

Title

Shifts in the thermal niche of fruit trees under climate change: the case of peach cultivation in France.

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Abstract

Climate influences plant phenological traits, thus playing a key role in defining the geographical range of crops. Foreseeing the impact of climate change on fruit trees is essential to inform policy decisions to guide the adaptation to new climatic conditions. To this end, we propose and use a phenological process-based model to assess the impacts of climate change upon the phenology, the suitability and the distribution of economically important cultivars of peach (*Prunus persica*), across the entire continental France. The model combines temperature dependent sub-models of dormancy, blooming, fruit survival and ripening, using chilling units, forcing units, frost occurrence and growing degree days, respectively. We find that climate change will have divergent impacts upon peach production. On the one hand, blooming will occur earlier, warmer temperatures will decrease spring frost occurrence and fruit ripening will be easily achieved before the start of fall. On the other hand, milder winters will impede the plant buds from breaking endodormancy, with consequent abnormal patterns of fruit development or even blooming failure. This latter impact will dramatically shift the geographic range of sites where peach production will be profitable. This shift will mainly be from the south

of France (Languedoc-Roussillon, Rhône-Alpes and Provence-Alpes-Côte d'Azur), to north-western areas where the winter chilling requirement will still be fulfilled. Our study provides novel insights for understanding and forecasting climate change impacts on peach phenology and it is the first framework that maps the ecological thermal niche of peach at national level.

Key words: Process-based suitability model; global warming; blooming time; plant dormancy; peach (*Prunus persica*); plant phenology

1 Introduction

Climate plays a key role in defining the geographic range of plants (Whittaker, 1975) and climate change is expected to severely influence plant distributions in the forthcoming decades (Lenoir et al., 2008; Morin et al., 2008; Chuine, 2010; Gritti et al., 2013; Zhao et al., 2018). Plant phenology is strongly responsive to temperature and, indeed, phenological changes have been among the first documented fingerprints of climate change (Menzel & Fabian, 1999; Körner & Basler, 2010; Lee et al., 2013; Wolkovich et al., 2017). Climate change will therefore have an impact on agricultural production by altering the geographical distribution of economically important crops (Tao et al., 2006; Duchêne et al., 2010; Teixeira et al., 2011; Ghrab et al., 2014). Foreseeing this impact is essential to alert stake holders, inform decisions to implement adaptation strategies and to alleviate damaging consequences. Although some of the impacts can be mitigated in agricultural settings (e.g. irrigation, frost protection), the challenge will be greater for perennial crops. These are subject to climate impacts throughout the year and their decades-long lifespan makes the choice of where to plant an orchard critical (Lobell & Field, 2012).

To date, spatial shifts in the distribution of crops have been assessed using empirical Species Distribution Models (SDMs) (Machovina & Feeley, 2013), as well as process-based Suitability Models (SMs) (e.g. White et al., 2006; Keenan et al., 2011). Both modeling frameworks have merits and shortcomings. However, as noticed by Parker and Abatzoglou (2017), “unlike SDMs, SMs can provide information on specific climatic limitations and crop phenology, and are not limited by the correlative approach”. When dealing with perennial plant phenology and climate, the blooming process is probably the most studied. Until one decade ago, most studies

23 provided consensus on the earlier occurrence of plant blooming. For example, Estrella et al.
24 (2007) reported that phenological events such as emergence and blooming “are significantly
25 earlier now than 53 years ago, with a mean advance of 1.1-1.3 days per decade”. According to
26 Chmielewski et al. (2004) “phenological phases of the natural vegetation as well as of fruit trees
27 and field crops have advanced clearly in the last decade of the 20th century”. Nevertheless, in the
28 last years some authors theorized a possible trend reversal due to a subtle, process-dependent
29 cause: strong warming in winter could slow the fulfillment of chilling requirements, which may
30 delay spring phenology (Hänninen & Tanino, 2011). These scenarios were also confirmed in
31 recent field studies (Yu et al., 2010; Laube et al., 2014). Accordingly, models that explicitly
32 considered the fulfillment of both chilling and forcing requirements, also called two-phase or
33 sequential Chilling/Forcing (CF) models, were proposed to better assess the date of blooming.
34 For the genus *Prunus*, this has been accomplished for apricot and peach trees (Chuine et al.,
35 2016) as well for cherry trees (Chmielewski & Götz, 2016).

36 To design adaptation strategies, the occurrence of blooming is a necessary yet not sufficient
37 condition to make an area suitable for fruit production. An area is considered suitable if the
38 environmental conditions allow all the processes leading to full fruit ripeness to occur. Follow-
39 ing this rationale, Parker and Abatzoglou (2017) assessed possible shifts in the thermal niche of
40 almond trees under climate change in the Western United States. Santos et al. (2016) assessed
41 values of chilling and heat accumulation over Portugal and discussed possible related shifts
42 of the thermal niche of several fruit classes (from carob to lemons and from olives to vines).
43 Similarly, Ahmadi and Baaghdeh (2018) assessed the impact of climate change on apple tree
44 cultivation in Iran. They assumed that a given area would be suitable if temperatures remained
45 within certain boundaries in any plant development stage.

46 Here, we propose a novel modeling approach that combines temperature-dependent phenologi-
47 cal models of blooming, fruit survival and ripening to assess suitability for peach *Prunus persica*
48 tree cultivation. We optimize the models for nine different peach cultivars and demonstrate
49 our approach for a reference past period (1996-2015). Then, we use it to project shifts in the
50 peach cultivation range in continental France under different scenarios, that are Representa-
51 tive Concentration Pathways, RCPs 4.5 and 8.5 (IPCC, 2013) of average and minimum daily
52 temperature change in the near (2021-2040) and far (2081-2100) future. Globally, the peach is

53 the third most cultivated plant of the *Rosaceae* family (Obi et al., 2018). This fruit has been
54 extensively studied in both field and modeling works (Génard & Huguet, 1996; Ziosi et al.,
55 2003; Allen et al., 2006; Miras-Avalos et al., 2011) and its sensitivity to climate change has
56 already been established (Litschmann et al., 2008; Ghrab et al., 2014). The French territory
57 is a paradigmatic case for studying peach cultivation, because it covers four climatic zones
58 (Mediterranean, continental, oceanic and mountain) in less than 600,000 km².

59 **2 Materials and Methods**

60 **2.1 Phenological dates**

61 From the database TEMPO (National Network of Phenology Observatories) (INRA, Seguin,
62 2004), we obtained 159 pairs of data that report blooming and harvest dates for nine peach
63 cultivars (Snowqueen, M. Sundance, Springlady, OHenry, Alexandra, Flavorglod, Benedicte,
64 Flavorcrest and Emeraude) in the period 1987-2008. These concern three experimental sites in
65 southern France: Bordeaux (long. 0° 34' W, lat. 44° 46' N), Balandran (long. 4° 28' E, lat.
66 43° 45' N) and Étoile-sur-Rhône (long 4° 53' E, lat. 44° 49' N) (see Supplementary Information
67 Figure S1).

68 **2.2 Temperature: records, assessments and projections**

69 Daily average temperature data (1987-2008) were obtained from meteorological stations in the
70 proximity of the experimental sites (see Supplementary Information Table S1) and provided
71 by the INRA CLIMATIK platform (https://intranet.inra.fr/climatik_v2). We obtained assess-
72 ments of daily minimum and average temperatures for the entire French continental area for
73 the period 1996-2015. Assessments were generated by the EUROCORDEX CNRM-CERFACS
74 CM5 model and downscaled with a spatial resolution of around 8 × 8 km² (~ 0.11° × 0.11°).
75 They have been made available by the project Drias developed by Météo-France (Drias, 2013).
76 The same project provided projections of the daily minimum and average temperature for two
77 Representative Concentration Pathways scenarios, RCP 4.5 and RCP 8.5 (IPCC, 2013), in the
78 period 2020-2100, downscaled at the same spatial resolution, for the entire French territory.
79 The RCPs refer to different emission scenarios providing an estimated increase of global mean

80 surface temperatures at the end of the 21st century, is likely to be in the ranges of 1.1-2.6°C
81 for RCP 4.5 and 2.6-4.8°C for RCP 8.5. To analyze the extent of climate warming in France,
82 we selected four cells of 24×24 km², one for each climatic zone (Mediterranean, continental,
83 oceanic and mountain), and we evaluated the daily temperature anomalies in the the near
84 (2021-2040) and far (2081-2100) future with respect to a reference time period (1996-2015).

85 **2.3 A model to predict blooming date**

86 First, we performed a 1-way ANalysis Of VAriance (ANOVA, with a significance level of
87 $\alpha = 0.05$) to test if observed blooming dates significantly differed between the nine consid-
88 ered cultivars. Second, we used the two-phases model proposed by Chuine et al. (2016) to
89 predict blooming dates. That model assumes that endodormancy break occurs at the time t_C ,
90 when the state of chilling $S_c(t)$, resulting from the sum of the daily rates of chilling R_c (as
91 detailed below, see eq. 5), reaches the critical value C^* . According to Chuine et al. (2016),
92 thus, we compute

$$S_c(t) = \sum_{\xi=t_0}^t R_c(\xi) \quad (1)$$

$$S_c(t_C) = C^* \quad (2)$$

93 with t measured in days and t_0 fixed to September 1st. Blooming is assumed to occur at time
94 t_B when the state of forcing S_f , *i.e.* the sum of the daily rates of forcing R_f (also detailed
95 below, see eq. 6), reaches the critical value F^* . According to Chuine et al. (2016), we have

$$S_f(t) = \sum_{\xi=t_c}^t R_f(\xi) \quad (3)$$

96

$$S_f(t_B) = F^* \quad (4)$$

97 with t_C determined by equation 2. Note that the endodormancy break is a necessary condition
98 to enter the ecodormancy phase that precedes blooming. Both daily rates of chilling R_c and
99 forcing R_f are functions of the average daily temperature $T(t)$, itself varying with time t . We

100 computed R_c using a symmetrical and unimodal function proposed by Chuine (2000):

$$R_c(T(t)) = \frac{1}{1 + \exp [a(T(t) - T_c)^2 + b(T(t) - T_c)]} \quad (5)$$

101 where a (in units of $^{\circ}\text{C}^{-2}$), b ($^{\circ}\text{C}^{-1}$) and T_c ($^{\circ}\text{C}$) are species-specific parameters that describe
102 the accumulation of chilling units. According to Chuine et al. (2016), we computed R_f as a
103 logistic function of temperature:

$$R_f(T(t)) = \frac{1}{1 + \exp [s(T(t) - T_f)]} \quad (6)$$

104 where the parameter s ($^{\circ}\text{C}^{-1}$) shapes the steepness of the curve and T_f ($^{\circ}\text{C}$) the logistic midpoint.
105 As for the chilling function, we used the parameters optimized by Chuine et al. (2016) for peach
106 ($a = 3.53$, $b = -25.85$, $T_c = 1.52$, $C^* = 49.6$), while we optimized the parameters of the forcing
107 function, *i. e.* s , T_f and the required F^* , using our data set by minimizing the Root Mean
108 Square Error ($RMSE$, see Burnham and Anderson, 2002):

$$RMSE = \sqrt{\frac{\sum_{i=1}^n (\hat{x}_i - \bar{x}_i)^2}{n}} \quad (7)$$

109 where $\hat{x}_i = \hat{t}_{Bi}$ is the estimated blooming date, $\bar{x}_i = \bar{t}_{Bi}$ is the observed one, and n the number
110 of observations over three different sites. We performed the $RMSE$ minimization via the
111 Nelder–Mead simplex algorithm (Lagarias et al., 1998), using the built-in MATLAB optimizer
112 “fminsearch”.

113 2.4 A model to predict ripening duration

114 We performed a 1-way ANalysis Of VAriance (ANOVA, with a significance level of $\alpha = 0.05$)
115 to test if the observed ripening duration d_R (*i.e.* the time from blooming = t_B to ripening
116 = t_R) was significantly affected by the factor “cultivar”. In case of significance, we performed
117 a multiple t-test using Scheffé’s procedure (Savin, 1980), with a significance level $\alpha = 0.05$, to
118 evaluate if cultivars could be classified in groups. If this was the case, we calibrated a different
119 parameter set of the ripening model for each group (see below). We assume that a fruit is ripe
120 when the state of ripening S_R , *i.e.* the sum of the daily rates of heating R_R (see eq. 10 below),

121 reaches the critical value R^* (Miller et al., 2001; Kenealy et al., 2015):

$$S_R(t) = \sum_{\xi=t_B}^t R_R(\xi) \quad (8)$$

122

$$S_R(t_R) = R^* \quad (9)$$

123 where t_B is the observed blooming time for each cultivar and year. $R_R(t)$ is calculated as

$$R_R(t) = \begin{cases} T(t) - T_a & \text{if } T(t) \geq T_a \\ 0 & \text{if } T(t) < T_a \end{cases} \quad (10)$$

124 where T_a is the base activation temperature, which we set equal to 7°C (Miller et al., 2001;
125 Kenealy et al., 2015). Thus, R^* is the sum of Growing Degree Days (GDD). Then, for each
126 cultivar group we estimated the value of R^* minimizing the $RMSE$ (see eq. 7), where $\hat{x}_i = \hat{d}_{Ri}$
127 is the estimated ripening duration, $\bar{x}_i = \bar{d}_{Ri}$ is the observed one, and n is the number of
128 observations. If ripening is not achieved by the 21st of September, when the plant is assumed
129 to enter dormancy (Battey, 2000; Gauzere et al., 2017), the yield is considered to be entirely
130 lost. In other words, the environmental conditions are not suitable for cultivation.

131 2.5 Thermal niche for fruit tree cultivation

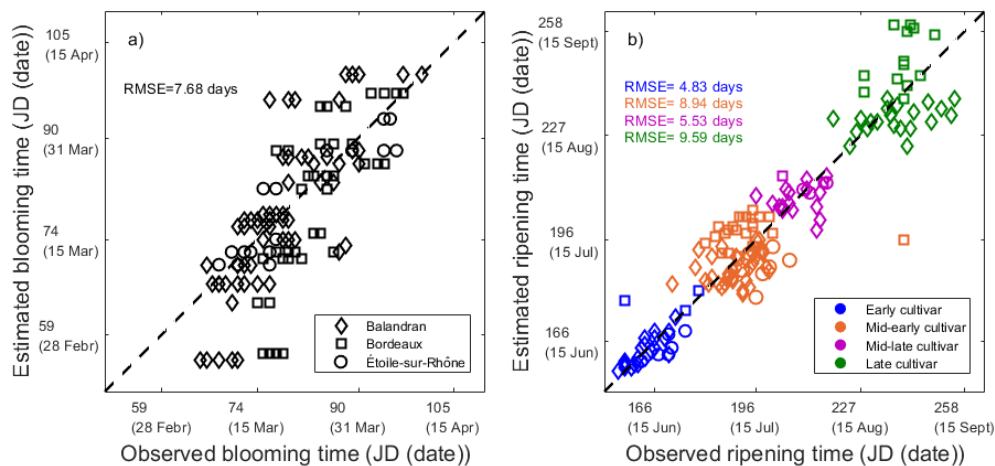
132 We assume that an area is suitable for peach cultivar if all three necessary conditions are
133 sequentially met: *i*) blooming can occur, *i.e.* both chilling and forcing requirements are satisfied;
134 *ii*) no frost events occur at the turn of blooming; and *iii*) ripening can occur. To model the fact
135 that flowers (considered here as newborn fruits) are sensitive to frost in their first week after
136 blossom (Rodrigo, 2000), we assume that fruits do not survive if there is at least one day, in the
137 period from $t_B - 3$ to $t_B + 3$, with minimum daily temperature below a critical temperature (T_x).
138 For peach, according to Snyder and Melo-Abreu (2005), this critical temperature corresponds
139 to -4.9 °C. We assumed that orchards can be irrigated so that water requirements are met
140 independently from climatic conditions. We assessed the spatial suitability for different varieties
141 of peach cultivation across France (8412 map cells of 8×8 km²) over three different periods,
142 *i.e.* a reference (1996-2015), a near future (2031-2050) and a far future (2081-2100) period.

143 We considered a map cell as suited for cultivation if all three conditions described above are
144 accomplished for at least 18 years out of 20.

145 3 Results

146 3.1 Models for blooming and ripening date

147 We found that the factor “cultivar” has no significant effect on the observed blooming dates
148 ($p = 0.61$). The values of parameters of the blooming model optimized against our data set,
149 namely s , T_f and F^* (see eq. 5), are $-0.5 \text{ }^\circ\text{C}^{-1}$, $14.4 \text{ }^\circ\text{C}$ and 4.19 , respectively. Although
150 the model is quite hyperactive, *i.e.* it tends to underestimate earliest blooming dates and to
151 overestimate the latest ones, it reproduces the observed data well (Figure 1a). We evaluated
152 if the model residuals are cultivar-dependent via an ANOVA and the resulting p-value of 0.65
153 suggests that the model simulates the peach blooming time in an equivalent way for all the
different cultivars.



154
155 *Figure 1: Blooming and ripening dates. Observed (x-axis) vs predicted (y-axis) dates in Julian Day (JD) and in calendar dates (within brackets) of (caption a) blooming and (caption b) ripening. Diamonds, squares and circles refer to the sites of Balandran, Bordeaux and Étoile-sur-Rhône, respectively. Ripening date is represented in blue for early, in orange for mid-early, in purple for mid-late and in green for late cultivars.*

156 observed ripening duration ($p < 0.01$). We therefore refined the analysis for the ripening dura-
157 tion applying Scheffé's procedure and performing a multiple t-test (Figure S2). The procedure
158 identified four cultivar clusters that we can name: "early cultivar" (Alexandra and Springlady),
159 "mid-early cultivar" (Snowqueen, Flavorgold, Flavorcrest and Emeraude), "mid-late cultivar"
160 (Benedicte) and "late cultivar" (M.Sundance and OHenry). We then optimized the parameter
161 of forcing requirement Growing Degree Days for each cultivar clustering, obtaining a GDD
162 value equal to 678, 1026, 1371 and 1772, respectively, for the "early", "mid-early", "mid-late"
163 and "late" cultivars. Adherence of model predictions (y-axis) to observations (x-axis), using
164 our model with calibrated parameters for each cluster, is quite satisfactory, as shown in Fig-
165 ure 1b. We also evaluated how the error in the estimate of blooming time propagates to the
166 ripening period duration. We used the estimated blooming time as initial time for the ripening
167 time model and we compared the estimated ripening period duration with the observed data
168 (Figure S3). Our analysis reveals that the predicted date of ripening does not vary significantly
169 whether we used the described model (estimated) or the observed blooming date (early cultivar
170 $p= 0.95$, mid-early cultivar $p= 0.44$, mid-late cultivar $p= 0.24$ and late cultivar $p= 0.87$).

171 **3.2 Peach thermal niche in the reference period 1996-2015**

172 Average temperature conditions over the reference period for the four considered cells repre-
173 sentative of climatic zones are reported in the Supporting Information (Figure S4a-d and Table
174 S2). Spring temperatures are warmer in the Mediterranean region [12-15 °C], with respect to
175 those in the continental and oceanic regions [10-12 °C], which are comparable between them,
176 while they are lower in the mountain areas [4-5 °C]. Similarly, the warmest summers were regis-
177 tered in the Mediterranean region [20-22 °C], the coldest ones in the mountain areas [12-16 °C]
178 and intermediate summers were found in oceanic and continental regions, where temperatures
179 are more variable. Autumns and winters were characterized by milder temperatures in the
180 Mediterranean [9-11 °C and 3-7 °C] and oceanic regions [8-12 °C and 4-7 °C].

181 As mapped in Figure 2a, our process-based suitability model estimates that for the reference
182 period 1996-2015 the chilling requirement was achieved first (by the end of December) in the
183 mountain regions (Alps, Pyrenees and in the Massif Central) and last (by mid February) in
184 the Mediterranean and the Southern Atlantic (MSA) regions. Also, in a few map cells of these

185 regions (notably, nearby the cities of Montpellier, Touloun, Sain Tropez, Perpignan et Bayonne)
186 some winters were so mild that the chilling requirement was not satisfied and the endodormancy
187 break was compromised. In contrast, blooming time (Figure 2b) occurred late (by end of April)
188 in the mountain regions and first (by mid March) in the MSA regions. This occurs because
189 in mountain regions, although the plants achieve chilling requirement first, they experience
190 high temperatures triggering forcing rates much later in the season. Likewise, in some cells of
191 mountain regions (Prealps and Massif Central), blooming was not effective because newborn
192 fruits were injured by frost. Note that this was not the case for map cells at the highest
193 altitudes where blooming occurred late enough to avoid frosting days. When considering the
194 date of ripening (Figure 2c), a mid-early cultivar, which is expected to ripe around mid July
195 in the MSA regions, has no time to ripe in the mountain regions. In the northern part of the
196 country, instead, ripeness is achieved only at the end of August. Such a difference in the date of
197 ripening is mostly due to blooming occurring later, rather than to lower summer temperatures.
198 Mid-late and late cultivars need significantly more days to attain ripeness, so they can only be
199 cultivated in those regions where blooming occurs first (Figure S5). For this reason, in terms
200 of peach suitability (Figure 2d), the MSA regions were the most suited for peach cultivation,
201 as both early to late cultivars could be grown and produce peach from mid June to the end of
202 August. On the other hand, the mountain regions (Pyrenees, Massif Central and Alps, black
203 in Figure 2d) were the only regions where no peach variety could be cultivated due to spring
204 frost damages, late blooming and the consequent lack of time to attain ripeness.

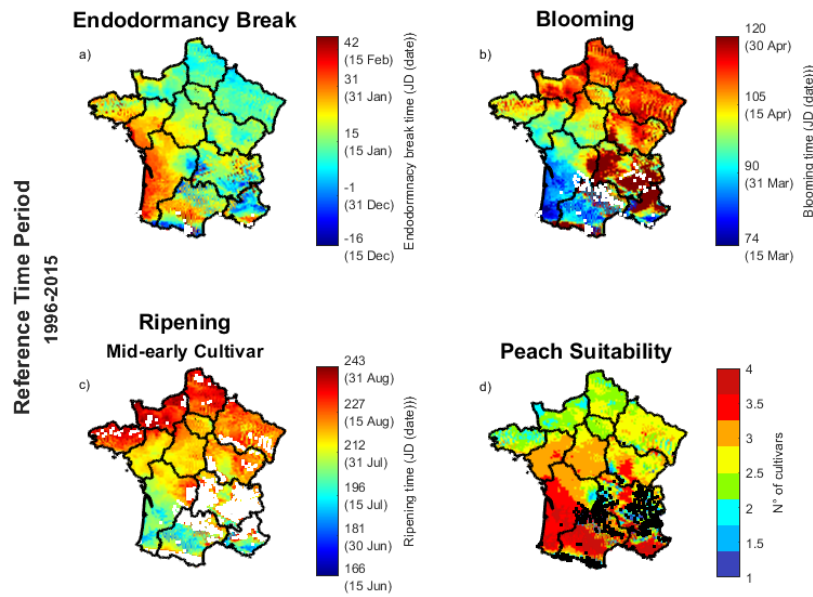


Figure 2: Peach phenological times and thermal niche for cultivation: hindcast in the French continental regions for the reference period (1996-2015). (a) Endodormancy break, (b) blooming, (c) ripening of mid early cultivars dates, in Julian Days and calendar dates (within brackets), and (d) peach suitability. For each map cell (8×8 km²), the average value over the considered 20 years is reported. White map cells (in captions a, b and c) represent those areas where the phenological event did not occur for at least two years. Likewise, black cells (in caption d) represent those areas where no peach cultivar could be cultivated for at least two years (out of 20).

205 3.3 Peach thermal niche in the future

206 According to the used scenarios (Figure S4 and Table S2), climate warming is expected to
 207 be more severe in the Mediterranean and mountain regions. Particularly, with the highest
 208 anomalies expected in winters for the Mediterranean regions, and in both summer and winters
 209 for the mountain areas. In the near future (2021-2040), the RCP 4.5 and RCP 8.5 scenarios are
 210 characterized by similar temperature anomalies. It is worthy to note that, for that period higher
 211 winter anomalies are generally expected for RCP 4.5 rather than for RCP 8.5. Conversely, in the
 212 far future (2081-2100), RCP 8.5 anomalies are expected to be consistently higher throughout
 213 the year, with the highest predicted warming in the mountain areas in summer (up to +5.1 °C).

214 Our mapping in Figure 3 reveals that the most suitable areas for peach cultivation in France
 215 are expected to significantly change in the near future (Figure 3a-d) and even more at the end
 216 of the century (Figure 3e-h). Some of the historically suitable zones in the MSA regions are
 217 predicted to become unsuitable because of blooming failure, being the chilling requirement

218 impossible to achieve (Figure 3a,e). In those areas where the chilling requirement will still be
219 satisfied, endodormancy break will be delayed. However, this will not directly reflect into a
220 delayed blooming time, as the forcing requirement will be achieved more quickly. This earlier
221 achievement of the forcing requirement will cause blooming time to occur 7-17 days in advance
222 (Figure 3b,f). In other words, blooming is expected either to be impaired or to occur earlier.
223 As a consequence of early blooming and warmer springs and summers, there will be an earlier
224 occurrence of the ripening date (Figure 3d,g). In comparison with the reference period, the
225 failures caused by not meeting the chilling requirement will cause numerous areas that are
226 currently productive to become unsuitable. This is expected mainly in the Mediterranean
227 regions and will be an important issue in the far future (see Figure 3e), especially for the
228 RCP 8.5 scenario (see Figure S6e). In contrast, spring frost events and ripening failures will
229 decrease, making continental areas in the northern part of the country more suitable for peach
230 cultivation. Changes will be exacerbated in the far future of the RCP 8.5 scenario (Figure
231 S6). In this far future scenario, there will be a paradoxical geographical divide, whereby the
232 most suitable areas will be close to those that will not be able to host any cultivar because of
233 the inability to bloom, while conditions would be optimal for fruit survival (*i.e.* no frost) and
234 ripening.

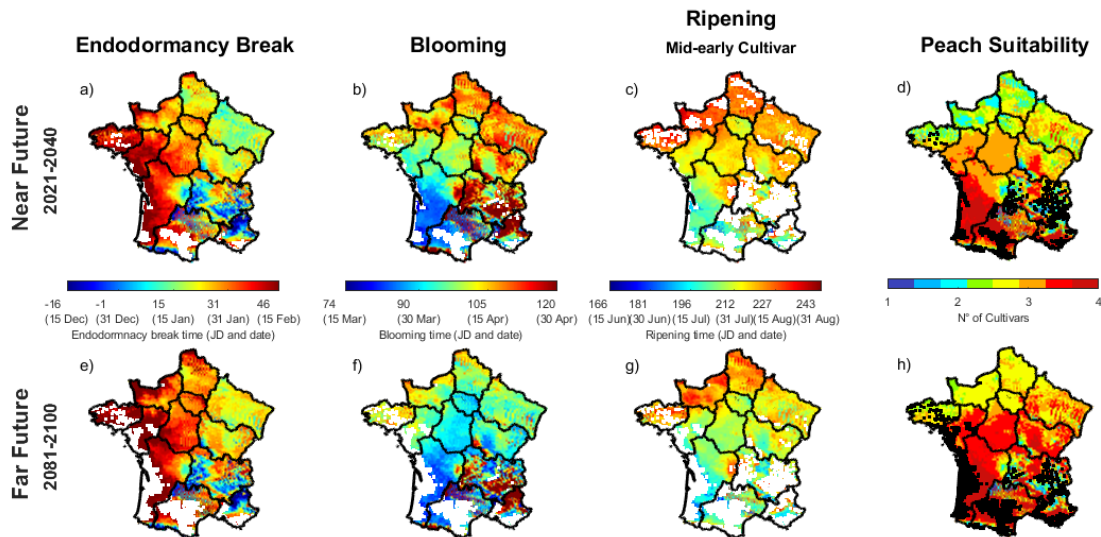


Figure 3: Peach phenological times and thermal niche for cultivation: scenarios forcing our model with RCP 4.5 in the French continental regions for the near (2021-2040, upper panels) and far (2081-2100, lower panels) future. (a,e) Endodormancy break, (b,f) blooming, (c,g) ripening of the mid-early cultivars dates, in Julian Days (JD) and calendar dates (within brackets), and (d,h) peach suitability. For each map cell ($8 \times 8 \text{ km}^2$), the average value over the considered 20 years is reported. White map cells (a-c and e-g) represent those areas where the phenological event did not occur for at least two years. Likewise, black cells (d and h) represent those areas where no peach cultivar could be cultivated for at least two years (out of 20).

235 4 Discussion

236 4.1 Peach thermal niche in the reference period 1996-2015

237 Chill accumulation has historically not been considered as a limitation for peach cultivation in
 238 the northern basin of the Mediterranean. This is consistent with our results indicating that
 239 chilling requirement was systematically satisfied for the reference period 1996-2015 in the entire
 240 French territory between mid-December and mid-February, with the milder regions satisfying
 241 it later. On the other hand, frost damage and insufficient heat accumulation are known to be
 242 primary limitations to the cultivation in Europe of peaches and other *Prunus* cultivars (e.g.
 243 apricot and cherry, see Julian et al., 2007; Reig et al., 2013; Chmielewski et al., 2017; Vitasse and
 244 Rebetez, 2018). For peaches, this was reflected in our study for those areas outside the current
 245 French peach production regions, such as the north-western areas of Occitanie and the alpine
 246 part of Auvergne Rhône-Alpes. Note that even if an area is suitable for the cultivation of very
 247 few cultivars, generally such an area cannot be considered cost-effective for peach production
 248 because the product would be available only for a limited time and late in the season (Layne

249 & Bassi, 2008). Our model identifies the locations of Occitanie, Auvergne-Rhône-Alpes (ARA)
250 and Provence-Alpes-Côte d'Azur (PACA) as the most suitable area for peach cultivation in
251 France. This outcome of our model is consistent with the fact that they currently provide more
252 than 90% of the France's total peach production (Talpin, 1954; Ministère de l'Agriculture et
253 de l'Alimentation, 2019).

254 **4.2 Peach thermal niche in the future**

255 Frost damage and heat accumulation constraints are predicted to wane in northern regions of
256 continental France, namely in Pays de la Loire, Centre-Val de Loire and Bourgogne-France-
257 Comté by the end of the 21st century, a result that is in line with projections of the northward
258 expansion of other cultivars due to global warming (Chmielewski & Rötzer, 2001). If warmer
259 winter temperatures determining earlier blooming are not followed by warmer spring temper-
260 atures at blooming time, an higher risk of spring frost can occur (Liu et al., 2018; Vitasse &
261 Rebetez, 2018). On the other hand, such risk can be neglected when also spring temperatures
262 increase below the frosting threshold (Eccel et al., 2009; Chmielewski et al., 2017). Our re-
263 sults suggest that, due to the specific interplay between peach phenology and French predicted
264 climate change the second situation will likely occur in France.

265 According to our findings, the decline in winter chill will become the major limitation for
266 peach cultivation. Blooming failure due to mild winters was first observed in northern Africa
267 (Ghrab et al., 2014) and it is likely to become the norm in southern France by the end of the
268 century. The limitation of the production area for peach appears to be even more severe using
269 our model under the RCP 8.5 scenario (Figure S6), with a further shift of the suitable area
270 towards the alpine and northern continental regions of the country. Our results are consistent
271 with the growing concern regarding the impact of global warming on the endodormancy phe-
272 nological phase. Several authors already provided evidence of abnormal patterns of bud break
273 and fruit development in Europe (Legave et al., 1983; Erez & Couvillon, 1987; Viti et al., 2010)
274 and relevant economic issues (Jackson & Hamer, 1980; Baldocchi & Wong, 2008; Luedeling
275 et al., 2009; Ghrab et al., 2014).

276 Beyond shifting the geographies of the thermal niche of peaches, climate change is expected
277 to alter peach phenology and physiology. We predict that blooming will occur between 5 and 20

278 days earlier than usual by the end of the century, depending on the area and the emission sce-
279 nario. Earlier blooming has also been predicted by other authors for other species (Chmielewski
280 et al., 2004; Dose & Menzel, 2004; Menzel et al., 2006; Estrella et al., 2007; Primack et al.,
281 2009; Fujisawa & Kobayashi, 2010; Jochner et al., 2016; Parker & Abatzoglou, 2017). When
282 dealing with commercial species, it is worth noting that faster fruit ripening due to increased
283 temperatures affects fruit growth physiology (Lescourret & Genard, 2005). This might impair
284 the quality, in terms of organoleptic properties, of the final product (Peiris et al., 1996).

285 Needless to say, all predicted changes in peach phenology and thermal niche depend on
286 methodological choices regarding both the climate change scenarios and the phenological mod-
287 els. Reliable models to estimate blooming dates are still lacking due to the complex interplay
288 of the processes governing endo- and eco-dormancy breaks (Bartolini et al., 2018) and dis-
289 crepancies between different models are unfortunately still the norm rather than the exception
290 (Chmielewski et al., 2012; Andreini et al., 2014). The main challenge is to achieve a better
291 understanding of the chill accumulation and endodormancy break processes (Luedeling, 2012).
292 New empirical data that provide measures of endodormancy break dates, such as those studied
293 by Chuine et al. (2016), are to our view urgently needed so as to conceive, calibrate and vali-
294 date models aimed at testing further hypotheses. In our model we assumed that plant water
295 needs would be met through irrigation. However, climate changes are expected to increase the
296 frequency of both drought and heavy precipitation events (IPCC, 2013), factors that surely
297 will impact water availability for agriculture (Elliott et al., 2014). Moreover, shifts in the plant
298 phenological dates and geographical range may result in another possible mismatch with pol-
299 linator phenology (Scaven & Rafferty, 2013) and photoperiod (Hänninen & Tanino, 2011; Way
300 & Montgomery, 2015). This latter will not change as the climate warms, yet the potential
301 break of the synchrony in the plant-climate integrated system remains highly probable. Like-
302 wise, plants may be exposed to new parasites and consequent diseases (Chakraborty & Newton,
303 2011). Only when a transdisciplinary and broader approach will provide further evidence, the
304 inclusion of these processes, and possibly others, in our framework will be possible.

305 4.3 Adapting to climate change

306 Despite the uncertainty which is inherent in any projection, there is no doubt that climate
307 changes will affect peach phenology and the geographical range of the thermal niche. Our study
308 corroborate the intuition that current suitable areas can turn into unsuitable and viceversa.
309 This will occur to the extent that adaptation strategies will be required. As insufficient winter
310 chill will be the major issue, some research breeding studies have been started to search for
311 cultivars with low or no chilling requirements. Such cultivars are already being utilized in
312 tropical and subtropical climates (Layne & Bassi, 2008). However, chill requirements evolved
313 to protect plants from frost damages and “artificially” lowering them might increase the impacts
314 of frost damages. Some chemicals have been shown to be effective in forcing flowering and bud
315 burst in dormant plants (Erez et al., 2008; Ashebir et al., 2010). However, they are effective only
316 under given environmental conditions (Campoy et al., 2011) and they make use of phytotoxic
317 compounds (Luedeling, 2012). It has also been shown that winter agronomic practices (like
318 sprinkling water over shoots) can reduce the buds temperature by evaporative cooling (Erez,
319 1995).

320 An alternative adaptation strategy could consist in moving the peach production into new
321 suitable areas. This strategy has three obvious drawbacks: *i*) the costs of crop translocation is
322 substantial, *ii*) additional costs can be incurred in transporting fruit to centralized processing
323 and distribution facilities (Parker & Abatzoglou, 2017), and *iii*) the sociological implications of
324 the growers’ behavior should be considered as they will ultimately govern the reality of peach
325 cultivation in new regions.

326 5 Conclusions

327 Our study provides a model to produce data-informed maps that quantitatively allow the anal-
328 ysis of the geographical limitations and challenges that the peach industry may face under
329 climate change in France. Here, we used the best currently available information about phe-
330 nology of fruit trees. We developed a novel phenological model that is not site specific, thus it
331 can be applied to other systems where climate change scenarios are available. It is clear that
332 global warming will impact the economic viability of peach production and models can support

333 actions to be taken now to adapt to change.

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566 **Supporting Information**

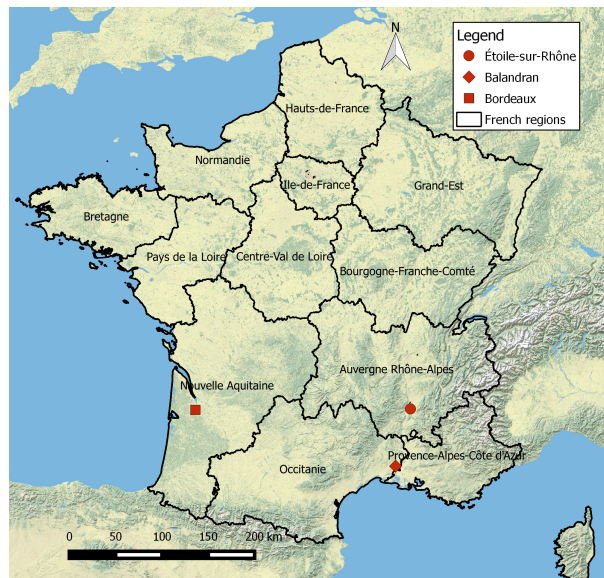


Figure S1: French continental regions and sampling sites of blooming and ripening dates.

Table S1: Meteorological stations. Meteorological station coordinates used for the period of interest in Balandran, Bordeaux and Étoile-sur-Rhône sites

Phenological data site	Years of interest	Longitude	Latitude
Balandran	1986-2004	4° 26' E	43° 47' N
	2005-2008	4° 28' E	43° 45' N
Bordeaux	1986-1999	0° 35' W	44° 47' N
Étoile-sur-Rhône	1996-2005	4° 44' E	44° 35' N

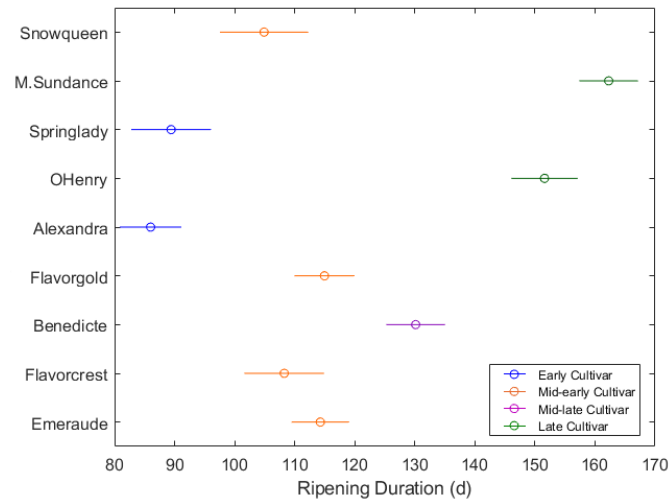


Figure S2: Peach cultivar and ripening duration. Cultivar clustering of the ripening duration period (multiple t-test, Scheffé's procedure). In blue, orange, purple and green are represented early, mid-early, mid-late and late cultivars, respectively.

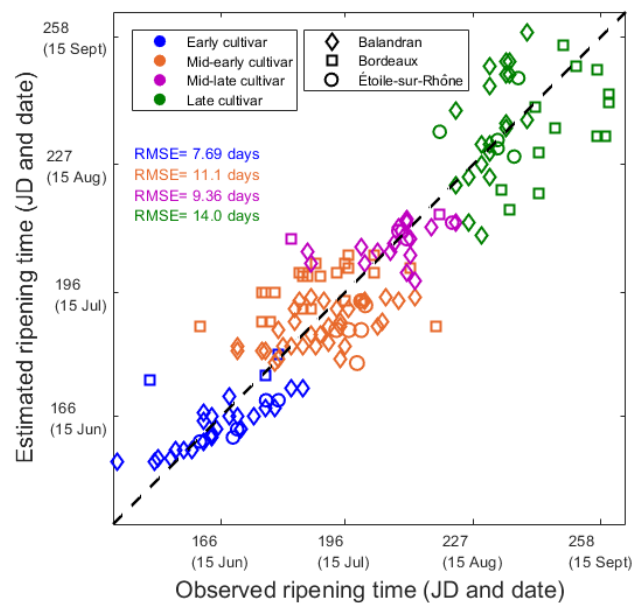


Figure S3: Observed and predicted dates of ripening. The ripening date has been predicted by summing the predicted ripening duration to the predicted (not observed as in fig. 1) date of blooming. Diamonds, squares and circles refer to the sites of Balandran, Bordeaux and Étoile-sur-Rhône, respectively. Ripening date is represented in blue for early, in orange for mid-early, in purple for mid-late and in green for late cultivars.

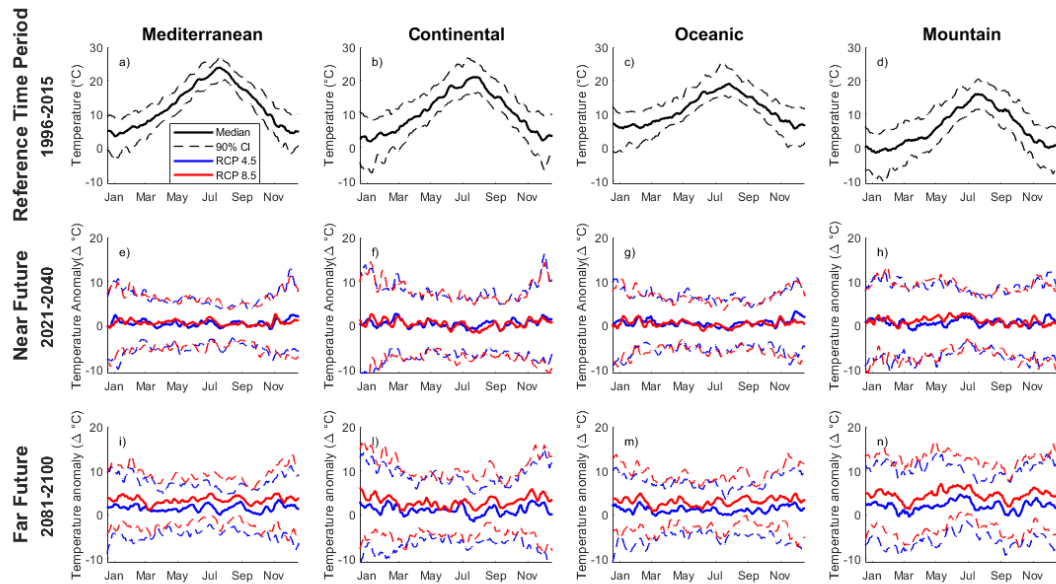


Figure S4: Temperatures in Mediterrean, continental, oceanic and mountain areas of the French territory: hindcast and forecast. Mean daily temperature, averaged with a moving window of 10 days, in the reference period (1996-2015) in black (a, b, c, d) and relevant anomalies in the near (2021-2040) and far (2081-2100) future in four areas of $24 \times 24 \text{ km}^2$ representative of different climatic zones: Mediterrean ($43^\circ 53' \text{ N } 5^\circ 4' \text{ E}$), continental ($48^\circ 48' \text{ N } 4^\circ 14' \text{ E}$), oceanic ($48^\circ 11' \text{ N } 2^\circ 59' \text{ W}$) and mountain ($42^\circ 53' \text{ N } 0^\circ 2' \text{ E}$). In black the temperatures in the reference period, in red and blue the temperature anomalies, compared to the reference period, in RCP 4.5 and in RCP 8.5, respectively. Solid lines represent the median daily temperatures, while dashed ones the 5% and 95% percentile.

Table S2: Seasonal temperature anomaly in Mediterranean, continental, oceanic and mountain areas of the French territory: forecast. Average seasonal temperature ($^{\circ}\text{C}$) in the reference period (1996-2015) and temperature anomaly ($\Delta^{\circ}\text{C}$) in the near (2021-2040) and far (2081-2100) future, in four different French climatic zones (Mediterranean, continental, oceanic and mountain). Relative 90% confidence interval are reported in square brackets. See Figure S4 for details about the climatic areas.

		Average Temperature ($^{\circ}\text{C}$)			
Time period	Season	Mediterranean	Continental	Oceanic	Mountain
Reference	Spring	11.2 [12.3;14.7]	11.6 [10.2;12.3]	11.3 [9.9;12.0]	5.4 [3.5;5.1]
	Summer	21.1 [19.6;22.2]	18.6 [17.1;21.0]	17.4 [16.0;19.2]	13.5 [11.7;15.8]
	Autumn	10.0 [8.7;11.3]	8.1 [6.3;9.7]	9.8 [8.3;11.5]	4.8 [2.7;6.2]
	Winter	5.2 [3.3;6.8]	3.8 [0.83;5.35]	6.5 [4.3;7.3]	-0.24 [-2.4;1.2]
		RCP 4.5-Temperature Anomaly ($\Delta^{\circ}\text{C}$)			
Time period	Season	Mediterranean	Continental	Oceanic	Mountain
Near future	Spring	-0.071 [-2.3;2.7]	0.22 [-2.4;2.2]	0.23 [-2.0;2.2]	0.044 [-3.0;5.2]
	Summer	0.44 [-1.5;2.2]	0.39 [-2.3;2.2]	0.052 [-1.4;2.4]	1.2 [-1.6;3.5]
	Autumn	0.39 [-2.0;3.2]	-0.045 [-1.2;3.8]	0.52 [-0.93;3.5]	0.015 [-2.4;5.0]
	Winter	1.3 [-2.2;3.1]	1.2 [-3.0;4.3]	0.93 [-1.3;3.2]	1.5 [-1.6;3.4]
Far future	Spring	1.2 [-0.78;0.87]	0.70 [-1.6;3.7]	0.92 [-1.3;3.6]	2.0 [-1.5;6.5]
	Summer	1.4 [0.36;3.1]	0.51 [-2.24;3.1]	0.75 [-0.70;2.4]	2.5 [0.47;4.3]
	Autumn	1.4 [-1.1;3.2]	1.6 [-1.3;4.0]	1.6 [0.94;4.0]	1.7 [-1.3;5.1]
	Winter	2.1 [-1.0;4.6]	1.3 [-2.6;5.1]	1.1 [-2.0;4.2]	2.2 [-1.0;5.1]
		RCP 8.5-Temperature Anomaly ($\Delta^{\circ}\text{C}$)			
Time period	Season	Mediterranean	Continental	Oceanic	Mountain
Near future	Spring	0.39 [-2.3;2.2]	0.35 [-1.8;3.1]	0.18 [-1.5;2.2]	1.4 [-1.9;4.9]
	Summer	0.21 [-1.9;2.5]	-0.28 [-4.4;3.0]	-0.51 [-2.6;2.2]	1.4 [-1.2;3.6]
	Autumn	0.22 [-0.99;2.3]	0.42 [-4.4;2.6]	0.33 [-1.6;2.7]	0.30 [-2.4;3.8]
	Winter	1.0 [-2.1;3.1]	0.71 [-2.9;4.2]	0.78 [-2.0;2.4]	0.83 [-1.6;2.9]
Far future	Spring	2.63 [1.2;5.6]	1.7 [0.41;5.3]	1.8 [0.94;4.7]	4.6 [0.84;9.1]
	Summer	3.6 [2.0;5.9]	2.9 [-0.41;5.3]	3.0 [0.47;5.6]	5.1 [3.1;7.5]
	Autumn	3.3 [1.4;5.4]	3.50 [0.66;6.0]	3.4 [1.2;5.9]	4.2 [1.7;6.6]
	Winter	3.2 [1.2;6.1]	3.04 [1.3;8.3]	2.8 [2.0;7.1]	4.1 [1.6;7.6]

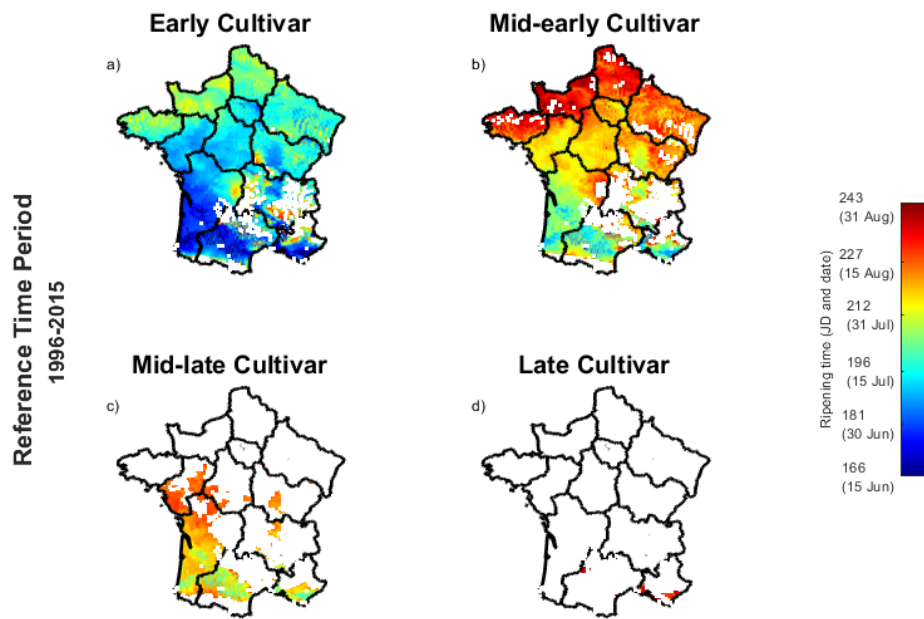


Figure S5: Peach cultivar ripening time in the French continental regions in the reference period (1996-2015). Predicted ripening date, in Julian Days (JD) and calendar dates (within brackets), in the reference time period (1996-2015) for (a) early, (b) mid-early, (c) mid-late and (d) late cultivars. White cells represent those areas that are not suitable for the cultivar production. For each cell (8×8 km²), the average value over the considered 20 years is reported.

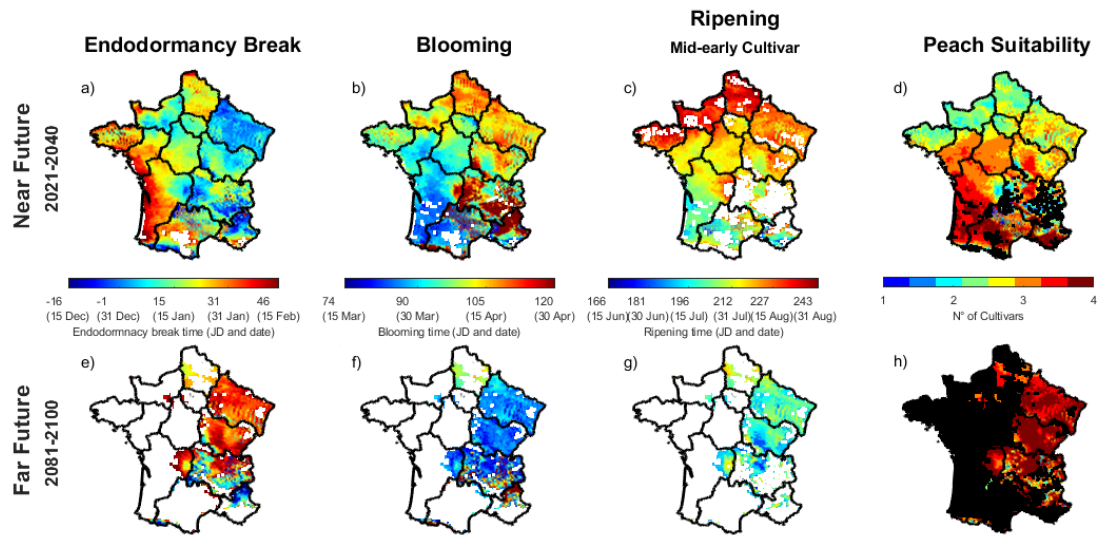


Figure S6: Peach phenological times and thermal niche for cultivation: scenarios forcing our model with RCP 8.5 in the French continental regions for the near (2021-2040, upper panels) and far (2081-2100, lower panels) future. (a,e) Endodormancy break, (b,f) blooming, (c,g) ripening of the mid-early cultivars dates, in Julian Days (JD) and calendar dates (within brackets), and (d,h) peach suitability. For each map cell ($8 \times 8 \text{ km}^2$), the average value over the considered 20 years is reported. White map cells (a-c and e-g) represent those areas where the phenological event did not occur for at least two years. Likewise, black cells (d and h) represent those areas where no peach cultivar could be cultivated for at least two years (out of 20).