizations of the nitrogen limitation factor and soil hydrology
412 schemes, respectively (~~in detail see~~ Table 3). Observed climate data and crop type
413 at each site were used to drive the models (in ORCHIDEE, winter wheat is ~~assigned~~
414 described by the C3 crop ~~standard-standard~~ parameters and maize by the standard C4
415 crop ones). The same mean soil depth and soil water holding capacity were prescribed
416 for the seven7 sites, and were averaged from the Harmonized World Soil Database
417 (HWSD), [http://webarchive.iiasa.ac.at/Research/LUC/External-World-soil-](http://webarchive.iiasa.ac.at/Research/LUC/External-World-soil-database/HTML/)
418 [database/HTML/](http://webarchive.iiasa.ac.at/Research/LUC/External-World-soil-database/HTML/)). ~~At each site with rotation~~ For each site, we selected one year of

419 observation during which winter wheat or maize was cultivated. The sowing date was
420 ~~prescribed to~~inputted into the model for each crop-site according to the management
421 data (Table 2). ~~However, but~~ the harvest date in ORCHIDEE-CROP was ~~calculated~~
422 ~~by the model~~determined by crop development processes. The observed nitrogen
423 fertilization and irrigation information for each crop-site were used in STICS
424 experiment STI-WN (Table 2 and Table 3). In STICS, the real date and quantity of
425 applied irrigation and nitrogen fertilization can be introduced into the model, which-
426 ~~and are involved~~affects into the water balance and nitrogen transformation modules,
427 respectively, ~~from the irrigation and fertilization calendar~~ (Brisson et al., 2008).

428 All simulations based on ORCHIDEE and ORCHIDEE-CROP started from an
429 equilibrium state of carbon pools ~~with where the~~ climate was obtained ~~with using a~~
430 model spin-up. For this spin-up, site-specific meteorological ~~1/2~~half-hourly data was
431 repeatedly cycled for 300 years to force ORCHIDEE and ORCHIDEE-CROP until the
432 soil water reached a steady state (data not shown). Then, simulations were conducted
433 for the period of evaluation, starting ~~with starting from the~~ initial conditions at the end
434 of model spin-up. Notably, C input from manure ~~input applications was~~ not taken into
435 account in this study, due to a lack of data for historical manure applications.

436 The same cultivar choice (represented by the ~~parameters of~~ “Soissons” and “DK250”
437 varietyies parameters in STICS for winter wheat and maize, respectively), rather than
438 site-year specific varieties ~~choice~~, ~~choice~~ was made ~~in the model~~ at all sites for winter
439 wheat and maize, ~~respectively~~ (see Table 3). This may lead to some discrepancies
440 between simulated and observed values, but our main purpose was to evaluate the

441 improvements achieved by ORCHIDEE-CROP in a generic way, without having to
442 calibrate the model for each site. Sensitivity tests were conducted to evaluate the effects
443 of nitrogen limitation and water stress on crop development, carbon, and energy
444 ~~balances budgets~~. The ~~experimental detailed ensemble of details~~ ~~experiments~~ ~~are~~
445 shown in Table 3.

446

447 2.3 Metrics for evaluating model performance

448 Three metrics were used to evaluate the model-data agreements. ~~These were with~~
449 ~~available observations~~ at a daily resolution for ~~different fluxes~~ (NEE, H, and LE fluxes,
450 and ~~the~~ LAI, AGB, and grain yield biometric variables, ~~at the different crop sites~~
451 ~~where these observations are available~~.

452 First, we calculated the index of agreement (IOA) (Willmott et al., 1985), given by

$$453 \quad \text{IOA} = 1.0 - \frac{\sum_{i=1}^n (O_i - P_i)^2}{\sum_{i=1}^n (|P_i - \bar{O}| + |O_i - \bar{O}|)^2} \quad (4)$$

454 where P_i is ~~the~~ modelled data, O_i is ~~the~~ observed data, \bar{O} is ~~the~~ observed mean and
455 n is the number of ~~data~~. The IOA, with values ranging from 0.0 to 1.0, is more
456 sensitive than correlation-based metrics to differences in the observed and modelled
457 means and variances (Willmott et al., 1985).

458 We also calculated the Pearson's product-moment correlation coefficient for ~~different~~
459 ~~all~~ sites. This metrics estimates the proportion of total variance in the observed data that
460 can be explained by model, ~~and is~~ given by

$$461 \quad r = \frac{\sum_{i=1}^n (P_i - \bar{P})(O_i - \bar{O})}{\sqrt{\sum_{i=1}^n (P_i - \bar{P})^2} \sqrt{\sum_{i=1}^n (O_i - \bar{O})^2}} \quad (5)$$

462 where ~~P_i is modelled data, O_i is observed data, \bar{P} is the modeled mean, \bar{O} is~~
463 ~~observed mean, and n is the number of data.~~

464 Third, the root mean square error (RMSE) and normalized root mean square error
465 (NRMSE) were used to quantify the model-observation agreement in absolute terms,
466 expressed as:

$$467 \text{ RMSE} = \sqrt{\sum_{i=1}^n (P_i - O_i)^2 / n} \quad (6)$$

$$468 \text{ and NRMSE} = \sqrt{\sum_{i=1}^n (P_i - O_i)^2 / n} / (O_{\max} - O_{\min}) \quad (7)$$

469 where P_i and ~~O_i is modelled and observed data, respectively, and n is the number of~~
470 ~~data.~~ O_{\max} and O_{\min} are observed maximum and minimum data.

471

472 3. Results

473 3.1 Crop phenology, plant development stages and productivity

474 Comparison of the seasonal evolution of observed and modelled LAI for winter wheat
475 and maize at different sites ~~is iswas~~ shown in ~~Fig. 2~~[Fig. 2](#). The modelled
476 seasonality ~~for~~ LAI ~~was~~has been markedly improved by ORCHIDEE-CROP (ORC-
477 CP1, Table 3) compared ~~with to~~ ORCHIDEE, for both winter wheat and maize. The
478 correlation coefficient between observed daily LAI and modelled daily LAI ~~showed a~~
479 ~~marked~~ ($p < 0.05$) ~~increas~~ed from ~~ORCHIDEE to ORCHIDEE-CROP (ORC-CP1)~~
480 ~~from~~ 0.44 ± 0.22 to 0.83 ± 0.17 for winter wheat and from 0.64 ± 0.22 ~~to~~s 0.79 ± 0.10
481 for maize ~~using from ORCHIDEE and to ORCHIDEE-CROP (ORC-CP1), respectively.~~
482 The IOA ~~increas~~ed from 0.47 ± 0.11 to 0.82 ± 0.12 (winter wheat) and from $0.57 \pm$

483 0.15 to 0.85 \pm 0.08 (maize), with a significant decrease ~~in~~ RMSE (2.71 \pm 0.49 vs.
484 1.12 \pm 0.36 and 2.06 \pm 0.86 vs. 1.04 \pm 0.31 for winter wheat and maize, respectively)
485 (~~Fig. 2~~~~Fig. 2~~~~Fig. 2~~, Table 4, ~~Figs. 5a–b~~). Despite its overall good performance for LAI,
486 ORC-CP1 (under moderate nitrogen limitation ~~to~~~~of~~ leaf growth) ~~could~~ ~~cannot~~
487 reproduce ~~the amplitude of the~~ observed LAI within the measurement uncertainty
488 (personal communications with PIs ~~in 2014~~) at a few sites (~~Fig. 2~~~~Fig. 2~~~~Fig. 2~~). For
489 example, maximum LAI ~~was~~ underestimated by 49% and 28% for winter wheat ~~at~~
490 FR-Gri and FR-Lam, respectively. Reducing the nitrogen limitation ~~for~~~~of~~ leaf growth
491 (ORC-CP3) at these two sites ~~could~~~~can~~ improve the modelled maximum LAI and bring
492 it ~~into~~ agreement with the observations (Fig. S3, Table 4). The modelled growing season
493 length (defined as the period ~~going~~ from crop sowing to harvest) by ORC-CP1 for all
494 crop ~~sites~~ ~~was~~ in good agreement with the observations (~~with~~-IOA = 0.96 and RMSE
495 = 25.4 days) (~~Fig. 3~~~~Fig. 3~~~~Fig. 3~~).

496 ~~Along with the an~~The accurately simulated timing and amplitude of LAI ~~improved~~, the
497 seasonal evolution of aboveground biomass (AGB) ~~was~~ ~~got~~ ~~ets~~ ~~improved~~ in
498 ORCHIDEE-CROP (ORC-CP1) compared ~~with~~~~to~~ ORCHIDEE for both winter wheat
499 and maize, except at BE-Lon for winter wheat and at NL-Lan for maize (Fig. 4, ~~Fig.~~
500 ~~5~~~~Fig. 5~~~~Fig. 5~~). In general, the bias of ~~the~~ modelled AGB ~~was~~ attributable to the bias
501 ~~in the~~ modelled LAI, as indicated by ~~the~~ a significant ($p < 0.005$) relationship between
502 ~~them~~ ~~AGB~~ and LAI for all ~~crop~~ sites (Fig. S4). However, the daily change rate of
503 above-ground biomass in the late growing season between the start of grain filling and
504 yield harvest ~~was~~ systematically and significantly ($p < 0.05$) underestimated for both

505 winter wheat (change rate of AGB underestimated by 36%—74%) and maize (~~by~~ 18%—
506 -70%), especially at the sites where LAI ~~is was~~ underestimated (e.g., winter wheat at
507 FR-Gri and FR-Lam) (Fig. 2, Fig. 4, Fig. S5). ~~In the~~ The reality observation data did not
508 show a; ~~the~~ decrease in above-ground biomass ~~does did not start~~ until harvest (Fig. 4
509 Fig. 4).

510

511 ORCHIDEE-CROP (ORC-CP1) ~~couldan~~ capture the timing of grain filling and yield
512 harvest well compareding to the observations and STICS simulations (Fig. S6).
513 Comparisons of modelled and observed crop yields for winter wheat and maize in FR-
514 Aur and FR-Lam showed ed that there was around a ~19%-% to 30% underestimation of
515 crop yields in ORC-CP1 without fertilization (Fig. 6~~Fig. 6~~), compareding to a
516 good match (NRMSE ~~=~~ ~8.8%) between STICS with real fertilization (STI-WN) and
517 the observed dataations (Fig. S6). However ORCHIDEE-CROP with real fertilization
518 (ORC-CP4) ~~can~~ could produce a better estimation of crop yields for these two sites than
519 ORCHIDEE-CROP without fertilization (ORC-CP1), leading to a ~~~50%~~ reduction in
520 the NRMSE (47% vs. 23% for ORC-CP1 and vs. ORC-CP4, respectively) (Fig. 6).

521 Considering the measurement uncertainties of FR-Aur and FR-Lam for crop yields
522 (personal communications with PIs in 2014), ORCHIDEE-CROP, with the-its simple
523 nitrogen fertilization, parameterization, ~~on crop productivity~~ generally conserves
524 conserved showed reasonable performance compared to STICS, ~~that~~ because it which
525 has has a full fledge complete nitrogen cycle, ~~that~~ and captures both the timing and
526 amplitude of crop yields.

527

528 3.2 CO₂ and energy fluxes

529 ORCHIDEE-CROP had a more realistic simulated seasonality and amplitude for
530 NEE at most of the winter wheat sites than ORCHIDEE (significant increase in IOA
531 and r and decrease in RMSE from $2.9 \pm 0.2 \text{ g C m}^{-2} \text{ day}^{-1}$ in ORCHIDEE to 1.9
532 $\pm 0.5 \text{ g C m}^{-2} \text{ day}^{-1}$ in ORC-CP1). Improved performances of ORCHIDEE-CROP
533 over ORCHIDEE were also found at the maize sites in humid regions (Fig. S1, Fig.
534 Fig. 7). Along with leaf area development (LAI) during the growing season, the
535 model produced a CO₂ sink until shortly before harvest, when most leaves
536 were senescent and crop photosynthesis could not compensate for respiration,
537 which was consistent with the observed data (Fig. 7).
538 ORCHIDEE-CROP could also capture the observed peak in CO₂ release to
539 atmosphere shortly (ranging from 10 to 20 days, Fig. 7) after harvest for both winter
540 wheat and maize, which was mainly due to increased litter
541 decomposition.

542

543 However, there was a mismatch between the simulations and observations regarding
544 the temporal evolution of NEE for winter wheat in BE-Lon, where there was a
545 weaker and earlier termination of CO₂ uptake in the model (Fig. 7). The
546 underestimated LAI and earlier cessation of crop growth in ORC-CP1 at this site
547 resulted in a negative bias for GPP during the late growing season (~170 days after
548 sowing) (Fig. 2, Fig. S7), which contributed to the

549 underestimation of NEE uptake ~~during the same period~~ (Fig. 7 Fig. 7 Fig. 7, Fig. S8).
550 Notably, ORC-CP1 ~~overestimates overestimated~~ the NEE peak uptake of CO₂ for maize
551 at sites with drier climates ~~regimes~~ in Europe (e.g., FR-Lam and IT-Bci). The
552 overestimation of NEE at these summer-dry sites ~~is was~~ probably ~~mainly~~ (~68%–85%
553 of explained variance ~~as~~ revealed by ~~the a~~ Generalized Linear Model) caused by ~~the~~
554 an overestimation of GPP rather than by an underestimation of ecosystem respiration
555 in ORC-CP1 (Fig. S7, Fig. S8). Further analysis ~~shows showed~~ a much higher ($p < 0.05$)
556 rate ~~for~~ GPP per unit LAI in ORC-CP1 than observ~~ations~~~~ed~~ at these southern
557 ~~European~~ maize sites (Fig. S9). Notably, ~~the~~ ORCHIDEE-CROP with the 11-LAY
558 hydrological scheme (ORC-CP5) ~~improves improved~~ the modelled NEE ~~largely~~ for
559 maize at these sites; ~~with a~~ because it showed a ~40% decrease in the NRMSE (Fig. 7).
560 Despite the improved seasonality of H for most of the crop-sites over Europe (Fig. S10),
561 ORCHIDEE-CROP ~~combined~~ with the 2LAY hydrological scheme generally
562 ~~overestimates overestimated~~ H for winter wheat sites, especially in the early- and mid-
563 growing season (from sowing to ~160–200 days after sowing) and ~~shows showed~~ a
564 more realistic simulation of H for maize sites (NRMSE of ~9%–13%). The
565 overestimation of H at wheat sites ~~occurs occurred~~ during the early- and mid-growing
566 season (Fig. 8) when the plants were grow~~ing~~ng slowly with a low canopy cover, ~~and it~~
567 ~~is was~~. This could be partly attributed to the underestimation of soil water content in
568 the top soil during that period (data ~~not~~ shown) or to the insufficiently deep roots
569 prescribed in the model. Notably, the ORC-CP5 with the 11LAY soil hydrological
570 scheme, which ~~has had~~ a more realistic representation of soil water infiltration after rain

571 and ~~allows allowed to~~could simulate the vertical profile of soil moisture with
572 ~~desiccation~~desiccation of the surface soil during dry episodes, ~~improves~~ improved the
573 simulation of H during this period, with the NRMSE being brought down from ~~~7%~~
574 10% in ORC-CP1 to ~~~5%~~~8% in ORC-CP5 (Fig. 8). Notably, however, the 11LAY
575 hydrological scheme usually ~~overestimates~~ overestimated the bare soil evaporation
576 (data not shown), which ~~will~~ would result in drier top soil conditions and lead to a
577 higher H, ~~which. This can~~ could partially explained the residual overestimation of H,
578 even in ORC-CP5 (Fig. S10).

579 Consistent with the overestimation of H in ORC-CP1, ~~ORCLE was~~ CP1 generally
580 ~~underestimates~~ underestimated LE ~~among at the~~ wheat sites (Fig. 9). A more realistic
581 estimation of LE was ~~observed~~ obtained by in ORC-CP5 for a majority of the ~~crop~~ sites
582 ~~simulations than ORC-CP1~~, showing a ~32% decrease in NRMSE from ORC-CP1 to
583 ORC-CP5. ~~The eE~~ Exceptions were the winter wheat and maize simulation ~~on at the~~
584 DE-Kli site, which could be attributed to a considerable energy balance gap (with an
585 energy closure of ~73%) in at this site (Fig. 9). For ~~the~~ maize, ~~simulation at DE-Kli,~~
586 ORC-CP5; ~~overestimates~~ overestimated the LE at DE-Kli; ~~for by~~ ~110% compared to
587 ~~with the observed data.~~ ations. The LE values were also overestimated At this site, was,
588 ~~while ORC-CP5 also overestimates~~ overestimated the LE for wheat at DE-Kli during
589 the early- and mid-growing season (from sowing to 230 days ~~since~~ after sowing). The
590 overestimation of LE at DE-Kli this site for both winter wheat and maize is was not
591 neither likely to not raise be explained from by the the simulated a bias of LAI bias
592 (good estimation, see above) nor ~~from by the~~ a systematic error in LE due to the effects

593 of rainfall events (with daily rainfall ≥ 3 mm) (Figs. 8–9), but was possibly due to
594 some other factors, such as soil water holding capacity ~~since in our study we used the~~
595 ~~same mean value among different sites despite the great difference~~. The slightly
596 negative (~~–16% of RMSE~~) bias of in LE simulated in by ORC-CP5 at the wheat site
597 FR-Lam during the peak leaf growth (~~during 210–250 days after planting~~) is was due
598 to an underestimation of the LAI (Fig. 9, Fig. 2). The slight overestimation of – LAI for
599 maize during periods of peak leaf growth (e.g., FR-Lam and NL-Lan) ~~does did~~ not
600 ~~appear however to~~ translates into a related overestimation of LE. This illustrates
601 illustrated the divergent responses of LE to changes of in LAI between ORCHIDEE-
602 CROP and the observations, which can could be due to several factors, such as the
603 parameterization of soil water stress (Fig. S11). The episodes of LE with low biases
604 (during LE peaks of LE) ~~are were symmetrical to episodes of~~ coincided with high H
605 biases, even though net radiation appears to be realistic, except for the maize site IT-
606 Bci in Italy (Fig. S12).

607

608 ORCHIDEE-CROP ~~also has had good ability could also to~~ capture the spatial
609 gradients of carbon and energy fluxes across different crop – sites in Europe. There are
610 were significant correlation coefficients between the observed and modelled GPP, NEE,
611 H₂ and LE data, with r ranging from 0.75 – to 0.90. Evaluation of IOA ~~reveals revealed~~
612 a generally good agreement between the observed and modelled GPP, NEE, H₂ and LE
613 data, with IOA ranging from 0.70 – to 0.90 (Fig. 10, Fig S14–S16).

614

615 4. Discussion

616 4.1 General performance of ORCHIDEE-CROP

617 ~~The ORCHIDEE-CROP has been developed here as an Agro-LSM by adopting and~~
618 ~~adopts a generic framework through to integrate the crop processes from STICS~~
619 ~~into the DGVM model ORCHIDEE LSM. Managed and natural vegetation share the~~
620 ~~same fundamental biophysical and biochemical functions within the DGVM~~
621 ~~framework. Given its generic structure, ORCHIDEE-CROP, tested in this study~~
622 ~~focusing wheat and maize in this study, can simulate many different kinds of other crop~~
623 ~~types over the global with a generic crop development structure. The crop phenology,~~
624 ~~developments, carbon allocations and crop grain filling are with a relatively complete~~
625 ~~scope and simulated explicitly driven by can be calculated from climate~~
626 ~~conditions forcing data, include and is mediated by limiting factors (e.g., nitrogen,~~
627 ~~extreme temperatures, and low soil moisture).~~

628 ~~Marked A significant improvement is was achieved obtained by using in~~ ORCHIDEE-
629 CROP ~~comparing with to~~ ORCHIDEE for the simulated ~~timing and amplitudes of~~
630 ~~plant developments (crop phenology and development) for winter wheat and maize at~~
631 different ~~winter wheat and maize sites investigated in Europe. It showed showing~~
632 ~~agreements with observations within 65%–95% (IOA) for biometric data and 78%–~~
633 98% (IOA) ~~agreement with the observed data~~ for all turbulent fluxes, despite the lack
634 of detailed crop management (e.g., irrigation, fertilization) parameterization (Figs. 2–
635 9), and the lack of an explicit calculation ~~for the of~~ nitrogen cycle in the croplands.

636 Remarkably, ORCHIDEE-CROP has a good ability to reproduce the observed spatial

637 gradients ~~foref~~ carbon and energy fluxes across different climate zones in Europe, even
638 using a fixed variety parameter setting ~~among for~~ different sites. ~~This,~~ impl~~ied~~ying that
639 these spatial gradients in biophysical and biochemical variables ~~should beare~~ mainly
640 ~~driven by~~ climate ~~driven~~ rather than ~~by~~ crop variety.

641 Improvements ~~inof~~ crop phenology and carbon allocation lead to a general good match
642 of the seasonality between modelled and observed AGB (with NRMSEs of 11% ~~–~~54%),
643 ~~crop yields, and, as well as~~ carbon and energy fluxes (NRMSEs of ~~~~~9.0% ~~–~~20.1% and
644 ~~~~~9.4% ~~–~~22.3% for NEE and sensible and latent heat fluxes, respectively). Comparisons
645 between the 2LAY and 11LAY hydrological schemes revealed that the 11LAY
646 hydrological scheme can improve the modelling of soil water dynamics and hence lead
647 to a better simulation of leaf growth and ~~the consequently~~ biochemical and biophysical
648 variables, especially for the C4 crops planted in ~~the~~ drier climate zones of Europe (Fig.
649 ~~7–9),). which This~~ in turn exerts great effects on the estimations of carbon balances in
650 these regions, especially in ~~the~~ context of the projected increasing climate variability
651 and extremes (e.g., heat waves and drought events) (Beniston et al., 2007; Ciais et al.,
652 2005; Stocker et al., 2013). Yet, parameterization of ~~the~~ water stress also depends on
653 the ~~profile distribution~~ of active roots, which is considered as fixed in all ~~the versions~~
654 ~~of ORCHIDEE versions. Such~~ The use of a static root profile is one limit on theation to
655 calculation ofe water stress, but the use of a 11-Layer hydrology allows to us to simulate
656 shifts in root uptake from the surface to deeper horizons as the soil dries out during
657 drought. distribution scheme in ORCHIDEE CROP could undoubtedly add some bias
658 in the simulation of crop growth, water and energy (e.g., the latent heat) fluxes. An

659 important ~~direction task for future improvement area for further research would could~~
660 be ~~to have~~ a more mechanistic parameterization of the root profile in the model.

661 Notably, the simple function of ~~the~~ additive nitrogen fertilization on crop productivity
662 can lead to better agreement between the observed and modelled crop yields in
663 ORCHIDEE-CROP, ~~showing which showed~~ a ~50% decrease in ~~the~~ NRMSE (Fig. 6).

664 The remaining discrepancies ~~in of~~ simulated crop yields, and energy fluxes are generally
665 within the observed uncertainties ~~for of~~ measurement and energy ~~balance~~ closure. More
666 importantly, ORCHIDEE-CROP has ~~good the~~ ability to capture ~~the~~ spatial gradients of
667 crop-related fluxes variables, such as GPP, NEE, H_e and LE_e , across the studied sites in
668 ~~the~~ different European climate zones ~~of Europe~~ (Fig. 10, Figs. S14–S16). This is
669 important for further applications of this model using gridded data over Europe, or even
670 ~~the~~ globe, ~~to when attempting to~~ investigate regional/global yield variations, and the
671 interactions between croplands and ~~the~~ climate system. ~~C, while the croplands has have~~
672 ~~the~~ potentially crucial climate feedbacks ~~from regarding the increasing increased~~
673 intensification of agricultural activities ~~and as well as~~ land use changes (Pitman et al.,
674 2009; Ramankutty et al., 2002; Sacks and Kucharik, 2011).

675 Failure of the model to ~~capture~~ the peak LAI at some crop ~~sites~~ (e.g., winter wheat
676 at FR-Gri and FR-Lam) under ORC-CP1 is ~~at~~ partly attributed to ~~the~~ simplified
677 representation of nitrogen limitation on crop growth and fertilization effects (~~in detail~~
678 ~~see~~ section 2). Alleviation of nitrogen limitation on leaf growth ~~on at these those~~ sites
679 can improve the simulated amplitudes of LAI and capture the maximum LAI (Fig. S3).

680 ~~Actually, n~~Nitrogen limitation has a strong influence on the seasonal evolutions of crop

681 growth (Fig. S3), ~~and a~~ more realistic representation of intra-seasonal nitrogen
682 processes (results based on STICS with an explicit nitrogen cycle) leads to a generally
683 much better match between the modelled and observed LAI, except ~~for the~~ NL-Lan ~~for~~
684 and maize (Fig. S13).

685 ~~Lack of~~ The failure to modelling of the irrigation effects ~~ss~~ — can also ~~contribute~~ introduce
686 some bias to the simulated LAI and other variables. Soil water stress on GPP and LE,
687 which also impact ~~ing~~ affects carbon allocation ~~as well~~, plays an important role in
688 controlling crop developments, especially for summer crops (e.g., maize) planted in
689 regions with dry summer episodes (Fig. S1, Table 1), ~~where~~ Those regions are
690 currently suffering from intensive irrigation managements ~~currently~~ (Table 2) and
691 ~~possibly there will possibly be an~~ increase ~~of in~~ irrigation requirements as the climate
692 warm ~~along with the climate warming~~ (Döll, 2002). As illustrated by our results ~~that~~
693 the lack ~~ing~~ of irrigation managements in the current version of ORCHIEE-CROP leads
694 to a lower LAI~~s~~ in the later crop season at FR-Lam for maize in drier climate zones
695 (Fig. 2 and Fig. 7), ~~which~~ in turn ~~in turn~~ affect NEE and the energy budget (Fig. 7–9). More
696 importantly, the projected increas ~~ed~~ ing drought stress for ~~current~~ cultivated croplands
697 (Dai, 2012), ~~with a more intense and longer lasting droughts~~ in drier climate zones
698 (Davin et al., 2014; Trenberth et al., 2014), ~~—~~ challenges the representations of soil
699 hydro-logical processes and their interactions with other factors for existing Agro-
700 LSMs.

701

702 4.2 Model limitation and uncertainty

703 ~~I~~The irrigation (as discussed above) effect~~s~~ on the crop developments and yields are
704 not accounted for in ~~this study~~the current version of ORCHIDEE-CROP, ~~and it is~~
705 ~~yet~~but it is important ~~for when attempting to~~ investigate~~ing~~ the historically long-term
706 changes ~~in~~ crop yields ~~during over~~ recently ~~past~~ decades, as ~~the~~ intensive human
707 management has tended to s-occur~~mainly~~ since approximately the middle of the 20th
708 century.

709 Several studies have shown that the spatial differences in crop managements contribute
710 significantly to the tempo-spatial patterns of crop yields (Licker et al., 2010; Lobell and
711 Field, 2007), ~~besides as well as~~ the impacts of climate and soil fertility (Rosenzweig et
712 al., 2013). Adaptive improvements in agricultural managements are regarded as a
713 potential way to close the “yield gaps” in a relative~~ly~~ sustainable manner ~~for social-~~
714 ~~environmental system~~ (Licker et al., 2010). How the model handles human
715 management factors (e.g., irrigation and fertilization) and their interactions with
716 changing CO₂ and climate variations could have significant impacts on the ~~simulations~~
717 ~~of crop evolutions and~~ production simulations and the consequent land surface carbon
718 budgets (Prescher et al., 2010). Additionally, our current crop development module
719 embodies a number of simplifications for ~~the~~ pests, diseases, and weeds, which ~~are we~~
720 assumed to be controlled. ~~Besides, the e~~Extreme soil conditions (e.g., high salinity or
721 acidity) are also crudely assumed to ~~exert~~have little effects on crop growth. ~~T~~All these
722 factors can also introduce great uncertainties into the biophysical and biochemical
723 simulations ~~over of~~ croplands.

724 Therefore, explicit nutrition dynamics and a human management (e.g., irrigation,
725 fertilization, ~~application-introduction~~ of new crop varieties, and pest management, etc.)
726 module ~~are with primary priority~~ need to be included in the updated version of
727 ORCHIDEE-CROP to improve our ability to understand and project the roles of
728 croplands in food security, environmental footprints and ecosystem services in response
729 to climate change.

730

731

732 **5. Conclusions**

733 ORCHIDEE-CROP, by integrating a generic process-based crop development and yield
734 harvest module into a generic LSM--ORCHIDEE program, allow us to assess the
735 spatial and temporal dynamics of the important biophysical and biochemical
736 interactions within the soil-vegetation-atmosphere continuum for temperate crops.
737 Comprehensive evaluations show thea generally good performance of ORCHIDEE-
738 CROP at predictingin crop phenology, productivity, as well asand the biosphere-
739 atmosphere carbon and energy exchanges in pan-Europe temperate crop sites covering
740 different climate zones, even without the explicit human management module. It
741 bBenefitsing from the a generic strategy in the crop module, which makes ORCHIDEE-
742 CROP can be widely applicable at the regional and global scale. Explicit
743 parameterizations of crop development processes in ORCHIDEE-CROP can improve
744 the simulations of both the seasonality and magnitudes of LAI for croplands, which in
745 turn affectsaffect the consequent surface roughness, surface albedo, water, energy, and

746 ~~carbon budgets for land surfaces. Therefore, Moreover, w~~with respect to future climate
747 change, ORCHIDEE-CROP will allow us ~~not only~~ to predict the footprints of climate
748 variations ~~in on~~ food security, ~~but also to and to~~ simultaneously account for feedbacks
749 ~~caused by~~ changes in crop behaviors to the atmosphere by coupling ~~it to~~ a general
750 atmospheric circulation model (e.g., LMDz).

751 Nevertheless, ~~a~~ further improvement, especially ~~the with regards to~~ explicit nutritional
752 dynamics and human management, is ~~with a~~ primary priority and could significantly
753 improve our ability to understand and predict the role of croplands in the biosphere-
754 atmosphere continuum, in ~~the~~ context of the increasing global demand for food and the
755 urgent requirement to reduce the environmental footprints (Godfray et al., 2010;
756 Mueller et al., 2012).

757

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763

764 **7. Code availability**

765 The ORCHIDEE-CROP is still undergoing development, especially for human
766 management processes, and the code is modified frequently. Therefore, the codes are

767 not ready for fully public access. However, the source codes of ORCHIDEE-CROP at
768 an early version (V0) can be requested from Xiuchen Wu (xiuchen.wu@bnu.edu.cn) or
769 Nicolas Vuichard (nicolas.vuichard@lsce.ipsl.fr).

770

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982

983 **Tables**

984

985 Table 1. Basic geography and climate information for different crop sites.

986

Crop type	SiteID	Country	MAP [*]	MAT [†]	Longitude	Latitude	Altitude (m)	KGCC [‡]
Winter wheat	FR-Lam	France	702	12.55	1.24	43.49	180	Cfb
	FR-Gri	France	579	11.5	1.95	48.84	125	Cfb
	FR-Aur	France	700	12.9	1.11	43.55	242.5	Cfb
	DE-Kli	Germany	674	7.1	13.52	50.89	478	Cfb
	Be-Lon	Belgium	800	10	4.74	50.55	165	Cfb
Maize	FR-Lam	France	702	12.55	1.24	43.49	180	Cfb
	FR-Gri	France	700	11.5	1.95	48.84	125	Cfb
	DE-Kli	Germany	674	7.1	13.52	50.89	478	Cfb
	NL-Lan	Netherland	786	9.8	4.9	51.95	-0.7	Cfb
	IT-Bci	Italy	900	15.5	14.96	40.52	20	Csa

987

Note:

988

* MAP: mean annual precipitation;

989

† MAT: mean annual temperature;

990

‡ KGCC, the Koppen-Geiger climate classifications.

991

992 Table 2. Management information for different crop-sites.

993

Crop type	SiteID	Year (sowing)	Sowing date		Irrigation (mm)				Fertilization (Kg N/ha)			
Winter wheat	FR-Lam	2006	291	0	0	0	0	0	46.5 (8 Jan 2007)	48.2 (4 May 2007)	\	\
	FR-Gri	2005	301	0	0	0	0	0	55.0 (15 Mar 2006)	55.0 (14 Apr 2006)	\	\
	FR-Aur	2005	300	0	0	0	0	0	50.0 (25 Jan 2006)	40.0 (23 Mar 2006)	33.5 (12 Apr 2006)	\
	DE-Kli	2006	269	0	0	0	0	0	74.3 (8 Apr 2007)	53.8 (4 May 2007)	35.8 (4 Jun 2007)	43.1 (22 Jun 2007)
	Be-Lon	2006	286	0	0	0	0	0	40.0 (17 Mar 2007)	60.0 (12 Apr 2007)	94.5 (8 May 2007)	\
Maize	FR-Lam	2006	121	25.0 (13 Jun 2006)	33.0 (3 Jul 2006)	27.8 (15 Jul 2006)	18.0 (26 Jul 2006)	44.0 (10 Aug 2006)	91.0 (8 Jun 2006)	\	\	\
	FR-Gri	2005	129	0	0	0	0	0	140.0 (9 May 2005)	\	\	\
	DE-Kli	2007	118	0	0	0	0	0	17.3 (22 Apr 2007)	67.2 (13 Jun 2007)	\	\
	NL-Lan*	2005	138	0	0	0	0	0	\	\	\	\
	IT-Bci	2004	129	21.8 (24 Jun 2004)	27.2 (2 Jul 2004)	20.3 (15 Jul 2004)	25.7 (18 Jul 2004)	23.4 (20 Jul 2004)	22.5 (8 May 2004)	142.0 (11 Jun 2004)	\	\
			22.1 (27 Jul 2004)	19.3 (31 Jul 2004)	22.9 (5 Aug 2004)	22.1 (12 Aug 2004)	15.0 (21 Aug 2004)					

994 Note: * There is strong organic fertilization. \ indicates no fertilization records.

995

996

997 Table 3. Description of the ensemble of simulations.

998

Name of experiments	Description of experiments	Irrigation	Nitrogen processes *		Soil water scheme †	Stlevdrp (GDD) ¶	Strpmat (GDD) !!
STI-NN	STICS without fertilization during crop development [†]	NO	DY	\		540/990	750/600
STI-WN	STICS with actual fertilization based on management records [†]	NO	DY	\		540/990	750/600
ORC-ST0	Standard version of ORCHIDEE without crop development module, no fertilization	NO	NO		LAY2	540/990	750/600
ORC-CP1	ORCHIDEE-CROP with moderate nitrogen limitation, no fertilization	NO	NO, innlai = 0.5		LAY2	540/990	750/600
ORC-CP2	ORCHIDEE-CROP with high nitrogen limitation, no fertilization	NO	NO, innlai = 0.2		LAY2	540/990	750/600
ORC-CP3	ORCHIDEE-CROP with low nitrogen limitation, no fertilization	NO	NO, innlai = 0.9		LAY2	540/990	750/600
ORC-CP4	ORCHIDEE-CROP with moderate nitrogen limitation, real fertilization	NO	ND, innlai = 0.5		LAY2	540/990	750/600
ORC-CP5	Same to ORC-CP1, but with 11 layer soil hydrological scheme, no fertilization	NO	NO, innlai = 0.5		LAY11	540/990	750/600

999 Note:

1000 * DY, with dynamic nitrogen processes, NO, without nitrogen processes, ND, without dynamic nitrogen processes but with a simplified additive
 1001 nitrogen response of crop productivity to fertilization. For ORCHIDEE-CROP, we introduced a fixed nitrogen limitation factor for leaf growth
 1002 (innlai, ranging 0.0-1.0) during the whole crop growing season.

1003 † Two soil hydrological schemes (the 2 layer soil scheme, referred as 2LAY, and the 11 layer soil diffusion scheme, referred as 11LAY, in detail
 1004 see Guimberteau *et al.*, (2014)) are available in ORCHIDEE and ORCHIDEE-CROP.

1005 ¶ The accumulated growing degree days (GDD) from crop emergence to start of grain filling for winter wheat (C3 crop) and grain (C4 crop),
 1006 respectively.

1007 !! The accumulated growing degree days (GDD) from start of grain filling to crop mature for winter wheat (C3 crop) and grain (C4 crop),
 1008 respectively.

1009 || JavaStics (v11.0) used here was obtained from <http://www6.paca.inra.fr/stics>.

1010 ! The detailed crop managements for each crop-site were shown in Table 2.

1011

1012

1013 Table 4. Comparisons between observations and different simulations.

1014

Crops	SiteID	IOA				R				RMSE (m ² m ⁻²)				NRMSE (%)			
		ORC-ST0	ORC-CP1	ORC-CP2	ORC-CP3	ORC-ST0	ORC-CP1	ORC-CP2	ORC-CP3	ORC-ST0	ORC-CP1	ORC-CP2	ORC-CP3	ORC-ST0	ORC-CP1	ORC-CP2	ORC-CP3
Winter wheat	BE-Lon	0.37	0.65	0.52	0.63	0.15	0.92**	0.98***	0.73	3.30	1.53	1.78	1.74	93.52	52.81	61.41	60.14
	FR-Lam	0.48	0.88	0.67	0.88	0.30	0.79*	0.83**	0.86**	2.68	0.90	1.48	1.21	60.72	20.44	33.52	27.56
	FR-Gri	0.66	0.87	0.63	0.97	0.74	0.96**	0.92*	0.97**	1.86	1.34	2.45	0.73	30.44	22.01	40.09	11.93
	FR-Aur	0.40	0.95	0.77	0.75	0.51	0.95**	0.91*	0.89*	3.06	0.52	0.85	1.58	107.47	18.42	29.84	55.61
	DE-Kli	0.46	0.74	0.56	0.62	0.49	0.55	0.47	0.56	2.68	1.31	1.17	2.07	101.02	49.26	44.06	77.96
Maize	DE-Kli	0.65	0.89	0.64	0.81	0.77	0.80*	0.74	0.89*	1.66	1.05	1.94	1.90	35.78	22.62	41.93	40.94
	FR-Lam	0.50	0.86	0.69	0.57	0.92*	0.76*	0.88*	0.55	2.46	1.00	1.31	2.58	74.95	30.42	40.08	78.75
	FR-Gri	0.58	0.96	0.64	0.91	0.45	0.95**	0.92**	0.97***	2.04	0.68	2.07	1.34	44.69	14.86	45.32	29.43
	NL-Lan	0.77	0.80	0.63	0.39	0.80	0.71	0.83*	0.45	0.79	0.89	1.34	2.52	24.82	27.98	42.28	79.18
	IT-Bci	0.38	0.74	0.49	0.73	0.42	0.70*	0.84*	0.65	3.37	1.60	2.62	1.98	85.37	40.59	66.33	50.11

1015

1016 Note: IOA, index of agreement; R, Pearson's product-moment correlation coefficients; RMSE and NRMSE are the root mean square error and
 1017 normalized root mean square error, respectively.

1018 *, ** and *** indicates statistically significant at 5%, 1% and 1‰ level, respectively.

1019

1020 **Figure captions**

1021

1022 Figure 1. Model structures of the ORCHIDEE-CROP. The crop development module
1023 ~~is~~ (based mainly on STICS, (Brisson *et al.*, 1998)) is integrated into the STOMATE
1024 module of ORCHIDEE (Krinner *et al.*, 2005). The crop development module simulated
1025 the phenology, developments and grain yields for crop PFTs. ORCHIDEE-CROP
1026 consists in the coupling of two modules. SECHIBA simulates the vegetation
1027 photosynthesis, water and energy budgets, STOMATE is a carbon module and
1028 calculates carbon allocation in different carbon pools and fluxes to the atmosphere.

1029

1030

1031

1032 Figure 2. Temporal changes of daily leaf area index (LAI) since planting from
1033 observations (green dots), standard ORCHIDEE (ORC-ST0, grey line) and
1034 ORCHIDEE-CROP (ORC-CP1, orange line). The upper and lower panel shows the
1035 results for different sites of winter wheat and maize, respectively.

1036

1037

1038 Figure 3. Comparisons of the observed and modelled (ORC-CP1, in detail see Table 3)
1039 growing season lengths (from sowing to maturity) for winter wheat and maize in

1040 different sites. Different colors indicate data for different crop-sites.

1041

1042

1043 Figure 4. Comparisons of the observed (green dots) and modelled daily aboveground
1044 biomass from ORCHIDEE-CROP (ORC-CP1, orange line) and ORCHIDEE (ORC-
1045 ST0, grey line) for winter wheat and maize in different sites. The upper and lower panel
1046 shows the results for different sites of winter wheat and maize, respectively.

1047

1048 Figure 5. Scatter plots of the modeled (ORC-CP1, in detail see Table 3) and observed
1049 daily LAI and aboveground biomass (AGB) for different sites of winter wheat (a) and
1050 c)) and maize (b) and d)), respectively. The units for RMSE of LAI and AGB are $m^2 m^{-2}$
1051 2 and $g C m^{-2}$, respectively. Different colors indicate different crop-sites with red, orange,
1052 light green, green and dark green for winter wheat (-W) at BE-Lon, DE-Kli, FR-Aur,
1053 FR-Gri and FR-Lam, respectively, and with light blue, medium blue, blue, purple and
1054 violet for maize (-M) at DE-Kli, FR-Gri, FR-Lam, IT-Bci and NL-Lan, respectively.

1055

1056

1057 Figure 6. Comparisons of the observed (blue bars) and modelled (green bars for ORC-
1058 CP1 and brown bars for ORC-CP4, see Table 3) harvested crop yields in different sites
1059 for winter wheat a) and maize b).

1060

1061 Figure 7. Temporal changes of daily net ecosystem exchanges (NEE) derived from
1062 observations (black line) and ORCHIDEE-CROP (ORC-CP1, blue line; ORC-CP5,
1063 brown line) since planting. The green and blue stems represent the fertilization (kg N
1064 ha⁻¹) and irrigation (mm) events during the selected growing season. The dotted orange
1065 line indicates the harvest date since planting. The upper and lower panel shows the
1066 results for different sites of winter wheat and maize, respectively.

1067

1068 Figure 8. Comparisons between the observed (black line) and modeled daily sensible
1069 heat fluxes (H) from ORCHIDEE-CROP (ORC-CP1, blue line; ORC-CP5, brown line)
1070 for different crop-sites. The grey stems represent the relative large rainfall events (with
1071 daily summed rainfall ≥ 3 mm) during the modelled growing season. The upper and
1072 lower panel shows the results for different sites of winter wheat and maize, respectively.

1073

1074

1075 Figure 9. Same to Figure 8 except for latent heat fluxes (LE).

1076

1077 Figure 10. Comparisons between the observed and modelled (based on ORC-CP5)
1078 mean growing season GPP among different crop sites for winter wheat (circle, -W) and
1079 maize (cross, -M). Different colors indicate different sites.

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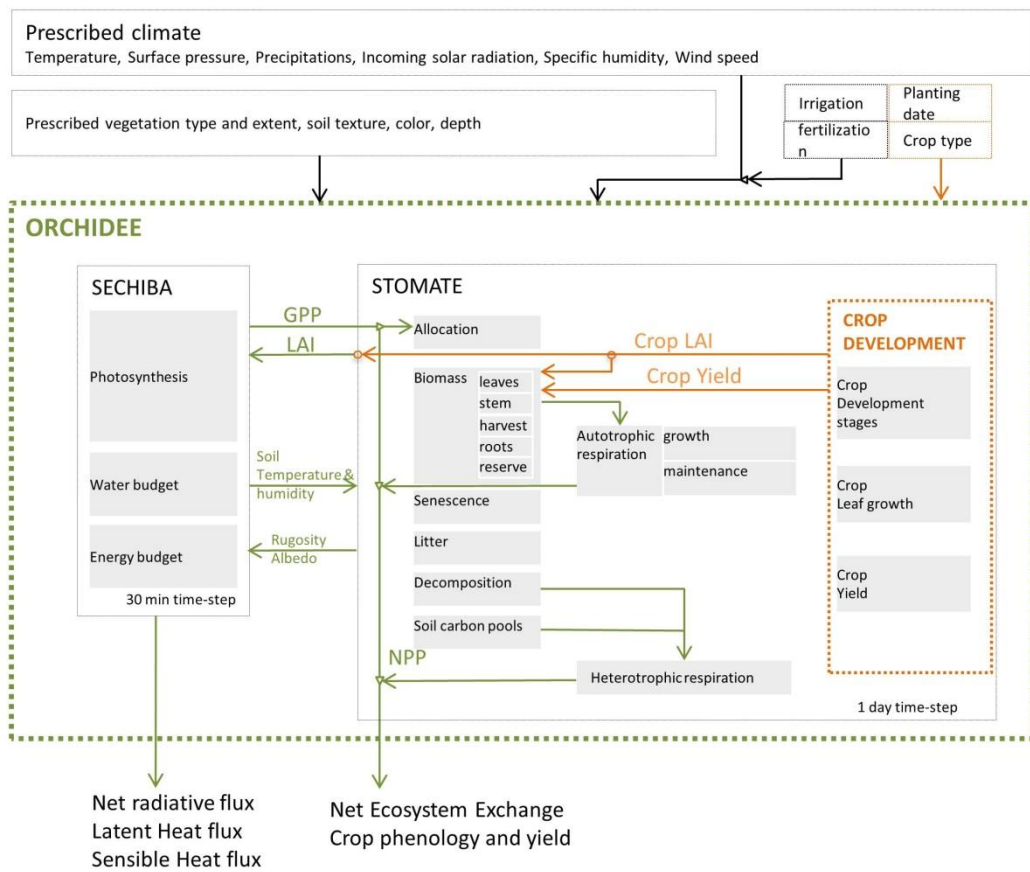
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1083 **Figures**

1084

1085 Fig. 1

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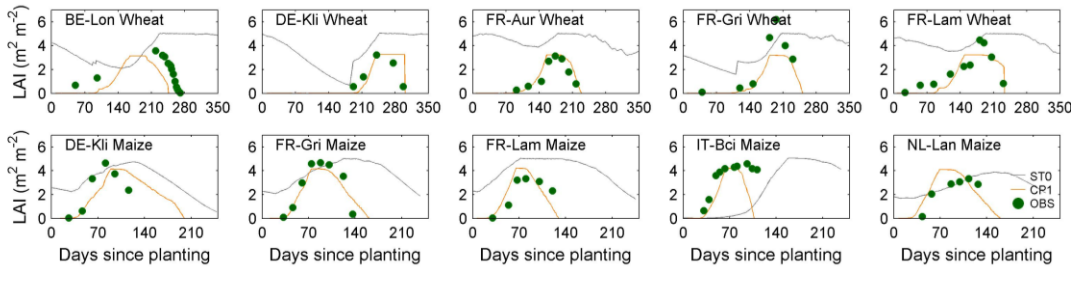
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1090 Fig. 2

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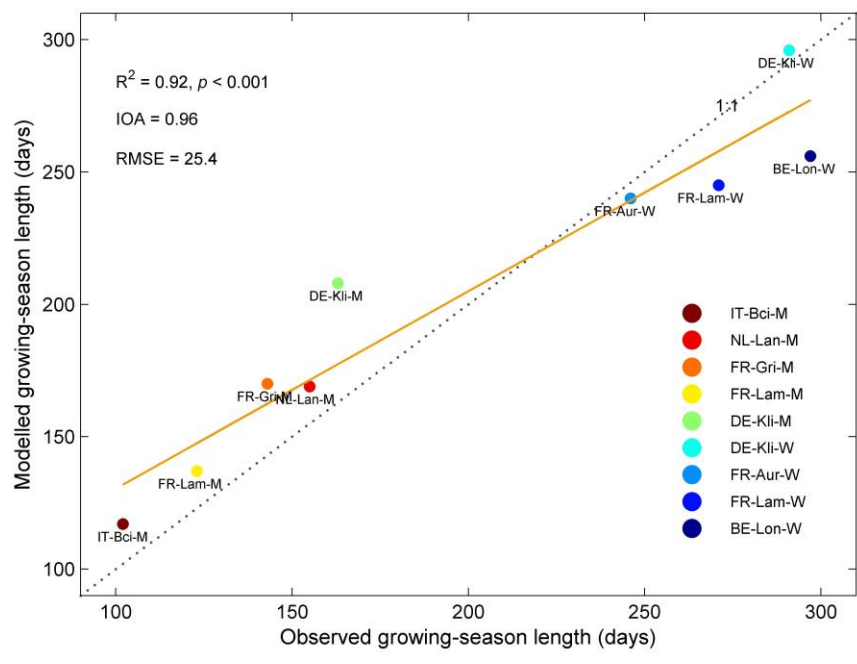
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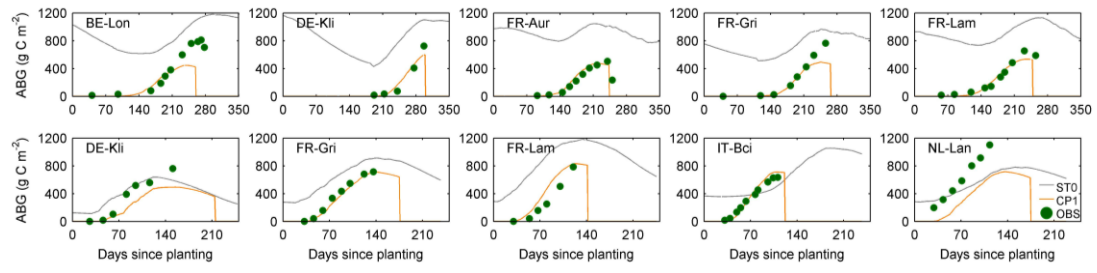
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1105 Fig. 4

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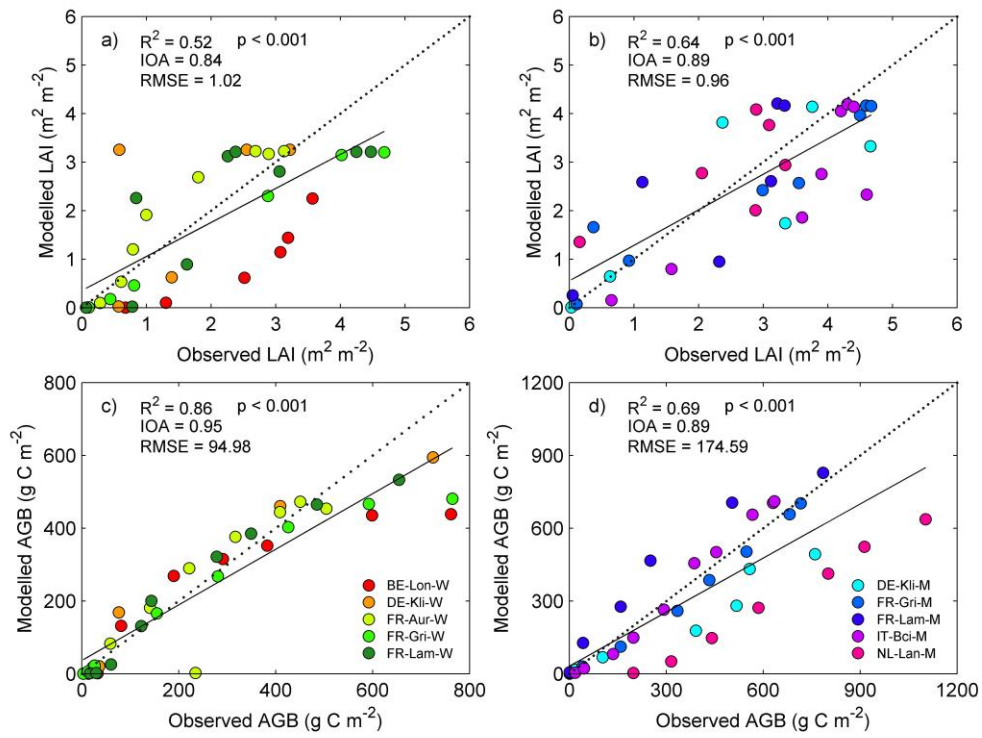
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1111 Fig. 5

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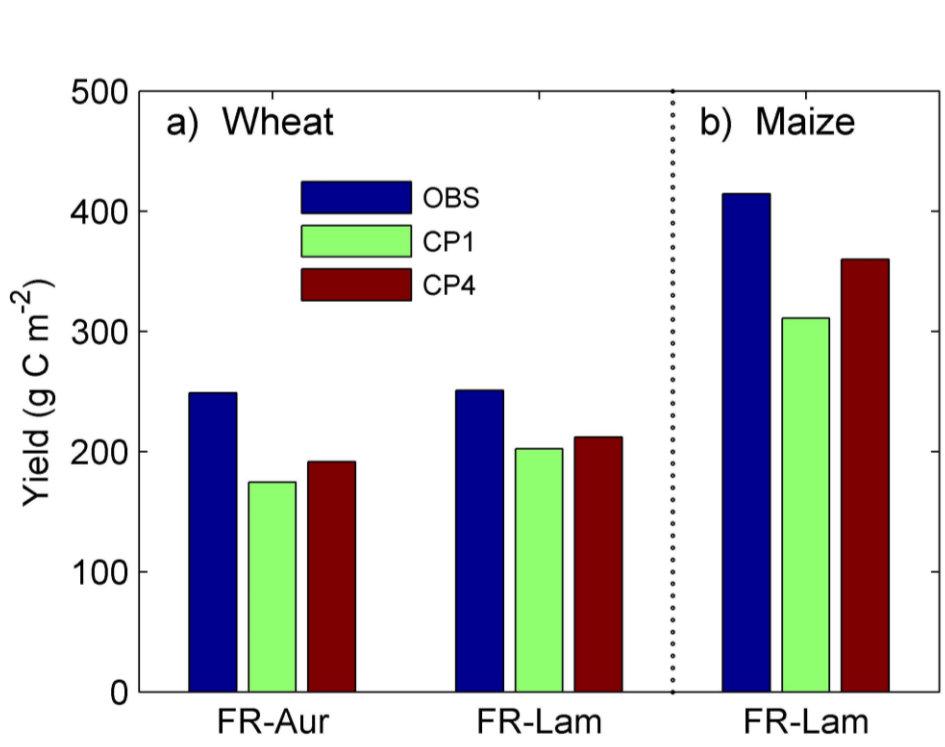
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1119 Fig. 6

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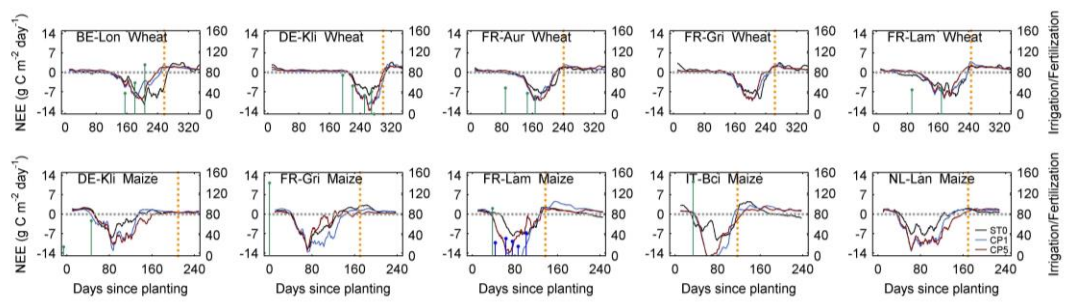
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1126 Fig. 7

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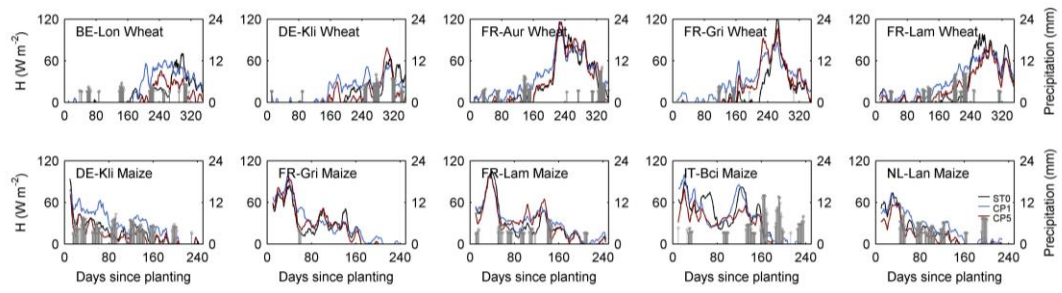
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1132 Fig. 8

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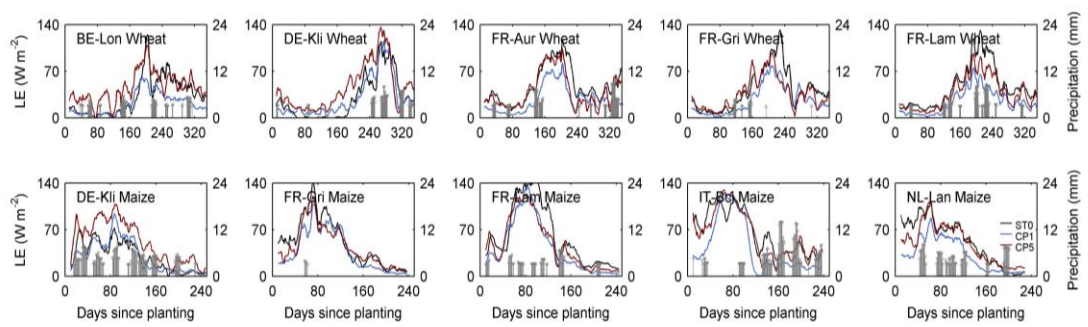
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1142 Fig. 9

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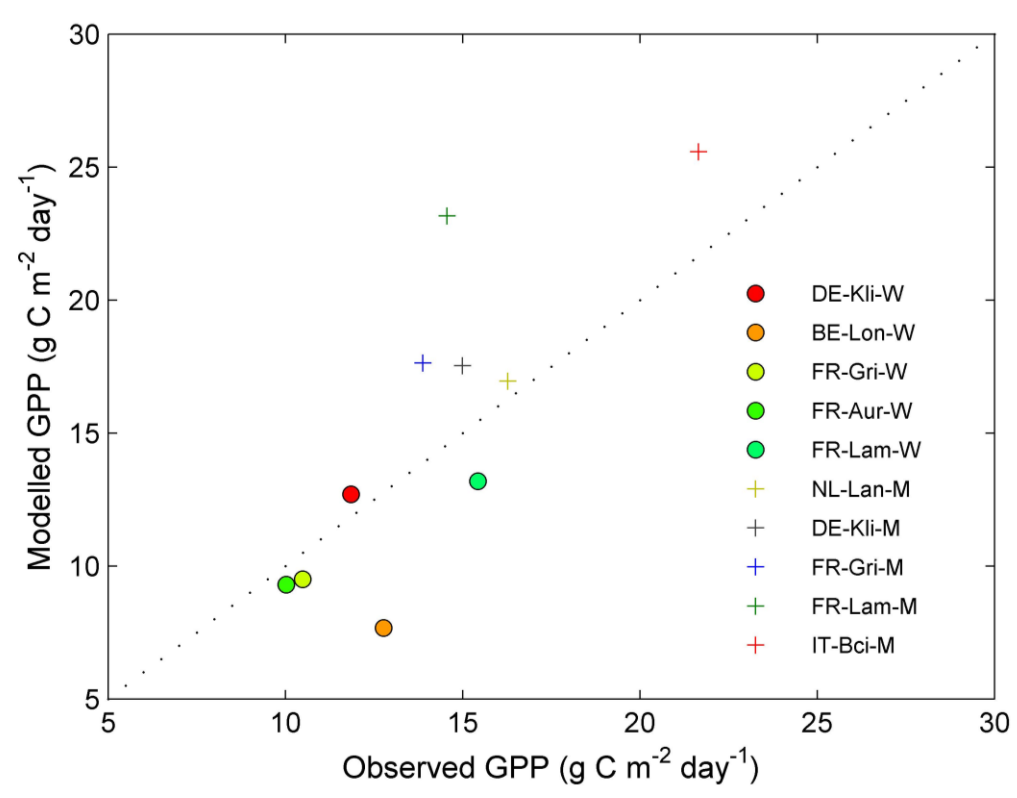
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1148 Figure 10.

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