



Diversity buffers winegrowing regions from climate change losses

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Agrobiodiversity—the variation within agricultural plants, animals, and practices—is often suggested as a way to mitigate the negative impacts of climate change on crops [S. A. Wood *et al.*, *Trends Ecol. Evol.* 30, 531–539 (2015)]. Recently, increasing research and attention has focused on exploiting the intraspecific genetic variation within a crop [Hajjar *et al.*, *Agric. Ecosyst. Environ.* 123, 261–270 (2008)], despite few relevant tests of how this diversity modifies agricultural forecasts. Here, we quantify how intraspecific diversity, via cultivars, changes global projections of growing areas. We focus on a crop that spans diverse climates, has the necessary records, and is clearly impacted by climate change: winegrapes (predominantly *Vitis vinifera* subspecies *vinifera*). We draw on long-term French records to extrapolate globally for 11 cultivars (varieties) with high diversity in a key trait for climate change adaptation—phenology. We compared scenarios where growers shift to more climatically suitable cultivars as the climate warms or do not change cultivars. We find that cultivar diversity more than halved projected losses of current winegrowing areas under a 2 °C warming scenario, decreasing areas lost from 56 to 24%. These benefits are more muted at higher warming scenarios, reducing areas lost by a third at 4 °C (85% versus 58%). Our results support the potential of in situ shifting of cultivars to adapt agriculture to climate change—including in major winegrowing regions—as long as efforts to avoid higher warming scenarios are successful.

agrobiodiversity | resilience | phenology | agriculture | climate change adaptation

The potential adverse effects of climate change on agriculture, including shifts in growing areas, decreased yields, and crop failures (1–6), are a major concern to practitioners, policymakers, scientists, and consumers alike (7). Forecasts predict a future where regional climates will become increasingly mismatched with crops currently cultivated in those regions (e.g., ref. 8), unless there are large shifts in agricultural practices.

Practices that increase the resilience of agricultural regions would foster growing regions that can maintain normal processes and function—including in yields and quality—despite increases in stress or disturbance (9) from climate change. Research has especially focused on exploiting intraspecific crop diversity (10–15) because of its potential to increase resilience without requiring agricultural regions or the crops they grow to shift. As expansion of agriculture is one of the primary drivers of biodiversity loss, keeping agricultural regions in place and thereby preventing natural lands from being lost to new agricultural regions is a major international conservation goal (16, 17).

To increase resilience with climate change, intraspecific diversity must link to the traits most needed to adapt to future

climate regimes (2). Such traits include a cultivar's heat and drought tolerance and its phenology—the timing of recurring developmental stages, such as budburst and maturity. Variation in phenology may be a particularly important trait for developing agricultural systems resilient to climate change, as differences in cultivar phenology (e.g., an early versus late-ripening cultivar) can translate to very different climatic conditions during critical developmental phases, such as fruit maturation.

Given enough variation in traits—such as phenology—across cultivars, growers could select and plant cultivars suited to their current climate, then shift to more appropriate cultivars over time as the climate shifts, a process we refer to as “turnover.” Cultivar turnover is expected to increase the resilience of agricultural systems and thus lead to improved agricultural forecasts

Significance

Increasing diversity within crops may be a powerful way to reduce agricultural declines from climate change. As such, it has garnered increasing attention, especially in documenting within-crop diversity through different cultivars or wild relatives. Yet, there are few tests of whether this diversity can mitigate losses with warming. Here, using European (predominantly French) databases to forecast winegrape phenology, we test if shifting cultivars changes predictions of future growing regions. We find that cultivar diversity halved potential losses of winegrowing regions under a 2 °C warming scenario and could reduce losses by a third if warming reaches 4 °C. Thus, diversity—if adopted by growers locally—can mitigate agricultural losses, but its effectiveness will depend on global decisions regarding future emissions.

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Data deposition: Analyses other than phenological parameterization and cross-validation utilized custom computer R code, freely available at GitHub, <https://github.com/MoralesCastilla/PhenoDiversity>.

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(18). Yet, this basic assumption, which underlies much of the current research, has rarely been tested.

Here, focusing on winegrapes (*Vitis vinifera* subspecies *vinifera*), we quantify how much in situ cultivar turnover affects forecasts of suitable growing regions with climate change. We selected winegrapes, given their high diversity and extensive records, which make testing the importance of intraspecific diversity to forecasts possible. Growers today plant over 1,100 different *vinifera* cultivars of winegrapes (19), called varieties, which are geographically and morphologically diverse. Different varieties possess important trait variation related to climate, including variation in phenology of 6 to 10 wk across varieties grown in the same climate (20).

Winegrape diversity is well documented, allowing us to combine winegrape phenology and global variety-level planting data with projections of daily temperature and precipitation from a large ensemble of a state-of-the-art climate model (Community Earth System Model [CESM]; *SI Appendix, Fig. S5*; ref. 21) to forecast climatic suitability of 11 globally planted varieties of winegrapes (Cabernet-Sauvignon, Chasselas, Chardonnay, Grenache, Merlot, Monastrell [synonym Mourvèdre], Pinot noir, Riesling, Sauvignon blanc, Syrah, and Ugni blanc). These varieties make up 35% of the area planted globally, reaching 64 to 87% in many important winegrowing countries (e.g., Australia, Chile, France, New Zealand, Switzerland, and the United States; ref. 22).

Our approach to model future winegrowing regions provides an important advance on previous efforts. Studies to date have generally ignored intraspecific diversity (forecasting only one or few varieties) and have used species-distribution models or simple linear phenological models, which fail to adequately include nonlinear developmental responses to temperature (23). Instead, our approach fits nonlinear process-based models for multiple

varieties, which can predict expected phenological delays due to heat stress, and characterizes specific climatic conditions during maturation. Using predominantly French long-term phenology records (*SI Appendix, Tables S1 and S2*), we developed and validated models to forecast budbreak, flowering, and the onset of ripening (veraison) in each region for two warming scenarios, +2 °C and +4 °C, and a 0 °C reference scenario of no warming (*SI Appendix, Fig. S5*; see also Warming Scenarios for more details). Next, using global data on winegrape variety plantings (22), we predicted the climatic suitability of each region during the ensuing maturation stage—a period that controls whether a variety can be harvested in a particular region each year (24–26)—for our reference and warming scenarios.

To quantify the change (including gains and losses) in areas suitable for winegrowing, and resulting cultivar turnover, we compare our results relative to: 1) current winegrowing regions, and 2) areas identified as climatically suitable (estimated as supporting at least one of the 11 cultivars modeled to maturity in most model years) under our 0 °C reference scenario (*Calculating Climatic Suitability*).

Results

Without cultivar turnover, our results predict major global gains and losses in future winegrowing regions (Fig. 1). Under a 2 °C warming scenario, 51% of all areas we identified as climatically suitable for winegrowing under our 0 °C reference scenario would be lost. At 4 °C, losses reach 77% (*SI Appendix, Fig. S14*). Losses were higher when focusing only on regions that currently grow winegrapes: At 2 °C, 56% of current growing regions were lost; at 4 °C 85% were lost (Fig. 2).

When we allowed turnover of cultivars, however, losses declined by up to 57% (Fig. 2). With cultivar turnover included

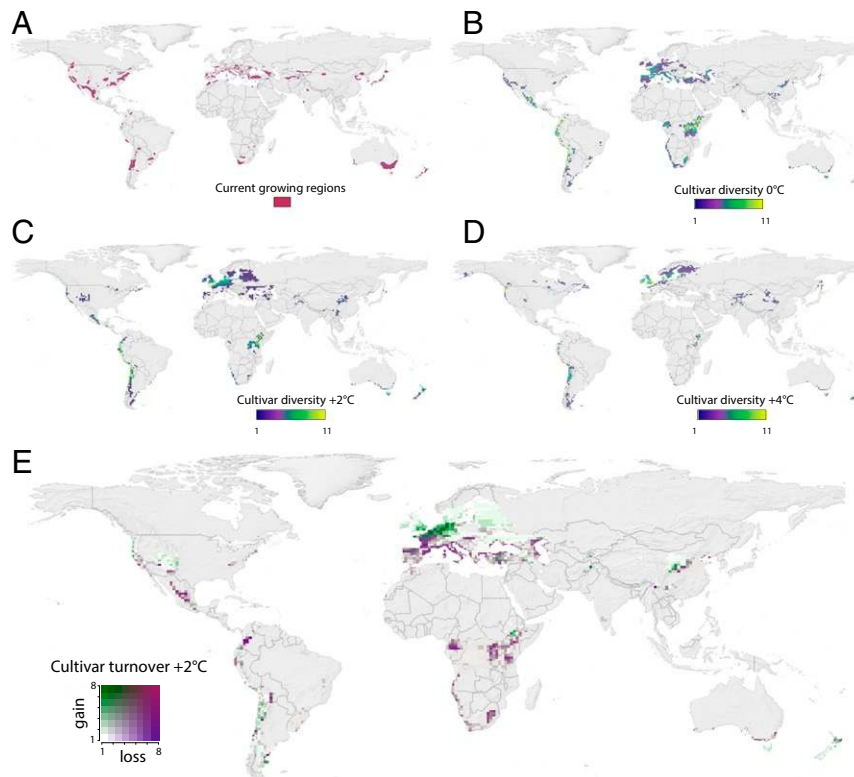


Fig. 1. Global maps of winegrowing. (A) Current winegrowing regions (see *SI Appendix, Fig. S1* for more detail). (B) Modeled predictions of cultivar diversity (total number of varieties) under our 0 °C reference scenario—all colored pixels show areas predicted as climatically suitable (*Calculating Climatic Suitability*). (C–E) Predicted effects of climate change on cultivar diversity and distribution under 2 °C warming (C), 4 °C warming (D), and cultivar turnover (E; cultivar gains and losses shown simultaneously via a bivariate color scale). See *SI Appendix, Fig. S13* for 4 °C turnover.

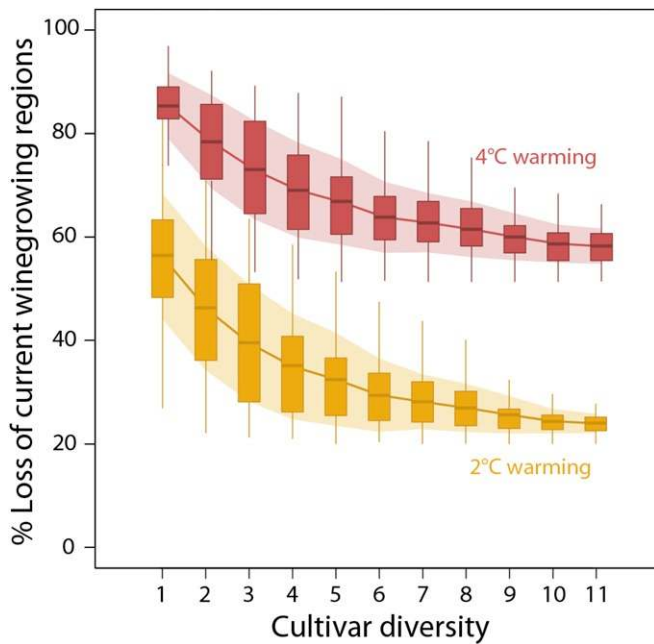


Fig. 2. Winegrape cultivar diversity can impact the loss of current winegrowing regions (see *SI Appendix, Fig. S14* for losses within all climatically suitable areas). Predictions of loss are shown for scenarios of 2 °C warming (yellow bars) and 4 °C warming (red bars) relative to a 0 °C reference scenario. Shaded areas illustrate ± 1 SD around the mean loss for each number of cultivars, combining two sources of uncertainty: 1) variability according to all possible combinations of n cultivars (e.g., at one cultivar, that cultivar could be any of the 11 considered and each covers a different area), and 2) modeled climatic suitability under each climate change scenario (e.g., one model member may predict suitability of an area, while another does not). These results are based on climatic suitability calculated with all eight climate variables (*Modeling Maturity*).

under the 2 °C warming scenario, our models predicted a loss of 24% of current winegrowing regions (compared to 56% without cultivar turnover, yielding: $(56 - 24)/56 = 57\%$ decline in areas lost). Thus, exploiting cultivar diversity more than halved the potential losses with warming. Similar improvements were seen when considering all climatically suitable areas (losses declined by a quarter to 38% under the 2 °C warming scenario). However, the benefits of including cultivar diversity were muted at higher warming: Under the 4 °C scenario, loss of current winegrowing regions was 58%, including turnover, an improvement of 32% over predictions without cultivar turnover (considering all climatically suitable areas losses declined to 64%).

The importance of using different varieties to maintaining current winegrowing regions, versus expanding into new areas, was related to variation in phenology. Many later-ripening varieties—e.g. Grenache and Monastrell—were critical to maintaining current growing areas (Fig. 3), while early ripening varieties dominated new regions (Fig. 3); in particular, at 4 °C, Pinot noir and Chasselas showed large increases (Fig. 3B) as they moved far north into large Northern Hemisphere land masses (Fig. 1D).

Geographical shifts were accompanied by phenological advances of veraison, which changes the timing of the ripening period for grapes and is important for winegrape quality potential (24–27). Early varieties—Pinot noir, Chasselas, and Riesling—advanced veraison date by over 15 d under a 4 °C warming scenario when averaged over all regions predicted as climatically suitable (*SI Appendix, Fig. S12*). This change, however, was smaller than shifts that could occur within regions

if growers do not shift varieties. For example, at 4 °C warming, Pinot noir in the area including Burgundy was predicted to advance 24 d (*SI Appendix, Fig. S11*). Advances were similarly large for some later-ripening varieties (e.g., advance of 28 d for Cabernet-Sauvignon in the area including Bordeaux). However, other late-ripening varieties showed delays of up to 10 d with 4 °C warming (*SI Appendix, Fig. S12*). Such delays were generally caused by a nonlinear response to temperature: While higher temperatures usually accelerate phenology, too-high temperatures can produce heat stress and slow phenology (*SI Appendix, Fig. S22* and ref. 28). Advances and delays in veraison date impacted the suitability of different cultivars by changing the timing, and, thus, climate, of the veraison-maturity window. These delays would likely impact wine quality, both

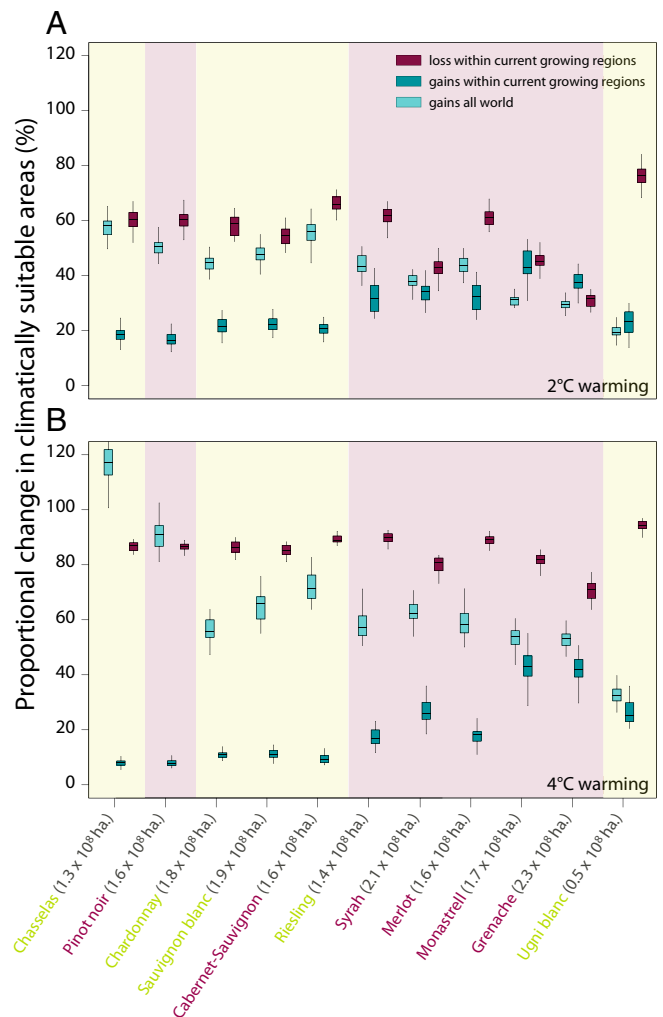


Fig. 3. Predicted gains (turquoise) and losses (purple) of climatically suitable growing areas for each of the 11 studied winegrape varieties (cultivars) under scenarios of 2 °C (A) and 4 °C (B) warming. To facilitate comparison of varieties and warming scenarios, we show gains and losses as the proportional change for each variety with warming (i.e., the area predicted with warming relative to the area predicted under our reference scenario of 0 °C). Gains are shown relative to both current winegrowing regions (darker turquoise) and all areas identified as climatically suitable under our reference scenario (pale turquoise). Background shading and variety name coloring differentiates red from white varieties; for each variety, we also give the total hectares predicted to be climatically suitable under our reference scenario of 0 °C within current winegrowing regions parenthetically. For a version of this figure showing gains and losses as absolute change in hectares, see *SI Appendix, Fig. S23*.

through the high temperatures that produce the delays (29) and through later-season effects by changing the climate during maturity (24).

Losses of climatic suitability across regions and varieties were mainly due to shifting temperature regimes during the veraison-maturity window (*SI Appendix, Figs. S16 and S17*), particularly increases in minimum and maximum temperatures and greater accumulations of temperatures above 25 °C. In the 4 °C warming scenario, the number of days above 40 °C was also commonly related to the loss of climatic suitability. These results are in accordance with increasing evidence that higher temperatures can desynchronize the development of sugars, acid, and other berry components important to quality during ripening (29). Predicted shifts in precipitation (maximum and/or summed precipitation) were less frequently associated with a loss of suitability for winegrowing (*SI Appendix, Figs. S16 and S17*). Including precipitation in our predictions, however, did have an important moderating effect: Models that excluded precipitation variables yielded higher estimates of varieties lost and showed a reduced capacity of cultivar turnover to buffer regions from such losses (*SI Appendix, Fig. S14 C and D*). While our models integrate over diverse winegrowing regions, we expect that effects of precipitation may be particularly location-specific, as irrigation—which can decouple a region from local precipitation regimes—is prohibited in some winegrowing regions and widespread in others. Even in irrigated areas, however, irrigating vines may not be a sustainable practice due to increasingly scarce water resources.

Discussion

Our results show that cultivar diversity can decrease the loss of agricultural areas by over 50%—highlighting the critical role that human decisions play in building agricultural systems resilient to climate change. We show that cultivar turnover—if adopted by growers locally—can reduce the negative outcomes of climate change on winegrapes, with implications for other crops with high diversity. However, we also find that the benefits of cultivar turnover decline under greater warming. Without global efforts to reduce emissions sufficiently to stabilize temperatures at or below 2 °C, our results suggest that half of current global winegrowing regions would become climatically unsuitable for today's major winegrapes.

These findings do not extend to all regions—in some regions, we find that cultivar diversity alone may not be enough to prevent declines. As seen in other studies (e.g., ref. 16), gains and losses of varieties are distributed unequally across the globe (Fig. 1), with warmer regions suffering the greatest losses and cooler regions seeing higher gains. Currently, even if growers exploit cultivar diversity, top-producing countries, particularly in Southern Europe, are predicted to sustain major declines of suitable winegrowing areas, with minimal gains (Fig. 1 C and D). For example, Spain and Italy are expected to lose 65% and 68%, respectively, of climatically suitable areas, under a 2 °C warming scenario (*SI Appendix, Fig. S15*), with gains of only 5% and 9% (respectively). France is projected to see balanced losses (22%) and gains (25%; *SI Appendix, Fig. S15*). In contrast, wine-producing regions in the Pacific Northwest (United States) or New Zealand expand in climatically suitable area for the latest-ripening varieties by 20 to 100% and 15 to 60%, respectively (*SI Appendix, Fig. S15*). Further, losses increase dramatically under a 4 °C warming scenario (*SI Appendix* and Figs. 1d and 2), while gains decrease. Losses at 4 °C are predicted to be particularly high in countries that are already warm; this includes losses reaching ~90% in Spain and Italy (*SI Appendix, Fig. S15*).

For regions where our results suggest that cultivar diversity may be most critical, growers must choose to actively shift varieties—which requires overcoming legal and cultural hurdles.

Currently, traditional practices, alongside regulations at local, regional, and higher levels, limit how much and where growers can shift varieties easily (19). This, coupled with other considerations, such as the temporal and related financial cost of replanting or regrafting a vineyard, may lead many growers to prefer alternative options that keep a particular variety tied to a region. For example, local management practices to reduce microclimatic temperatures or adjust the pace of development (e.g., shade cloth, reduced leaf area to fruit weight ratio, or longer-cycle rootstocks)—may help some growers (30–32), but generally work best for lower amounts of warming, especially compared to changing varieties.

Growers who want to exploit cultivar diversity would benefit from improved climate and crop-diversity data. For winegrapes, an immediate need is data on a greater number of varieties at a vineyard-relevant spatial scale. Our modeling approach requires projected climatic data at a high temporal resolution (e.g., simulated daily temperature values), which are only available at a low spatial resolution (e.g., circa 100 km² pixels), and thus cannot capture the unique microclimates of many vineyards. Our results could be expanded to finer spatial resolutions, given climate data downscaled with attention to the important climatic attributes of a particular viticultural region (e.g., coastal influences and/or cold air pools in complex terrains). Additionally, our approach requires sufficient phenological data, which we obtained for 11 varieties from a narrow geographical range (i.e., mainly France). These varieties span a diversity of phenologies (*SI Appendix, Figs. S4 and S14E*), yet they still represent less than 1% of known winegrape diversity, suggesting that benefits from cultivar diversity could be higher if more varieties were included.

Our results apply clearly to winegrowing regions, but have implications for many of the world's agricultural regions. We focused on winegrapes given their diversity and extensive data resources: winegrape harvest dates are some of the longest written records on earth (33); major repositories collect, preserve, and document the crop's diversity (20); and newly available data describe the planted geographic distribution of winegrape varieties across the globe (22). Such resources make winegrapes an excellent crop to test how intraspecific diversity may help agriculture adapt to a changing future, but many other crops also harbor high genotypic and phenotypic (e.g., morphological) diversity. Some of this diversity is obvious to consumers (e.g., historical and new cultivars of apples; ref. 13), while other diversity is hidden, present mainly in the wild or in research collections (e.g., banana and orange; refs. 34 and 35, respectively). Gathering sufficient data for tests similar to ours will be critical to identifying the full potential of cultivar turnover, but we expect that our results extend to many other crops, if growers have the flexibility and resources to shift in step with climate change.

Materials and Methods

Phenological Data for Parameterization and Validation. We assembled historical data for 50% budbreak, 50% flowering, and 50% veraison dates from 62, mostly French, locations between 1956 and 2015 (see list of data sources in *SI Appendix, Table S1*). Most observations are for the 1995 to 2007 period and, secondarily, the 1975 to 1994 period. The dataset included 517 observations of budbreak, 757 observations of flowering, and 688 veraison observations. Requirements for phenological observations were as follows: Budbreak was identified as stage 4 on the modified Eichhorn and Lorenz (E-L) scale (36); flowering was identified as the 50% flowering date corresponding to stage 23 on the modified E-L scale; and veraison corresponded to stage 35 on the modified E-L scale, where 50% of berries softened or changed from green to translucent for white cultivars, or changed color for red cultivars. These data represent a portion of the original database collected in ref. 37, which was subsequently released for use within this project and includes matched local meteorological data. To conduct an independent validation of our phenological model predictions, we used further phenological observations

from three locations in Europe and two in North America (*SI Appendix, Table S2*).

Viticulture Data—The Geography of Winegrowing. To analyze climatic suitability in current winegrowing regions, we digitized the global distribution of the major winegrowing regions of the world according to a published atlas of winegrowing (38). We considered our results within this limited region (*SI Appendix, Fig. S1*) and, also, all areas that our models classified as suitable (see below).

Climate Data. We used two different sources of climate data to build and validate our models. First, meteorological local data from weather stations (situated not more than 5 km away and not more than 100 m of difference in elevation from each vineyard), providing daily minimum and maximum temperatures (*SI Appendix, Table S1*). The average daily temperature was calculated as the arithmetic mean of the daily maximum and minimum temperature. Second, a gridded reconstruction of daily minimum and maximum land-surface air temperatures (Berkeley Earth Surface Temperatures, BEST; <http://berkeleyearth.org/data/>), based on 37,000 climate records spanning the period 1880 to 2013. We used both the BEST and local climate data to parameterize the phenological models and to test for any major parameter differences across the two datasets. We further used the BEST data to test our phenological models against validation datasets (see above in Phenological Data for Parameterization and validation), save for one site (Davis, CA), where BEST did not provide overlapping years and we instead used local station data (<http://atm.ucdavis.edu/weather/ucdavis-weather-climate-station/>). BEST data are very strongly correlated with weather-station data ($r = 0.982$; rms error [RMSE] = 1.829) and, where biased, tend to underestimate warming trends (39).

Warming Scenarios. For our climate projections, we used output from the National Center for Atmospheric Research Large Ensemble (LENS; ref. 21). The LENS is a multimember ensemble of a single general circulation model (GCM), the CESM. Each member starts from its own unique initial condition in the atmosphere, and all members are simulated with the same scenario of historical climate forcings (1920 to 2005) and a high-emissions/high-warming scenario for the 21st century (2006 to 2100; Representative Concentration Pathway [RCP] 8.5). We chose this model and ensemble because it provided some of the highest spatial-temporal resolution output available from climate models ($\sim 1^\circ$ latitude/longitude, daily projections), similar to the resolution of the BEST data on which our models were ultimately calibrated.

The LENS ensemble is also well-designed for our warming threshold approach, allowing us to sample a large number of model-years (300) at different warming levels above preindustrial temperatures to force our phenology models: $+2^\circ\text{C}$ (2039 to 2048) and $+4^\circ\text{C}$ (2076 to 2085), in addition to a 0°C (1970 to 1979) reference scenario that corresponds to a recent period where the temperature was the same as our preindustrial baseline (*SI Appendix, Fig. S5*). The median estimates for the $+2^\circ\text{C}/+4^\circ\text{C}$ warming scenarios are $+2.03/+3.99^\circ\text{C}$, with interquartile ranges of 1.99 to 2.06°C and 3.97 to 4.04°C , respectively. We used this temperature-based approach because we believe that it is easier to interpret and link to current global agreements on climate change and that it provides relevant information on potential losses and gains of climatic suitability for winegrowing at different plausible future warming levels, without tying those predictions to any time horizon in the future. Our projections, therefore, do not depend on any singular future greenhouse-gas forcing scenario and are not intended to; however, we note that the $+2^\circ\text{C}$ and $+4^\circ\text{C}$ temperature thresholds we use in this study correspond closely with the broad warming estimates (relative to preindustrial) in surface air temperature for the end of the 21st century found for the RCP 4.5 ($+2.55\text{ K}$) and RCP 8.5 ($+4.39\text{ K}$) scenarios (40).

Data and Code Availability. Raw data were generated at several large-scale facilities (see *SI Appendix* for details). Derived data supporting the findings of this study are available from the corresponding author upon request. Phenological parameterization and cross-validation was implemented with the software PMP (Version 5.0; ref. 41). All other analyses utilized custom computer R code, freely available at GitHub, <https://github.com/MoralesCastilla/PhenoDiversity> (42).

Phenological Modeling. We modeled winegrape phenology for each of the 11 varieties according to a phenological process-based sequential model, considering only pixels where each 10-y scenario had no more than 2 d below -20°C or 1 d below -30°C , which represent lower lethal temperatures for most winegrape buds (43, 44). Our approach combined model

estimates of three key stages of grapevine development: budbreak, flowering, and veraison. After comparison against alternative models (*SI Appendix, Table S3*), the budbreak stage was simulated by combining the Smoothed-Utah model (45, 46), to simulate dormancy break (accumulating chilling units), and the Wang and Engel model (47), to simulate the postdormancy phase until budbreak. Then, the Wang and Engel model was also used to simulate the accumulation of forcing units until flowering and from flowering until veraison. The curvilinear structure of both the Smoothed-Utah and Wang and Engel models reproduce the known effect of a developmental slowdown at high temperatures (48). We fixed the minimum- and maximum-temperature thresholds for development at 0 and 40°C (respectively), based on physiological thresholds well established in the literature of winegrapes (49, 50) and other species (51, 52). We calculated a single optimum temperature for each phenological stage and parameterized the chilling and forcing units required to reach each stage for each variety. See *SI Appendix* for equations and further details, including a discussion of our 40°C maximum.

To evaluate model accuracy and performance, we calculated 1) the RMSE as described by ref. 53 and 2) model efficiency; and 3) we performed a leave-one-out cross-validation calculating RMSE of prediction (RMSEP) as described by ref. 54 and 4) residual prediction deviation (RPD), which can increase comparability of metrics such as RMSEP (55).

Following this validation, we further validated our phenological models against independent observations of winegrape phenology recorded at other geographical locations (e.g., Germany, Portugal, Serbia, and the United States; *SI Appendix, Tables S2 and S8*). Our analyses show good accuracy of the fitted phenological models to other regions (*SI Appendix, Figs. S6–S9*).

Modeling Maturity. We modeled the veraison to harvest phenophase using bioclimatic envelope models (56, 57) based on a suite of bioclimatic variables and the recorded climatic conditions experienced by each variety under pre-climate change conditions. We selected eight bioclimatic variables relevant to winegrape ripening (but necessarily excluded some potentially relevant variables, see also *SI Appendix, Diurnal Temperature Range*); more details on the selection of these variables and references are provided in *SI Appendix, Modeling Maturity*. Bioclimatic envelope models are often used to characterize the climatic niche or climatic conditions under which a studied species can survive. We chose bioclimatic envelope models over alternative algorithms—e.g., MaxEnt (58) or Random Forests (59)—because they allow for direct traceability of which climatic variables are responsible for changes in suitability, leading to either gains or losses as climate changes (*SI Appendix, Figs. S16 and S17*), which was a goal of our analysis.

We fitted these models according to the climatic conditions recorded during the veraison-harvest temporal window over a 30-y normal period (1950 to 1980), within existing winegrowing regions where each of the 11 modeled varieties is cultivated (22). We chose a 30-y normal period for fitting our models to capture a period before significant anthropogenic warming with robust global climate data; this period is longer than our reference scenario (0°C), as a longer time series allows us to better characterize the climate envelope for each winegrape variety.

Calculating Climatic Suitability. Our estimates of climatic suitability are the outcome of a multistep process, characterized, first, by sequential phenological models (explained above and in detail in *SI Appendix*), ensuring that veraison occurs within adequate dates—e.g., prior to October 1 in the Northern Hemisphere (this cutoff had a negligible effect on suitability; *SI Appendix, Fig. S17*); and, second, by comparison of the bioclimatic envelope forecasted under climate change with the envelope fitted during the 30-y normal period (1950 to 1979) for each variety (explained in detail above). Then, third, we considered the percentage of model-years (using 30 members from the CESM GCM, with 10 y each compiled into 300 model-years) for each grid cell and warming scenario where our models predicted suitability for a given variety (see also *SI Appendix, Fig. S11*). All figures and calculations—unless otherwise noted—used a cutoff of 75% or more of model-years predicting that a grid cell is climatically suitable for a given variety. Thus, some regions we suggest as climatically suitable may be suitable in only some years, and other regions we suggest are not suitable may be suitable, but in fewer model-years than this cutoff.

Our modeled estimates of climatically suitable winegrowing areas under our reference scenario overlap substantially with current regions (76%), but do not capture all growing regions, and include areas where winegrapes are not currently grown. This is expected given our modeling framework (e.g., we consider only 11 varieties) and its assumption that not all potential growing areas will be—or are currently—exploited.

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Supplementary Information for

Cultivar diversity buffers winegrowing regions from climate change losses

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Supporting Information Text

Data

Phenological data for parameterization & validation

We assembled historical data for 50% budbreak, 50% flowering and 50% veraison dates from 62, mostly French, locations between 1956 and 2015 (see list of data sources in Table S1). Most observations are for the 1995-2007 period and secondarily the 1975-1994 period. The dataset includes 517 observations of budbreak, 757 observations of flowering and 688 veraison observations. Requirements for phenological observations were: budbreak was identified as stage 4 on the modified Eichhorn and Lorenz (E-L) scale (1); flowering was identified as the 50% flowering date corresponding to stage 23 on the modified E-L scale; veraison corresponded to stage 35 on the modified E-L scale where 50% of berries softened or changed from green to translucent for white cultivars, or changed color for red cultivars. These data represent a portion of the the original database collected in Parker *et al.* (2), which was subsequently released for use within this project, and include matched local meteorological data.

Our phenology dataset included data for 11 common winegrape cultivars (varieties): Cabernet-Sauvignon, Chardonnay, Chasselas, Grenache, Merlot, Monastrell (syn. Mourvedre), Pinot noir, Riesling, Sauvignon blanc, Syrah and Ugni blanc. These varieties were selected based on their widespread cultivation (all are globally planted and many are considered international varieties, 3, 4), and ensuring that the set of selected varieties encompassed representative phenological diversity (Fig. S4). All are 100% *V. vinifera* subsp. *vinifera* varieties.

The above-described phenology dataset was utilized to parameterize models designed to forecast phenology across space and time, and then validate them via cross-validation methods (see *Methods* below). To further validate phenological models we used data independent from the data used to parameterize the models. We assembled a validation dataset of phenological observations from the following sources: (a) published articles on winegrape phenology containing observations for budbreak, flowering and veraison (i.e., 5-7); (b) phenological observations extracted from a report sponsored by the Napa Valley Vintners (8) (see https://napavintners.com/about/docs/nvv_climate_exec_summary.pdf; data not publicly available); and (c) our own phenological data collected at the University of California, Davis (9). These data sources comprise a dataset for five sites (which we refer to as the ‘independent validation dataset’), containing all analyzed varieties except for Grenache, and included observations from 1949 to 2016. Note that the dataset does not contain observations for all varieties and all years in each site (see Table S2).

Finally, we also used phenological data from INRA Domaine de Vassal Grape Collection to put the phenology of our selected cultivars into context by comparing them with a wider diversity of cultivars (130 others, selected to capture major varieties and the phenological diversity of winegrapes). The Domaine de Vassal Grape Collection, currently located in Marseillan-plage (France), is dedicated to the conservation, characterization and valorization of grapevine genetic resources. As such it grows many varieties of *Vitis vinifera* subsp. *vinifera* and records phenology relative to a standard variety (Chasselas, 10). For more information visit: INRA Vassal-Montpellier Grapevine Biological Resources Center at www6.montpellier.inra.fr/vassal and French Network of Grapevine Repositories https://bioweb.supagro.inra.fr/collections_vigne/Home.php?l=EN.

Viticulture data—The geography of winegrowing

To analyze climatic suitability in current winegrowing regions we digitized the global distribution of the major winegrowing regions of the world according to a published atlas of winegrowing (11). We considered our results within this limited region (Fig. S1) and, also, all areas our models found were climatically suitable (see below). The dataset on major winegrowing regions is available at <https://knb.ecoinformatics.org/view/doi:10.5063/F1SX6BH1> (12).

Climate data

See the main text *Materials & Methods*.

Data availability

Most data used in the paper are already published and publicly-available, including: climate projections (13), historical climate data from BEST <http://berkeleyearth.org/data/>, winegrowing regions distributions (3, 12), phenological data (2, 5-9); these data can be made available upon request of the authors of each publication. Derived data supporting the findings of this study are available from the corresponding author upon request.

Methods

General workflow & rationale

To forecast winegrape phenology we followed a multi-step procedure. In our first step, using our phenological dataset (Table S1) and BEST climate data, we parameterized phenological models for 11 winegrape varieties. (Note that we did not find major differences between models based on BEST data compared to the meteorological observations, see Table S6.) We chose BEST to parameterize the phenological models for several reasons. First, BEST is one of the few datasets that provide global, gridded, and quality-controlled estimates of daily maximum and minimum temperatures—i.e. using daily temporal resolution is a requirement for our phenological models. Second, the BEST data are gridded at a similar resolution to the LENS, facilitating

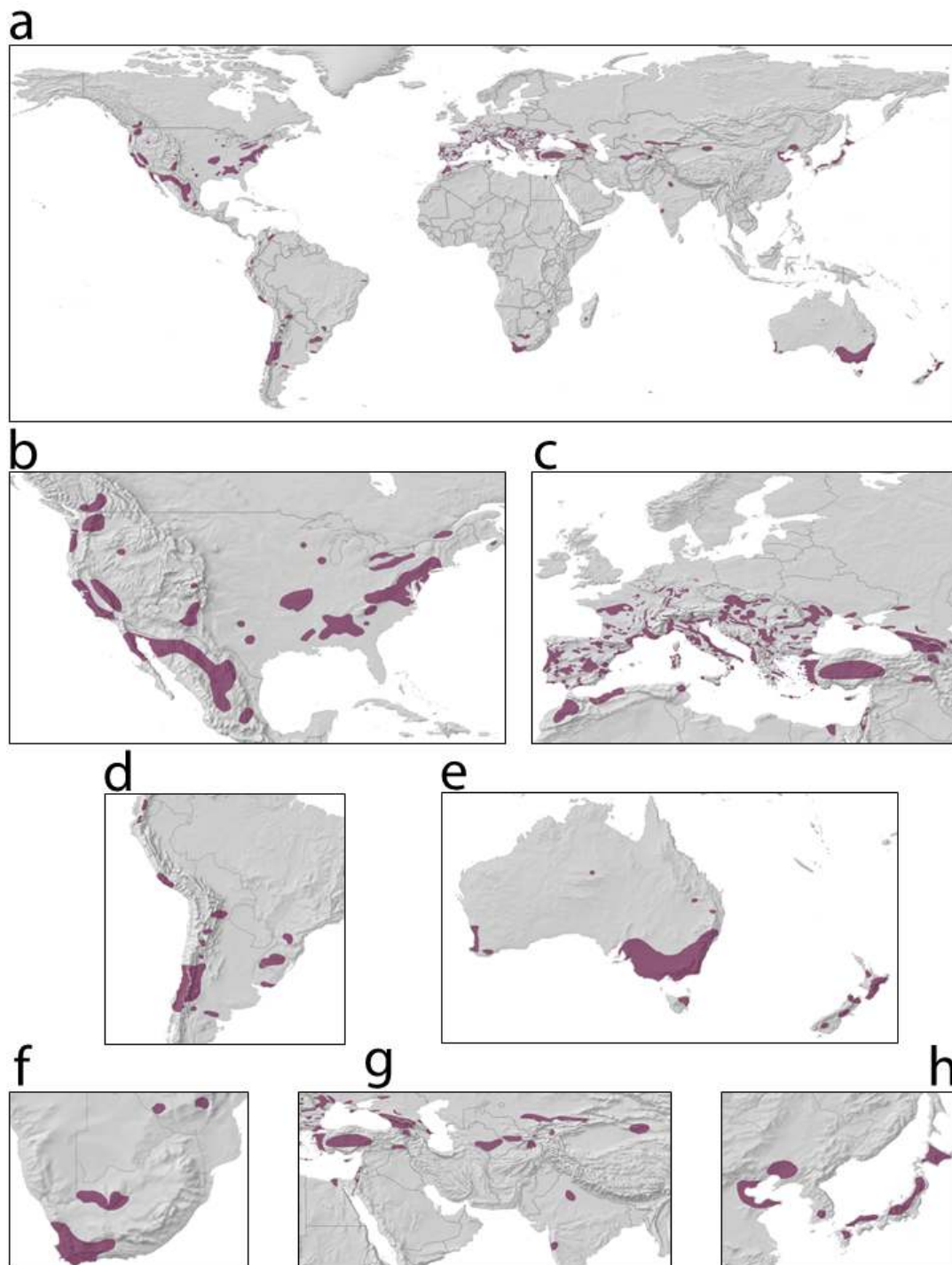


Fig. S1. Maps of current major winegrowing regions of the world (digitized from (11)). The panels show the distribution of winegrowing regions globally (a), in North America (b), Europe and Northern Africa (c), South America (d), Australia and New Zealand (e), Southern Africa (f), Southeast Europe, Middle East and South Asia (g), and Eastern Asia (h). Winegrowing regions are shaded in purple. Note that the map is not comprehensive, it lacks regions developed after 2007 and that some regions depicted only have marginal winegrape production (e.g., southeastern US, central Mexico, areas of Mongolia, north island of Japan).

1) straightforward bias correction of the model climate information and 2) re-gridding of the BEST datasets to the same spatial resolution as the LENS climate projections. We fitted phenological models to estimate the dates at which budbreak, flowering and veraison occur, sequentially—i.e. each phenological stage starts on the predicted date for the preceding stage. In our second step, we validated the models by comparing results of observed vs. modeled winegrape phenology for each stage, variety and region (using the independent validation dataset described above). In our third step, using the validated models, we projected phenology through to veraison for all 11 varieties during our reference scenario (0°C), and the two warming levels of 2°C and

4°C (see *Climate data*; Fig. S5). Finally, we used bioclimatic niche models to predict the climatic suitability to reach maturity of each variety in each region and during the 45 days following veraison (more details given below in *Modeling maturity*). Given these results we could then compare predictions for the climatic suitability of each variety across sites and warming scenarios.

Modeling multiple varieties' specific responses to future climate has two major advantages for growers. First, beyond forecasting geographic shifts in climatic suitability for viticulture, which has already been done (e.g., 14, 15), it allows establishing which particular regions will become most suitable to grow which variety. This information could guide a grower's decisions by, for example, focusing on a popular variety such as Pinot noir, and forecasting whether the grower's land will present suitable conditions to grow it under different warming scenarios (Figs. 1 and S13). While this approach has been done previously for a single variety (e.g., 16), it is rarely done for more than one at a time (but see 17, for an example using growing season mean temperatures). Second, having multiple variety-specific results allows us to build variety diversity maps (Figs. 1b and S13b). These maps yield more explicit information of the potential of winegrape diversity with climate change: the more varieties forecasted to reach maturity in a given region, the more resilient that region will be to losing climatic suitability for viticulture. Finally, modeling multi-variety responses to climate change is required to quantify the adaptive potential of within-crop agrobiodiversity.

Phenological modeling

We modeled winegrape phenology for each of the 11 varieties according to a phenological process-based model (which treated each stage sequentially), considering only pixels where each 10-year scenario had no more than two days below -20°C or one day below -30°C (18, 19). Freezing temperatures below -20°C have been shown to generate tissue damage in grapevine (18), and thus, these thresholds are necessary because as climate warms high latitude locations are predicted to start providing the necessary heat during the growing season for winegrowing, but winters may still be too cold.

Our approach combined model estimates of three key stages of grapevine development: budbreak, flowering and veraison. The budbreak stage was simulated using a process-based sequential model combining the Smooth-Utah model (20, 21), which simulates dormancy break (accumulating chilling units) and the Wang and Engel model (22), which simulates the post-dormancy phase until budbreak. Then, the Wang and Engel model was used to simulate the accumulation of forcing units until flowering, and after flowering, it was used to simulate accumulation of forcing units until veraison. The curvilinear structure of both the Smoothed-Utah and Wang and Engel models reproduce the known effect of a developmental slowdown at high temperatures (23).

The Smoothed-Utah model (20, 21) assumes that chilling can only occur within a given range of temperatures, and allows for negative chilling in warm days:

$$F_{SmoothedUtah} = \begin{cases} \frac{1}{1 + e^{-4 \frac{T_d - T_{m1}}{T_{opt} - T_{m1}}}} & \text{if } T_d < T_{m1} \\ 1 + \frac{-0.5(T_d - T_{opt})^2}{(T_{m1} - T_{opt})^2} & \text{if } T_{m1} \leq T_d < T_{opt} \\ 1 - \left((1 - min) \frac{(T_d - T_{opt})^2}{2(T_{n2} - T_{opt})^2} \right) & \text{if } T_{opt} \leq T_d < T_{n2} \\ min + \left(\frac{1 - min}{1 + e^{-4 \frac{T_{n2} - T_d}{T_{n2} - T_{opt}}}} \right) & \text{if } T_d \geq T_{n2} \end{cases} \quad [1]$$

Where T_{m1} is decrease in cold efficiency for bud endodormancy, T_{opt} is the optimal mean daily temperature, T_{n2} is the temperature with half the efficiency of T_{opt} to induce endodormancy, and min is the negative impact of high temperatures. This model yields the amount of chill units based on daily temperatures (T_d), which are accumulated until a threshold of accumulated chilling action (C^*) is reached defining the end of endodormancy.

The Wang and Engel model (22) belongs to the family of beta functions, is asymmetric and has four parameters:

$$f_{Wang \& \text{ Engel}} = \begin{cases} \frac{2(T_d - T_{min})^\alpha (T_{opt} - T_{min})^\alpha - (T_d - T_{min})^{2\alpha}}{(T_{opt} - T_{min})^{2\alpha}} & \text{if } T_{min} < T_d < T_{max} \\ 0 & \text{if } T_d \leq T_{min} \text{ or } T_d \geq T_{max} \end{cases} \quad [2]$$

$$\alpha = \frac{\ln(2)}{\ln\left(\frac{T_{max} - T_{min}}{T_{opt} - T_{min}}\right)} \quad [3]$$

Where T_{min} , T_{opt} , T_{max} represent the minimum, optimum and maximum temperature values, respectively, that are included to calculate the amount of forcing needed until a threshold of accumulated forcing units (F^*), is reached for a given phenological event—i.e., budbreak, flowering or veraison—to occur. Following many other crops and approaches in winegrapes (2, 24), we consider T_{min} , T_{opt} , T_{max} to be species-level characteristics and thus define one value for each parameter that is repeated across varieties. This approach is consistent with the observation that the precocity hierarchy across varieties is generally constant

within a site across years (as opposed to shifting as expected if $T_{min}, T_{opt}, T_{max}$ vary strongly across varieties), allows us to compare varieties directly, and prioritizes robustness over precision given our data limitations (i.e., at high temperatures we have very few observations for any one variety). Our estimated parameters (see Tables S4-S6) were in line with values estimated by previous research (25). In our modeling approach here, the thresholds for minimum and maximum temperatures (T_{min} at 0°C and T_{max} at 40°C for the Wang & Engel model) were fixed according to physiological thresholds well-established in the literature of winegrapes (26, 27) and other species (24, 28). We set T_{max} at 40°C after comparison of model fit against models using alternative T_{max} values of 36, 37, 38, and 39°C. Values of T_{max} comprised between 35 and 40°C are common in the literature (26, 27, 29, 30), which often shows a drastic reduction in photosynthetic activity above 40 °C (31–33), and thus in phenological development. We acknowledge that phenological models relying upon threshold parameters such as T_{min} and T_{max} may predict different responses above and below the thresholds—e.g. in our example, forcing units are accumulated at T_{max} of 39.9 but not at 40.1°C. However, research on winegrape physiology seems to support the existence of temperature thresholds above which damage may be too high for further plant development (31–33). As new data becomes available it will be interesting to see how phenological models accommodate uncertainty around their threshold parameters. While T_{min}, T_{opt} and T_{max} were fitted or set at the species level, the accumulated thresholds for chilling (C^*) and forcing (F^*) corresponding to each phenological stage were parameterized at the variety-level (see Tables S5-S7), thus allowing for phenological variation across varieties.

To evaluate model accuracy and performance we calculated: 1) the root mean square error (RMSE, providing a classical evaluation between observed and simulated values) as described by (34), 2) model efficiency and, 3) we performed a leave-out-one cross-validation calculating RMSEP (root mean square error of prediction) as described by (35, 36):

$$N_{PS}(i) = f(X(i), \theta(i)) \quad [4]$$

RMSE in our approach yields the number of days our predictions deviate from the average observed (thus RMSE = 0 would correspond to a perfect fit).

$$RMSEP = \sqrt{\frac{1}{N} \sum_{i=1}^N (N_{PS}(i) - N_{PO}(i))^2} \quad [5]$$

RMSEP represents the quadratic distance between the observed (N_{PO}) and the simulated (N_{PS}) phenological stages (here, budbreak, flowering and veraison). RMSEP and RMSE are calculated the same but the result of the former specifically informs of the error of prediction in the cross-validation and the latter computes the calibration error; f represents the model (Smoothed Utah or Wang and Engel) and $X(i)$ and $\theta(i)$ are the vectors of the input data and parameters for the observation i . As with RMSE, the lowest RMSEP are indicative of higher accuracy.

The model efficiency (37) can be defined as:

$$EF = 1 - \frac{\sum_{i=1}^N (N_{PO}(i) - N_{PS}(i))^2}{\sum_{i=1}^N (N_{PO}(i) - \overline{N_{PO}})^2} \quad [6]$$

Where $\overline{N_{PO}}$ is the mean value of the observed phenological stages.

Additional accuracy metrics have been proposed to standardize RMSE with respect to the standard deviation of observed values. Here we calculated the Residual Prediction Deviation (RPD), defined as:

$$RPD = \frac{\sqrt{\frac{1}{N} \sum_{i=1}^N (N_{PS}(i) - N_{PO}(i))^2}}{RMSE} \quad [7]$$

RPD is applicable to observed and predicted values that are normally distributed, and can increase comparability of metrics such as RMSE. It is also fairly easy to interpret as prediction accuracy increases with increasing RPD (38).

The combination of the above models was selected after rigorous tests of alternative modelling approaches. The alternative models tested for the dormancy phase were the BRIN model (39) and the Chuine model (40); alternative models for post-dormancy included the Richardson model (20) both in its constrained and unconstrained version and the sigmoid model (41). Our choice of the Smoothed Utah and Wang & Engel models for budbreak was made according to accuracy metrics (e.g. RMSE, EF and AIC) showing that their combination outperformed alternative options in terms of fit and parsimony (see Table S3). Although the Smoothed-Utah + Sigmoid combination performed similarly well, we favored the Wang & Engel model as it is more realistic biologically—e.g. it does not accumulate forcing units at temperatures below 0°C. The Wang & Engel model has the additional benefits of reducing uncertainty (42), and being more realistic biologically for budbreak, flowering and veraison by replicating the common plant biology of slowed development at higher temperatures (24, 42). We note that while

this approach may not be perfectly accurate it provides a reasonable trade-off between prediction accuracy, strong physiological foundations and simplicity to upscale computationally.

Following the above described validation metrics, we additionally validated our phenological models against independent observations of winegrape phenology recorded at other geographical locations (e.g. California in the USA, Germany, Portugal, Serbia, see Table S2 for data sources) from the locations where calibration data were extracted (e.g. predominantly France in Table S1; see Tables S8-S9 for validation results). For locations in California, where we had raw data on phenological observations—i.e. yearly values of any phenophase for a given variety—validation was straightforward—we calculated the RPD and RMSE based on observed vs. the simulated values by our phenological models. For European locations where we had only the mean \pm standard deviation values of observed phenology, validation was restricted to comparing these means and errors to those resulting from our predictions for each given period and dataset (see Table S8).

Our cross-validation and independent validation datasets provide estimates of model uncertainty for our global projections, which encompass many regions for which we did not have phenological data. Our core validation method—cross validation of our main dataset (Table S1)—show RMSE of 5.54 ± 1.64 days and RPD of 2.43 ± 0.84 for BEST climate data (all estimates are mean \pm SE unless otherwise noted). These values are well within margins generally considered a good fit (2, 39, 43–45). While we note that growers would benefit from an even lower RMSE, these values are limited by how accurately phenology can be observed, especially when integrating across diverse datasets, as we do here. Moreover, our additional validation with independent observations from North American sites with high resolution data—i.e. yearly observations of each phenophase—showed equal or reduced uncertainty (average RMSE of only 2.55 ± 1.04 days in Napa Valley, see Table S8, and RMSE of 2.50 ± 0.71 days in Davis). Correlations between the phenology simulated by our models and independently observed winegrape phenology were high for both the North American and European locations ($R^2 > 0.98$; see Figs. S6-S8). Even for European locations with low temporal resolution data—i.e. we had only mean and SD of phenology across a set of years—but measuring the same phenophase (i.e., 5, 6), we found that our averaged predictions overlap with average observed values (see Fig. S8). Our independent validation results are, however, inherently limited in their scope—i.e. they are generated for 5 locations where not all modelled cultivars are planted—given the current availability of phenological data across winegrowing regions. Yet we argue that our validation analyses show that our fitted phenological models can predict phenology in other regions accurately enough (Figs. S7-S6) to meet our research goals: compare phenological forecasts across cultivars, regions and warming scenarios.

Modeling maturity

Phenological modeling of winegrape maturity is particularly challenging for several major reasons. First, date of harvest responds to a mix of decisions related to the chemical composition of the berries—e.g. sugars, acids, phenolic content, aroma precursors and aromas—that mainly develop during the veraison to harvest period. Thus, while measures of budbreak, flowering and veraison are strongly related to physiological processes, harvest timing is often closely linked to growers and winemakers' preferences in winestyle, or even historical and cultural determinants. Models of maturity are thus often based on sugar measurements (e.g., 46), but data for such models are extremely limited, and recent studies suggest sugar may change separately from acids and other juice compounds with warming (47, 48). Further, while temperature alone can suffice to model the preceding phenological stages (49), more complex interactions among climatic variables, including those related to vine water status (39, 50), are often needed to correctly characterize winegrape maturation.

Thus, we modeled the veraison to harvest phenophase using bioclimatic envelope models (51, 52) based on 8 bioclimatic variables we expected were relevant globally to grape ripening (see also *Diurnal Temperature Range* below) and the recorded climatic conditions experienced by each variety under pre-climate change conditions. Such models are often used to characterize the climatic niche or climatic conditions under which a studied species can survive.

We used the following set of bioclimatic variables (indicators) known to be relevant for winegrowing:

1. Growing Degree Days (GDD) – defines the accumulation of temperatures during which active grapevine growth is expected. Here we consider only temperatures above 10°C , which is the classical threshold to define GDD in grapevines (53, 54), and calculate it starting from the estimated veraison date.
2. Number of days on which maximum temperatures exceed 40°C (temperature above 40°C) – are generally assumed to severely retard or stop photosynthesis, growth and developmental activity (26, 27, 29, 30).
3. Number of days on which minimum temperatures are below 10°C (number of days below 10°C) – as for the previous indicators we consider there is less photosynthetic activity below 10°C (29, 30).
4. Minimum daily temperatures – minimum temperatures during maturation period are used to describe night temperatures, which are well related to good development of secondary compounds (e.g. aromas and color, 55). For each variety, we recorded the distribution of minimum temperatures corresponding to each day within the 45-day period following veraison, and across the 30-year normal period (1950-1980).

5. Maximum daily temperatures – these values are related to sugar content, acidity degradation, physiological problems and *cooked fruit* aromas (if temperatures are too high, 56, 57). For each variety, we recorded the distribution of maximum temperatures corresponding to each day within the 45-day period following veraison, and across the 30-year normal period (1950-1980).
6. Growing Degree Days above the optimal developmental temperatures. Here we consider 25°C as the base temperature, which is coherent with the optimal value of the phenological models (see *Phenological modeling* above; and (58)).
7. Total summed precipitation (mm) – this value has been often linked to impacts on the final yield, and is associated with pests and diseases (59) but also with overall wine quality (50).
8. Maximum daily precipitation: similar information to total summed precipitation, but also helps differentiate between high precipitation due to many smaller events versus one large precipitation event (calculated as the maximum daily precipitation registered within the 45-day period for each year of the 30-year normal period we considered).

Some of these indicators are known to be collinear—i.e. to varying degrees across the different regions where cultivars are planted—and yet we chose them because they can be linked to particular physiological and developmental processes which are relevant for ripening (see above). We found collinearity to be higher for precipitation indicators (see Table S10) but even so, their relative importance to determine loss of climatic suitability differed significantly for most cultivars (see Fig. S17). We present results for all the variables (and an additional analysis where we include diurnal temperature range, see below) as they are relevant to different aspects of the ripening process (e.g., GDD represents accumulated temperatures over the ripening window while maximum and minimum temperature represent extremes). We note, however, that analyses using only a subset of these variables may find different drivers of climatic suitability (e.g., an analysis with few temperature-related metrics outside of GDD would likely find a greater importance of GDD to climatic suitability). The choice of threshold-based indicators was robust to the specific thresholds used—e.g. the effects of the number of days above 40°C on climatic suitability loss were strongly correlated with the effects yielded by the number of days above 39°C ($R^2 = 0.93$).

The existing winegrowing regions for each of the 11 modeled varieties were extracted from (3) according to the following criteria: 1) a minimum of planted 100 *ha*, thus guaranteeing the ability of those regions to adequately support a given cultivar, 2) the target variety represented at least 2.5% of the total area planted globally for that variety, and 3) to minimize prevalence effects derived from certain varieties being distributed in more locations than others (60), we ensure that each variety was modeled based on a minimum of 20 locations (except for Monastrell, for which $n=10$) and to avoid over-representation of most common varieties—e.g. Cabernet-Sauvignon—we trimmed down the number of locations to a maximum of 120 (Fig. S10). Finally, we cross-validated the models by re-running the models based on randomized subsampled training (70%) and testing (30%) datasets, and comparing results against observations for the 30-year normal period (1950-1980). Selection of the geographical distribution for each variety could, potentially, improve if quality criteria was utilized (i.e. considering only regions where a given variety is known to produce high quality wines). Such data, however, are not available across all areas where each variety is planted. Our approach is thus conservative in focusing on climatic criteria exclusively: it identifies where a given variety would reach maturity regardless of production or quality criteria.

Our bioclimatic envelope approach characterized the distribution of each of the above climatic variables for each winegrape variety over the 45 days following veraison, given estimates over our 30-year normal period. Certain varieties may show shorter or longer ripening periods in different geographic locations, but a lack of available data prevented us from incorporating such variation in our envelopes. Thus we assume a 45-day ripening period across varieties and regions, and tested the sensitivity of our results to this assumption (Figs. S20-S21). Then, for each 16,000 land pixels of the world we characterized the forecasted daily distribution of each climatic variable for each year within each warming period—e.g. 2°C, and 4°C—and our 0°C reference period. Finally, we classified each climatic variable as suitable or unsuitable for the target variety to reach maturity according to whether or not forecasted values for that variety fell within the 90% quantiles of the distribution. We set a 75% threshold for a given pixel to be classified as suitable, meaning that at least 6 out of the 8 climatic variables should have forecasted values within their distribution over our 30-year normal period. This approach is rather permissive—i.e. high quantile allowance and low threshold—in order to account for inter-year climatic variability and to reduce the amount of false negatives. Given the 30 ensemble members of our GCM we had a total of 300 estimates ($300=30$ GCM members \times 10 years for each warming period) of climatic suitability for each pixel and each warming scenario.

Climatic suitability

See the main text *Materials & Methods* for how we calculated climatic suitability.

Our models of climatic suitability overlap significantly with current growing areas but exclude some areas; such areas may use unique practices to maintain vines, may use varieties very different than those we modelled (e.g., many areas in the southern and mid-west United States use hybrid vines that are not 100% *V. vinifera* subsp. *vinifera*), and/or may be areas our models suggested were suitable for winegrowing in too few years. Additionally, our estimates contain areas where winegrapes are not currently grown, including areas that have been highlighted in other studies of winegrowing areas (e.g., 17). We quantify gains and losses of climatic suitability for winegrowing by comparing the forecasted suitability under 2°C and 4°C with comparisons

to: (1) current growing regions and (2) areas our approach identified as climatically suitable under our 0°C reference period. Given that our models do not (and are not designed to) predict only areas currently growing winegrapes, estimates against current winegrowing areas could under or over-estimate changes. Thus, our two comparisons (current winegrowing regions and all climatically suitable regions under 0°C reference) capture both change relevant to current areas, and to our particular modeling approach.

Code availability and software used

Phenological parameterization and cross-validation was implemented with the software PMP v5.0 (61). All other analyses utilized custom computer R code, freely available at <https://github.com/MoralesCastilla/> under the GNU General Public License.

Additional simulations & tests

Sensitivity analyses of indicators included in bioclimatic niche models

To test for the sensitivity of our results to including precipitation variables—e.g. Total summed precipitation, Maximum precipitation—in the maturation stage, we also modelled maturity based only on temperature variables (see above section on *Modeling maturity*): Growing Degree Days, Number of days above 40°C, Number of days below 10°C, Minimum daily temperatures, Maximum daily temperatures, Growing Degree Days above a threshold of 25°C. The ability of cultivar diversity to decrease the loss of climatically suitable regions holds when only temperature variables are accounted for to model maturity, regardless whether all potentially suitable regions (see Fig. S14b), or current growing regions of the world are considered (see Fig. S14d). Notably, by not accounting for precipitation variables in the models, the total amount of regions loss for any one single variety is higher, reaching 70% of average losses under the 2°C warming scenario (see Fig. S14d).

We also assessed whether our estimates of regions lost were sensitive to including minimum and maximum temperatures in our models. We found that our absolute estimates of regions lost were sensitive to these two variables being included—predicted losses are higher if minimum and maximum temperatures are considered (Fig. S14c)—however, our relative estimates were not: the overall pattern of cultivar diversity decreasing the loss of climatically suitable regions by almost a half was robust to whether these parameters were included in the model (Fig. S14).

Diurnal Temperature Range

Our bioclimatic niche models account for a wide array of indicators relevant for viticulture, but ignore additional potentially important indicators such as the daily variation in temperature. Shifts in Diurnal Temperature Range (DTR), mostly during the ripening phase, have been linked to major viticultural properties such as winegrape quality (62, 63), and berry development and chemical characteristics (64). Evidence for consistent directional shifts in DTR with climate warming is, however, scarce. Based in our future climate projections for a 2°C warming scenario (see above *Climate data*) we found that, shifts in DTR with 2°C of warming will not exceed 0.5°C locally and may affect the world's winegrowing regions unevenly—e.g. slight decreases in central Europe vs. mild increases in Southern Europe (Fig. S2). Future work on winegrape phenology may benefit from accounting for DTR in models and projections.

Although we do not include this variable in our bioclimatic niche models, we tested for the sensitivity of our results to dismissing DTR. Specifically, we re-ran our bioclimatic niche models for maturity of all 11 cultivars globally, adding DTR to our set of 8 bioclimatic indicators (see *Modelling maturity* above) and projected shifts under a +2°C warming scenario (results were averaged across five randomly selected simulations of the LENS model—i.e. simulations 5,7,14,19 and 26). Results show that DTR is less important for region loss than most other indicators, including precipitation variables (Fig. S18). Although low, the influence of DTR in determining climatic suitability appears important and should be considered in future studies, especially those at higher spatial resolutions or in regions known to be more influenced by DTR (e.g., 62).

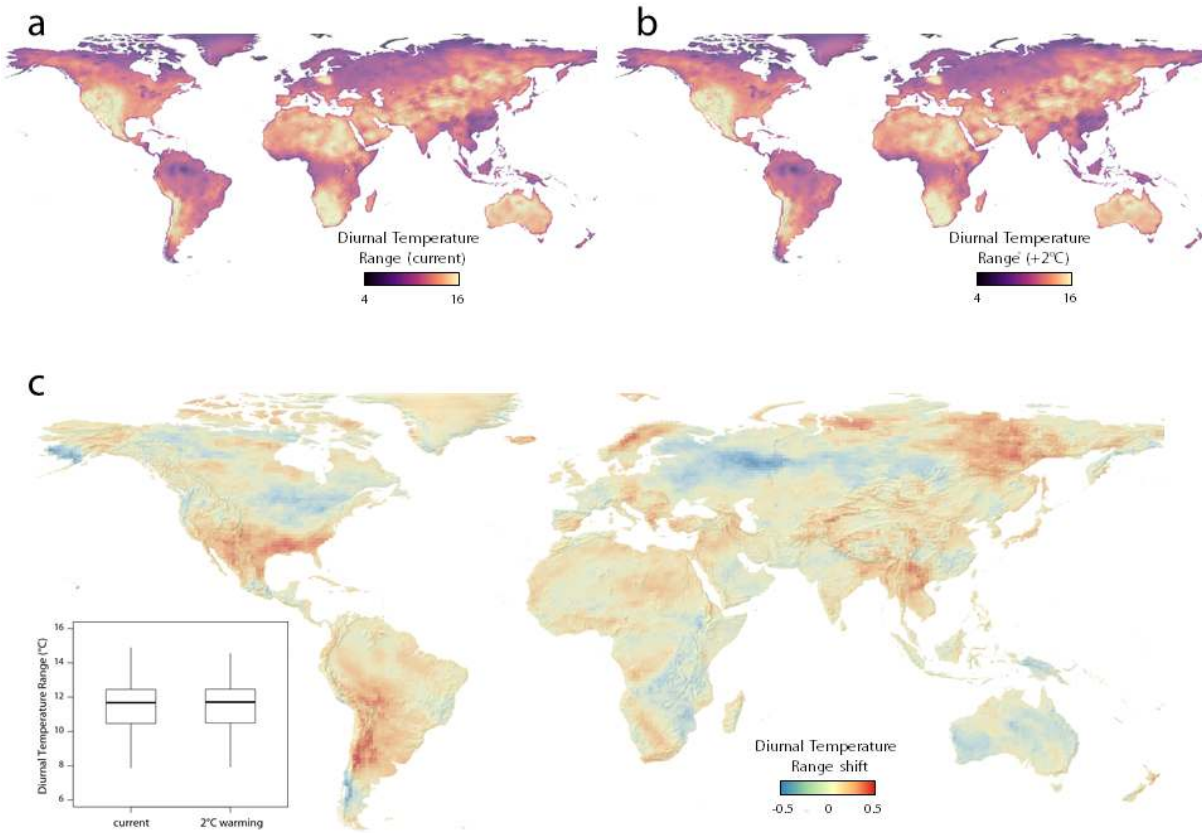


Fig. S2. Maps of Diurnal Temperature Range (DTR) measured globally for a 10-year period (2006-2015) representative of current climatic conditions (a), and the 10-year period (2039-2048) corresponding to the +2°C warming scenario (b). Shifts in DTR are calculated as the difference between current and projected DTR (c). The boxplot inset compares the distributions of DTR globally under both current and future climate, showing no significant differences in median DTR before and after warming. Across locations, however, recorded DTR shifts range between +0.5°C and -0.5°C with respect to current climatic conditions. These results are for one randomly selected GCM simulation of the LENS model.

Gains of climatic suitability

Our results identify a substantial impact of climate change in reducing the areas of both current winegrowing regions and other regions where wine is not cultivated but where climate would allow so. However, previous research (e.g., 17) has also highlighted that large extensions of territory may become suitable for winegrowing due to climate change. While gains of climatic suitability due to climate change are considerably lower than losses under both 2°C and 4°C warming scenarios (Fig. S14), these patterns are highly heterogeneous across geographic regions. For example, gains of climatic suitability were rather limited in Southern European countries like Spain, France or Italy (Fig. S15), and only occurred for varieties with later phenologies (e.g. Grenache, Monastrell or Ugni blanc, see Fig. S15). In contrast, gains of climatic suitability were highest in New World regions such as the Pacific Northwest (USA), New Zealand or California, not only for late varieties but also for earlier ones (e.g. Chardonnay, Riesling or Chasselas, see Fig. S15). Interestingly, gains of climatic suitability in the latter regions are highest under a 4°C warming scenario, but in European countries, most opportunities to gain climatic suitability coincide with the 2°C warming scenario. This information would be relevant for winegrowers from the regions with high potential to increase their suitability, by informing them of the varieties that may be best adapted to grow there in years to come.

Cultivar turnover also has a detectable effect on gains of climatic suitability (Fig. S19). The potential for gains of climatic suitability for any one variety is consistently higher under the 2°C than under the 4°C warming scenario (Fig. S19). Increasing diversity—i.e., allowing for cultivar turnover—from one to 11 varieties increases the amount of gains by 19.8% (i.e. from 80.2% of average gains when only one variety is considered to 100% gains when all are considered), under 2°C of warming (Fig. S19). A similar pattern, but showing a greater effect of turnover is found under 4°C of warming, with 43.1% increase in climatic suitability (i.e. from 56.9% of average gains when only one variety is consider to 100% gains when all are considered). These results thus show that our modeled varieties will have a larger potential to occupy novel growing regions under the 2°C scenario (i.e. climate suitability gains for winegrowing are greater under less warming), and also that the relative effects of cultivar turnover on suitability gains would be greater under 4°C of warming scenario.

Late-ripening composite (LRC) variety

To examine how including a much later-ripening cultivar impacts results, we parameterized and tested a hypothetical extremely late variety, built by combining our models with information on several late-ripening varieties. Our subset of 11 varieties captures significant phenological diversity (Fig. S4), but does not represent the full phenotypic variability existing within the genetic diversity of *Vitis vinifera* subsp. *vinifera*. In particular, our range of varieties fails to represent the later maturation exhibited by very late varieties (e.g. Nero amaro, Roditis or Maratheftiko). To include such phenotypic variability in our analyses, we simulated a virtual variety in an analogous way to previous approaches (65). To compute the parameters of this virtual variety, we find the maximum difference in C*, F*BB, F*Flo and F*Ver among the varieties included in our dataset, and either sum it or subtract it to the value of the variety with either maximum or minimum values for the parameters, respectively. Subsequently, for these set of parameters, we conducted all analyses, replicating the above described procedure for our 11 actual varieties.

The last step of our modeling workflow involves fitting bioclimatic envelopes to simulate maturity (see above), which requires data on the geographic distribution of each variety in order to estimate the climatic niche or the environmental space occupied by each variety over the 30-year normal period (1950-1980). Since we did not have distributional data for the hypothetical Late-ripening composite variety (LRC), we utilized as a surrogate the following varieties for which we had comprehensive enough distributional data, amongst those with the known later phenologies: Fogarina, Lambrusco Di Sorbara, Savatiano, Verdelho Tinto, Brun Argente, Amaral, Uva Cao and Verdicchio Bianco. Overall we selected 51 localities where these varieties are distributed, across Greece, Italy, France and Portugal (Fig. S3), that were later used to fit bioclimatic envelopes for the LRC variety.

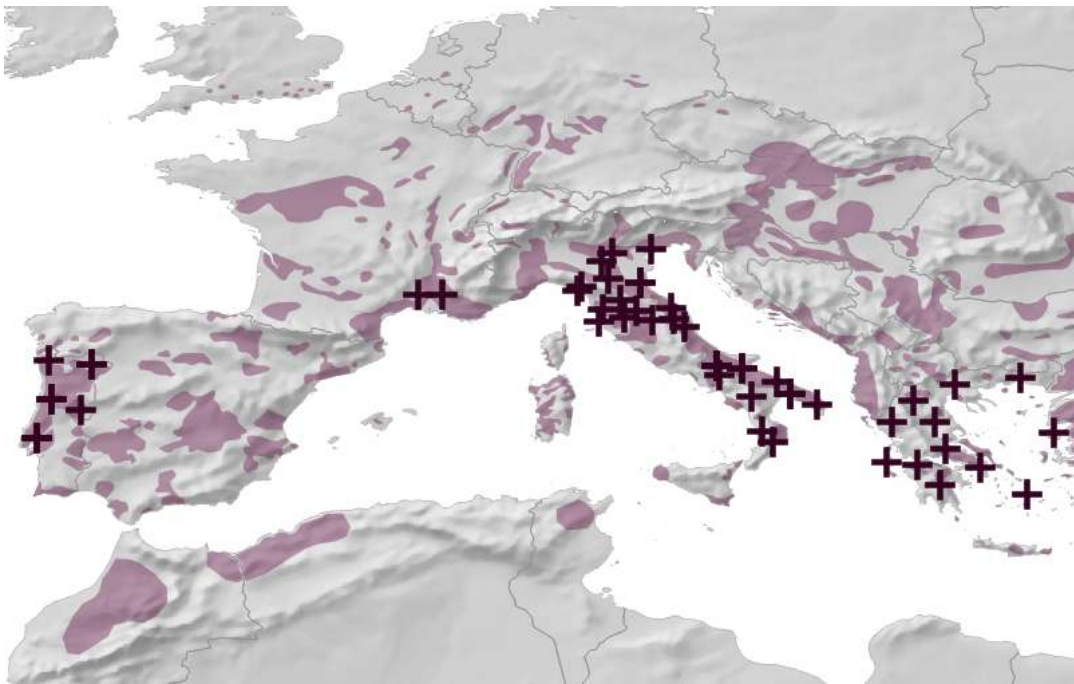


Fig. S3. Simulated geographical distribution of the Late-ripening composite (LRC) variety, based on the distribution of 8 varieties known to have late phenologies. The 51 points utilized to fit a bioclimatic envelope for the LRC variety are marked with crosses. In the background current winegrowing regions are shaded in purple.

Overall, including the LRC has an almost negligible effect on our results. Analyses of loss of climatic suitability within current winegrowing regions revealed that including the LRC decreased region loss by less than 1% (Fig. S14): under 2°C of warming, 23.6% out of all global current winegrowing regions would lose their climatic suitability for winegrowing even after including our simulated LRC. This figure does not represent a large improvement in comparison with the 24% that would be loss allowing for turnover across the 11 varieties in our study. This result supports that cultivar diversity can buffer loss of agricultural regions but such ability is not unlimited. However, modeling one virtual variety is a suboptimal option in comparison to modeling one (or several) actual late-ripening variety (e.g. Maratheftiko), which may potentially have a tolerance to high temperatures. Unfortunately, modeling further late-ripening varieties was unfeasible due to data limitations and thus, future work will benefit from systematic collection of phenological data for less-known, Southern European varieties.

Sensitivity analyses based on veraison-harvest temporal window

Our models for maturity (see above section on *Modeling maturity*) assumed for all varieties a temporal window of 45 days following veraison within which climatic variables were recorded to build the climatic envelopes. A 45-day period for winegrapes to ripen is empirically supported for most varieties, however, certain varieties have ripening periods shorter or longer than 45

days in some sites and years (25, 66). To provide some assessment of the effect that our choice of a 45-day veraison-harvest period would have on estimates for gains and losses of winegrowing regions we re-fitted climatic envelopes for *Chasselas*, the variety with the shortest veraison-harvest period in our dataset.

We first evaluated shifts in the distribution of the values of each climatic variable according to a 30-day period after veraison against a 45-day period (Fig. S20). Results show an overall close match among the climatic distributions of each variable regardless of whether the distribution is recorded over the 30 or the 45 days following the predicted veraison date. An exception to this pattern was found for Summed precipitation and Growing Degree Days, which is expected given that more precipitation and more Degree Days can accumulate in 45 than in 30 days.

In a second step we replicated our analyses quantifying climatic suitability to reach maturity for *Chasselas* within Europe based on the 30 days following veraison instead of the 45-day window employed in our main analyses. Climatic suitability was calculated under the same three scenarios that were used to estimate increases and decreases in suitable regions (by comparing the amount of climatically suitable regions at 0°C, with suitable regions at both 2°C and 4°C of warming). Results showed similar trends, but magnified estimates of both gains and losses of suitable regions (Fig. S21). Thus, if climatic envelopes were characterized across a shorter period, the proportion of climatic suitability loss would increase—e.g. from 71.2% to 81.9% on average under 2°C warming—and the proportion of current growing regions becoming more suitable to grow *Chasselas* would decrease—e.g. from 15% to 0.2% on average under 2°C warming—S21. These results confirm that our approach is conservative in estimating both losses and gains.

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Tables

Table S1. Summary of data used to parameterize phenological models. For each site for which we had data, we indicate its geographic coordinates, the minimum and maximum year of observation, the phenological stage (BB for budbreak, FL for flowering, VER for veraison) for which there was data (marked with x), and the winegrape varieties observed. Part of these data are from Phenoclim Database (INRA Vassal, Colmar, Angers, Bordeaux, Pech Rouge - <https://www6.inra.fr/soere-tempo/Ressources/Portail-de-donnees>) - Seguin (2004).

Site	Lon	Lat	Years	minYear	maxYear	BB	FL	VER	Varieties
1	4.58	48.05	4	1999	2002	x	x	x	Chardonnay, Pinot noir
2	5.14	43.55	1	2003	2003	x	x	x	Syrah
3	5.67	43.52	2	2006	2007	x	x	x	Grenache
4	5.43	43.66	1	2006	2006	x	x	x	Grenache
5	5.55	43.73	3	2005	2007	x	x	x	Grenache
6	2.65	43.18	3	2000	2002	x	x	x	Monastrell
7	4.99	47.27	2	2004	2005	x	x	x	Chardonnay, Pinot noir
8	4.41	43.78	2	2004	2005	x	x	x	Grenache, Syrah
9	4.56	44.30	5	1999	2003	x	x	x	Monastrell
10	4.49	44.14	2	2001	2003	x	x	x	Monastrell
11	4.68	43.94	4	1999	2003	x	x	x	Monastrell
12	4.70	44.01	4	1999	2003	x	x	x	Monastrell
13	4.56	43.88	5	1999	2003	x	x	x	Monastrell
14	4.43	43.80	5	1999	2003	x	x	x	Monastrell
15	-0.47	44.77	28	1974	2001	x	x	x	Merlot, Cabernet-Sauvignon
16	-0.56	44.75	1	2004	2004	x	x	x	Merlot
17	-0.16	44.89	4	2000	2003	x	x	x	Merlot
18	-0.20	44.93	3	2001	2003	x	x	x	Merlot
19	-0.19	44.92	12	1995	2006	x	x	x	Merlot, Cabernet-Sauvignon
20	-0.77	45.21	15	1990	2004	x	x	x	Sauvignon blanc
21	3.56	43.33	50	1956	2005	x	x	x	Chasselas, Monastrell, Riesling, Pinot noir, Chardonnay, Grenache, Sauvignon blanc, Syrah, Ugni blanc, Cabernet-Sauvignon, Tempranillo, Merlot
22	3.92	43.84	2	2004	2005	x	x	x	Grenache, Syrah
23	3.17	43.27	3	1997	1999	x	x	x	Cabernet-Sauvignon
24	3.17	43.61	1	2005	2005	x	x	x	Chardonnay, Syrah
25	3.32	43.34	3	1997	2006	x	x	x	Riesling, Chardonnay
26	3.10	43.58	1	2005	2005	x	x	x	Chardonnay, Syrah
27	2.91	43.56	1	2005	2005	x	x	x	Chardonnay, Syrah
28	3.42	43.46	8	1999	2006	x	x	x	Merlot, Monastrell, Chardonnay, Riesling
29	3.14	43.44	2	2003	2004	x	x	x	Grenache
30	3.28	43.48	6	2001	2006	x	x	x	Syrah, Merlot
31	3.04	43.37	5	2002	2006	x	x	x	Merlot, Syrah
32	3.20	43.52	1	2003	2003	x	x	x	Monastrell
33	3.03	43.50	1	2002	2002	x	x	x	Monastrell
34	3.87	43.80	3	2003	2005	x	x	x	Syrah
35	-0.16	47.13	22	1981	2005	x	x	x	Chasselas, Riesling, Grenache, Sauvignon blanc, Syrah, Ugni blanc, Cabernet-Sauvignon
36	3.95	49.05	7	1998	2004	x	x	x	Chardonnay, Pinot noir
37	3.98	49.02	6	1999	2004	x	x	x	Chardonnay, Pinot noir
38	7.36	48.20	40	1976	2015	x	x	x	Cabernet-Sauvignon, Chardonnay, Chasselas, Merlot, Pinot noir, Riesling, Syrah, Ugni blanc, Grenache, Sauvignon blanc
39	6.23	43.14	4	2002	2007	x	x	x	Grenache
40	6.07	43.24	1	1999	1999	x	x	x	Monastrell
41	6.53	43.25	3	2005	2007	x	x	x	Grenache
42	6.12	43.47	3	2005	2007	x	x	x	Grenache
43	5.78	43.20	1	2007	2007	x	x	x	Grenache
44	4.76	44.18	1	2004	2004	x	x	x	Grenache, Syrah
45	4.94	43.93	8	1997	2004	x	x	x	Grenache, Syrah
46	5.13	44.27	4	2000	2003	x	x	x	Grenache
47	5.06	44.08	8	1998	2005	x	x	x	Grenache, Syrah
48	4.96	44.16	4	1999	2002	x	x	x	Monastrell
49	4.80	44.21	4	2000	2003	x	x	x	Monastrell
50	5.07	44.17	4	1999	2002	x	x	x	Monastrell
51	5.24	43.86	4	2000	2003	x	x	x	Monastrell
52	5.06	43.86	4	1999	2003	x	x	x	Monastrell
53	4.81	44.08	4	1999	2003	x	x	x	Monastrell
54	4.93	44.23	3	1999	2001	x	x	x	Monastrell
55	4.93	44.23	2	2000	2002	x	x	x	Monastrell
56	5.05	44.09	2	2006	2007	x	x	x	Grenache
57	4.83	44.06	3	2005	2007	x	x	x	Syrah, Grenache
58	5.15	43.89	1	2006	2006	x	x	x	Grenache
59	4.93	43.96	3	2005	2007	x	x	x	Grenache, Syrah
60	5.18	44.12	3	2005	2007	x	x	x	Grenache, Syrah
61	5.05	43.80	2	2006	2007	x	x	x	Grenache
62	7.97	49.98	40	1974	2013	x	x	x	Pinot noir

Table S2. Summary of European and North American sources used to validate phenological models with independent observational data. For each site, we indicate its geographic coordinates, the minimum and maximum year of observation, the phenological stage for which there was data (marked with x), and the winegrape varieties observed. Data for European locations comes from published work (i.e. Bock et al. (2013); Malheiro et al. (2013); Ruml et al. (2015)) and reports means and standard deviations of the observations. Data for North American locations comes from a data report (Cayan et al. (2010) for Napa Valley) and own-collected data (Wolkovich et al. 2017), and includes raw observations of yearly timing of each phenological stage (i.e. BB for budbreak, FL for flowering, VER for veraison) for each variety.

Site.ID	Lon	Lat	Years	minYear	maxYear	BB	FL	VER	Varieties
Bock	9.87	49.83	38	1968	2010	x	x	x	Riesling
Malheiro	-9.20	39.00	22	1990	2011	x	x	x	Chasselas
Ruml	20.17	45.17	25	1986	2011	x	x	x	Cabernet Sauvignon, Pinot noir, Merlot, Chardonnay, Riesling
Napa	-122.29	38.30	19	1985	2007	x	x	x	Cabernet Sauvignon, Merlot, Chardonnay, Sauvignon blanc
Davis	-121.74	38.54	4	2013	2016	x	x	x	Cabernet Sauvignon, Chardonnay, Chasselas, Merlot, Monastrell, Pinot noir, Riesling, Sauvignon blanc, Syrah, Ugni blanc

Table S3. Comparison of combined dormancy-post-dormancy models for budbreak. Compared alternatives include the Smoothed-Utah (Richardson 1974), BRIN (García de Cortázar-Atauri et al 2009) and Chuine (Chuine et al. 1999) models for dormancy and the Richardson (Richardson 1974), sigmoid (Hanninen et al. 1990) and Wang-Engel (Wang & Engel 1998) models for the post-dormancy phase. Model accuracy was calculated through the Root Mean Square Error (RMSE), the Efficiency (EF) and the Akaike Information Criterion (AIC; Anderson et al. 1994). The number of fitted parameters is shown for the dormancy (Dorm.) and Post-dormancy (Post-dorm.).

Model	n	Dorm.	Post-dorm.	RMSE	EF	AIC
BRIN + Richardson	482	2	1	7.58	0.74	1958.70
Smoothed-Utah + Richardson	482	5	1	7.40	0.76	1941.21
Smoothed-Utah + Wang-Engel	482	5	2	7.19	0.77	1916.25
Smoothed-Utah + Sigmoid	482	5	3	7.16	0.77	1914.30
Chuine + Richardson	482	4	1	7.50	0.75	1953.03
Chuine + Richardson (free parameter)	482	4	3	7.44	0.75	1948.44
Chuine + Sigmoid	482	4	3	7.29	0.76	1929.01
Chuine + Wang-Engel	482	4	2	7.41	0.75	1943.16

Table S4. Parameterization of winegrapes to sequential phenological models from dormancy to budbreak (Smoothed-Utah model, SU), to flowering and veraison (Wang and Engel, WE). The SU model includes four parameters: 1) Tm1 or decrease in cold efficiency for bud endodormancy, 2) Topt optimal mean daily temperature, 3) Tn2 the temperature with half the efficiency of Topt to induce endodormancy, and 4) min the negative impact of high temperatures. The WE model includes three parameters: 1) Tmin, 2) Topt.1, and 3) Tmax, representing the minimum, optimum and maximum temperature values to calculate the amount of forcing, respectively (for details see Methods section above). The units for all parameters are in °C. Fitted parameters are identical across varieties.

SU.params	SU.values	WE.params	BB.values	BB.FL.values	FL.VER.values
Tm1	-6.70	Tmin	0	0	0
Topt	7.94	Topt.1	26.1	29.34	21.84
Tn2	40.58	Tmax	40	40	40
min	-0.17				

Table S5. Chilling (C*) and forcing (F*) parameters, used as thresholds above which a given phenological stage from dormancy to budbreak (C* BB and F* BB), from budbreak to flowering (F* BB-FL) and from flowering to veraison (F* FL-VER) is reached. The units are in number of days for all parameters.

Variety	C* BB	F* BB	F* BB-FL	F* FL-VER
Cabernet-Sauvignon	166.02	20.24	23.61	65.30
Chardonnay	199.01	12.50	22.26	63.28
Chasselas	196.68	12.40	24.46	53.40
Grenache	197.68	14.02	24.71	66.90
Merlot	152.14	19.91	24.96	64.45
Monastrell	216.56	7.64	28.04	62.07
Pinot noir	198.53	12.98	21.85	59.50
Riesling	185.38	14.08	22.13	64.63
Sauvignon blanc	203.48	13.06	25.34	59.27
Syrah	198.63	13.55	23.91	59.99
Ugni blanc	199.74	17.80	24.41	67.36
Late-ripening composite	232.67	23.39	29.59	70.85

Table S6. Comparison between the parameterization of winegrapes in Table S4, based on gridded climate data extracted from BEST (Berkley Earth Surface Temperatures) and a parameterization resulting from using local weather station data. As in Table S4 the models cover sequential phenological stages from dormancy to budbreak (Smoothed-Utah model, SU), to flowering and veraison (Wang and Engel, WE). Note that fitted parameters are identical across varieties. The units for all parameters are °C. Note that the Tmin and Tmax parameters were fixed at 0 and 40°C, respectively, for both climatic datasets. Values of Tmax of 40°C were selected after comparison against alternative values (e.g. 36, 37, 38, 39°C).

SU.params	SU.values	WE.params	BB.values	BB.FL.values	FL.VER.values
Tm1 local weather	-10.83	Tmin local weather	0	0	0
Topt local weather	11.13	Topt.1 local weather	28	30.92	25.08
Tn2 local weather	29.67	Tmax local weather	40	40	40
min local weather	-0.32				
Tm1 BEST data	-6.70	Tmin BEST data	0	0	0
Topt BEST data	7.94	Topt.1 BEST data	26.1	29.34	21.84
Tn2 BEST data	40.58	Tmax BEST data	40	40	40
min BEST data	-0.17				

Table S7. Comparison between the parameterization of winegrapes in Table S5, based on gridded climate data extracted from BEST (Berkeley Earth Surface Temperatures; <http://berkeleyearth.org/data/>) and a parameterization resulting from using local weather station data. As in Table S5 the (C*) and forcing (F*) parameters, used as thresholds above which a given phenological stage from dormancy to budbreak (C* BB and F* BB), from budbreak to flowering (F* BB-FL) and from flowering to veraison (F* FL-VER) is reached. The phenological stages parameterized are BB (budbreak), FL (flowering) and VER (veraison). The units for all parameters are the number of days.

Variety	C* BB	F* BB	F* BB-FL	F* FL-VER
Cabernet-Sauvignon Weather data	158.34	16.09	16.56	60.84
Cabernet-Sauvignon BEST data	166.02	20.24	23.61	65.30
Chardonnay Weather data	189.44	9.86	15.35	58.98
Chardonnay BEST data	199.01	12.50	22.26	63.28
Chasselas Weather data	193.20	10.14	17.24	49.53
Chasselas BEST data	196.68	12.40	24.46	53.40
Grenache Weather data	187.40	11.44	17.49	63.98
Grenache BEST data	197.68	14.02	24.71	66.90
Merlot Weather data	144.72	16.16	17.28	60.35
Merlot BEST data	152.14	19.91	24.96	64.45
Monastrell Weather data	171.13	15.42	20.34	60.43
Monastrell BEST data	216.56	7.64	28.04	62.07
Pinot noir Weather data	194.18	10.27	15.29	54.53
Pinot noir BEST data	198.53	12.98	21.85	59.50
Riesling Weather data	190.22	10.32	15.50	59.88
Riesling BEST data	185.38	14.08	22.13	64.63
Sauvignon blanc Weather data	192.22	10.26	17.76	55.16
Sauvignon blanc BEST data	203.48	13.06	25.34	59.27
Syrah Weather data	191.43	11.05	16.88	57.19
Syrah BEST data	198.63	13.55	23.91	59.99
Ugni blanc Weather data	216.88	10.18	17.23	62.47
Ugni blanc BEST data	199.74	17.80	24.41	67.36

Table S8. Validation of phenological modeling results against independent observations of winegrape phenology corresponding to dates of budbreak (BB), flowering (FL) and veraison (VER). Independent observations are extracted from Bock et al. (2011), Malheiro et al. (2013) and Ruml et al. (2015) in Europe, where only means and standard deviations of observations are provided. And from Cayan et al. (2010) in North America, where full yearly observations of each phenophase were available. Results are for Riesling (Rsl), Chasselas (Chs), Pinot noir (Pn), Cabernet-Sauvignon (C-S), Merlot (Mrl), Chardonnay (Chr) and Sauvignon blanc (S-b). Model predictions and observations from Bock et al. (2011) and Malheiro et al. (2013) for flowering and veraison were for when 50% of clusters were at stages BBCH 65 and 81, respectively (these correspond to modified Eichorn-Lorenz of 23 and 35); for Ruml et al. (2015), observations are for the beginning of these stages (while our model estimates are for the mid-point—thus our model would not be expected to accurately predict these observations). Budburst is defined as green shoot tips visible for all observations and model predictions. Values shown correspond to averages and associated confidence intervals (95%) across observations and model predictions over the period for which phenological data were available. RPD and RMSE values could be computed only for Napa, as it was the only dataset where yearly observations matched yearly predictions. For European data validation is restricted to comparison of means of observed and predicted values. Note that data from Davis in Wolkovich et al. (2017) was not evaluated here due to lack of long enough temporal resolution—i.e. 4 years of observations—but see Fig. S6.

Dataset	Phenophase	Observed	Predicted	RPD	RMSE
Bock (Rsl)	BB	122.00 ± 10.00	175.00 ± 9.00		
Bock (Rsl)	FL	236.00 ± 12.00	118.57 ± 16.50		
Bock (Rsl)	VER	177.10 ± 16.20	246.62 ± 18.20		
Malheiro (Chs)	BB	73.00 ± 8.00	140.00 ± 9.00		
Malheiro (Chs)	FL	208.00 ± 11.00	80.25 ± 7.20		
Malheiro (Chs)	VER	139.50 ± 13.20	195.06 ± 13.30		
Ruml (Pn)	BB	99.00 ± 10.50	148.00 ± 7.80		
Ruml (Pn)	FL	203.00 ± 9.70	108.07 ± 14.50		
Ruml (Pn)	VER	158.20 ± 11.50	219.87 ± 11.00		
Ruml (C-S)	BB	108.00 ± 6.70	151.00 ± 7.60		
Ruml (C-S)	FL	212.00 ± 8.50	114.67 ± 17.00		
Ruml (C-S)	VER	164.20 ± 11.90	231.83 ± 12.10		
Ruml (Mrl)	BB	104.00 ± 8.30	149.00 ± 8.10		
Ruml (Mrl)	FL	212.00 ± 10.50	112.52 ± 19.70		
Ruml (Mrl)	VER	165.50 ± 12.70	232.16 ± 12.90		
Ruml (Chr)	BB	97.00 ± 10.70	146.00 ± 8.30		
Ruml (Chr)	FL	205.00 ± 8.90	107.24 ± 14.80		
Ruml (Chr)	VER	158.50 ± 11.30	224.08 ± 10.90		
Ruml (Rsl)	BB	102.00 ± 9.30	149.00 ± 7.40		
Ruml (Rsl)	FL	211.00 ± 10.20	106.18 ± 16.00		
Ruml (Rsl)	VER	157.74 ± 12.20	224.86 ± 11.80		
Napa (C-S)	BB	90.27 ± 13.58	84.69 ± 15.91	1.06	2.26
Napa (C-S)	FL	143.86 ± 15.27	150.29 ± 15.23	0.97	2.37
Napa (C-S)	VER	214.66 ± 17.14	218.03 ± 15.61	1.68	1.40
Napa (Chr)	BB	81.16 ± 12.96	87.99 ± 13.27	0.72	2.54
Napa (Chr)	FL	135.85 ± 21.62	148.82 ± 15.32	0.43	4.90
Napa (Chr)	VER	208.51 ± 25.10	214.65 ± 15.63	0.69	3.02
Napa (Mrl)	BB	87.35 ± 16.62	76.24 ± 14.96	0.67	2.78
Napa (Mrl)	FL	143.27 ± 19.36	149.28 ± 15.25	0.95	2.00
Napa (Mrl)	VER	218.77 ± 19.52	216.20 ± 15.20	2.36	0.80
Napa (S-b)	BB	86.86 ± 13.13	92.33 ± 13.34	0.82	2.08
Napa (S-b)	FL	145.40 ± 16.71	157.16 ± 14.69	0.55	3.43
Napa (S-b)	VER	208.72 ± 18.63	218.42 ± 14.89	0.63	3.01

Table S9. Results of uncertainty, accuracy and cross-validation relative to calibrated parameters. Uncertainty around fitted parameters is measured as lower and upper 95% Confidence Intervals (CI). Accuracy of each calibration is shown by the RMSE and EF metrics. Cross-validation results following a leave-one-out procedure (analogous to that in Chuine et al. 2016) are shown by RMSEP. EF, RMSE and RMSEP are defined above in 'Phenological Modeling' section. Values for C* and F* are in the chilling and forcing units, respectively, necessary for a phenological event to occur. The units of threshold parameters (C*, F*) and RMSE and RMSEP are in days. RPD is included for parameters obtained with BEST to allow comparability with other validation results. Abbreviations are used for Cabernet Sauvignon (C-S), Chardonnay (Chr), Chasselas (Chs), Grenache (Gre), Merlot (Mrl), Monastrell (Mon), Pinot noir (P-n), Riesling (Rsl), Sauvignon blanc (S-b) and Ugni blanc (U-b). Weather or BEST indicate what climatic data was used to parameterize the data and BB, FL and VER indicate the phenological stage of budbreak, flowering and veraison, respectively.

Variety_Data_Stage	C*	C* CI.low	C* CI.up	F*	F* CI.low	F* CI.up	RMSE	EF	RMSEP	RPD
C-S.BEST_BB	166.02	157.91	180.47	20.24	17.86	21.50	5.70	0.91	6.38	3.35
C-S.BEST_FL				23.61	22.64	24.62	4.60	0.90	4.73	3.12
C-S.BEST_VER				65.30	63.95	66.62	5.36	0.86	5.44	2.36
C-S.Weather_BB	158.34	149.33	166.34	16.09	14.79	16.95	5.96	0.90	7.42	
C-S.Weather_FL				16.56	15.88	17.26	4.51	0.90	4.70	
C-S.Weather_VER				60.84	59.60	62.01	5.24	0.86	5.31	
Chr.BEST_BB	199.01	192.02	203.51	12.50	11.59	13.73	7.92	0.53	8.21	1.46
Chr.BEST_FL				22.26	21.55	22.95	4.05	0.85	4.19	2.61
Chr.BEST_VER				63.28	61.30	65.35	7.46	0.82	7.60	2.37
Chr.Weather_BB	189.44	187.17	195.42	9.86	9.26	10.22	6.85	0.65	7.53	
Chr.Weather_FL				15.35	14.90	15.82	3.78	0.87	3.90	
Chr.Weather_VER				58.98	57.45	60.50	5.92	0.89	6.07	
Chs.BEST_BB	196.68	192.24	205.71	12.40	11.14	14.11	6.15	0.55	7.72	1.49
Chs.BEST_FL				24.46	23.74	25.22	3.62	0.86	3.72	2.68
Chs.BEST_VER				53.40	52.07	54.77	5.12	0.92	5.21	3.44
Chs.Weather_BB	193.20	189.64	195.10	10.14	9.76	10.59	5.02	0.70	5.35	
Chs.Weather_FL				17.24	16.77	17.72	3.19	0.89	3.28	
Chs.Weather_VER				49.53	48.47	50.55	4.25	0.94	4.32	
Gre.BEST_BB	197.68	195.36	205.78	14.02	12.45	14.67	7.21	0.79	8.00	2.18
Gre.BEST_FL				24.71	24.18	25.34	3.64	0.94	3.71	4.21
Gre.BEST_VER				66.90	65.20	68.52	7.77	0.81	7.88	2.29
Gre.Weather_BB	187.40	162.40	191.70	11.44	10.57	15.14	7.24	0.79	8.89	
Gre.Weather_FL				17.49	17.06	17.92	3.45	0.95	3.50	
Gre.Weather_VER				63.98	62.72	65.21	6.26	0.88	6.36	
Mrl.BEST_BB	152.14	146.83	169.62	19.91	16.92	21.10	8.96	0.73	9.42	1.91
Mrl.BEST_FL				24.96	24.26	25.60	5.04	0.83	5.13	2.46
Mrl.BEST_VER				64.45	63.19	65.75	5.91	0.83	5.98	2.44
Mrl.Weather_BB	144.72	142.54	147.71	16.16	15.75	16.65	7.59	0.80	7.71	
Mrl.Weather_FL				17.28	16.84	17.73	4.50	0.87	4.56	
Mrl.Weather_VER				60.35	59.19	61.46	5.50	0.86	5.56	
Mon.BEST_BB	216.56	205.56	217.77	7.64	6.66	12.11	6.67	0.09	6.89	1.05
Mon.BEST_FL				28.04	27.49	28.67	3.18	0.43	3.24	1.32
Mon.BEST_VER				62.07	58.81	65.29	3.97	0.39	4.52	1.28
Mon.Weather_BB	171.13	167.90	184.08	15.42	13.08	16.18	5.65	0.34	6.23	
Mon.Weather_FL				20.34	19.91	20.80	3.01	0.49	3.08	
Mon.Weather_VER				60.43	57.03	63.78	3.58	0.50	4.06	
P-n.BEST_BB	198.53	197.54	221.51	12.98	8.09	13.49	6.28	0.32	7.51	1.22
P-n.BEST_FL				21.85	21.44	22.17	3.33	0.83	3.39	2.39
P-n.BEST_VER				59.50	58.17	60.86	7.11	0.77	7.17	2.07
P-n.Weather_BB	194.18	190.97	197.06	10.27	9.92	10.68	5.02	0.57	5.18	
P-n.Weather_FL				15.29	15.03	15.58	3.35	0.82	3.39	
P-n.Weather_VER				54.53	53.28	55.78	7.06	0.77	7.09	
Rsl.BEST_BB	185.38	181.37	194.30	14.08	13.10	14.89	6.13	0.74	7.01	1.97
Rsl.BEST_FL				22.13	21.59	22.66	3.62	0.91	3.72	3.39
Rsl.BEST_VER				64.63	62.54	66.47	7.96	0.69	8.16	1.79
Rsl.Weather_BB	190.22	183.23	191.45	10.32	10.10	10.95	4.90	0.83	5.53	
Rsl.Weather_FL				15.50	15.13	15.88	3.30	0.92	3.49	
Rsl.Weather_VER				59.88	58.58	61.15	5.34	0.85	7.99	
S-b.BEST_BB	203.48	199.19	207.29	13.06	12.17	14.53	4.22	0.92	4.78	3.62
S-b.BEST_FL				25.34	24.47	26.03	3.41	0.94	3.56	4.04
S-b.BEST_VER				59.27	57.93	60.56	4.80	0.90	4.90	3.16
S-b.Weather_BB	192.22	188.24	195.34	10.26	9.37	11.06	4.97	0.89	6.21	
S-b.Weather_FL				17.76	16.17	19.20	3.20	0.95	3.34	
S-b.Weather_VER				55.16	53.97	56.42	4.77	0.90	4.84	
Syr.BEST_BB	198.63	192.34	204.42	13.55	12.61	14.75	7.22	0.70	7.54	1.83
Syr.BEST_FL				23.91	23.25	24.56	3.79	0.91	3.85	3.40
Syr.BEST_VER				59.99	58.53	61.39	6.55	0.84	6.65	2.52
Syr.Weather_BB	191.43	185.55	196.34	11.05	10.41	12.05	6.68	0.75	7.14	
Syr.Weather_FL				16.88	16.42	17.34	3.55	0.92	3.64	
Syr.Weather_VER				57.19	56.20	58.18	4.77	0.92	4.87	
U-b.BEST_BB	199.74	196.64	219.76	17.80	12.83	18.60	5.33	0.52	6.66	1.45
U-b.BEST_FL				24.41	23.17	25.36	3.80	0.84	4.05	2.46
U-b.BEST_VER				67.36	65.40	69.48	6.80	0.88	6.96	2.92
U-b.Weather_BB	216.88	201.96	230.47	10.18	6.87	13.40	5.39	0.51	8.06	
U-b.Weather_FL				17.23	16.66	17.87	3.19	0.88	3.41	
U-b.Weather_VER				62.47	61.01	63.80	5.07	0.93	5.28	

Table S10. Correlations amongst indicators used to fit bioclimatic envelopes. Results show the average \pm standard deviation of correlations computed across 30 GCMs, 30 years (1951-1980), and 11 cultivars at locations utilized to fit their envelopes (see above). Precipitation variables are highly collinear, and yet they differ in their relative importance to determine loss of climatic suitability (see Fig. S17). Temperature-related indicators are not strongly collinear except for minimum and maximum temperatures, but each variable affects winegrape maturation differently.

	Days<10°C	Days>40°C	GDD	GDD>25°C	MaxTemp	MinTemp	MaxPrec	SumPrec
Days<10°C	1.00 \pm 0.00	0.01 \pm 0.04	-0.61 \pm 0.08	-0.16 \pm 0.07	-0.61 \pm 0.10	-0.79 \pm 0.10	0.21 \pm 0.25	0.25 \pm 0.23
Days>40°C	0.01 \pm 0.04	1.00 \pm 0.00	0.01 \pm 0.03	0.01 \pm 0.02	0.00 \pm 0.04	-0.00 \pm 0.03	0.01 \pm 0.06	0.01 \pm 0.07
GDD	-0.61 \pm 0.08	0.01 \pm 0.03	1.00 \pm 0.00	0.05 \pm 0.01	0.69 \pm 0.16	0.68 \pm 0.16	-0.14 \pm 0.15	-0.19 \pm 0.15
GDD>25°C	-0.16 \pm 0.07	0.01 \pm 0.02	0.05 \pm 0.01	1.00 \pm 0.00	0.45 \pm 0.19	0.55 \pm 0.16	-0.12 \pm 0.14	-0.19 \pm 0.16
MaxTemp	-0.61 \pm 0.10	0.00 \pm 0.04	0.69 \pm 0.16	0.45 \pm 0.19	1.00 \pm 0.00	0.74 \pm 0.11	-0.39 \pm 0.17	-0.44 \pm 0.17
MinTemp	-0.79 \pm 0.10	-0.00 \pm 0.03	0.68 \pm 0.16	0.55 \pm 0.16	0.74 \pm 0.11	1.00 \pm 0.00	-0.23 \pm 0.19	-0.30 \pm 0.17
MaxPrec	0.21 \pm 0.25	0.01 \pm 0.06	-0.14 \pm 0.15	-0.12 \pm 0.14	-0.39 \pm 0.17	-0.23 \pm 0.19	1.00 \pm 0.00	0.96 \pm 0.02
SumPrec	0.25 \pm 0.23	0.01 \pm 0.07	-0.19 \pm 0.15	-0.19 \pm 0.16	-0.44 \pm 0.17	-0.30 \pm 0.17	0.96 \pm 0.02	1.00 \pm 0.00

Additional Supplemental Figures

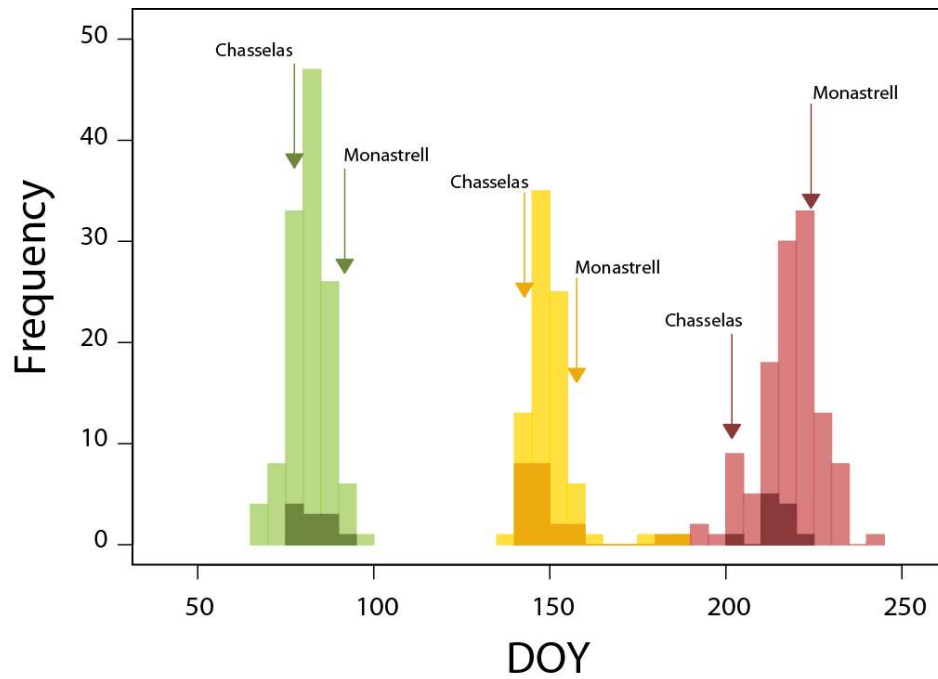


Fig. S4. We selected 11 winegrape cultivars (varieties), ensuring that they span enough diversity in their phenologies. Histograms depict the temporal distribution of the three modeled phenological stages—e.g. budbreak (green), flowering (yellow) and veraison (red)—across 130 varieties included in the Domaine de Vassal collection (lighter colors) and across the 11 varieties analyzed in this study (67). To illustrate the large differences in the phenologies among early and late varieties, we mark the average dates of each phenological stage for two characteristic winegrape varieties known to have either early—Chasselas—or late—Monastrell—phenologies. For more information see *Phenological data for parameterization & validation*.

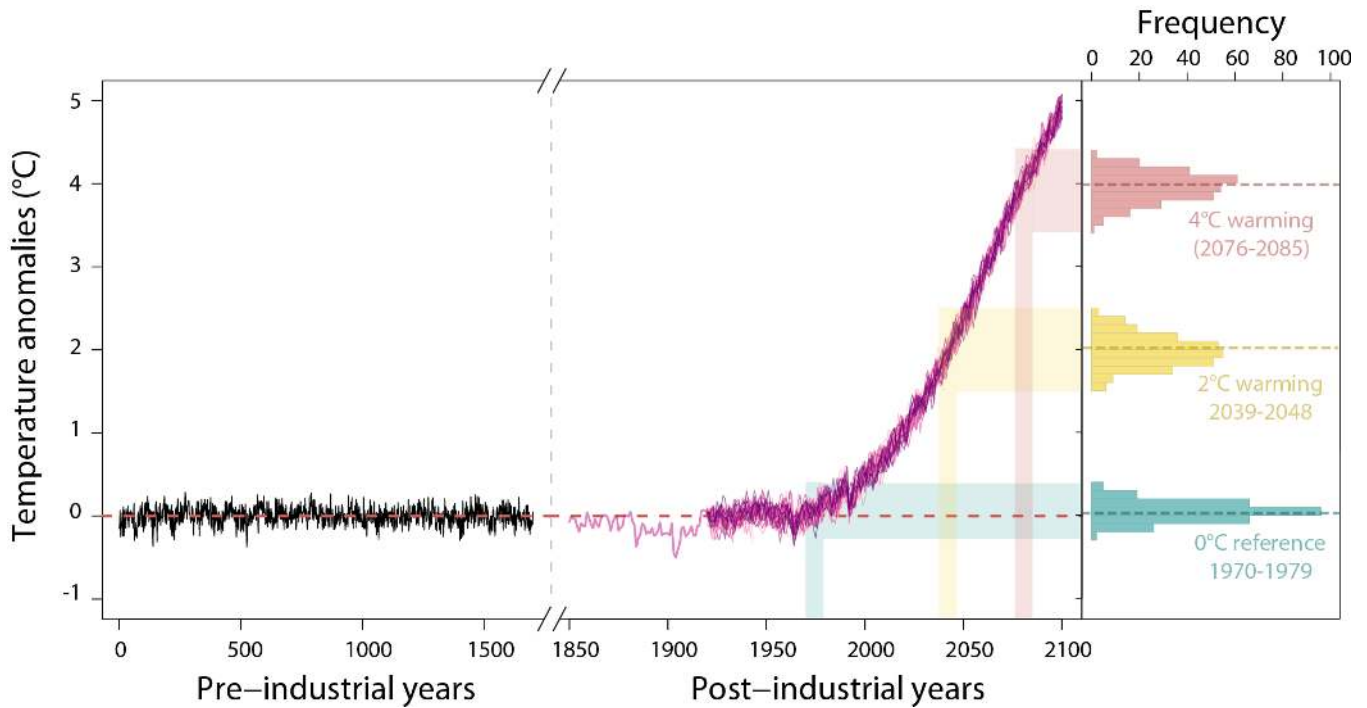


Fig. S5. Evolution of the anomalies in global average temperatures through time. The figure shows global anomalies in temperatures based on 1700 simulations for the pre-industrial era, prior to 1850 (black line), the global average in temperature anomalies for the 1850-1920 period based on one GCM simulation (purple line), and 30 simulated global averages in temperature anomalies for the 1921-2100 period based on 30 GCM simulations (light pink to dark purple lines). We examined two major warming scenarios: 2°C (2039-2048 in RCP 8.5, yellow), 4°C (2076-2085 in RCP 8.5, pink) against a 0°C reference period (1970-1979 in RCP 8.5, blue). Histograms in the right-hand panel show non-overlapping distributions of temperature anomalies across 300 simulations corresponding to each 10 year period and each of the 300 GCM simulations. Dotted lines on top of the histograms represent median values of each distribution of temperatures. The median estimates for the +2°C/+4°C warming scenarios are +2.03/+3.99°C, with interquartile ranges of 1.99-2.06°C and 3.97-4.04°C, respectively.

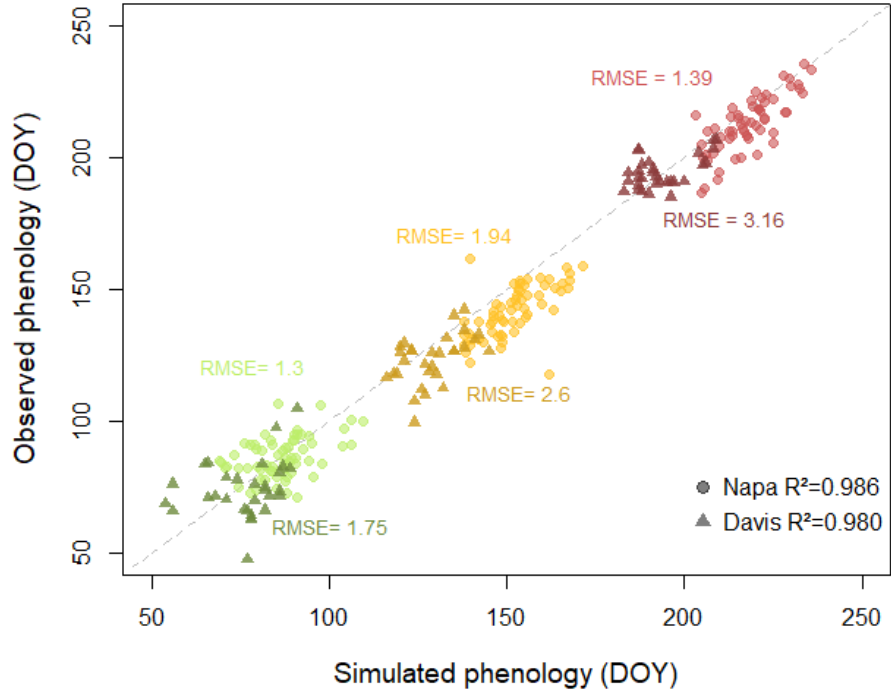


Fig. S6. Phenological modeling validation results comparing simulated dates of phenological stages budbreak (green), flowering (yellow) and veraison (red) for two locations in California (USA), Napa Valley (circles, light colors) and Davis (triangles, dark colors), against independent observations of the phenology therein (8, 9). Observations and simulations comprise the 1985 to 2007 period for Cabernet-Sauvignon, Sauvignon blanc, Chardonnay and Merlot in Napa and the 2014 to 2016 in Davis for all cultivars except for Grenache. Overall regression R^2 are noted for each location across phases and RMSE values for each phenophase and location are marked next to the corresponding cloud of points. Note that error values are rather low, particularly for the Napa dataset.

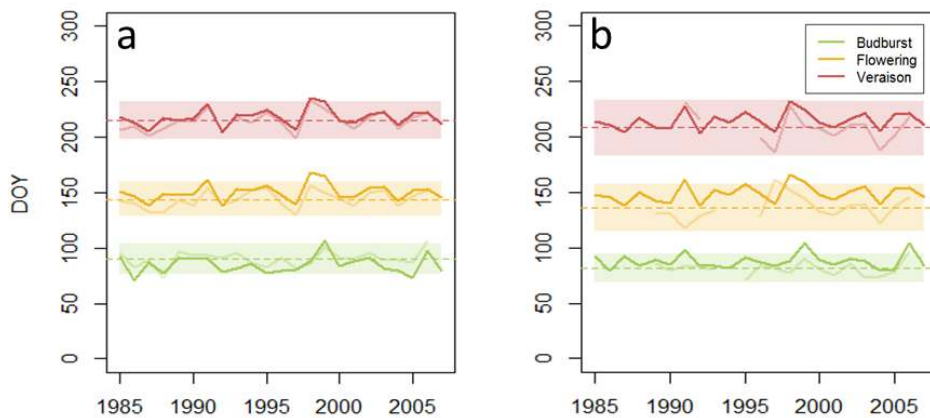


Fig. S7. Example of model validation with independent observations comparing our projections of the phenological stages budbreak (green), flowering (yellow) and veraison (red) for Napa Valley (California, USA) against independent observations therein (8, 9). Results in this example are for Cabernet-Sauvignon (a) and Chardonnay (b). Although projections are based on models calibrated with European data (see above), and thus there are notable differences between climate in the calibration and validation regions, phenological projections (solid lines) closely match the observed phenology (light lines) ($r = 0.994$ for Cabernet-Sauvignon; $r = 0.983$ for Chardonnay), and fall within 95% confident intervals for the variation of the observed phenology (shaded areas).

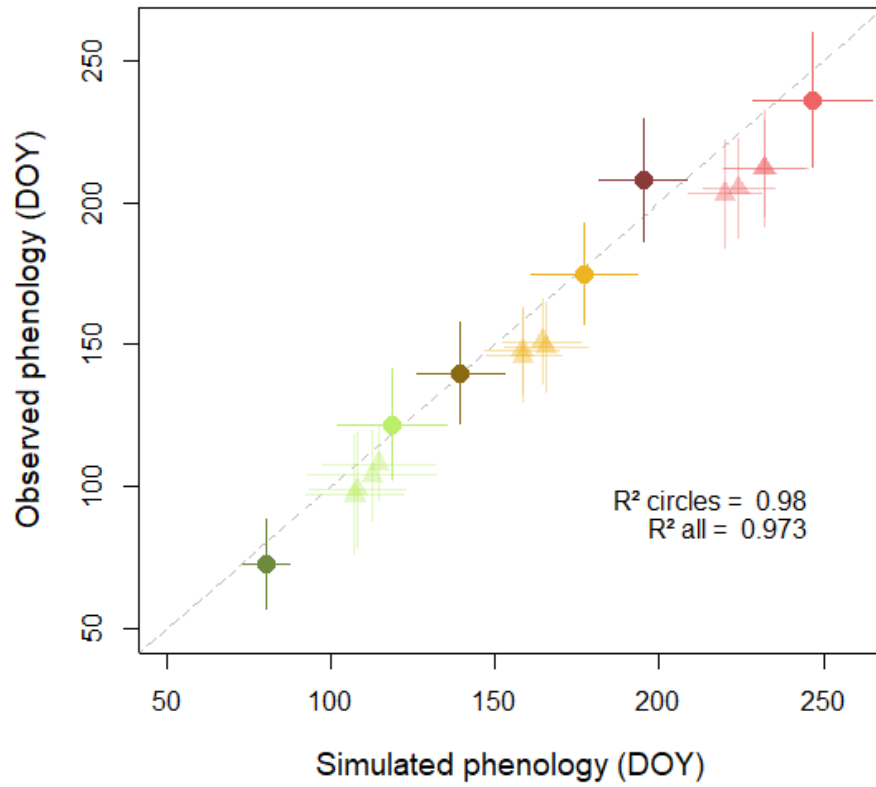


Fig. S8. Phenological modeling validation results comparing our projections of the phenological stages budbreak (green), flowering (yellow) and veraison (red) against independent phenological observations in (5) (light-colored circles), (6) (dark-colored circles), and (7) (light-colored triangles). The different symbol used for the Ruml *et al.* (2016) data stems from it reporting an earlier phenological stage than we modeled. Results show an overall strong correlation between observed phenology and the phenology simulated by phenological models. Error bars represent 95% CI around the mean.

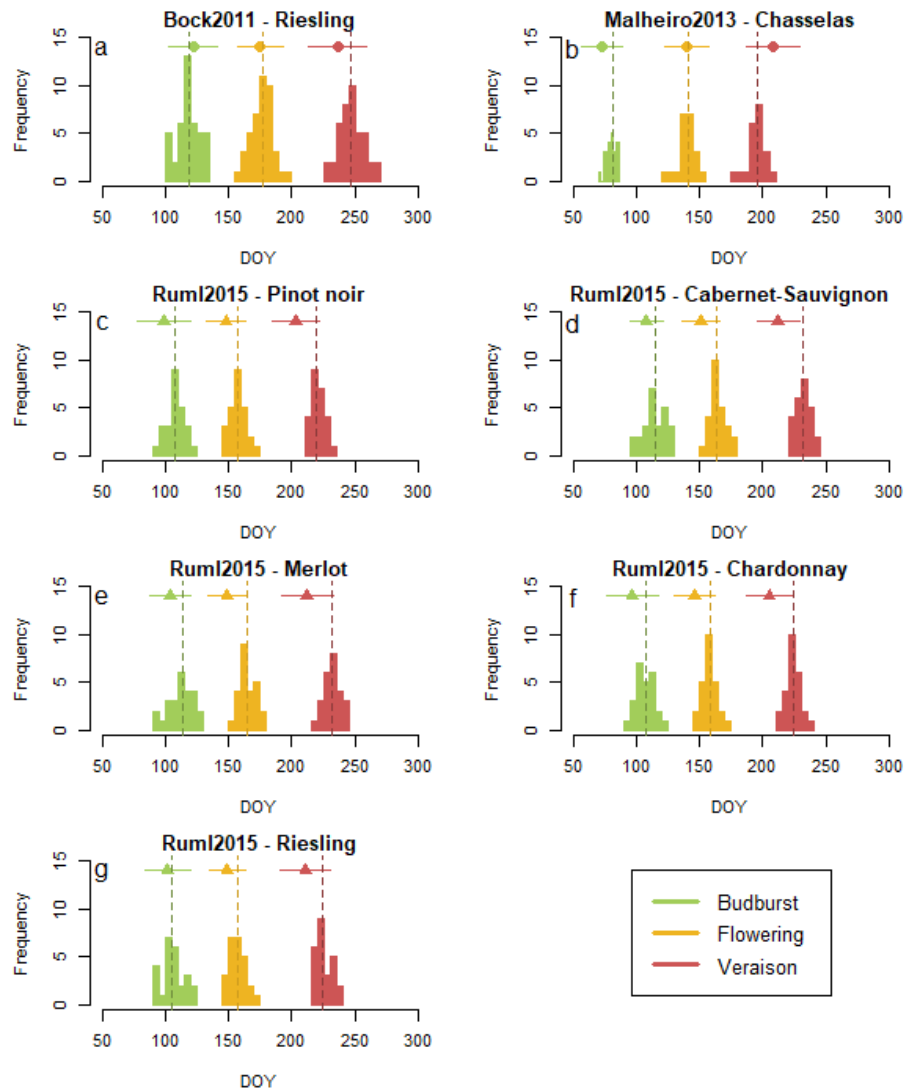


Fig. S9. Phenological modeling validation results comparing our projections of the phenological stages budbreak (green), flowering (yellow) and veraison (red) against independent observations in published work (5–7). The histograms summarize our phenological projections across the time period during which independent observations were recorded. The circles and triangles above the histograms represent the means and their error intervals represent the 95% confident intervals of observed phenology. Note that our projections for winegrape phenology consistently predict later than observed phenology for the Ruml *et al.* (2016) dataset, given that their results are for earlier stages than those parameterized by our models. This is the reason for the use of triangles instead of circles. Results in this example are for Riesling (a,g), Chasselas (b), Pinot noir (c), Cabernet-Sauvignon (d), Merlot (e), Chardonnay (f).



Fig. S10. Geographical locations utilized to model maturity for each of the 11 winegrape varieties analyzed in this study. Locations are marked by red dots and were extracted from (3), see also *Modeling maturity*.

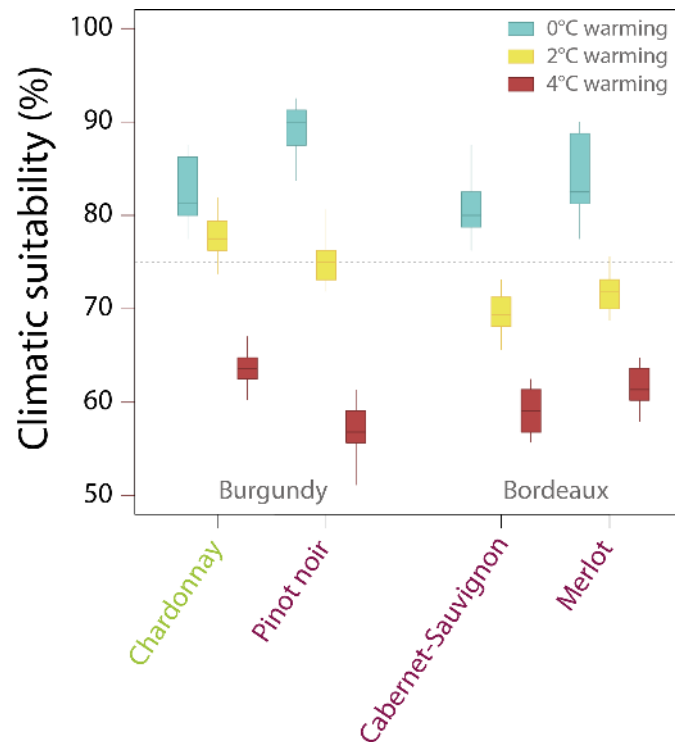


Fig. S11. Estimated climatic suitability for regions including Bordeaux and Burgundy under our 0°C reference scenario, 2°C, and 4°C warming scenarios. Our metric of suitability incorporates uncertainty in the climate that determines whether a location is good for growing a particular variety every year. Thus, suitability below 100%, in part, highlights that every year is not always ideal for a variety in a region (68). See also *Calculating climatic suitability*.

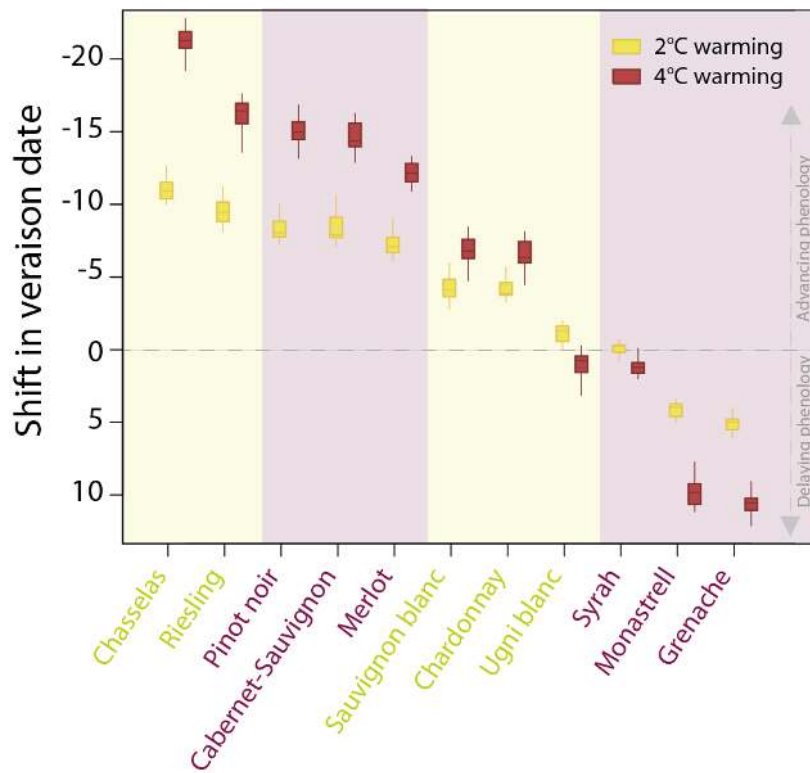


Fig. S12. Variation in the predicted shifts in phenology across the 11 winegrape varieties in our dataset. Shifts in phenology are shown for two warming scenarios: 2°C (yellow) and 4°C (red), and are calculated by comparing predicted veraison dates under these scenarios against predicted veraison during the 0° reference scenario. Winegrape varieties are ranked according to the magnitude of the veraison shifts.

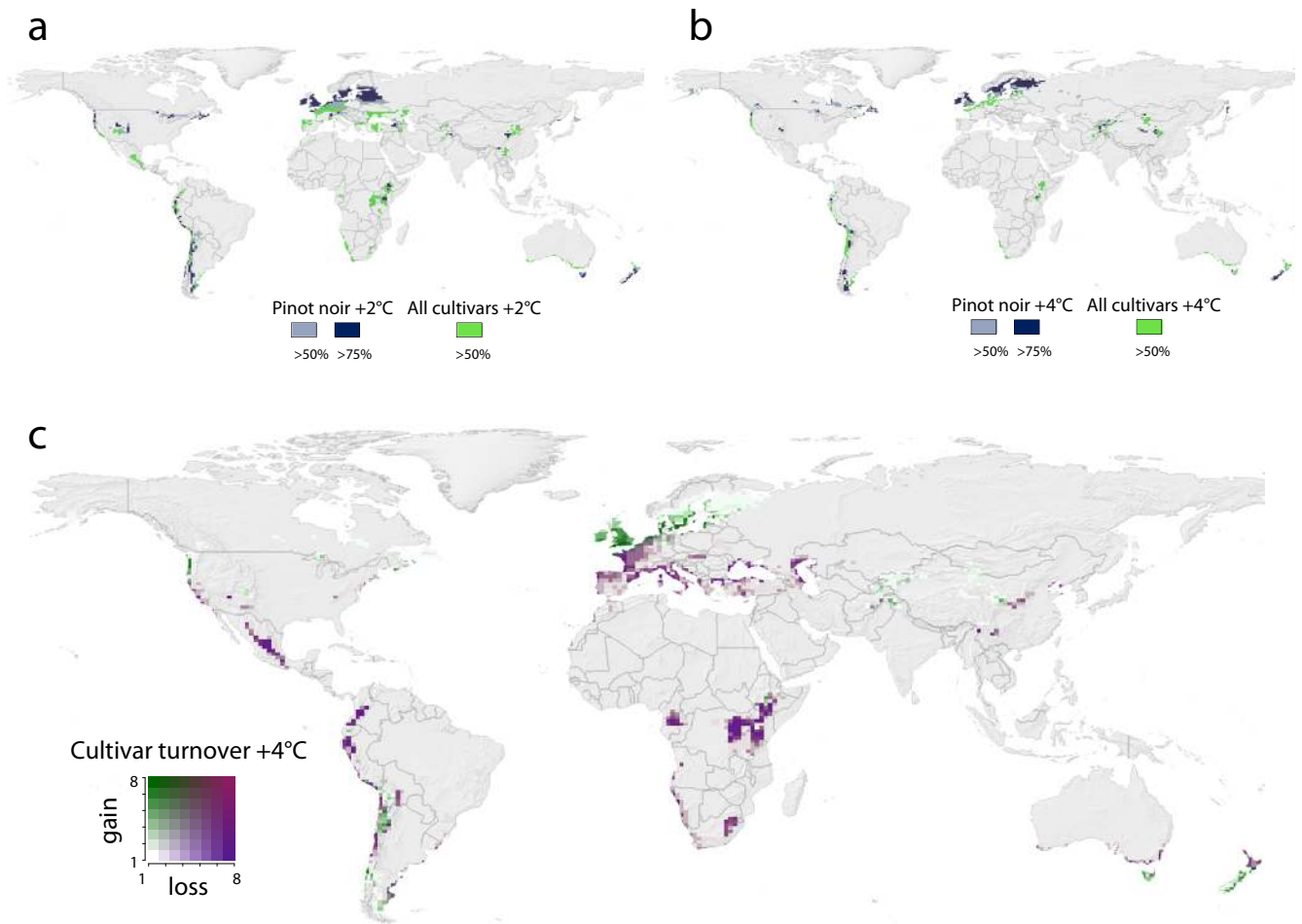


Fig. S13. Predicted effects of climate change on cultivar (variety) distribution. Panel (a) depicts the predicted areas suitable to grow Pinot noir under different levels of climatic suitability (over 50% and 75%) and areas of the globe our model predicts are climatically suitable for the remaining varieties, for a 2°C warming scenario. Similarly, panel (b) shows predicted suitable areas for Pinot noir vs. other varieties under the 4°C warming scenario. Panel (c) shows in a bivariate color scale simultaneously variety gains (white to green) and losses (white to purple) predicted under 4°C of warming. Climatic suitability is defined as the percentage of model-years (i.e., 300 years: 10 years x 30 ensemble members) when our modeling approach suggested a variety would reach maturity in a region (see above for details). Note that both gains and losses of climatic suitability (c) are calculated with respect to areas projected to be suitable under the 0°C, and thus, do not refer to current growing regions (see Fig. S1).

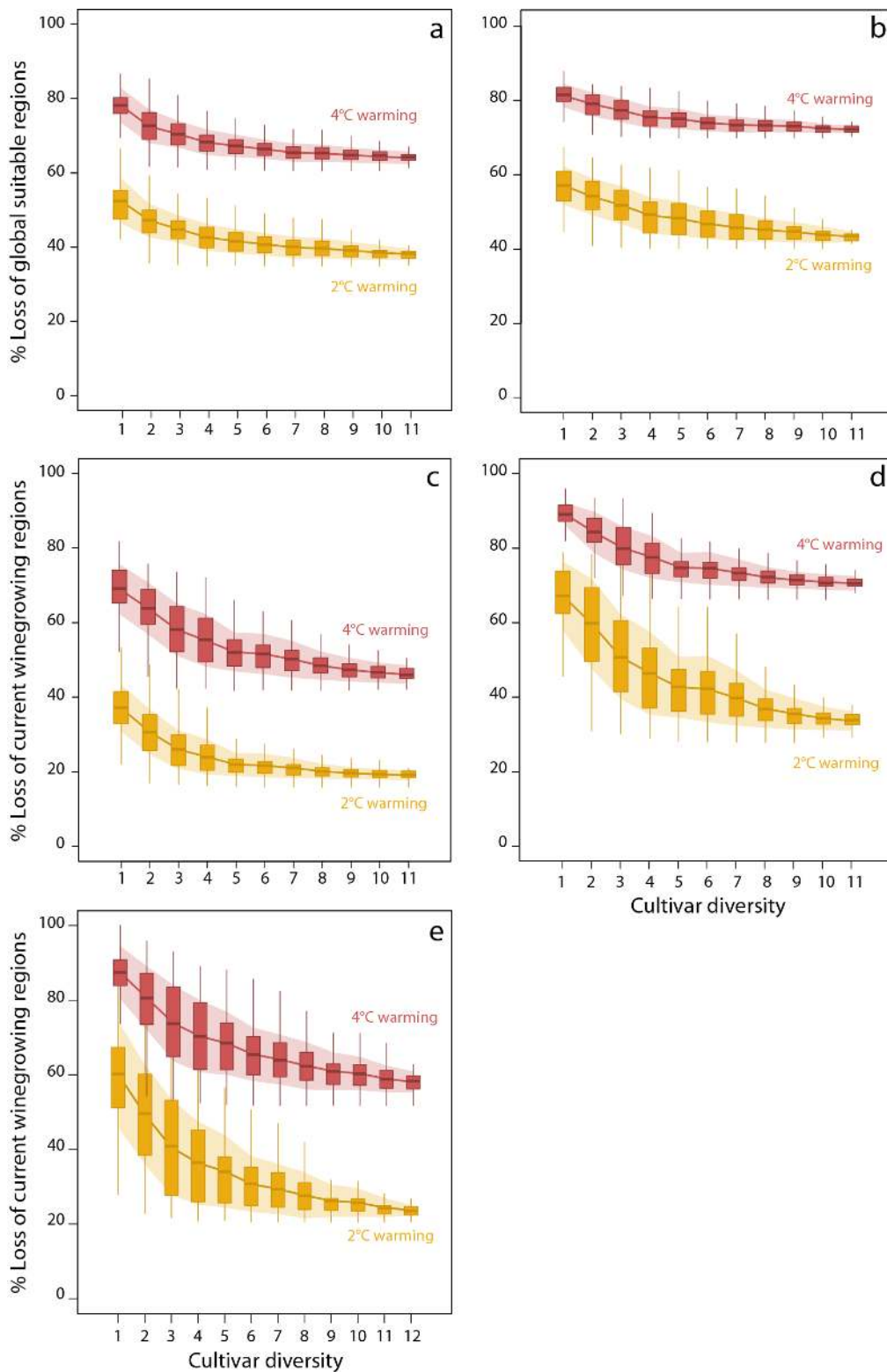


Fig. S14. Alternative analyses of the proportion of climatic suitability losses versus winegrape cultivar diversity (see also main text Fig. 2). Panels show decrease in loss of suitable regions with increasing diversity for both a scenario of 2°C warming (yellow bars) and one of 4°C warming (red bars) for: (a) all global regions potentially suitable for winegrowing using all 8 climatic variables, (b) all global regions potentially suitable for winegrowing using temperature variables only, (c) current growing regions excluding maximum and minimum temperatures, (d) current growing regions using temperature variables only and, (e) current growing regions using all 8 climatic variables and including the Late-ripening composite variety and thus, 12 winegrape varieties. Shaded areas illustrate ± 1 standard deviation around the mean climatic suitability loss for the corresponding number of sampled varieties.

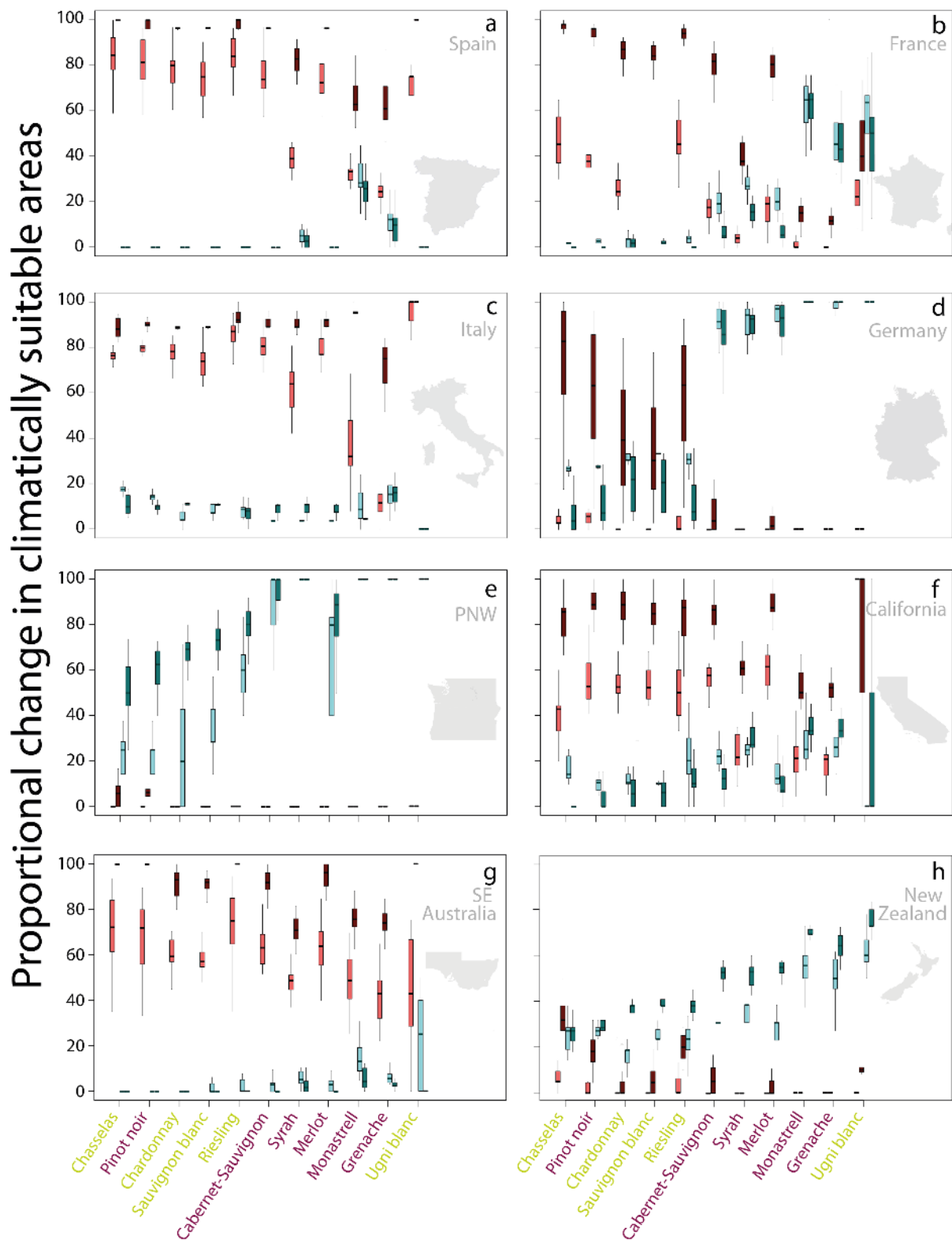


Fig. S15. Shifts in predicted climatic suitability for 11 studied varieties within 8 popular winegrowing regions (see also main text Fig. 3), including Spain (a), France (b), Italy (c), Germany (d), Pacific Northwest (USA) (e), California (USA) (f), South Eastern Australia (g) and New Zealand (h). Increases in suitability are represented by light turquoise (2°C warming) and dark turquoise (4°C degree warming) colors, and suitability decreases by light red (2°C warming) and dark red (4°C warming). Winegrape varieties are ranked according to their phenological order, following (2). Note that these estimates are based on modelled suitability and thus we may show low suitability for some varieties already grown in some regions (see *Calculating climatic suitability* in main text for more information).

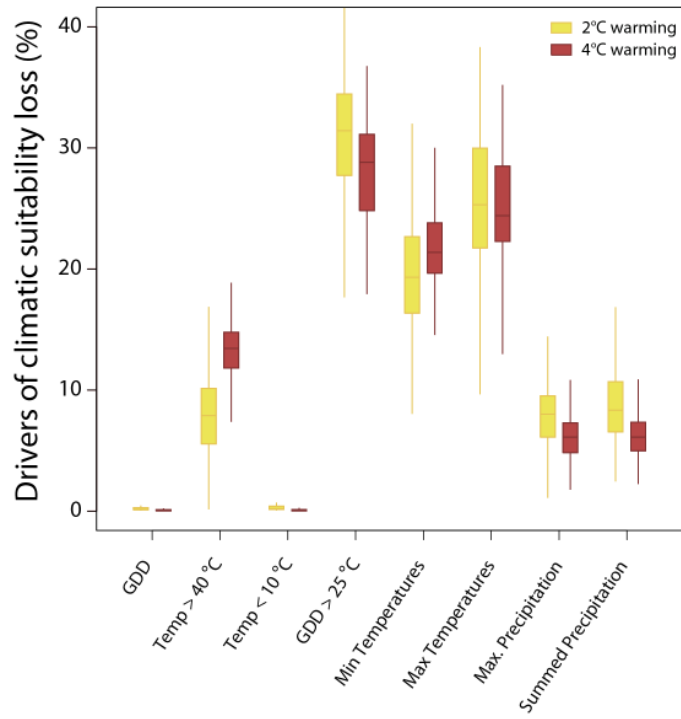


Fig. S16. Drivers of loss of climatic suitability within current winegrowing regions. Results for all eight climatic variables included as predictors in the bioclimatic envelopes (see *Modeling maturity*). Estimates were calculated as a proportion of losses for all varieties under scenarios of warming of 2°C (yellow bars) and 4°C (red bars). Error bars show variation in the proportion of losses due to each climatic variable across GCMs and years.

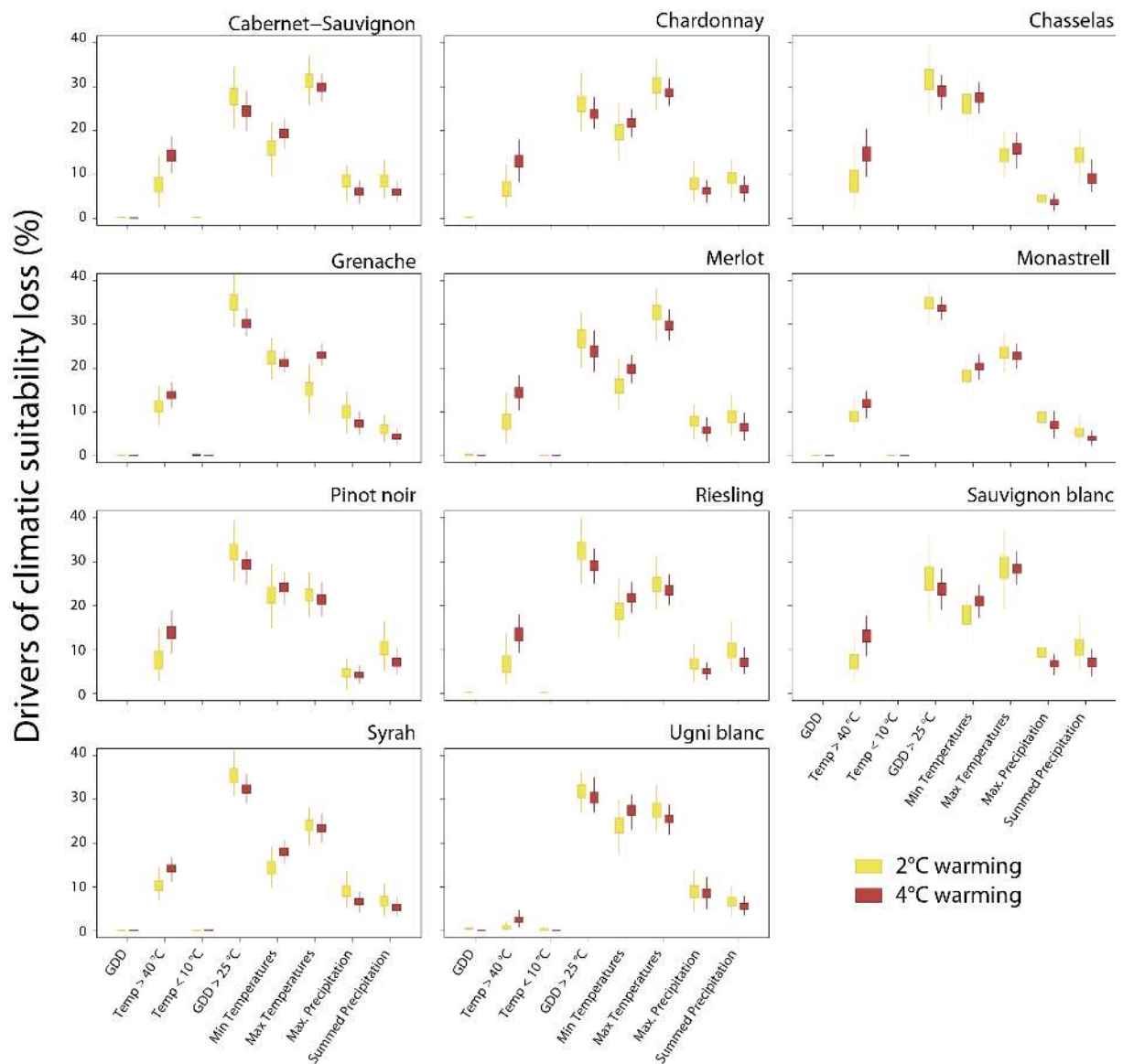


Fig. S17. Drivers of loss of climatic suitability within current winegrowing regions. Results for all eight climatic variables included as predictors in the bioclimatic envelopes to model climatic suitability to reach maturity. Estimates were calculated as a proportion of losses for each of the 11 studied varieties under scenarios of warming of 2°C (yellow bars) and 4°C (red bars). Error bars show variation in the proportion of losses due to each climatic variable across GCMs and years. Note that we do not show loss due to the veraison date cut-off, as no regions were lost due to this at 2°C, and at 4°C less than 1% of regions were lost for all varieties, except for Ugni blanc, which lost 2% of regions to veraison after October 1.

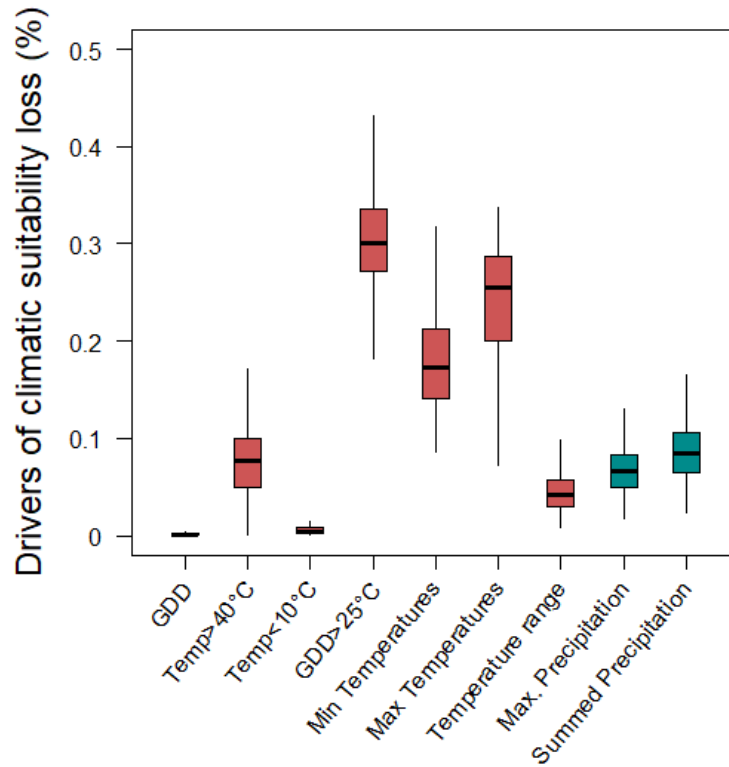


Fig. S18. Sensitivity analysis of the drivers of loss of climatic suitability within current winegrowing regions, including the influence of Diurnal Temperature Range amongst the temperature-related predictors (red bars). Results for all nine climatic variables included as predictors in the bioclimatic envelopes (see *Modeling maturity* and *Sensitivity analysis for Diurnal Temperature Range*). Estimates were calculated as a proportion of losses for all varieties under scenarios of warming of 2°C. Error bars show variation in the proportion of losses due to each climatic variable across 5 randomly selected GCM simulations (see above) and the 10 years corresponding to each 0°C and 2°C scenario for which region loss is computed.

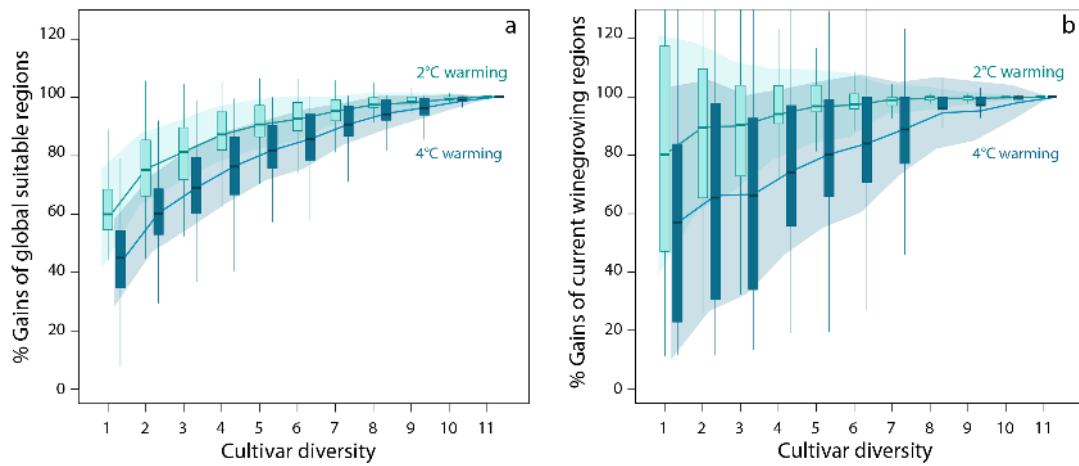


Fig. S19. Proportion of climatic suitability gains calculated across all potential climatically suitable regions of the world (a) and within current winegrowing regions (b), versus winegrape cultivar diversity, out of maximum suitability gains by the 11 varieties. Predictions of climatic suitability gains are shown for both a scenario of 2°C warming (green bars) and one of 4°C warming (blue bars). Depicted uncertainty shows variability in climatic suitability losses according to all possible combinations of n varieties, and their modelled suitability under each climate change scenario. Shaded areas illustrate ± 1 standard deviation around the mean climatic suitability gains for each number of varieties.

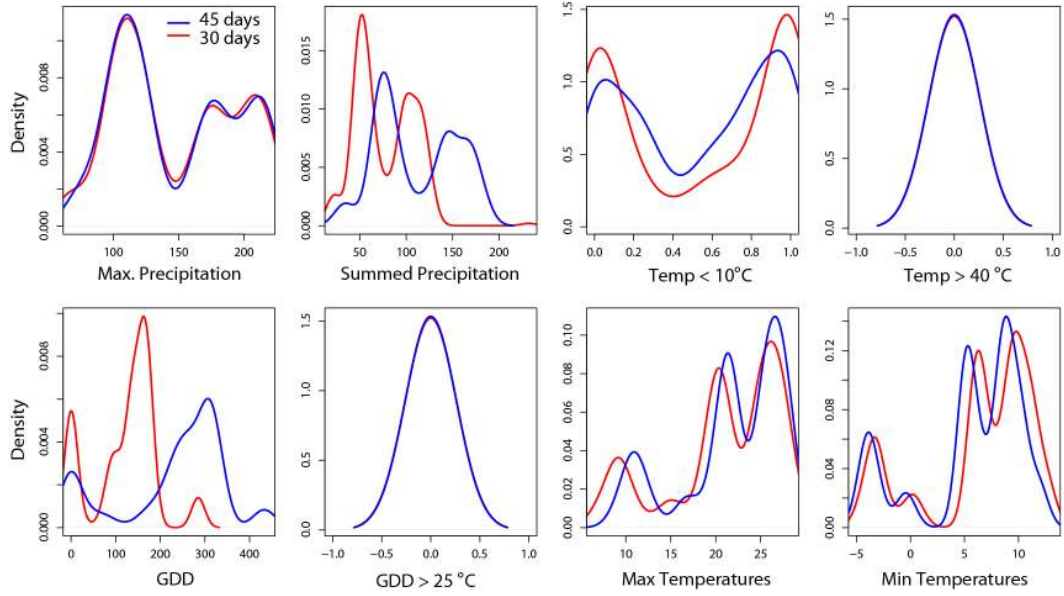


Fig. S20. Density plots comparing the distribution of values corresponding to each of the eight climatic variables utilized to fit climatic envelopes, recorded either over a 45-day period after veraison (blue) or over a 30-day period after veraison (red) for Chasselas across the 19 locations selected to fit the envelopes for this variety.

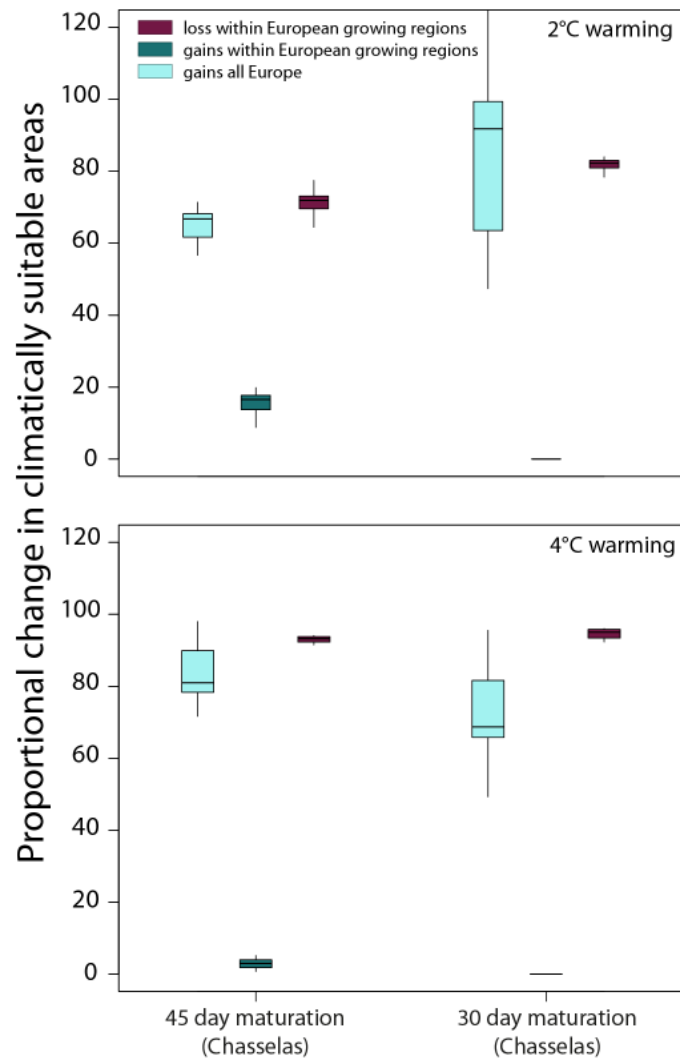


Fig. S21. Shifts in predicted climatic suitability for Chasselas in Europe (see also main text Fig. 3), according to climatic envelopes computed over either a 45-day period after veraison (left) or a 30-day period after veraison (right). Increases in suitability across all of Europe are represented by light green boxes, increases within current growing regions (see Fig. S1) are represented by dark green boxes, and decrease in climatic suitability within those regions are represented by dark red boxes. Both estimates of suitability gains and losses are presented for two warming scenarios, 2°C warming (above) and 4°C warming (below).

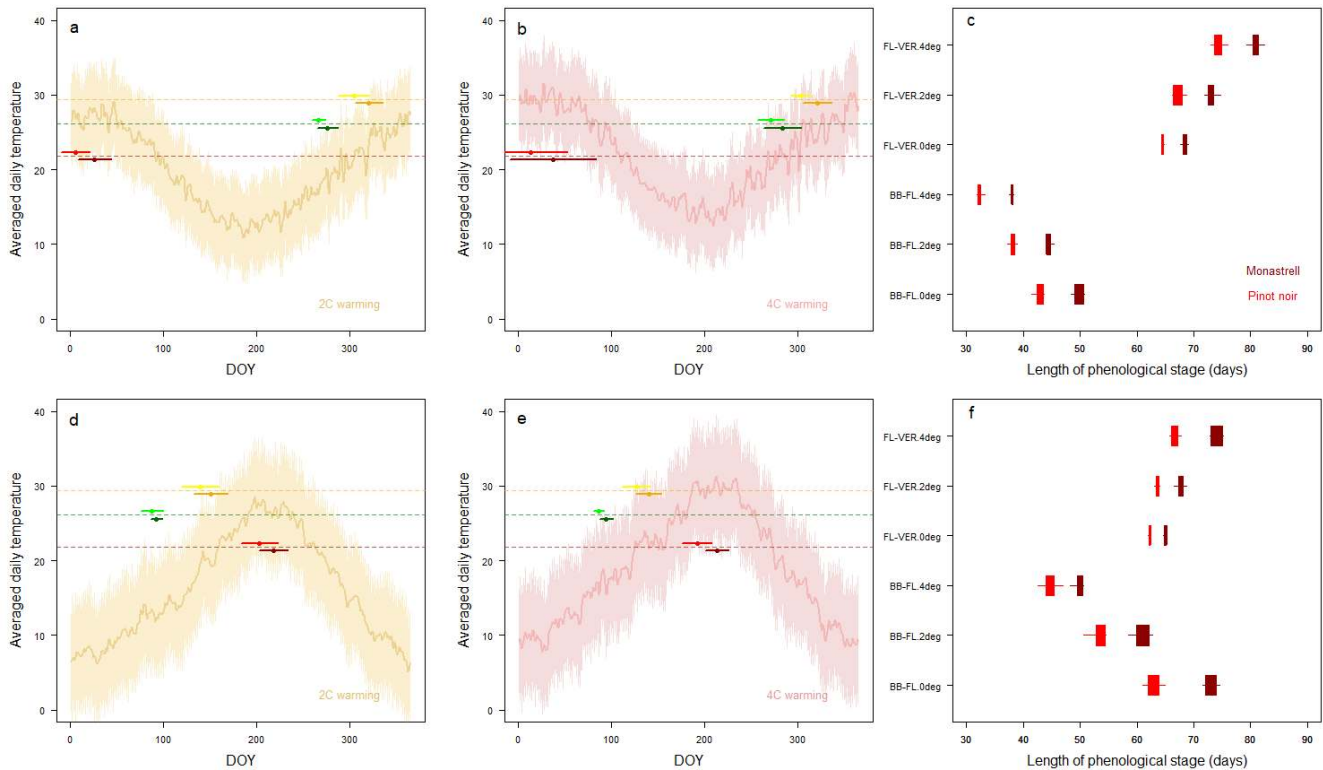


Fig. S22. Drivers of phenological advances and delays. Our analyses show that some winegrape cultivars with late phenology (e.g., Monastrell and Grenache; see Fig. S12) have delays in veraison under warming scenarios. In our non-linear phenological models, delays are expected when the temperatures experienced during a given phenological stage are above the optimal parameterized temperature for that stage (T_{opt} , shown here as dashed lines). The panels illustrate average daily temperatures for two example countries, Australia (a,b) and Spain (d,e), under the two warming scenarios of 2°C (yellow; a,d) and 4°C (red; b,e). Dates of predicted phenology are shown for budbreak (BB, green), flowering (FL, yellow) and veraison (VER, red) for Pinot noir, a variety with early phenology (lighter colors) and a variety with late phenology, Monastrell (darker colors), which often shows delayed phenology in our higher warming scenario (4°C). Delays are expected only when temperatures are above the parameterized optimum values (dashed lines). Thus, in both countries (c,f) the period between budbreak and flowering shortens as climate warms (BB-FL) because warming increases temperatures, but (in these cases) never beyond the parameterized optimum values (dashed lines). In contrast, the period of flowering to veraison (FL-VER) lengthens as climate goes from 0°C to 4°C warming (c,f), and temperatures are often higher than the fitted optimums—this leads to delaying phenology. This is seen in our models most commonly for late varieties such as Monastrell and Grenache because they take longer to move through phenological stages (because of their high requirements of forcing accumulation, F^*) – and thus experience the higher temperatures longer than early-ripening varieties.

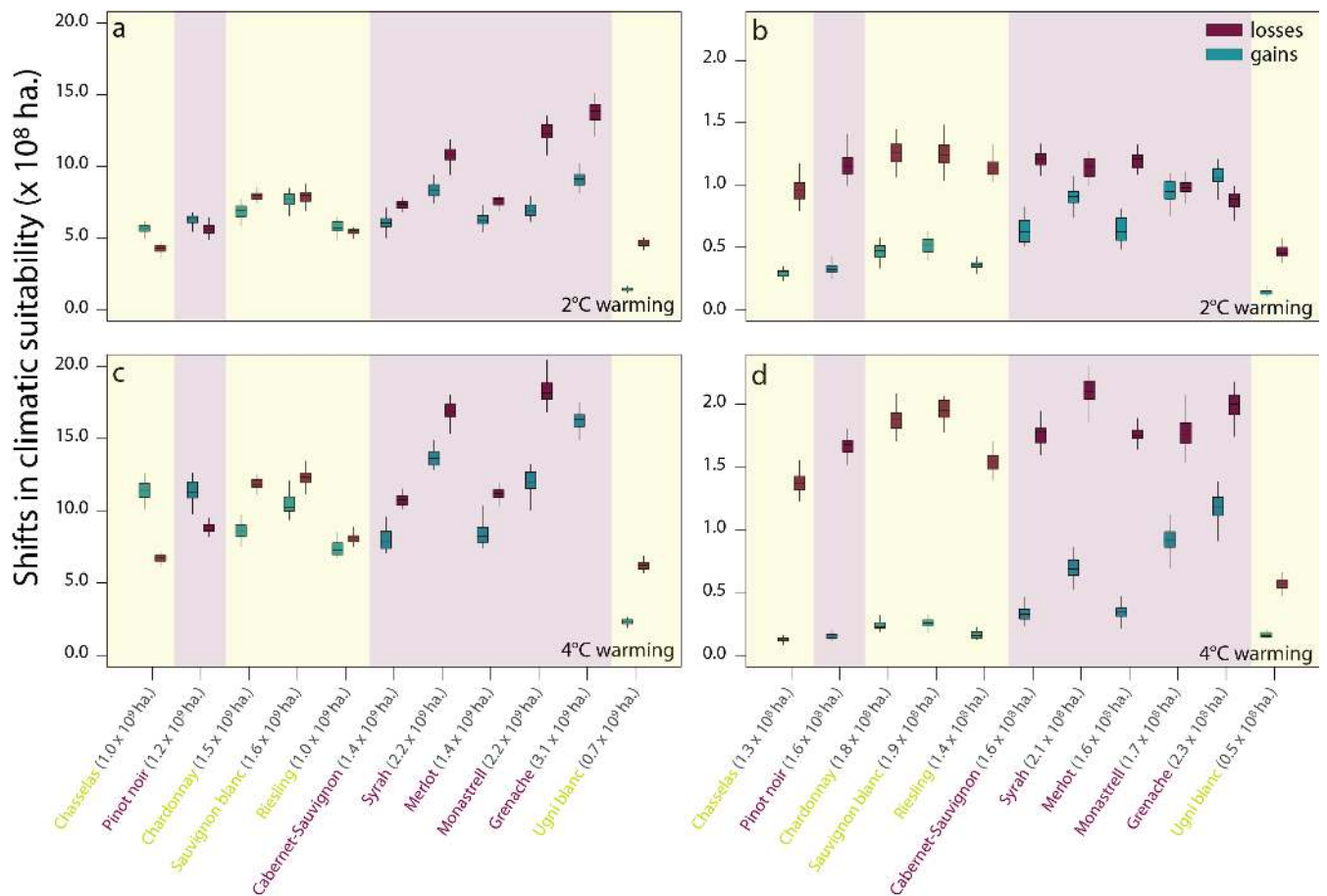


Fig. S23. Predicted gains (turquoise) and losses (purple) of climatically suitable growing areas for each of the 11 studied winegrape varieties (cultivars) under scenarios of (a, b) 2°C and (c, d) 4°C warming. We show gains and losses as absolute values (in hectares) for each variety with warming (i.e., the area predicted with warming relative to the area predicted under our reference scenario of 0°C). Both gains and losses are calculated relative to areas identified as climatically suitable under our reference scenario both globally (a, c) and within current winegrowing regions (b, d). Values in parenthesis in the x-axes report the amount of hectares with suitable climate to grow each variety under the 0°C reference scenario. These values differ among panels as suitable hectares are computed either across all the world (a, c) or within current growing regions (b, d). Note that both gains and losses computed for global suitable areas (a, c) should not be interpreted as actual gains or losses of suitability for winegrowing but as potential ones if winegrapes had been planted in all climatically suitable regions of the planet in our reference scenario of 0°C. Background shading and variety name coloring differentiates red from white varieties.