

Modelling forest management within a global vegetation model-part 2: model validation from a tree to a continental scale

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19 Abstract

20 The construction of a new forest management module (FMM) within the ORCHIDEE 21 global vegetation model (GVM) allows a realistic simulation of biomass changes during 22 the life cycle of a forest, which makes many biomass datasets suitable as validation data 23 for the coupled ORCHIDEE-FM GVM. This study uses three datasets to validate 24 ORCHIDEE-FM at different temporal and spatial scales: permanent monitoring plots, 25 yield tables, and the French national inventory data. The last dataset has sufficient 26 geospatial coverage to allow a novel type of validation: inventory plots can be used to 27 produce continuous maps that can be compared to continuous simulations for regional 28 trends in standing volumes and volume increments. ORCHIDEE-FM performs better than 29 simple statistical models for stand-level variables, which include tree density, basal area, 30 standing volume, average circumference and height, when management intensity and 31 initial conditions are known: model efficiency is improved by an average of 0.11, and its 32 average bias does not exceed 25%. The performance of the model is less satisfying for 33 tree-level variables, including extreme circumferences, tree circumference distribution 34 and competition indices, or when management and initial conditions are unknown. At 35 the regional level, when climate forcing is accurate for precipitation, ORCHIDEE-FM is 36 able to reproduce most productivity patterns in France, such as the local lows of 37 needleleaves in the Parisian basin and of broadleaves in south-central France. The 38 simulation of water stress effects on biomass in the Mediterranean region, however, 39 remains problematic, as does the simulation of the wood increment for coniferous

40	trees. These pitfalls pertain to the general ORCHIDEE model rather than to the FMM.
41	Overall, with an average bias seldom exceeding 40%, the performance of ORCHIDEE-FM
42	is deemed reliable to use it as a new modelling tool in the study of the effects of
43	interactions between forest management and climate on biomass stocks of forests
44	across a range of scales from plot to country.
45	Keywords: model validation; global vegetation model (GVM); ORCHIDEE-FM; forest
46	management; carbon cycle;

48 **1** Introduction

49 Global vegetation models (GVMs) simulate fluxes of carbon, energy and water in 50 ecosystems at the global scale, generally on the basis of processes observed at a plant 51 scale. Despite their correct ability to simulate hourly local (e.g., at flux tower sites) and 52 global seasonal to interannual (e.g., compared with atmospheric CO_2 observations) 53 variations in CO₂ fluxes, these models usually fall short of simulating biomass and soil 54 carbon pool dynamics within ecosystems (Desai et al., 2007; Viovy et al., in prep.; in 55 revision). This shortcoming has been attributed to forest age structure and 56 management, which are not simulated by most GVMs (Ciais et al., 2008). ORCHIDEE-FM, 57 a new GVM with an explicit representation of forest management practices typical of 58 European forests (Bellassen et al., this issue), addresses this challenge, but it has yet to 59 be validated against independent datasets. 60 Several of the many variables processed by GVMs can be measured and, thus, used for 61 validation: for example, leaf area index (Demarty et al., 2007), light absorption and light 62 use efficiency (Jung et al., 2007), carbon stocks (Masek and Collatz, 2006), 63 evapotranspiration (Thornton et al., 2002), and latent and sensible heat fluxes 64 (Abramowitz et al., 2008). However, validation exercises for GVMs most frequently 65 focus on carbon fluxes estimated with eddy-covariance techniques (Thornton et al., 66 2002; Krinner et al., 2005; Turner et al., 2005; Schaefer et al., 2008).

67 Flux towers have two major strengths: the flux data that they deliver have a very fine 68 resolution in time, often half-hourly, and their reasonably large footprint of 69 approximately 100 ha (Nagy et al., 2006; Reichstein et al., 2007) averages the variability 70 due to individual trees. With regard to GVM validation, they also have two major 71 drawbacks. First, their costly structure and maintenance limits their numbers: large 72 networks such as FLUXNET manage to have good coverage of the different land-use 73 types and climates (Baldocchi et al., 2001), but they seldom provide clear insights on 74 inter-site variability within a given climate and land-use type. Therefore, it is difficult to 75 generalise measurements that could be heavily influenced by local conditions (e.g., soil 76 fertility or hydrological parameters) or management (e.g., recent thinning) (Lindner et 77 al., 2004). Second, because eddy-covariance technology is quite recent, the time-series 78 are seldom longer than one decade (Urbanski et al., 2007). This precludes the validation 79 of full stand rotations, which commonly last between 100 and 200 years, on temperate 80 forest ecosystems (Bottcher et al., 2008) unless flux towers are smartly distributed to 81 measure chronosequences (Amiro et al., 2006).

Thus, in their review of terrestrial carbon models, Hurtt et al. (1998) concluded that GVMs need to be validated for a diverse range of spatial and temporal scales. Datasets of forest stand structure variables (e.g., height, basal area, and volume increment) are good candidates for this diversification because they are often available on wider spatial and temporal scales than eddy-covariance data, but these variables are not simulated by most GVMs. A new generation of GVMs that explicitly simulate forest management begins to bridge this gap: Desai et al. (2007) validated the regional forest biomass
simulated by Ecosystem Demography with data derived from forest inventories in the
midwest United States, and Sato et al. (2007) compared the local age structure
simulated by SEIB-DGVM with intensive monitoring plots. However, to date, no GVM
has been evaluated simultaneously at the diverse spatial and temporal scales relevant
to managed forests.

94 Beyond the assessment of model error, a model validation exercise also provides the 95 opportunity for a better understanding of the model's strengths and pitfalls. In 96 particular, it should be designed to attribute a share of model error to each model 97 component. One way of attributing this model error is to quantify the improvement of 98 model fit when a given component is switched on. This approach was used by Zaehle et 99 al. (2006) for a model component simulating the processes involved in the age-related 100 decline of net primary productivity (NPP). Because it requires a validation variable that is 101 simulated both in the absence and presence of the component, this method is not 102 always applicable. Another way of attributing modelling error is to force a model by 103 replacing the outputs of one model component by site measurements. By assimilating a 104 satellite-derived leaf area index (LAI) in ORCHIDEE, Demarty et al. (2007) showed that 105 the phenology component of the model is responsible for 25% of the lack of fit to flux 106 tower data. Because the new forest management module (FMM) of ORCHIDEE 107 generates a whole new set of variables and processes, this second approach was found

to be better suited to discriminate between its error and the error coming from thegeneral core of ORCHIDEE.

In this study, we use forestry datasets to further evaluate the performance of the
ORCHIDEE-FM simultaneously at several spatial and temporal scales, which are all
relevant to the novelty introduced by the forestry management module, namely the
simulation of stand structure and its evolution with age. Two requirements are set for a
validation dataset:

It should cover the diverse spatial and temporal scales necessary for exploring
 regional variations and the full lifespan of a forest from harvest to harvest.

It should provide the possibility to replace the input of ORCHIDEE to the FMM
 with actual field values so that a share of modelling error can be attributed to
 both the ORCHIDEE and the FMM components (see Figure 1).

120 No single dataset was found to match all of these requirements. Instead, we selected 121 three complementary datasets: permanent forest monitoring plots, yield tables, and 122 extensive forest inventory data (see Figure 2). Permanent plots provide long time series 123 and detailed within-plot measurements, but their spatial coverage is very limited. Yield 124 tables cover the entire European continent, but their precise area of relevance and 125 source data are often unknown. Inventory data is sufficiently abundant to create 126 spatially continuous maps of carbon stocks and stock changes (through surface tree 127 cores), but only one snapshot measurement is available for each plot. Here, these

128 datasets are successively compared to specific simulations to validate ORCHIDEE- and to

130	The notion of model validation is controversial (Oreskes et al., 1994). Confirmation that
131	models reproduce existing in situ measurements reasonably well is, nevertheless,
132	required of GVMs, the projections of which are used in the definition of climate change
133	mitigation and adaptation policy (IPCC, 2007). Therefore, we use the term of
134	"validation" with the cautionary requirements set by Rykiel (1996), clearly specifying the
135	model's purpose, its context of operation and the criteria that it must meet for being
136	considered "acceptable for use":
137	 Model purpose: to simulate the age-related dynamics of carbon stocks and
138	fluxes that are ignored in the standard version of ORCHIDEE.
139	\checkmark Context of operation: a tree to continental scale, limited to Europe.
140	\checkmark Validation criteria: whereas plot-scale models, calibrated with site- and species-
141	specific parameters, can be expected to fit local data series, the aim of a GVM is
142	to simulate a regional average CO $_2$ flux, typically using 0.5° resolution. The
143	performance of ORCHIDEE-FM is, therefore, assessed through its ability to cut
144	across a cloud of data points corresponding to different sites in the same region.
145	All abbreviations used in this paper are indexed in Annex 1.

129 identify the most important sources of model error.

All abbreviations used in this paper are indexed in Annex 1. 145

146 **2 Material & Methods**

147 **2.1 ORCHIDEE and its forest management module (FMM)**

148 **2.1.1 Description of ORCHIDEE-FM**

149 The ORCHIDEE global vegetation model (ORganising Carbon and Hydrology In Dynamic 150 Ecosystems) was designed to operate from regional to global scales (Krinner et al., 151 2005). ORCHIDEE typically represents an average mature forest at steady-state 152 equilibrium in a *big-leaf* manner. For a given climate, it simulates the carbon, water and 153 energy budget at the pixel scale. For carbon, ORCHIDEE computes its fixation (gross 154 primary productivity or GPP), allocates photosynthetates to the different biomass 155 compartments where they are respired or stored, and recycles carbon through constant 156 tree mortality and soil respiration. To simulate forest management, several processes 157 have been added to the standard version of ORCHIDEE, among which is a forest management module (FMM) inspired by the stand-level model FAGACEES (Dhôte and 158 159 Hervé, 2000). The key concept is to add to the "average tree" representation of 160 ORCHIDEE an explicit distribution of individual trees, which is the basis for a process-161 based simulation of mortality (see Figure 1). The above-ground plot-scale wood 162 increment simulated by ORCHIDEE is distributed among individual trees according to the 163 rule of Deleuze et al. (2004):

164
$$\delta ba_i = \frac{\gamma}{2} \times \left(\operatorname{circ}_i - m \times \sigma + \sqrt{(m \times \sigma + \operatorname{circ}_i)^2 - 4 \times m \times \sigma \times \operatorname{circ}_i} \right)$$
 (1)

165 where δba_i is the annual increase in the basal area of tree i in square meters, and *circ_i* is 166 the circumference of tree i in meters. γ , σ and m are the slope, the threshold and the 167 smoothing parameters, respectively (see Figure S1): trees whose circumference is lower 168 than σ grow very little; thus, γ is the slope of the δba_i vs. the *circ_i* relationship above σ . 169 Then, tree mortality processes, due to natural competition, anthropogenic thinning or 170 clear-cutting, rely on the self-thinning rule (Eq. 2) of Reineke (1933).

$$dens_{nex} = \frac{\alpha_{st}}{Dg^{\beta_{st}}}$$
171 (2)

where *dens_{max}* is the stand maximum density in ind ha⁻¹ (individuals per hectare); α_{st} and β_{st} are parameters; and *Dg* is the quadratic mean diameter in m.

174 For more information on the structure of ORCHIDEE-FM, see Bellassen et al. (this issue).

175 **2.1.2 Pedo-climatic inputs and model "spinup"**

176 The climate data used in this study to drive ORCHIDEE is from the 0.25°-resolution

177 REMO reanalysis covering the 1861-2007 period (Kalnay et al., 1996; Vetter et al., 2008).

178 Maps of soil depth and texture were derived from FAO and IGBP products (Vetter et al.,

179 2008). Following a standard method in GVM modelling, a model "spinup" is performed

180 before all simulations to define the initial conditions from which subsequent simulations

181 will be performed, in particular for soil carbon. For this "spinup", ORCHIDEE and

182 ORCHIDEE-FM are repeatedly run for the climate of the years 1861-1911 with a CO₂

183 concentration of 280 ppm until all ecosystem carbon and water pools reach a cyclical

- 184 (clearcut-regrowth) steady-state equilibrium. The conditions of the stand after the last
- 185 clearcut are used as initial conditions for all ORCHIDEE-FM simulations.
- 186 **2.2 Validation data**
- 187 Three complementary datasets are used to validate the FMM and its integration in
- 188 ORCHIDEE: permanent plots, yield tables and national inventory. All three are necessary
- to cover the three different scales of interest: tree scale (e.g., individual tree growth and
- 190 circumference distribution), stand scale (e.g., tree density and basal area), and
- 191 continental scale (e.g., inter-regional variations). The following paragraphs describe
- 192 each dataset and its specific use in our model validation assessment. Table 1
- 193 summarises the characteristics and aims of each simulation. Figure S 4, Figure S 5, and
- 194 Figure S 9 summarise measurements, simulations and validated variables for each of the
- three datasets. The uncertainty associated to each dataset in discussed in part 2 of the
- 196 Supplementary Materials.
- 197 2.2.1 Permanent plots
- 198 Description
- 199 Fifty-eight permanent plots (PP) were set by the Institut National de la Recherche
- 200 Agronomique (INRA) for long-term monitoring of the evolution of forest stands (Dhôte
- and Hervé, 2000). These plots contain either oaks or beeches. They all belong to even-
- 202 aged stands and were subject to different management intensities with a post-thinning

- relative density index (*rdi*, see Eq. 3) ranging from 0.4 (heavy thinning) to 1 (no
- anthropogenic thinning = unmanaged).

$$205 \quad rdi = \frac{dens}{dens_{max}}$$
(3)

- where *rdi* is the relative density index, and *dens* and *dens_{max}* are the actual and maximal
- tree densities, respectively, of the stand in ind.ha⁻¹.
- 208 *dens_{max}* is derived from Eq. 2. All trees in the plots are marked, and for each
- 209 measurement year, the status of trees is recorded (dead, alive, or thinned), as is their
- circumference at breast height. Measured ages span 37-203 years, with an average
- 211 measurement frequency of 4.2 years.
- A summary of permanent plot characteristics is provided in Annex 2.
- 213 Estimation of non-measured variables
- 214 The key variables of interest available at each plot for the validation of the FMM-
- 215 simulated counterparts are as follows:
- circumference distribution variables: minimum, average, and maximum
- 217 circumference as well as number of trees in a given circumference class
- stand variables: tree density and basal area.
- 219 Other variables, such as standing volume, standing biomass, tree height, wood
- increment, and individual tree growth indicators (the σ and γ of Equation 1) can be
- estimated. For detailed information on the estimation method, see part 1 of the
- 222 Supplementary Materials.

224	The <i>PP_f</i> (permanent plot-forced) simulation is aimed at validating tree-scale and stand-
225	scale characteristics between two measured states: the state of a stand at first
226	measurement and its state at last measurement. To validate the FMM separately from
227	the rest of ORCHIDEE, the annual increase in aboveground woody biomass (<i>wood_{inc}</i>) is
228	forced by the <i>in situ</i> estimate instead of the value simulated by the core of ORCHIDEE.
229	The initial conditions of PP_f are the conditions of each permanent plot at its first
230	measurement regarding tree circumferences and, therefore, aboveground biomass. The
231	other biomass compartments (e.g., leaves, roots, and soil) are not used as inputs in the
232	FMM and, therefore, do not need to be accounted for when the FMM is forced.
233	The aim of the PP_c (permanent plot-coupled) simulation is to assess the additional error
234	brought to the FMM outputs by an initial error in the simulation of stand-scale wood
235	increment by ORCHIDEE. PP_c is therefore similar to PP_f , except that the FMM is no
236	longer forced by data-derived wood _{inc} . Instead, the coupled ORCHIDEE-FM model is run
237	over the measurement period of each plot using the corresponding climate forcing to
238	provide a simulated <i>wood</i> _{inc} . Whereas the differences between PP_f and the data reveal
239	the error of the FMM in simulating management and growth distribution, those
240	between PP_c and PP_f reflect the error due to the simulation of wood _{inc} by ORCHIDEE.
241	Comparing the performance of PP_f vs. <i>data</i> to that of PP_c vs. PP_f allows us to attribute a
242	share of the total error (PP_c vs. data) to each of the model's components. Comparing PP_c
243	directly to the data would be confusing because the error of each component might

- cancel each other out and, consequently, be wrongly interpreted as a high modellingefficiency.
- 246 *PP_{fi} simulation: validation of initial distribution*
- 247 The *PP_f* (permanent plots initial conditions) simulation complements the *PP_f* simulation
- 248 by assessing the model's ability to reproduce the state of each plot at its first
- 249 measurement, starting from the default model initial conditions of 10,000 trees per
- 250 hectare, with circumferences following a decreasing exponential distribution (Bellassen
- 251 et al., this issue). The FMM is, therefore, run on each permanent plot from its date of
- regeneration until the first measurement year. For the *PP_f* simulation, the FMM is
- 253 decoupled from the rest of ORCHIDEE: the annual increase in aboveground woody
- biomass between year one and the first measurement is forced by the average annual
- wood increment estimated from field data over the measured period.
- 256 **2.2.2 Yield tables**
- 257 Description

258 More than a thousand forest yield tables have been compiled by the Joint Research 259 Centre (JRC, 2009). They cover 26 European countries and 23 genuses. Forest yield 260 tables give the evolution of typical stand variables, including tree density, basal area, 261 dominant or average height, average circumference, standing volume and thinned 262 volume, with age. All of these variables will be tested against FMM simulations for 263 validation. Yield tables are usually established based on either permanent plots

264	monitored over an entire rotation, or temporary plots of different ages monitored once.
265	Their aim is to reproduce the average growth pattern of a tree species in a given region,
266	which sometimes declines in yield classes representing different levels of treatment or
267	local fertility. Because the FMM simulates the growth of an average coniferous or
268	broadleaf species managed as a high stand, coppices and fast-growing poplars and
269	eucalypts were discarded from the database. When needed, cormometric volume
270	(merchantable volume) was converted into dendrometric volume (whole tree) using a
271	branch to total volume ratio of 0.25 for needleleaf species and 0.38 for broadleaf
272	species (Loustau, 2004).
273	Testing the effect of climate and management in the dataset
274	Yield tables complement permanent plots by providing a presumably much more
275	diverse range of climates, species and management conditions. However, neither
276	management style nor climate are represented by explicit indicators as is the case with
277	permanent plots for which accurate location, plot age and the targeted relative density
278	index play that role. A first step in testing the FMM against this assumed variety of
279	climates and management conditions is therefore to test whether climate and
280	management effects can indeed be detected in the dataset. To test the climate effect,
281	an analysis of variance was performed using the mixed linear model of Eq. 4.
282	$vol_{tot}(i,j,k) = \alpha + \beta_i + \gamma_j + \varepsilon(i,j,k)$ (4)
283	where α is the intercept; β_i and γ_j are the coefficients associated with plant functional

type (PFT) i and country j, respectively, and $vol_{tot}(i,j,k)$, and $\varepsilon(i,j,k)$ are the total volume

produced at year 80 and the error term associated with yield table k of PFT i and country j, respectively. The error terms $\varepsilon(i,j,k)$ are assumed to be dependent upon the

yield table k, justifying the use of a linear mixed model with PFT and country as fixedeffects.

Because the total biomass (standing biomass + thinned biomass) produced by a plot is
largely independent of the management style (Lanier, 1994), this variable was not suited
to test the diversity of management styles in the data. For this purpose, a second
analysis of variance based on tree density was performed using the linear mixed model
of Eq. 5.

294
$$dens(i, j, k) = \alpha + \beta_i + \gamma_i + \delta \times vol_{tot}(i, j, k) + \varepsilon(i, j, k)$$
(5)

295 where α is the intercept; β_i , γ_i and δ are the coefficients associated with plant functional 296 type (PFT) i, country j, and total volume produced at year 80, respectively; dens(i,j,k), 297 $vol_{tot}(i,j,k)$ and $\varepsilon(i,j,k)$ are the density, the total volume produced at year 80 and the 298 error term associated with yield table k of PFT i and country j, respectively. The error 299 terms $\varepsilon(i,j,k)$ are assumed to be dependent upon the yield table k, justifying the use of a 300 linear mixed model. 301 Density is highly dependent on management for a given productivity level, which is 302 embedded in the random factor voltot, and management is, therefore, likely to explain

303 most of the variance attributed to country and PFT when productivity is already

304 captured by another variable (here vol_{tot}). In this model, β and γ can thus be interpreted

305 as indicators of the country- and PFT-specific variation in management style.

306 *YT_f* simulation: validation of stand-scale characteristics across Europe



316

French national inventory

317 Description

2.2.3

318 The French National Forest Inventory (NFI) conducts yearly field measurement 319 campaigns covering the entire French metropolitan territory. Each intersection of a 320 systematic grid of 10 x 10 km² is photo-interpreted to determine land cover and land-321 use. Of these intersections, every other forested point, totalling about 8 000 points per 322 year, is visited and inventoried following the NFI protocol (IFN, 2006): circumference at 323 breast height, width of the last five rings, height and species are recorded for a 324 representative sample of trees. NFI allometric rules are used to estimate tree volume 325 and annual volume increment, and all of these data provide the basis for an estimate of

320	piot-scale tree density, basal area, dominant neight, standing volume and the annual
327	volume increment. For even-aged stands, a few trees are cored to the stem centre to
328	estimate stand age. For this study, we pooled together the results of three campaigns
329	(2005, 2006 and 2007). Because the FMM only represents even-aged high stands, all
330	other management types were excluded from the analysis. Our sample size was,
331	therefore, reduced to 11,222 sites. The raw data are available on the IFN website:
332	www.ifn.fr.

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333 Interpolation

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334 Both permanent plots and yield tables are unsuitable to test the ability of ORCHIDEE-FM 335 to simulate regional trends in carbon stocks and fluxes. Thus, a spatially continuous 336 dataset is needed. With its high spatial density, the NFI dataset presents the opportunity 337 to build continuous maps: for the category of broadleaf plots of the 80-100 years age 338 class alone, half the French territory has at least 10 plots within a distance of 0.5° (55 339 km), and only 24% of the territory has less than 5 plots. However, this dataset is 340 heterogeneous: the order of magnitude of the standard deviation of the volume 341 increment within a radius of 0.5° is 30%. Therefore, a smoothing is necessary to 342 eliminate the local variations due to topography, soil fertility and species composition 343 and to retain only the regional climate-related variations in carbon stocks and fluxes. 344 Several interpolation techniques were tested to obtain these smoothed data-derived 345 maps, resulting in the joint use of the following two methods:

346	•	Large footprint interpolation technique: the data were interpolated with a
347		minimum footprint radius of 0.5° and no distance weighting. Where necessary,
348		the footprint was extended to include a minimum of 10 plots. The result is a
349		0.05° resolution map for which each final pixel represents the average of all plots
350		within a 0.5° radius of the centre of the pixel.
351	•	Data density mask: density masks were created to distinguish pixels with more
352		than 10 plots within a 0.5° radius. Applying these masks restricts model-data
353		comparisons to the areas where the uncertainty in the data is lowest.
354	NFI _{std}	and NFI _{fmm} regional simulations
355	Two ty	pes of simulations were conducted to assess the model's ability to reproduce the
356	trends	observed in the NFI data.
357	•	The <i>NFI_{std}</i> simulation aims at representing an average forest at steady-state
358		equilibrium, typical of GVMs. Thus, the standard version of ORCHIDEE was run
359		between 1956 and 2006, which is the average measurement year for the dataset
360		(2005-2007).
361	•	The three NFI _{fmm} simulations aim at validating the coupled version of the
362		ORCHIDEE-FM. The model was run for 50, 90, and 130 years with all runs ending
363		in 2006. The resulting NFI _{fmm50} , NFI _{fmm90} , and NFI _{fmm130} results can, thus, be
364		compared to NFI plots of three selected age-classes: 40-60 years, 80-100 years

366

and 120-140 years. In both cases, the CO_2 concentration follows its historical increase from 290 ppm in 1876 to 378 ppm in 2006.

367 The wood increment estimated in the NFI data comes from surface cores of live trees. It

is a gross commercial wood increment, and it does not account for woody losses from

369 artificial thinning or natural mortality. From the simulation of tree-level growth and

370 mortality, a similar variable can be extracted from the ORCHIDEE-FM simulations,

allowing its validation. The commercial wood increment is converted to the total wood

increment using the relevant PFT-specific branch expansion factor (BEF) of IPCC (2003).

373 "Difference maps" present the relative difference between each pixel of data-derived

374 maps and its closest simulation point. These maps are limited to pixels complying with

the data density masks, which are pixels with at least 10 inventory plots within a 0.5°

- 376 radius of the centre of the pixel.
- 377 NFI_{opt} and NFI_{st} simulations for error attribution

Unlike permanent plots, it is not possible to estimate the history of productivity in each
NFI plot. The only data point available is the average tree-ring width over the previous
five years that is obtained from a surface core. Therefore, attributing error to the
management or productivity simulation is not straightforward when one is looking at
cumulative variables such as standing biomass. To do so, two additional types of
simulations are performed:

For the *NFl_{opt}* simulations, we replaced the default values of the photosynthesis
 efficiency parameters (vc_{max}, the maximum capacity of the Rubisco enzyme, and

386	$v j_{\text{max}}$, the maximum regeneration speed of the Rubisco enzyme) with the values
387	of Santaren (2006), who optimised their ORCHIDEE model based on eddy-
388	covariance measurements from six European sites. Broadleaves were
389	unaffected, but the photosynthesis efficiency of needleleaves was increased by
390	20%. <i>NFI_{opt}</i> simulations are a sensible variant of <i>NFI_{fmm}</i> simulations for
391	productivity.
392 •	For the NFI _{st} simulations, artificial thinning is disabled, and only self-thinning
393	occurs, thus representing the minimal level of management. When a lack of fit
394	between the ORCHIDEE-FM and data for standing biomass comes from overly
395	intensively simulated management, NFI_{st} provides a comparison with the most

396 extensive type of management.

397 **2.3** Criteria of model performance

398 Two common criteria are used to evaluate model performance: *EF*, model efficiency,

and *AB*, model average relative bias (Soares et al., 1995; Smith et al., 1997). Their

400 definition is given by Eq. 6 and 7.

401
$$EF = 1 - \frac{\sum_{i} (mes_{i} - sim_{i})^{2}}{\sum_{i} (mes_{i} - \overline{mes})^{2}}$$
 (6)

402
$$AB = \frac{1}{n} \sum_{i=1}^{n} \frac{(sim_i - mes_i)}{mes_i}$$
 (7)

respectively; \overline{mes} is the data average; and n is the number of data points.					
EF reflects the ability of the model to reproduce the data: the closer it gets to 1, the					
better the fit. AB indicates whether the model has a systematic bias. Whereas an					
efficient model necessarily has a small systematic bias, the reverse is not always true.					
However, when a large-scale model such as ORCHIDEE is compared to plot-scale					
measurements, avoiding systematic bias may be more important than scoring high					
efficiency: large-scale models are not expected to reproduce each stand specifically but					
rather to simulate an "average stand" within the gridcell of interest.					
To improve the interpretation of these criteria, we undertook three complementary					
analyses:					
• "Shadow models": for each simulation, we built a "shadow model" for the					
ORCHIDEE-FM. These "shadow models" are simple statistical models using the					
same input variables as ORCHIDEE-FM. For the stand-scale variables of the PP_f					
simulation, for example, the main input variables of ORCHIDEE-FM are total					
volume, initial conditions (initial median circumference), and management					
intensity (post-thinning relative density index). The shadow model thus follows					
Eq. 8.					

421
$$mes(i) = a \times vol_{tot}(i) + b \times med_{circ}(i) + c \times rdi_{target}(i) + \varepsilon(i)$$
 (8)

422	where mes is the measured variable of interest (e.g., tree density and standing
423	volume); <i>vol_{tot}</i> is the total volume of the stand at the last measurement; <i>med_{circ}</i>
424	is the median circumference of the stand at the first measurement; rdi_{target} is the
425	post-thinning relative density index; <i>a</i> , <i>b</i> , <i>c</i> , and <i>d</i> are regression coefficients; <i>i</i> is
426	the permanent plot number; and $\varepsilon(i)$ is the error term associated with <i>mes(i)</i> .
427	"Shadow models" are calibrated on one half of the dataset, and their efficiency
428	(EF_{stat}) is assessed on the other half. The details of each model and its calibration
429	are presented in the Supplementary Materials.

Systematic vs. unsystematic error: to assess the importance of the average bias,
we computed the systematic (*RMSEs*) and unsystematic (*RMSEu*) errors of
Willmott (1982), defined by Eq. 9 and 10, respectively:

433
$$RMSEs = \frac{1}{n} \sqrt{\sum_{i=1}^{n} (pred_i - mes_i)^2}$$
 (9)

434
$$RMSEu = \frac{1}{n} \sqrt{\sum_{i=1}^{n} (pred_i - sim_i)^2}$$
 (10)

435where *RMSEs* and *RMSEu* are the systematic and unsystematic root mean square436error, respectively; n is the number of the measurement; sim is the simulated437variable; mes is the measured variable; i is the measurement number; and pred438is the value predicted by the linear regression sim = f(mes): pred_i = a + b x mes_i,439where a and b are the regression coefficients.440RMSEs represents the error due to a systematic bias in the model, and RMSEu

- 443 *RMSEs/RMSEu* ratio is lower than one.
- Error share of a given model component: an index (*ES*_{fmm}) of the share of the
- 445 total error of ORCHIDEE-FM that can be attributed to the FMM component was

446 computed based on the permanent plots data as well as the *PP_f* and *PP_c*

447 simulations (see Eq. 11).

448
$$ES_{fmm} = \frac{1 - EF_{PPf}}{(1 - EF_{PPf}) + (1 - EF_{PPc})}$$
(11)

449 where ES_{fmm} is the error share of the FMM model component (0 when all of the 450 error comes from ORCHIDEE and 1 when it comes entirely from the FMM), and 451 EF_{PPf} and EF_{PPc} are the efficiency of PP_f to reproduce the data and the efficiency 452 of PP_c to reproduce the PP_f simulation, respectively.

453 **3 Results**

454 **3.1 Stand scale: stand characteristics**

455 **3.1.1 Permanent plots**

456 *PP_f* and *PP_{fi}* simulations: good performance of the FMM under controlled conditions

- 457 Average stand characteristics such as tree density, basal area, average circumference
- 458 and standing volume are efficiently simulated under the control conditions of *PP_f* (Figure
- 459 3 and Figure 4). All of these characters have modelling efficiencies higher than 0.5 and

460 average biases below 20%. This average bias is negligible because the systematic error is 461 smaller than the unsystematic error: all RMSEs to RMSEu ratios are lower than 0.6 462 (Table 2). The model is not as accurate for extreme circumferences: both have lower 463 efficiencies, and the minimum circumference is consistently underestimated with an 464 average bias of -25% and a systematic error component overtaking the unsystematic 465 component. These deficiencies essentially occur for plots with large trees (Figure 3). 466 With the approximations necessary for the PP_{fi} simulation (default model initial 467 distribution and average growth rate), the fit of all variables deteriorate. Except for 468 standing volume and average circumference, all model efficiencies become negative. 469 From the results of the *PP_{fi}* simulation, we conclude that the model could not correctly 470 reach the initial state of the *PP_f* simulation. Average biases are also higher than for the 471 PP_f simulation, although none exceeds 45%. However, because all RMSEs to RMSEu 472 ratios remain below 0.7, the default initial conditions of ORCHIDEE-FM can be considered to induce no strong systematic bias to the simulations. For both simulations, 473 474 the FMM is more efficient than its simple statistical "shadow models". For stand-scale 475 variables, its efficiency is on average of 0.11 higher for PPf and 0.6 higher for PPf. 476 PPf and PPc simulations: a minor share of modelling error for the FMM component 477 The (inaccurate) simulation of wood increment by ORCHIDEE is a more important source 478 of error than the processes simulated by the FMM. For most variables, the forced FMM 479 (PP_f) is more efficient at reproducing the data than ORCHIDEE-FM (PP_c) is at reproducing 480 the forced FMM (Figure 4). For basal area, which is the variable most commonly

estimated by forest inventories, the efficiency of the forced FMM to reproduce the data is three times higher than that of ORCHIDEE-FM to reproduce the forced FMM, giving an ES_{fmm} value of only 35%. Because the efficiency of the coupled PP_c remains quite high for standing volume, the error for this variable is, therefore, split evenly between ORCHIDEE and the FMM ($ES_{fmm} = 48\%$).

486 **3.1.2 Yield tables**

487 Statistically significant effect of climate and management practices in the dataset

488 The statistical model of Equation 4 explains 64% of the total variance, and both country

and PFT predictors have a significant effect (p-value < 0.001) on the total volume

490 produced (the detailed statistics are provided in the Supplementary Materials).

491 Therefore, the effect of climate is present, though blurred, in the yield table dataset.

492 This result can be ascertained visually from Figure 5: the estimated coefficients for

493 country (γ_j), representing the relative effect of each country corrected for PFT effects,

494 present a climatic pattern with lower values in arid Spain and the cold Russo-

495 Scandinavian countries. This pattern is clearly blurred over western and central Europe,

496 where the differences between countries are difficult to explain based on climate alone.

497 The statistical model of Equation 5 explains 47% of the total variance, and all

498 explanatory variables (country, PFT, and total volume produced) have a significant effect

499 (p-value < 0.01) on stand density. The effect of the total volume produced is, as

500 expected, more important than that of PFT and country (F-value is about 50 times

501 higher for total volume). Because management style is expected to vary between PFTs 502 and countries, this result points to a detectable effect of management style on tree 503 density, although other explanations for the effect of PFT and country cannot be 504 discarded (e.g., ecophysiological differences between PFTs and differences in methodology for establishing yield tables between countries). Similar results are 505 506 obtained if density is replaced by basal area or standing volume in Equation 5, showing 507 that management styles also affect these two variables. 508 YT_f simulation: validation across a variety of management and climate conditions 509 Except for tree density, average biases do not exceed 55% for the YT_f simulation, and 510 most modelling efficiencies are higher than 0.3, with the exception of average height 511 and average circumference (Figure 6 and Figure 7). The FMM performs particularly well 512 for standing volume with an EF value of 0.83 and an average bias of only -2%. This value is slightly better than the "shadow model" (EF_{stat} = 0.82, AB_{stat} = 16%, see Table 2). 513 514 Because standing volume varies little for a given level of total volume produced, the 515 linear regression is indeed more sensitive to extreme values, which may differ between 516 the calibration and test subsamples and produce a higher average bias in the shadow 517 model. 518 For most variables, however, the performance of the FMM is lower for YT_f than under 519 the highly controlled conditions of *PP_f*: Efficiencies are lower and average biases are 520 higher, as is the systematic to unsystematic error ratio; however, it remains below 1 for

521 all variables except average height.

522 The FMM does not efficiently simulate tree density (EF = -8). In particular, it

523 overestimates high densities. However, the average bias of +160% is not uniform: Figure 524 6 shows that the fit is best for low densities (around 600 trees.ha⁻¹), meaning that the 525 average bias comes from the high number of data points from the high densities where 526 the bias is particularly high, rather than a systematic bias spanning the entire density 527 range. The average bias of +96% in the shadow model shows that reproducing the tree 528 density trends from the yield tables is not easy to accomplish. This difficulty could 529 originate from a specific treatment effect or measurement errors for the higher tree

- 530 densities.
- 531 3.1.3 French national inventory

532 Interpolated NFI plots and NFI_{fmm} simulations: regional trends

The interpolation technique unearths regional differences in volume increments (Figure 8a and Figure 8c), most of which are bolstered by a large number of plot measurements. For broadleaves, the range of the volume increment is from 2 to 18 m³.ha⁻¹yr⁻¹, half that of the needleleaves, which can grow as fast as 30 m³.ha⁻¹yr⁻¹ in northeastern France. In particular, regional lows of -48% and -59% in the Mediterranean region (2)¹ can be observed, extending somewhat inland toward south-central Toulouse to the west for

¹ To help readers unfamiliar with French geography, numbers between brackets refer to the regional markers of Figure 8d. The exact boundaries of these "regions" are given in **Erreur ! Source du renvoi introuvable.** of the Supplementary Materials.

broadleaves (3, -21%), and from the mid-Atlantic coast (7, -12% and -26%) to the
Parisian basin for needleleaves (1, -10%). Robust regional highs occur in northeastern

541 France (4, +36% for both) for both functional plant types, in Britanny (5, +5%) for

542 needleleaves and at the southwestern tip (6, +15%) for broadleaves.

543 The sign of these regional trends in volume increment is generally correctly simulated

544 (see Table 4). However, the amplitude of these variations is often underestimated; in

545 particular, the regional high in the north-eastern region (4) and the regional low for the

- 546 Mediterranean (2) are both underestimated in the simulations (Figure 8b and Figure
- 547 8d).
- 548 Model fit for different age classes

549 Leaving the Mediterranean region aside, the simulated broadleaf increments are

generally within the 20% uncertainty associated with the data-derived map (Figure 9).

551 The increment is, nevertheless, slightly underestimated around Paris and in the

southwest, by 20% and 50%, respectively. On the contrary, needleleaf increments are

systematically underestimated by at least 20% and often by more than 50% with the

exception of the southwest (6). For both plant functional types, the volume increment is

- 555 largely overestimated for the Mediterranean region.
- 556 Improvement in the simulation of biomass

557 For 50-year-old broadleaves, the standard version of ORCHIDEE (*NFI*_{std50}) overestimates

standing volume, which is directly related to above ground biomass stocks through wood

density, by an average of 60% (Figure 10a and Figure 10c). ORCHIDEE-FM (*NFI_{fmm50}*) is

560 much closer to the data (Figure 10b), with an average underestimate of -16%. This 561 pattern is also true for needleleaves at the southwestern tip of France (Figure 11a). For 562 the rest of the country except for the Mediterranean region, the standing volume is 563 systematically underestimated. When productivity is optimised in NFI option, model fit 564 improves in some regions at the expense of others (Figure 11b). The same happens 565 when management is made more extensive with no artificial thinning (Figure 11c). Only 566 when productivity optimisation is combined with reduced management intensity in 567 NFlopt st50 can the high volumes measured in central and northeastern France be 568 reproduced in the model (Figure 11d). This result reflects the lesser intensiveness of 569 management in these mountainous areas. A similar pattern in data-derived rdi confirms 570 this interpretation (Figure S 8).

571 **3.2** Tree scale: individual tree growth and circumference distribution

572 **3.2.1** Individual tree growth

The FMM model imperfectly reproduces individual tree growth variables as measured on the permanent plots (Figure 12). Both σ and γ have low model efficiencies of 0.1 and -0.3, respectively, and γ is even significantly overestimated. However, both simulated variables vary within the correct range of values, and their average biases of around 15% are not alarmingly high given the low efficiencies. The relevant "shadow models" are also very inefficient, suggesting that the current input variables are not sufficient to correctly predict these variables. Thus, the simulation of σ and γ will be difficult to 580 improve without a more detailed representation of inter-tree competition processes.

581 This representation would require an additional level of complexity and site-specificity

- in the FMM, which is not compatible with the aimed generality of ORCHIDEE-FM.
- 583 **3.2.2 Circumference distribution**

584 When permanent plots are sorted by increasing the simulated proportion of trees in the 585 greater than 1.4-m circumference category (Figure 13), a similar trend towards larger 586 circumference classes appears in the observed circumference distributions. This trend 587 shows that the model can capture the inter-plot differences in circumference 588 distribution. The trend in the data however, is blurred by several plots with a high 589 proportion of narrower trees than simulated. Some of these are merely attenuated in 590 the simulations (e.g., plots n°14-21-26), suggesting that circumference distribution is 591 essentially driven by the volume increment, with the FMM slightly overestimating tree 592 growth for lower values of the volume increment. In other cases, the high proportion of 593 narrower trees is not simulated at all (e.g., plots N°6-46-47-55). In these cases, 594 circumference distribution is probably driven by other factors that are not modelled in 595 the FMM (e.g., a high level of competition for light due to local topography or a "from 596 above" thinning strategy).

597 4 Discussion

598 **4.1 Effect of climate and management on carbon stocks and fluxes**

599 4.1.1 Regional assessment of carbon fluxes

600 The introduction of management and tree-level mortality into a GVM allows us to 601 validate carbon stocks and stock changes on continuous maps derived from the spatially 602 abundant inventory data. To our knowledge, this type of validation is a first for a GVM. 603 It complements the validation of short-term CO_2 fluxes at flux towers. Although the 604 inventory data only has a five-year resolution in time, it uncovers regional variations in 605 carbon fluxes that are very difficult to capture with flux towers. In particular, the low 606 productivity of the Parisian basin and the high productivity of northeastern France that 607 were detected in the data are correlated with pockets of low and high precipitation, 608 respectively, in particular for the five years before 2006. The mixed performance of 609 ORCHIDEE-FM in simulating these pockets is partly due to the mediocre accuracy of the 610 climate forcing data: while the REMO reanalysis clearly shows a regional low in 611 precipitation over the Parisian basin, it does not reproduce the pockets of higher 612 precipitation in the northeast (Meteo-France, 2009). This shortcoming combined with a 613 similar one in soil data (depth and texture) explains that simulations have a lower 614 amplitude of spatial variation than averaged measurements. Another reason for this 615 lower amplitude is the structure of the model itself. ORCHIDEE probably underestimates 616 water stress in the Mediterranean context (Morales et al., 2005; Gervois et al., 2008). In

617 northeastern France, an area with high nitrogen deposition, the model's inability to 618 simulate the high observed values in the volume increment partly comes from the 619 absence of an explicit simulation of the nitrogen cycle. 620 Some larger-scale patterns, however, can be found in both the inventory and eddy-621 covariance data. Using eddy-covariance data, Luyssaert et al. (2007) found that 622 precipitation drives NPP when average yearly temperature is higher than 10°C. Because 623 only a few mountainous grid cells (less than 10%) have an average temperature lower 624 than 10°C in France, this rule is consistent with our previous observation that 625 precipitation drives most regional trends in the country, both in data-derived maps and 626 simulations. 627 These comparisons between eddy-covariance-based and inventory-based validations 628 must, nevertheless, be made cautiously. Flux towers measure whole-stand NEE (and 629 GPP through flux-separation algorithms), while forest inventories estimate the share of 630 NPP allocated to above-ground woody growth (wood_{inc}) over a time period of several 631 years. Both variables are strongly correlated, but a model with a faulty allocation 632 scheme could perform well for total GPP and badly for *woodinc*. However, the joint use 633 of both methods presents new opportunities for the separate validation of production 634 and allocation processes.

4.1.2 Optimisation of biophysical parameters

Another key result from this spatially continuous validation is the rescaling from an 636 637 optimisation of photosynthesis efficiency parameters for needleleaves. Using the 638 optimised parameter values of Santaren (2006) improves model fit, but this does not 639 prove sufficient for all regions: rescaling allocation, plant respiration or management 640 intensity parameters also seem necessary. The model better reproduces the estimated 641 standing volume in southwestern France, which is not surprising: the parameters of 642 ORCHIDEE are based on published experimental studies, which are much more 643 abundant for southwestern Pinus pinaster than for northeastern Abies alba and Picea 644 abies. Our results question the generality of this parameterisation. Although this 645 optimisation of maximum photosynthesis rates is very coarse, the results are 646 qualitatively similar to the much finer GVM-oriented optimisation of vc_{max} using leaf 647 nitrogen content that was carried out by Kattge et al. (2009). This optimisation was 648 indeed an large upward correction from the original values of Beerling and Quick (1995) 649 for temperate needleleaves (see Table 5).

650

4.1.3 Management intensity maps

Management variability has also been shown to be an important driver of stand
characteristics and carbon stocks, both at regional (forest inventory) and continental
(yield tables) scales. In particular, management intensity has been shown to play a
comparable role to photosynthesis efficiency in explaining regional patterns of standing
volume. This result suggests that the performance of GVMs could be significantly

improved if management and photosynthesis efficiency were allowed to vary regionally
instead of having unique PFT-specific parameterisation. Such a regional
parameterisation would be feasible in Europe, where management intensity and species
distribution can potentially be mapped (Nabuurs et al., 2008).

660 **4.2 Simulating endogenous heterogeneity in a GVM**

661 In the field of ecological modelling, a common distinction exists between exogenous 662 heterogeneity, which arises from abiotic components such as climate or soil type, and 663 endogenous heterogeneity, such as the heterogeneity in individual tree circumferences, 664 which exists even in physically homogenous environments (Moorcroft et al., 2001). The 665 validation of ORCHIDEE-FM highlight the use of simulating such fine-scale processes in 666 large-scale GVMs. Although the endogenous heterogeneity now represented by the new 667 model structure with extreme circumferences, competition indexes, and circumference 668 distribution, is inefficiently simulated by ORHCIDEE-FM, the process-based model 669 outperforms simple statistical models for stand-level variables such as basal area and 670 standing volume. A similar pattern is found for the stand-level FORSKA model (Lindner 671 et al., 1997). This similarity suggests that a correct average representation of 672 endogenous heterogeneity is better than none at all, even if it poorly matches the data 673 on a plot per plot basis. Moreover, the process-based simulation of endogenous 674 heterogeneity presents the possibility to assess the impact of concrete management 675 decisions, such as short rotations, high thinning intensity, and high thinning frequency, 676 on carbon stocks and fluxes at large scales.

677 **4.3 Model strengths and limitations**

4.3.1 Model robustness for basal area and standing volume 678 679 The validation results for the yield tables show that despite a significant impact of 680 management styles in the data, the simulations are quite efficient and not strongly 681 biased, in particular, for standing volume and basal area. Similar conclusions can be 682 drawn from the *NFI_{fmm}* simulations for broadleaf standing volume. The performance of 683 ORCHIDEE-FM for basal area and standing volume probably has two explanations: first, 684 the model is robust to changes in management parameters for these variables, as 685 shown by the sensitivity analysis of Bellassen et al. (Bellassen et al., this issue), and second, basal area and standing volume are less heavily influenced by local conditions 686 687 than other variables such as average diameter or tree density and, therefore, respond 688 more directly to the large-scale climatic variations driving GVMs like ORCHIDEE (Wang et 689 al., 2006). Overall, this robustness justifies the rationale for management simulation in 690 GVMs, namely that simulating an "average" management is more realistic than not 691 simulating management.

692 **4.3.2 Tree density and self-thinning curves**

The performance of ORCHIDEE-FM to simulate tree density and average diameter is
much worse when the initial conditions and management style are unknown. The
proportion of systematic error for these two variables is twice as high as for basal area

696 or standing volume in the YT_f simulation, whereas they are comparable for all four 697 variables in the *PP_f* simulation. There are two likely reasons for this systematic error: 698 The self-thinning curves of ORCHIDEE-FM are not generic enough. Although 699 Reineke (1933) originally thought that site or species productivity made no 700 difference to his equations and would only accelerate the self-thinning process, 701 this has recently been questioned (Yang and Titus, 2002; Vacchiano et al., 2008). 702 Needleleaves tolerate higher densities than broadleaves, thus suggesting an 703 effect of at least plant functional type, if not species, on self-thinning curves 704 (Figure 14). For needleleaves, the default self-thinning curve of ORCHIDEE-FM 705 seems to be generic enough because it encompasses yield table data for the 706 entire productivity range (total volume produced at year 80). However, this is 707 not the case for broadleaves, for which many productive yield tables lie above the curve: some broadleaf species may, thus, be more tolerant to crowding than 708 709 the oaks and beeches on which the self-thinning curve was established (Dhôte, 710 1999). 711 The management style varies with density. Indeed, Figure 14 suggests that • 712 management may be more intense when the stand is dense. For both plant

functional types, the data-derived thinning curve cuts across the simulated
values as the stand grows sparser. Different management styles and intensities
between European countries could explain the important variability of the data.

These results highlight the important contributions that empirical studies of selfthinning and thinning curves could make to the performance of ORCHIDEE-FM, which
has already been shown to be very sensitive to these parameters (Bellassen et al., this
issue). Another approach would be to construct a data assimilation framework for
ORCHIDEE-FM to optimise the thinning and harvest parameters on existing wood
production datasets.

722 4.3.3 Bridging the gap with raw inventory data: basal area increment 723 One of the original ideas in ORCHIDEE-FM is its ability to put a process-based GVM on par with forest inventory data. In terms of proxy variables for productivity, ORCHIDEE-724 725 FM performs better for the volume increment than for the basal area increment. This is 726 mainly due to the lack of performance of ORCHIDEE-FM at the tree scale: the basal area 727 increment is very dependent on tree circumference distribution because many small 728 trees will show a higher basal area increment than a few large trees for the amount of 729 volume increment. Therefore in this study, we used the estimated volume increment 730 from the French inventory instead of the measured basal area increment. If the same 731 validation exercise was undertaken at the European scale, then the basal area 732 increment might be the only option because the methods for estimating volume vary 733 strongly between countries, and a comparison based on the compilation of European 734 forest inventories by Schelhaas et al. (2006) would be challenging without full 735 documentation of each inventory's method. In this case, a possibility for improving the 736 performance of ORCHIDEE-FM for the basal area increment would be to force the model

for its initial conditions. If the model were fed with the measured tree circumference
distribution before the productivity measurement by surface coring, ORCHIDEE-FM
would be more reliable in its simulation of the basal area increment and would,
therefore, provide a meaningful comparison with direct measurements.

741 **5 Conclusion**

742 The double aim of this study was to validate ORCHIDEE-FM at the various temporal and 743 spatial scales necessary for a GVM and to separate the modelling error due to the 744 simulation of management from that due to the simulation of productivity. We showed 745 that ORCHIDEE-FM performs reasonably well over long time-scales for most stand-level 746 variables (tree density, basal area, standing volume, average height, and average 747 circumference) and at spatial scales varying from local to continental with several 748 degrees of continuity between measurements. The performance of ORCHIDEE-FM is, 749 however, less satisfying for fine-scale processes such as competition between trees. In 750 terms of error separation, we showed that when initial conditions and management 751 style are controlled, the error from the FMM management component tends to be 752 lower than that of the ORCHIDEE productivity component. However, the volume 753 inventory data shows that both the management and the productivity components 754 need to be calibrated if we want the model to finely reproduce the conditions of a 755 specific region.

756 The validation of ORCHIDEE-FM also paves the way for its improvement. Specific 757 attention should be paid to thinning parameters, either through more empirical studies 758 or through an optimisation framework. The assimilation of initial conditions in the 759 model could also present the possibility of a comparison with the raw measurements of 760 forest inventories rather than the estimated volume and volume increment. Overall, 761 ORCHIDEE-FM is deemed reliable enough to carry out prospective studies on the large-762 scale impact of management on climate and on the impact of climate change on 763 business-as-usual management. In these applications, ORCHIDEE-FM will provide a 764 useful complement to inventory-based studies because it allows the separation of the 765 effects of CO₂, climate, and management on wood stocks and wood production.

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960 **Tables**

Simulation	Dataset	Model used	Source of woody NPP	Initial conditions	Time period	Validated variables
PP _f	Permanent plots	FMM	Data	Data	First measurement \rightarrow Last measurement	dens, ba, av _{circ} , circ _{min} , circ _{max} , distrib, σ, γ
PP _c	Permanent plots	ORCHIDEE-FM	Model	Data	First measurement \rightarrow Last measurement	dens, ba, av _{circ} , circ _{min} , circ _{max} , σ, γ, vol _{tot}
PP _{ic}	Permanent plots	FMM	Data	Model	Year 0 \rightarrow First measurement	dens, ba, av _{circ} , circ _{min} , circ _{max}
ΥΤ _f	Yield tables	FMM	Data	Model	Year $0 \rightarrow$ Year 80	dens, ba, dom _{height} , av _{height} , av _{circ} , vol _{main} ,
NFI_{std}	National Forest Inventory	ORCHIDEE	Model	Model	1876, 1916, 1956 → 2006	NPP_{woody}, vol_{main}
NFI _{fmm}	National Forest Inventory	ORCHIDEE-FM	Model	Model	1876, 1916, 1956 → 2006	NPP_{woody}, vol_{main}
NFI _{opt}	National Forest Inventory	ORCHIDEE-FM, optimized photosynthesis	Model	Model	1876, 1916, 1956 → 2006	NPP_{woody}, vol_{main}
NFI _{st}	National Forest Inventory	ORCHIDEE-FM, self-thinning only	Model	Model	1876, 1916, 1956 → 2006	NPP_{woody}, vol_{main}

961

962 Table 1. Simulations summary

963 Variable abbreviations: dens (tree density), ba (basal area), av_{circ} (average

964 circumference), circ_{min} (minimum circumference), circ_{max} (maximum circumference),

- 965 distrib (circumference distribution), σ (threshold circumference for basal area growth), γ
- 966 (competition index), voltot (total wood volume produced), volth (cumulated thinned
- 967 wood volume), NPP_{woody} (annual wood increment), vol_{main} (standing wood volume).

Validation	Variable name	EF	EF _{stat}	AB	AB _{stat}	RMSEs/RMSEu
PP _f vs data						
	dens	0.6	0.40	18%	1%	0.40
	ba	0.56	0.29	2%	10%	0.43
	vol _{main}	0.6	0.39	4%	9%	0.15
	av _{circ}	0.53	0.77	-5%	6%	0.57
	circ _{min}	-0.33	0.36	-25%	46%	1.30
	circ _{max}	0.36	0.67	10%	-3%	0.67
	σ	-0.11	-0.14	16%	72%	0.69
	γ	-0.27	0.01	15%	4%	0.91
PP _{fi} vs data						
	dens	-0.33	-0.29	30%	2%	0.02
	ba	-0.15	-0.90	-12%	13%	0.13
	vol _{main}	0.09	-1.59	-9%	18%	0.18
	av _{circ}	0.09	0.10	-3%	0%	0.10
	circ _{min}	-0.22	0.05	45%	10%	0.30
	circ _{max}	-0.45	-1.24	22%	30%	0.10
PP _c vs PP _f						
	dens	0.16	na	-13%	na	0.72
	ba	0.19	na	1%	na	0.68
	vol _{main}	0.57	na	6%	na	0.77
	vol _{tot}	0.08	na	22%	na	0.85
	av _{circ}	0.21	na	14%	na	0.81
	circ _{min}	0.72	na	10%	na	0.94
	circ _{max}	0.54	na	12%	na	1.09
	σ	0.22	na	10%	na	1.05
	γ	-1.12	na	27%	na	1.00
YT _f vs data						
	dens	-8	-0.08	161%	96%	0.83
	av _{circ}	-0.67	0.48	-18%	-7%	0.78
	ba	0.52	0.41	10%	13%	0.33
	dom _{height}	0.32	0.63	-9%	4%	0.62
	av _{height}	-0.85	0.60	-35%	0%	1.33
	vol _{main}	0.83	0.82	-2%	16%	0.44
	vol _{th}	0.45	0.81	54%	-3%	0.96

- 970 Table 2. Efficiencies and average biases of PP and YT simulations
- 971 Modelling efficiency of ORCHIDEE-FM (EF), modelling efficiency of the relevant
- 972 statistical model (EF_{stat}), average bias (AB), and systematic/unsystematic error ratio
- 973 (RMSEs/RMSEu) for three validations.
- 974 Variable abbreviations: dens (tree density), ba (basal area), av_{circ} (average
- 975 circumference), circ_{min} (minimum circumference), circ_{max} (maximum circumference),
- 976 distrib (circumference distribution), σ (threshold circumference for basal area growth), γ

978 wood volume), NPP_{woody} (annual wood increment), vol_{main} (standing wood volume).

979

Variable	ES _{fmm} (%)
dens	32%
ba	35%
vol _{main}	48%
av _{circ}	37%
circ _{min}	83%
circ _{max}	58%
σ	59%
γ	37%

⁹⁸⁰

981 Table 3. The FMM share of the total modelling error based on the permanent plots

- 982 dataset (ES_{fmm})
- 983 For the definition of ES_{fmm}, see Eq. 11.
- 984 Variable abbreviations: dens (tree density), ba (basal area), volmain (standing wood
- 985 volume), av_{circ} (average circumference), circ_{min} (minimum circumference), circ_{max}
- 986 (maximum circumference), σ (threshold circumference for basal area growth), γ
- 987 (competition index).
- 988

Region (label of Figure 9d)	Broadleaves				Needleleaves			
	Volume increment (m³.ha ⁻¹ .yr ⁻¹)		Relative difference to national average		Volume increment (m³.ha ⁻¹ .yr ⁻¹)		Relative difference to national average	
	Measured	Simulated	Measured	Simulated	Measured	Simulated	Measured	Simulated
Parisian Basin (1)	10.3	8.7	10%	-12%	15.2	6.8	-10%	-26%
Mediterranean (2)	3.8	6.2	-59%	-37%	8.8	9.0	-48%	-2%
Toulouse (3)	7.3	6.6	-21%	-34%	16.1	7.1	-5%	-23%
North-East (4)	12.7	12.0	36%	21%	23.0	11.3	36%	23%
Britanny (5)	9.2	14.2	-2%	44%	17.7	16.0	5%	74%
South-West (6)	10.8	9.0	15%	-9%	12.6	8.8	-26%	-4%
Mid-Atlantic (7)	8.3	8.6	-12%	-13%	12.3	7.3	-27%	-20%
France	9.4	9.9	0%	0%	16.9	9.2	0%	0%

990 Table 4. Regional breakdown of the measured and simulated (IFN_{fmm50}) annual volume

991 increment

992 The exact boundaries of these regions are given in Figure 6 of the supplementary

993 materials.

994

989

Plant functional type	Study	Improved Vc _{max}	Vc _{max} change from standard value
Temperate needleleaves			
	This study (ORCHIDEE model)	41.7	19%
	Kattge et al., 2009 (BETHY)	62.5	116%
Temperate broadleaves			
	This study (ORCHIDEE model)	55.0	0%
	Kattge et al., 2009 (BETHY)	57.7	65%

996 Table 5. Improved average vc_{max} values

997

998 Annexes

- 999 AB: Average Bias
- 1000 EF: Modelling Efficiency
- 1001 FMM: Forest Management Module
- 1002 GPP: Gross Primary Productivity
- 1003 GVM: Global Vegetation Model
- 1004 HR: Heterotrophic Respiration
- 1005 LAI: Leaf Area Index
- 1006 NEP: Net Ecosystem Productivity
- 1007 NFI: National Forest Inventory
- 1008 NPP: Net Primary Productivity
- 1009 ORCHIDEE-FM: Name of the new version of the ORCHIDEE GVM, which includes a forest
- 1010 management module
- 1011 PFT: Plant Functional Type
- 1012 PP: Permanent Plot
- 1013 YT: Yield Table
- 1014 Annex 1. List of abbreviations
- 1015

ID	Departement	Species	Age at last measurement year	First measurement year	Last measurement year	Post thinning relative density index	Size (ha)
1	Loir-et-Cher	oak	201	1927	2006	0.8	1
2	Allier	oak	153	1931	2003	0.5	1
3	Allier	oak	153	1931	2003	0.8	1
4	Meurthe-et-Moselle	mixed oak/beech	151	1904	2006	0.7	1
5	Meurthe-et-Moselle	oak	122	1959	2007	0.5	0.5
6	Meurthe-et-Moselle	oak	122	1959	2007	0.8	0.5
7	Seine-Maritime	beech	124	1931	2004	0.35	1
8	Seine-Maritime	beech	124	1931	2004	0.5	1
9	Seine-Maritime	beech	124	1931	2004	0.35	1
10	Seine-Maritime	beech	124	1931	2004	0.7	1
11	Orne	oak	165	1934	2005	0.8	1
12	Orne	oak	165	1934	2005	0.5	1
13	Meurthe-et-Moselle	beech	140	1904	1995	0.8	0.2
14	Meurthe-et-Moselle	beech	140	1904	1995	0.5	0.2
15	Meurthe-et-Moselle	beech	142	1904	1997	0.8	0.25
16	Meurthe-et-Moselle	beech	142	1904	1997	0.5	0.25
17	Meurthe-et-Moselle	beech	151	1904	2006	1	0.25
18	Aisne	beech	118	1922	1978	0.4	0.25
19	Aisne	beech	118	1922	1978	0.8	0.25
20	Aisne	beech	118	1922	1978	1	0.25
21	Aisne	beech	118	1922	1978	0.6	0.5
22	Allier	oak	183	1931	2003	0.7	1
23	Orne	oak	145	1933	2005	0.4	1
24	Meurthe-et-Moselle	beech	151	1904	2006	0.7	0.25
25	Meurthe-et-Moselle	oak/beech	151	1904	2006	0.5	0.25
26	Meurthe-et-Moselle	beech	151	1904	2006	1	0.25
27	Aisne	beech	121	1922	2006	0.5	0.2
28	Aisne	beech	121	1922	2006	0.5	0.2
29	Aisne	beech	121	1922	2006	0.5	0.2
21	AISHE	beech	121	1922	2000	0.5	0.2
22	Mourtho at Macalla	Deech	202	1923	2007	0.7	1
32	Meurthe-et-Moselle	oak	137	1920	2007	0.5	1
34	Meurthe-et-Moselle	heech	142	1904	1997	0.0	0.25
35	Orne	oak	112	1951	2005	0.7	0.25
36	Orne	oak	113	1951	2005	0.6	1
37	Orne	oak	113	1951	2005	0.4	1
38	Orne	oak	138	1934	2005	0.8	1
39	Orne	oak	138	1934	2005	0.5	1
40	Orne	oak	200	1934	1960	0.7	2
41	Loir-et-Cher	oak	181	1928	2006	0.8	1
42	Loir-et-Cher	oak	181	1928	2006	0.5	1
43	Loir-et-Cher	oak	146	1928	2006	0.8	1
44	Loir-et-Cher	oak	146	1928	2006	0.5	1
45	Allier	oak	98	1992	2003	0.8	0.5
46	Allier	oak	98	1992	2003	0.5	0.5
47	Aisne	beech	173	1922	1968	0.5	1
48	Aisne	beech	146	1922	2006	0.5	0.53
49	Allier	oak	203	1931	2003	0.7	2
50	Loir-et-Cher	oak	116	1966	2006	0.4	0.83
51	Loir-et-Cher	oak	116	1966	2006	0.4	0.47
52	Loir-et-Cher	oak	116	1966	2006	1	0.47
53	Loir-et-Cher	oak	116	1966	2006	0.8	1
54	Loir-et-Cher	oak	116	1966	2006	0.6	1
55	Orne	oak	188	1934	2005	0.9	1
56	Allier	oak	123	1959	2003	0.9	1
57	Allier	oak	123	1959	2003	0.7	1
58	Allier	oak	123	1959	2003	0.5	1

1017 Annex 2. Summary of permanent plot characteristics

- 1018 Annex 3. Details on the estimation of individual tree growth variables (σ and γ)

1022 Figure legends



- 1024 Figure 1. ORCHIDEE-FM: coupled or forced with field measurements
- 1025 Wood increment can be simulated by the core of ORCHIDEE or derived from site
- 1026 measurements. The management module simulates its allocation to individual trees in
- 1027 the stand and computes mortality (self-thinning, anthropogenic thinning or clear-
- 1028 cutting) based on a density index.



- - Vieldtables

NE

1029

- 1030 Figure 2. Characteristics of the three datasets used
- 1031 Eight French forests containing 1 to 10 permanent plots with broadleaves, amounting to
- 1032 58 plots over the metropolitan territory.



- 1035 Figure 3. Validation of stand characteristics: *PP_f* simulation
- 1036 Each blue dot corresponds the state of one permanent plot at its last measurement. The
- 1037 dotted blue line represents their linear regression. AB and EF are average relative bias
- 1038 and model efficiency, respectively. An "*" indicates that the systematic error is higher
- 1039 than the unsystematic error (*RMSEs* > *RMSEu*).
- 1040



- 1042 Figure 4. Summary diagrams of model performance for *PP* simulations
- 1043 These diagrams represent model efficiency and average bias for a selected set of stand
- 1044 variables. Red circles indicate the ability of the FMM forced with the local wood
- 1045 increment to reproduce data (*PP_f* vs. data and *PP_{fi}* vs. data), whereas black diamonds

- 1046 indicate the ability of the FMM coupled with the wood increment from ORCHIDEE to
- 1047 reproduce the "forced" simulation (*PP_c* vs. *PP_f*). An "*" indicates that the systematic
- 1048 error is higher than the unsystematic error (*RMSEs* > *RMSEu*).
- 1049 1. tree density / 2. basal area / 3. standing volume / 4. total volume / 5. average
- 1050 circumference / 6. minimum circumference / 7. maximum circumference / 8. sigma / 9.
- 1051 gamma
- 1052



1054 Figure 5. Country productivity index based on the European yield table dataset

1055 The "average relative productivity" index corresponds to the country-specific coefficient 1056 of the linear mixed model of Equation 4. The studied variable is the volume increment at 1057 the age of 80, and the two explanatory variables are country and plant functional type. 1058



1059



1061 Each blue dot corresponds the state of one permanent plot at its last measurement. The

- 1062 dotted blue line represents their linear regression. AB and EF are the average relative
- 1063 bias and model efficiency, respectively. An "*" indicates that the systematic error is
- 1064 higher than the unsystematic error (*RMSEs* > *RMSEu*).
- 1065



- 1067 Figure 7. Summary diagrams of model performance for the *YT_f* simulation
- 1068 This diagram represents the model efficiency and average bias for a selected set of
- 1069 stand variables. An "*" indicates that the systematic error is higher than the
- 1070 unsystematic error (*RMSEs* > *RMSEu*).
- 1071 1. tree density / 2. basal area / 3. standing volume / 5. average circumference / 10.
- 1072 dominant height / 11. average height / 12. thinned volume



- c. Interpolated broadleaf plots of the 40-60 age class
- Figure 8. Validation of the volume increment at a regional scale
- "Interpolated data" maps (a and c) are derived from National Forest Inventory plots, and
- "NFI_{fmm50} simulation" maps (b and d) represent the output of ORCHIDEE-FM simulations
- for 50-year-old stands.



1081 Figure 9. Model fit for the volume increment: evolution with age

1082 These 6 maps represent the model fit
$$\left(\frac{NFI_{fmm} - data}{data}\right)$$
 for the volume increment for

1083 three age classes of needleleaves (a, b, c) and broadleaves (d, e, f). White areas

- 1084 represent less than 10 data plots within a 55-km radius. Thus, the interpolation is
- 1085 considered too weak to assess model fit.



- 1087
- 1088 Figure 10. Standing volume of 50-year-old broadleaf stands
- 1089 The "interpolated data" map (a) is derived from National Forest Inventory broadleaf
- 1090 plots of the 40-60-year age class and the "NFI_{fmm50} and NFI_{std50} simulation" maps (b and
- 1091 c) represent the output of ORCHIDEE-FM and ORCHIDEE simulations for 50-year-old
- 1092 broadleaf stands.
- 1093



1095 Figure 11. Model fit for standing volume of 50-year-old needleleaf stands

1096 These four maps represent the model fit
$$\left(\frac{NFI_{fmm} - data}{data}\right)$$
 for standing volume for four

1097 different simulation options: ORCHIDEE-FM (a), ORCHIDEE-FM with optimised

1098 photosynthesis efficiency (b), ORCHIDEE-FM without anthropogenic thinning (c) and

- 1099 ORCHIDEE-FM both with optimised photosynthesis efficiency and without
- 1100 anthropogenic thinning (d). White areas represent less than 10 data plots within a 55-
- 1101 km radius, and the interpolation is, therefore, considered too weak to assess model fit.



- 1103 Figure 12. Validation of individual tree growth variables (σ and γ): *PP_f* simulation
- 1104 Each blue dot corresponds to the state of one permanent plot at its last measurement.
- 1105 The dotted blue line represents their linear regression. *AB* and *EF* are average relative
- bias and model efficiency, respectively. An "*" indicates that the systematic error is
- 1107 higher than the unsystematic error (*RMSEs* > *RMSEu*).
- 1108



- 1109
- 1110 Figure 13. Validation of circumference distribution: *PPf* simulation
- 1111 The different hues indicate the repartition of trees between 11 circumference classes
- 1112 (ordinates) for the last measurement of each permanent plot (abscissa). Permanent
- 1113 plots were sorted by increasing the simulated proportion of trees in the greater than
- 1114 1.4-m category.
- 1115



- 1117 Figure 14. Self-thinning curves based on yield tables
- 1118 Full dots represent values of tree density against the total volume produced from the
- 1119 *YT_f* simulation (orange for a "managed" scenario and red for a "self-thinning only"
- scenario). Empty blue dots represent the corresponding data from yield tables, with a
- 1121 dashed line for the log-log linear regression.