

Classification of wine grape cultivars in Chile and France according to their susceptibility to *Botrytis cinerea* related to fruit maturity

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Keywords:	Botrytis bunch rot, Grape maturity, Resistant, Susceptibility Index, <i>Vitis vinifera</i>

Reviewer 1

Comment	Answer
<p>The authors did a good job for classification based on climate, phenology and their observations as well as others authors observations already published. However, there is nothing in the index concerning the yield of the production for each variety that should be taken into account as well as the thickness of the skins of each grapes variety. In my opinion this 2 parameters should be also taken into account if the authors want to be closer to the reality and to the different conditions that can apply for each vines variety growing. Can the authors add elements on this aspect and in the new index?</p>	<p>We agree with the reviewer that these two parameters are important to be closer to the reality under the different conditions. In our study, we did not measure the yield or thickness of the berry skins. Therefore, unfortunately, we were not able to add and analyse more data related to these two parameters. However, after considering the importance of some bunch morphological aspects, such as the compactness (more compact clusters may be associated with higher yields and thinner berry skins, thus increasing berry susceptibility to <i>B. cinerea</i>), we have added in the discussion (lines 522-545) a new paragraph on the possible relationships among BBR intensity, cluster compactness and other key agronomic factors. Thus, future field investigations should be conducted to better determine the relationships between these parameters and the classification of cultivars according to the susceptibility to the disease.</p>
<p>There is different type of <i>Botrytis cinerea</i> strains and some of them can conduct to more severity than some others. How the authors deals with these aspects ? They should include possible modulation concerning the index on this aspect.</p>	<p>Yes, there are different types of <i>B. cinerea</i> strains, notably based on the transposon genotypes differing in virulence in grapevine, and this might have affected the results to a certain extent. In our study, we did not consider the phenotypic variability of the pathogen. Thus, we cannot include this aspect in the index, as suggested by the reviewer. Nevertheless, it has been demonstrated that the two major sympatric transposon genotypes in <i>B. cinerea sensu stricto</i> (excluding <i>B. pseudocinerea</i>) are present similarly in vineyards in Chile and France. Moreover, they seem to have similar key phenotypic features in both countries. A paragraph on this aspect was added to the Discussion (lines 465-482).</p>

Reviewer 2

Comment	Answer
<p>The main objective of the study was to set up a procedure to classify the wine grape cultivar according to the susceptibility to <i>B. cinerea</i> despite contrasting climatic conditions and cropping.</p> <p>The idea is good but no evidence on the "validation" of the method are reported.</p>	<p>The main goal of this work was not only to set up a procedure to classify the cultivars but also to compare the susceptibility of different wine-grape cultivars to <i>B. cinerea</i> under contrasting climatic and cropping conditions. For this purpose, it was necessary to objectively group and classify the cultivars and then to question the already published classifications, which were made based on experience rather than experimental data. Our method used to classify the cultivars was based on a methodological reference, the index calculated by Boso et al. (2014) (see Materials and methods lines 161-164).</p> <p>It should also be noted that an overall validation of our cultivar classification is the fact that most of the cultivars were finally classified in accordance with the literature (even if there are a few discrepancies).</p> <p>Furthermore, a secondary objective of our study was to demonstrate how a key potential explanatory factor, <i>i.e.</i>, fruit maturity, supports the observed differences in susceptibility to <i>B. cinerea</i>. This is clarified in the text (see the Abstract, lines 7-9, and the Introduction, lines 83-87).</p>
<p>English must be improved and also the Authors must apply all the standard detailed in the guidelines for the authors (<i>i.e.</i> citation in the text and references).</p>	<p>English editing was performed by a professional from the "American Journal Experts" service.</p>
<p>The results herein presented are sometimes conflicting with those reported in other researches and it is not well supported.</p>	<p>Our results come from experimental fields in which the measurement protocols (e.g. numbers of sampled vines, assessment stages/dates, and observation form) were standardized; thus, the results are as objective as possible. In contrast, the classifications reported in the literature are mostly based on professional experience rather than experimental data, as stated in the introduction (see lines 58-64). Thus, it may be expected, to a certain extent, that our results differ from some previously reported ones. Even the data from different sources in the literature may also be in conflict, and some of them differ in cv ranking, occasionally to a large extent (Table 1).</p> <p>When our results were very different from those reported in the literature, a possible explanation has been put forward in the discussion section (see Discussion, lines 391-426).</p>
<p>No sprays to control downy mildew are reported. Is it true?</p>	<p>The main viticulture areas in Chile (located at the centre of the country) are usually not favourable to downy mildew development; thus, no specific spray was included in the</p>

	<p>phytosanitary program. This is the case at the Panguilemo experimental station, where no specific fungicide spray to control downy mildew was applied in any year. In contrast, in France, four fungicide applications were used to control downy mildew. A paragraph addressing this issue has been added to the Materials and methods section (see Materials and methods, lines 132-139)</p>
<p>No canopy and bunch management is detailed. Is it true?</p>	<p>Neither in Chile nor in France were leaf-removal or bunch thinning performed during the studied seasons. This information has been added to the Materials and methods section (see Materials and methods, lines 130-131).</p>
<p>BBR was assessed observing the "surface" of the clusters. Is it true? No information on BBR inside the cluster? Can it twist the result?</p>	<p>Yes, BBR was assessed by observing the surface of the clusters, and it was confirmed by looking more precisely, if possible, within the grapevine bunches.</p> <p>BBR developing within the cluster could possibly affect the results, but only slightly. Usually, both parts of the clusters, <i>i.e.</i>, the surface and the inside part, are attacked by the pathogen, and there are generally no BBR attacks that affect only the bunch surface. We also preferred evaluating the surface because this methodology has been used in most of the works reported in the literature (e.g. Valdés-Gómez et al., 2008, González-Domínguez et al., 2015). This allowed us to better compare all of the available results. A paragraph addressing this issue has been added to the Materials and methods section (see Materials and methods, lines 154-157).</p>
<p>None relation with morphological and structural aspects has been reported by the Authors. All them can contribute to the assessed result</p>	<p>We agree with the reviewer, but identifying the various relationships with morphological and structural aspects was not the main objective of the present study. A paragraph addressing this issue has been added to the discussion section (see the Discussion, lines 522-545).</p>

Bibliography

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(i) **Title:**

Classification of wine grape cultivars in Chile and France according to their susceptibility to *Botrytis cinerea* related to fruit maturity

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All authors contributed significantly to this manuscript, and they are in agreement regarding the content of it. This article or one substantially similar has not been published or submitted for publication elsewhere.

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18 This study has been performed in the framework of the French Cluster of
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20 Excellence COTE, Bordeaux.
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24 **(vii) Disclosure statement:**

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26 We declare the absence of any conflict of interest in the submitted manuscript.
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29 **(viii) Short running title:**

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31 Susceptibility of wine grape cvs to *Botrytis cinerea*
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5 24 **Keywords:** Botrytis bunch rot, Grape maturity, Resistant, Susceptibility Index, *Vitis*
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7 25 *vinifera*.
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10
11 27 **Introduction**

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13
14 28 *Botrytis cinerea* is a polyphagous fungus that infects more than 1400 species of cultivated
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16 29 plants (Elad et al. 2016). On grapevine, this fungus causes one of the most serious
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18 30 diseases, namely, Botrytis Bunch Rot (BBR). The pathogen can reduce drastically both
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20 31 the yield and quality of wine (Ribéreau-Gayon et al. 1998), especially sensory qualities
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22 32 such as colour, taste and odour (Pszczolkowski et al. 2001). Important organoleptic
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24 33 negative consequences are perceived in the wine from a threshold of 5% fruit infection at
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26 34 harvest (Ky et al. 2012). Thus, this fungus causes substantial economic losses in
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28 35 grapevines, which have been estimated to be approximately 2 billion \$US per annum
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30 36 (Elmer and Michailides 2004).

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33 37 To control this disease, fungicides have long been used (Rosslenbroich and Stuebler
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35 38 2000), leading to the generation of site-specific fungicide resistant strains (Hahn 2014)
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37 39 and harm to both human health and the environment (Damalas and Eleftherohorinos
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39 40 2011). Therefore, new control strategies that allow growers to reduce the application of
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41 41 pesticides should be developed based on the principles of Integrated Pest Management
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43 42 (IPM) (IOBC 2007). In this context, some cropping practices aiming at BBR control
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45 43 should contribute to decrease the favourable conditions for the pathogen's development.
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47 44 This development depends on three major factors: i) climatic and microclimatic
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49 45 conditions, ii) the presence/amount and characteristics of the pathogen inoculum, and iii)
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51 46 the susceptibility of the host, i.e., grapevine. Climatic and microclimatic conditions,
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53 47 specifically temperature and humidity, are key factors for *B. cinerea* infection, notably in
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4 48 grapevine (Savage and Sall 1984, Thomas et al. 1988, English et al. 1989, Nair and Allen
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7 49 1993, Broome et al. 1995, Fermaud et al. 2001, Valdés-Gómez et al. 2008, Ciliberti et al.
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9 50 2016). Favourable climatic conditions are temperatures between 15 and 25°C and wetness
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11 51 duration between 12 and 24 h (Thomas et al. 1988). Concerning the pathogen, the
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13 52 population genetic structure is also a key factor to consider in the epidemiological
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15 53 development (Giraud et al. 1997, 1999, Levis et al. 1997, Beever and Weeds 2004,
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17 54 Martinez et al. 2003, 2008, Walker 2016). Regarding the host, the disease development
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19 55 depends on various genetic and phenotypic traits, such as the cluster compactness and
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21 56 morphological, anatomical, and chemical features of the berry skin (Latorre 2015), which
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23 57 are highly dependent on the grapevine cultivar.
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28 58 Grapevine cultivar susceptibility to *B. cinerea* can be considered an essential management
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30 59 indicator in IPM. Although different cultivar classifications according to their
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32 60 susceptibility to the pathogen are available in the literature (Orffer 1979, Brocuher-
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34 61 ACTA-ITV 1980, Robinson 1986, Jackson and Schuster 1987, Galet 1988, Dry and
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36 62 Gregory 1990, Marois et al. 1992, Dubos 2002), they sometimes differ greatly from one
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38 63 another (Table 1). This situation may have come to be because the proposed
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40 64 classifications are based mostly on professional experience rather than experimental data.
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43 65 Additionally, there are some gaps in these classifications: i) few studies compare the
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45 66 cultivars under the same environmental and management conditions, and ii) no study has
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47 67 proposed a cultivar susceptibility ranking that considers contrasting climatic and cropping
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49 68 conditions, e.g., northern vs southern hemisphere.
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54 69 The cropping conditions include agronomic factors, such as the canopy and/or foliar
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56 70 density, water and mineral nutrition, grape training systems and winter pruning, which
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58 71 also predispose grapevine berries to *B. cinerea* infection (Latorre 2015). Several studies
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5 72 have investigated the relationship between *B. cinerea* development and these factors
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7 73 (Barbetti 1980, Savage and Sall 1984, Marois et al. 1986, Gubler et al. 1987, English et
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9 74 al. 1989, Vail and Marois 1991, Zoecklein et al. 1992, Percival et al. 1994, Ferree et al.
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11 75 2003, Mundy 2007, Valdés-Gómez et al. 2008, Hed et al. 2009, Molitor et al. 2011,
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13 76 Pereira de Bem et al. 2015), but most often by taking into account and investigating only
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15 77 one model cultivar. Similarly, some works have studied the correlation between maturity
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17 78 and disease infection (Kosuge and Hewitt 1964, Blakeman 1975, Coley-Smith et al. 1980,
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19 79 Doneche 1986, Padgett and Morrison 1990, Vercesi et al. 1997, Mikota et al. 2003,
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21 80 Deytieux-Bellau et al. 2009), but none of them have related a classification of many
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23 81 cultivars with an explanatory factor of sensibility to the pathogen, such as the grape
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25 82 maturity.
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30 83 Thus, the main objective of this work was to compare and classify the susceptibility to *B.*
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32 84 *cinerea* between different grapevine cultivars in two contrasting climatic and cropping
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34 85 conditions, in Central Chile and Western France. Additionally, the fruit maturity was
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36 86 simulated, and we analysed the extent to which this factor may account for the
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38 87 susceptibility rankings.
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44 89 **Materials and methods**

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47 90 This study evaluated the susceptibility to Botrytis Bunch Rot (BBR) of different *Vitis*
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49 91 *vinifera* L. cultivars under contrasting conditions. The analysis was performed in three
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51 92 grapevine collections, two of them located in France and one in Chile. A total of 33 and
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53 93 22 cultivars were evaluated in both grapevine collections located in Aquitaine Region in
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55 94 France, in the sites “Tour Blanche” (Bommes 44°32'33.81" N, 0°21'02.17" W, 57 m.a.s.l)
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57 95 and “Grande Ferrade” (Villenave d’Ornon 44°47'15.4"N, 0°34'37.43"W, 22 m.a.s.l),
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5 96 respectively (Table 2). In contrast, 19 cultivars were evaluated in Maule Region in Chile,
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7 97 in the site “Panguilemo” (Panguilemo, 35°22.24’ S, 71°35.62’ W, 125 m.a.s.l). A total of
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9 98 13 common cultivars were evaluated in both countries. The experimental trials were
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11 99 performed during three seasons in the “Tour Blanche” site (2011, 2012, 2014), one season
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13 100 in the “Grande Ferrade” site (2011) and two seasons in Panguilemo site (2013-14, 2014-
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20 21 103 *Climatic characterization*

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23 104 The climatic conditions are different in the two regions. The sites located in France are
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25 105 characterized by an Oceanic climate with mild temperatures and annual rainfall of 890
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27 106 mm, with approximately 55 and 45% falling during the autumn-winter and spring-
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29 107 summer periods, respectively. In contrast, the site in Chile has a Dry Mediterranean
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31 108 climate with an annual rainfall of 600 mm, with more than 500 mm (80%) falling during
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33 109 the autumn-winter period. To characterize the climatic conditions for the study seasons
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35 110 of both sites, an automatic weather station (AWS) (Adcon Telemetric, A730,
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37 111 Klosterneuburg, Austria in Chile and Cimel Electronique S.A.S, CimAGRO, Paris in
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39 112 France) were installed 50 m from the trial plots and provided data about the air
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41 113 temperature, relative humidity and precipitation at 15-min intervals.

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43 114 Since Chilean climatic conditions were not favourable to *B. cinerea* development, we
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45 115 moistened the vines during the second season (2014-15) to promote the pathogen
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47 116 development. For this, the vines were water sprayed using a knapsack sprayer (Solo 435).
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49 117 At two consecutive days, close to harvest (approximately 25°Brix), a total of 2 L of water
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51 118 was applied per vine, every 2 hours from 8 pm (day 1) to 9 pm (day 2), resulting in the
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53 119 fruit being moistened for a period of 36 hours.
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7 121 *Experimental conditions*

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9 122 The characteristics of the experimental fields are summarized in Table 3. The main
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11 123 differences between experimental sites are the irrigation and rootstock. The use of
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13 124 irrigation is typical in vineyards in central Valley in Chile but not in Western France. In
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15 125 contrast, vines were grafted in French sites, but in Chile, the vines were planted on their
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17 126 own roots. Concerning disease management and with the aim to study the cultivar
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19 127 susceptibility to *B. cinerea*, no fungicide was applied to control this pathogen. For the
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21 128 others crop managements, conventional agricultural practices as used in commercial
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23 129 vineyards in Central Chile and Western France were used throughout the study period.

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25 130 Neither in Chile nor in France were leaf removal and/or cluster thinning performed during
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27 131 the studied seasons. The vineyards were protected against European Grapevine Moth, and
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29 132 sulphur sprays were applied to control Powdery Mildew in both countries. Additionally,
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31 133 one application of quinoxyfen (Legend ®), one of tebuconazol (Corail ®) and one of
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33 134 trifloxystrobin (Natechez ®) were used to control Powdery Mildew in France, whereas
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35 135 one application of flusiolazol (Nustar ®) and one of penconazol (Topas ®) were
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37 136 performed in Chile. Downy Mildew was controlled only in France with four fungicide
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39 137 applications per season, corresponding to two applications of cymoxanil (Option ®) and
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41 138 two copper applications. In Chile, due to the unfavourable conditions for grapevine
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43 139 Downy Mildew, no sprays were applied in any season and site.

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45 140 Regarding the experimental design at both sites, in the “Tour Blanche” site (France), each
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47 141 cultivar was replicated two times in a random design, and each replication consisted of a
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49 142 total of 6 adjacent vines. For the site “Grande Ferrade” (France), the cultivars were
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51 143 repeated in a randomized block design (4 blocks), and each block consisted of a total of
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4 144 10 vines. Finally, in “Panguilemo” (Chile), each cultivar was replicated four times in a
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6 145 randomized block design (to remove the effect of the soil slope), and each block consisted
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9 146 of a total of 15 vines.
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13 148 *Disease susceptibility assessment*

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16 149 To determine the susceptibility of the different cultivars, the incidence and severity of
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19 150 BBR were evaluated at harvest (approximately 25° Brix) in each study season. In France,
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21 151 the surface of all clusters from 3 vines per cultivar, corresponding to environ 70 clusters,
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23 152 was visually evaluated. In Chile, 5 and 20 vines per cultivar, corresponding to
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26 153 approximately 110 and 500 clusters, were evaluated in 2013-14 and 2014-15,
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28 154 respectively. BBR was assessed by observing the surface of the clusters because this
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30 155 methodology has been used in most published works (e.g., Valdés-Gómez et al., 2008,
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32 156 González-Domínguez et al., 2015), thus allowing more direct comparisons of the results
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34 157 from different sources. The incidence was obtained by dividing the number of clusters
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37 158 infected by the total number of clusters. The severity was calculated in each cluster as the
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40 159 percentage of the rotted and/or sporulating area. Both the incidence and severity were
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42 160 expressed as percentages.
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45 161 Additionally, to classify the 13 common cultivars in both countries, a susceptibility index
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47 162 (SI) was calculated using the severity data. The SI was calculated using as reference the
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49 163 index calculated by Boso et al. (2014). Thus, the SI values were calculated for all cultivars
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51 164 at each season and site as specified in equation (1):
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$$56 \quad SI = \frac{\text{Severity (\% for cultivar in question)}}{\text{Highest severity (\%) recorded in the season and in the most rotted cultivar}} \times 100 \quad (1)$$

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5 168 The cultivars were then classified into 5 categories of susceptibility: Highly Resistant
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7 169 (HR) = 0-3.5%; Resistant (R) = 3.51-10%; Intermediate (I) = 10.1-25%; Susceptible (S)
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9 170 = 25.1-50% and Highly Susceptible (HS) = 50.1-100%.

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14 172 ***Maturity assessment***

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16 173 A maturity index (F_{Mat}) was calculated to relate the berry maturity to the disease
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18 174 susceptibility of the 13 common cultivars in France and Chile. The index was calculated
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20 175 for each season and site using the Grapevine Flowering Veraison model (GFV) of Parker
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22 176 et al. (2011, 2013) and weather data for each study season, as indicated in equation (2).
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25 177 This phenological model was chosen because it was developed under similar conditions
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28 178 as observed in France and it was calibrated at the Panguilemo site, Chile (data not shown).

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$$F_{Mat} = F_{B.c \text{ assessment}} - F_{veraison} \quad (2)$$

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35 181 where $F_{B.c \text{ assessment}}$ is the timing of the *B. cinerea* assessment in each study season and
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37 182 $F_{veraison}$ is the timing of veraison for each cultivar, using the model proposed by Parker et
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39 183 al. (2011, 2013). Both variables were estimated as the critical degree-day sum (above
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41 184 0°C) calculated from the 60th and 242th day of the year in France and Chile, respectively,
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43 185 to the dates of *B. cinerea* assessment ($F_{B.c \text{ assessment}}$) and veraison ($F_{veraison}$). In Chile, the
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45 186 $F_{veraison}$ was corrected according to the results of calibration process by subtracting 100
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47 187 from the $F_{veraison}$ value proposed by Parker et al. (2013).

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50 188 Finally, to prevent the effect of the different dates of assessment depending on the season,
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52 189 the F_{Mat} was adjusted (F_{Mat_adj}) in both countries by removing the value of F_{Mat} of the latest
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54 190 cultivar, i.e., Petit Verdot, among the 13 cultivars studied, as shown in equation (3):
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$$F_{Mat_adj} = F_{Mat} \text{ for each cultivar} - F_{Mat} \text{ Petit Verdot} \quad (3)$$

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9 194 ***Statistical analyses***

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11 195 To determine differences of disease incidence and severity among the cultivars, an
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13 196 analysis of variance (ANOVA) was performed using the PROC GLM procedure for each
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15 197 experimental site. The variable “Cultivar” was considered as a fixed factor, whereas the
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17 198 variable “season” was considered as a random factor. When significant differences were
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19 199 found, a least significant difference (LSD) test at a significance level of 95% ($p = 0.05$)
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21 200 was used to compare cultivars. Additionally, a cluster analysis was performed for each
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23 201 site using the disease severity data. In this analysis, the furthest neighbour method and
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25 202 the squared euclidean distance metric were used. Furthermore, to establish a classification
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27 203 for the 13 common cultivars according to their susceptibility to *B. cinerea*, a box plot
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29 204 analysis was performed using together the SI data from all sites and all studied seasons.
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31 205 Moreover, a Kruskal-Wallis analysis and a Student-Newman-Keuls test at a significance
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33 206 level of 5% ($p = 0.05$) were performed on the SI data to compare the cultivar
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35 207 susceptibility. Finally, for the 13 common cultivars, the relationship between maturity of
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37 208 cultivars and their susceptibility to the pathogen was studied and modelled using the SI,
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39 209 F_{Mat} and F_{Mat_adj} data in all sites and study seasons. To build this relationship, a nonlinear
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41 210 model based on the equation $SI = a * (F_{mat_adj})^b$ was chosen. In both analyses using SI data
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43 211 (Box Plot and modelling), we did not include the values of cv. Roussanne in 2011 because
44
45 212 the disease was difficult to assess due the presence of sour rot. All statistical analyses
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47 213 were performed using the Statistical Software Statgraphics Plus 5.1 (StatPoint Inc.,
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49 214 Warrenton, Virginia, USA).
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216 **Results**

217 *Climatic conditions*

218 In all years studied in France, spring and summer were characterized by humid and
219 temperate conditions, which favoured the growth and development of *B. cinerea* (Figure
220 1a, c). From budbreak to harvest, the mean air temperature fluctuated between 8 and 27
221 °C and was rather similar in all seasons, except in 2011, which was characterized by
222 slightly higher temperatures. From April to October, i.e., during spring and summer in
223 France, a total rainfall of 418 mm and 439 mm were recorded in 2012 and 2014,
224 respectively, whereas a total rainfall of only 240 mm was registered in 2011. However,
225 in the last year, half of this total rainfall fell from veraison to harvest, notably in August
226 and September (124 mm), leading to favourable conditions for disease development.
227 Chilean conditions were characterized by dry and temperate spring and summer periods,
228 in both studied seasons, which were not conducive to disease development (Figure 1b, d).
229 From budbreak to harvest, the mean air temperature in both seasons ranged from 10 to 27
230 °C, similar to France. However, the total rainfall was much lower than in France: from
231 October to April, only 22 and 36 mm were recorded in 2013-14 and 2014-15, respectively
232 (Figure 1b). In the 2014-15 season, the rain periods were mostly concentrated before
233 veraison.

235 *Disease incidence and severity under field conditions*

236 *Experiments in France*

237 In the “Tour Blanche” site for the different *Vitis vinifera* cultivars evaluated, the mean
238 values of disease incidence and severity for the three studied years fluctuated from 0 to
239 98% and from 0 to 66%, respectively (Table 4). In contrast, for disease incidence, in 2011,

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4 240 the cultivars Riesling, Semillon, Muscat Petit Grain, Chenin, Folle Blanche, Roussanne
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7 241 and Negrette showed the highest values (> 83%). In contrast, Gros Manseng, Petit Verdot,
8
9 242 Petit Manseng and Cabernet Franc showed the lowest values (< 16%). In 2012, the
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11 243 cultivars Sauvignon Blanc, Chardonnay, Folle Blanche, Riesling, Muscadelle, Muscat
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13 244 Petit Grain, Grenache Blanc and Semillon showed the greatest incidence values (> 84%).
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16 245 However, Grenache Noir, Carignan, Tannat, Cabernet Sauvignon, Petit Verdot, Merlot,
17
18 246 Cabernet Franc and Petit Manseng showed the lowest values (< 18%). In 2014, the
19
20 247 cultivars Semillon, Folle Blanche and Pinot Noir showed the highest incidence values (>
21
22 248 74%), whereas Cabernet Franc, Syrah, Grenache Noir, Gros Manseng and Petit Manseng
23
24 249 showed the lowest values (< 14%).
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26
27 250 In contrast, for disease severity, in 2011, Riesling showed the highest value (66%),
28
29 251 followed by Semillon and Chenin (39%), consistent with the incidence levels. Moreover,
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31 252 the cultivars Gros Manseng, Petit Manseng, Cabernet Franc, Colombard, Cabernet
32
33 253 Sauvignon, Tannat, Merlot and Petit Verdot showed the lowest severity values (< 1.3%).
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35
36 254 In 2012, Riesling again was the most rotted cv, with a severity value reaching 47%,
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38 255 followed by Folle Blanche and Sauvignon Blanc (approximately 31%). Grenache Noir,
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40 256 Petit Verdot, Gros Manseng, Carignan, Cabernet Sauvignon, Petit Manseng, Cabernet
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42 257 Franc, Rolle, Tannat, Mourvèdre, Colombard, Ugni Blanc and Merlot were the least
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44 258 attacked, showing the lowest severity values (< 1.2%). In 2014, Folle Blanche showed
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46 259 the highest disease severity (30%), followed by Pinot Noir (22%). Gros Manseng, Petit
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48 260 Manseng, Cabernet Franc, Grenache Noir, Petit Verdot, Tannat, Cabernet Sauvignon,
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50 261 Carignan, Mourvèdre and Alicante Bouchet showed the lowest severity values (< 1.2%).
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53 262 In the “Grande Ferrade” site, mean incidence and severity values, for the studied season,
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55 263 fluctuated from 65 to 100% and from 5 to 51%, respectively (Table 5). The cultivars
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4 264 Cabernet Franc, Cot, Muscadelle, Petit Verdot, Roussanne, Sauvignon Blanc, Semillon,
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7 265 Tempranillo and Touriga Nacional showed the highest disease incidence, greater than
8
9 266 98%. However, Mourvèdre showed the lowest value (65%). The cultivar Roussanne
10
11 267 showed the highest disease severity value (51%), whereas the cultivars Marselan and
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14 268 Mourvèdre showed the lowest values (< 8%).

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19 270 *Experiments in Chile*

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21 271 The *V. vinifera* cultivars evaluated showed disease incidence and severity values lower
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23 272 than in France in both years (Table 6). The cultivars Cabernet Franc, Cabernet
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25 273 Sauvignon, Cot, Merlot, Mourvèdre and Petit Verdot did not develop any BBR symptom
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27 274 in any year, even when the vines were sprayed with water in the 2014-15 season in Chile.
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29
30 275 Thus, these cultivars are considered not susceptible to the pathogen under Chilean
31
32 276 conditions. In addition to these cultivars, Carménère, Grenache, Syrah and Tempranillo
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34 277 were not affected by the disease in 2013-14. In this season, the cultivars Gewürztraminer
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36 278 and Sauvignon Blanc showed the highest incidence values, reaching 5 and 8%,
37
38 279 respectively. In 2014-15, the cultivars Sauvignon Gris, Sauvignon Blanc and
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40 280 Gewürztraminer exhibited the greatest incidence, with values fluctuating between 12 to
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42 281 38%.

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46 282 Regarding the disease severity, in 2013-14, the cultivars Gewürztraminer and Sauvignon
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48 283 Blanc showed the highest values (approximately 0.2%), followed by Pinot Gris (0.12%).
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50 284 In 2014-15, the cultivar Sauvignon Gris exhibited the highest disease severity (9.8%),
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52 285 followed by Sauvignon Blanc and Gewürztraminer, with 3.9 and 2.3%, respectively.
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59 287 *Classification of cultivars according to the disease severity*
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288 *Situation in France*

289 In the “Tour Blanche”, the cluster analysis classified the cultivars tested into 7 groups
290 according to the disease severity (Figure 2a). The groups obtained were classified as
291 follows: resistant-intermediate "R-I" (group 1), susceptible "S" (groups 2 to 4) and highly
292 susceptible "HS" (groups 5-7) cultivars. The first group comprised 17 cultivars (Alicante
293 Bouschet to Syrah) that showed a mean severity value of 1.6% for all of the three seasons.
294 The disease severity for these cultivars was stable between seasons, i.e., the mean severity
295 fluctuated from 0.1 to 5.3% through the 3 years. The second group from the cluster
296 analysis included 3 cultivars (Gamay to Viogner) presenting a mean severity value of
297 9.8%. The third group was composed of 6 cultivars (Chenin to Negrette) presenting a
298 mean severity value of 13.8% for the three seasons. The severity values for these cultivars
299 were similar in 2011 and 2012 but lower in 2014. The fourth group, with a mean severity
300 value of 17.4%, included 3 cultivars (Chardonnay through Gewürztraminer). The fifth
301 group comprised 2 cultivars (Pinot Noir and Semillon), which showed a mean severity
302 value of 23.3%. Finally, the cultivars Folle Blanche and Riesling were classified in the
303 sixth and seventh categories showing mean severity values of 30.7 and 39.3%,
304 respectively. A particular case was the cultivar Riesling, which was classified in the most
305 susceptible category and presented a very high severity for the 2011 and 2012 seasons
306 but a relatively low severity value in 2014.

307 Furthermore, a classification was established based on all the databases from France. A
308 cluster analysis was performed with the common cultivars present in La Tour Blanche
309 and Grande Ferrade sites. The groups obtained in this analysis were classified as follows:
310 resistant-intermediate (group 1), susceptible (group 2) and highly susceptible (groups 3
311 and 4) cultivars (Figure 2b). The first group was composed of 9 cultivars (Cabernet Franc

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4 312 through Mourvèdre), with a mean severity value of 6.8%. The disease severity for these
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7 313 cultivars was similar in the “Tour Blanche” site during all the three seasons but higher at
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9 314 the “Grande Ferrade” site. The second group included 8 cultivars (Chardonnay through
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11 315 Roussanne), which were characterized by a mean disease severity value of 21%.
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13 316 Similarly, the severity results were higher at “Grande Ferrade”. Finally, the cultivars
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15 317 Pinot Noir and Riesling were classified in the third and fourth categories, showing mean
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17 318 severity values of 22.2 and 36%, respectively.
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22 23 320 *Situation in Chile*

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25 321 In Chile, the cultivars were grouped into 6 groups (Figure 3) according to disease severity.
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27 322 The groups obtained were classified as follows: resistant-intermediate (group 1),
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29 323 susceptible (groups 2 to 5) and highly susceptible (group 6) cultivars. The first group was
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31 324 composed of 12 cultivars (Cabernet Franc through Sangiovese). Within this group, 6
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33 325 cultivars did not present any rot symptom in any season. However, the other cultivars
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35 326 showed a very low mean severity value of 0.1%. The second group comprised 3 cultivars
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37 327 (Chardonnay through Roussanne) that presented a mean rot severity value of 0.2%. The
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39 328 cultivars Pinot Gris and Gewürztraminer were classified in the third and fourth groups
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41 329 with mean disease severity values of 0.4 and 1.3%, respectively. Finally, the cultivars
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43 330 Sauvignon Blanc and Sauvignon Gris were ranked in the fifth and sixth groups with mean
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45 331 severity values of 2.0 and 4.9%, respectively.
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53 54 333 *Classification of common cultivars in Chile and France according to the susceptibility*

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56 334 *index*
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5 335 According to the susceptibility index (SI), we classified the common cultivars evaluated
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7 336 in Chile and France in 5 categories: i) highly resistant (HR), ii) resistant (R), iii)
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9 337 intermediate (I), iv) susceptible (S) and v) highly susceptible (HS) cultivars (Figure 4).
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11 338 Five cultivars – Grenache Noir, Cabernet Franc, Petit Verdot, Cabernet Sauvignon and
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13 339 Mourvèdre – were highly resistant ($SI \leq 3.5$). Three cultivars were included in the
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15 340 resistant category (Merlot, Syrah and Cot). Only Roussanne was classified as an
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17 341 intermediate cultivar. Finally, the cultivars Chardonnay and Pinot Noir were identified as
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19 342 susceptible, whereas Gewürztraminer and Sauvignon Blanc were highly susceptible (SI
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21 343 > 50). This classification was corroborated with a non-parametric statistical analysis. This
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23 344 analysis demonstrated that the cultivars classified as HR and HS were stable between
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25 345 seasons and sites, in contrast with the R, I and S cultivars, which showed significant
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27 346 variability.

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348 ***Relationship between the cultivar susceptibility ranking and fruit maturity***

349 An exponential relationship between the susceptibility to the pathogen, as indicated by
350 the SI value, and the fruit maturity (F_{Mat}) of cultivars studied in France and Chile was
351 observed (Figures 5 and 6). For every combination "country x season" (experimental
352 condition), the relationship between the two variables was positive, thus showing clearly
353 that the cultivars with more mature berries were the most susceptible. This pattern was
354 very similar in all experimental conditions, but it was noticeable that the F_{Mat} values
355 differed to a large extent from one experimental condition (combination "country x
356 season") to the next (Figure 5).

357 To prevent the effect of the different dates of assessment depending on the season, the
358 F_{Mat} was adjusted (F_{Mat_adj}) in both countries by removing the value of F_{Mat} of the latest

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4 359 cultivar among the 13 cultivars studied. The relationship between F_{Mat_adj} and the SI value
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6 360 was positive and exponential in both countries (Figure 6). In France ($r^2 = 0.73$), the
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8 361 equation was $y = 3.2 \text{ E-}4 * x^{2.1}$ (Figure 6a), whereas in Chile ($r^2 = 0.55$), it was $y = 4.6\text{E-}$
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10 362 $11 * x^{4.78}$ (Figure 6b), with “y” representing the SI value and “x” the F_{Mat_adj} value. This
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12 363 pattern was quite similar in both sites, but with a steeper slope in Chile. Note that a change
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14 364 in cultivar susceptibility occurred for adjusted F-Maturity values of greater than
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16 365 approximately 250. In France, for higher F_{Mat_adj} values, the cultivars were classified as
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18 366 susceptible with an SI value higher than 25 (Figure 6a). In Chile, the cultivars with F_{Mat_adj}
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20 367 > 250 corresponded to those developing disease symptoms to some degree, whereas
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22 368 below this value, mostly no disease or very few rot symptoms were recorded (Figure 6b).
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24 369 The Roussanne cultivar was the exception in both sites, presenting a higher disease
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26 370 susceptibility in the 2012 and 2013-14 seasons, despite its low maturity (Figure 6a, b).
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372 **Discussion**

373 ***Cultivar classification according to disease susceptibility***

374 The results of this study showed that the cultivar classification according to the
375 susceptibility to *B. cinerea* was generally similar in the two countries, despite the
376 contrasting climatic conditions and cropping practices. Thus, on the one hand, the two *V.*
377 *vinifera* white cultivars Sauvignon Blanc and Gewürztraminer were classified as the
378 highest-susceptibility cultivars, followed by Chardonnay and Pinot Noir. On the other
379 hand, the four wine black cultivars – Petit Verdot, Cabernet Sauvignon, Mourvèdre and
380 Syrah – were identified as resistant or highly resistant. These classification features
381 confirm various previously published findings (Orffer 1979, Brocuher-ACTA-ITV 1980,
382 Robinson 1986, Jakson and Schuster 1987, Galet 1988, Dry and Gregory 1990, Marois

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4 383 et al. 1992, Dubos 2002) (Table 7). However, for the other cultivars tested, our results
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7 384 differ greatly from those published in the literature. We have classified the two black
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9 385 cultivars, Grenache Noir and Cabernet Franc, as highly resistant, yet they were considered
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11 386 as susceptible or highly susceptible by other authors (Robinson 1986, Galet 1988, Dry
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13 387 and Gregory 1990, Dubos 2002). Similarly, both the Merlot and Cot cultivars, which were
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16 388 identified as resistant in this study, appear in the literature as susceptible cultivars. Finally,
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18 389 we classified Roussanne as a cultivar intermediate in susceptibility, whereas it had been
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21 390 identified previously as a highly susceptible cultivar (Table 6).

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23 391 These differences observed between our results and those from the literature could be
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25 392 accounted for by possible changes in agronomic conditions that could affect the plant, the
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27 393 pathogen, the environment and/or the interactions between these epidemiological factors.
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29 394 Diverse studies have demonstrated the relationship between *B. cinerea* infection and/or
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31 395 BBR development and various environmental/agronomic factors, such as the following:
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33 396 first, climate and microclimate within the canopy (Savage and Sall 1984, Thomas et al.
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35 397 1988, English et al. 1989, Fermaud et al. 2001, Pieri and Fermaud 2005, Valdés-Gómez
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37 398 et al. 2008, Ciliberti 2015, 2016); second, canopy density and leaf removal after flowering
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39 399 (Gubler et al. 1987, English et al. 1989, Zoecklein et al. 1992, Valdés-Gómez et al. 2008,
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41 400 Molitor et al. 2011); third, cluster compactness and thinning (Barbetti 1980, Marois et al.
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43 401 1986, Vail and Marois 1991, Percival et al. 1994, Ferree et al. 2003, Hed et al. 2009,
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45 402 Molitor et al. 2011); fourth, mineral and water nutrition (Mundy 2007, Valdés-Gómez et
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47 403 al. 2008); fifth, grape training systems (Pereira de Bem et al. 2015); sixth, winter pruning
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49 404 (Savage and Sall 1984); seventh, cracks caused by biotic (insects, birds, snails, other plant
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51 405 pathogens) and abiotic (rain, hail, frost, sunburn, rapid water intake) factors (Nair et al.
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53 406 1988, Fermaud and Le Menn 1989, Coertze and Holz 1999, Becker and Knoche 2012a,
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5 407 b); and eighth, clone and rootstock (Bernard and Leguay 1988, Vail and Marois 1991,
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7 408 Derckel et al.1998, Vail et al. 1998).

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9 409 An important source of variation may be the clone effect, which may cause important
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11 410 susceptibility differences within one considered cultivar. From this point of view, Pinot
12
13 411 Noir is a model cultivar of interest. Significant differences in susceptibility to *B. cinerea*
14
15 412 between Pinot Noir clones have been attributed to variations in cluster compactness
16
17 413 (Bernard and Leguay 1988). Additionally, Derckel et al. (1998) also detected differences
18
19 414 in susceptibility to *B. cinerea* amongst the four Pinot Noir clones, suggesting that some
20
21 415 grape berry defences may play an important role in this interaction. Similarly, within the
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23 416 Chardonnay cultivar, variability in the susceptibility of different clones to *B. cinerea* has
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25 417 also been shown, although the variability attributable to the clone may be considered
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27 418 lower than the variability explained by the cultivars (Vail and Marois 1991, Vail et al.
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29 419 1998).

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34 420 The rootstock may also play an important role in the observed variability in the
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36 421 susceptibility to the pathogen among and within cultivars. For example, the SO4 rootstock
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38 422 induces higher disease infection in Pinot Noir cultivar because it promotes vine vigour,
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40 423 which is conducive to the disease (Dubos 2002). Additionally, the rootstock, by affecting
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42 424 depth of the root system and vine vigour, can influence significantly the cluster
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44 425 compactness, berry size and fruit maturity, which are known factors that modify the
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46 426 susceptibility to *B. cinerea* (Cordeau 1998).

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51 427 As a first conclusion, despite all the variations and differences possibly due to agronomic
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53 428 factors, the cultivar effect *per se* seems to be the most important for the extreme
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55 429 susceptibility groups of cultivars (highly resistant and susceptible), as defined and
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57 430 demonstrated in the present work.
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432 ***Stability of cultivar classification between years, sites and literature***

433 Our results suggest that the susceptibility of some cultivars is not stable and changes
434 depending on environmental, seasonal or management conditions. To compare the
435 differences in susceptibility and to know the stability of the cultivar classification, we
436 calculated the standard deviation corresponding to the literature results (Sdlit) and that
437 from our experimental data (Sdres) (Table 6). The susceptibility classification of Cabernet
438 Franc cultivar was not stable, neither in the literature nor in our study (Sdlit = 1.5; Sdres
439 =1.6). This could be due to the use of different clones because a great variability among
440 Cabernet Franc clones has been demonstrated to be related to key susceptibility factors,
441 notably, maturity, berry size, yield and tannin content (Van Leeuwen et al. 2013).
442 However, in our case, this difference appears to be due to the vegetative growth because
443 this cultivar was classified differently only at the “Grande Ferrade” site, at which the
444 vigour was higher. For the other cultivars, Petit Verdot and Grenache Noir, their
445 susceptibility rank was rather stable in the literature (Sdlit = 0.3 and 0.5), but it differed
446 according to the season and country in our work (Sdres = 1.6 and 1.0). For the cultivars
447 Merlot, Cot and Roussanne, the classification was the same in all other works (Sdlit = 0),
448 but it differed significantly under our conditions (Sdres = 1.5 and 1.2). Interestingly, the
449 four cultivars Grenache Noir, Petit Verdot, Merlot and Cot are susceptible to flower
450 abortion (Reynier 2011); consequently, they may present very different cluster
451 compactness depending on seasonal climatic conditions during bloom, leading to more
452 or less flower abortion (Keller 2015). Such a difference in compactness should account
453 for great variations in the susceptibility to *B. cinerea*, as has been often demonstrated in
454 the literature (Marois et al. 1986, Vail and Marois 1991, Percival et al. 1994, Ferree et al.

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4 455 2003, Hed et al. 2009, Molitor et al. 2011). Regarding the susceptibility classification, the
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6 456 cultivars Grenache Noir, Cabernet Franc, Merlot, Cot and Roussanne showed significant
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8 457 differences between literature works and our study (Table 6). To understand this
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10 458 difference, further studies about the clone and the vegetative growth related to the
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12 459 rootstock are necessary.

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16 460 It is important to note the effect of Chilean data, which decrease the average of the
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18 461 Susceptibility Index (SI) in the cultivars classification due to the existence of climatic
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20 462 conditions unfavourable to disease development. Even if the grapevines were water
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22 463 sprayed in Chile, this effect was temporary and did not allow the pathogen to develop to
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24 464 a large extent, as may occur under natural wet conditions such as e.g., under oceanic
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26 465 conditions. Finally, it may be discussed whether these results could have been affected
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28 466 by the phenotypic variability among *B. cinerea* strains, particularly in terms of difference
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30 467 in virulence. It has been demonstrated that the virulence of the two *B. cinerea* genetic
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32 468 types, *vacuma* and *transposa*, differed significantly in terms of disease incidence and
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34 469 severity, with *transposa* strains being more virulent than *vacuma* ones. This virulence on
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36 470 leaves or on berries was significantly and negatively correlated with the mycelial growth
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38 471 rate (Martínez et al. 2005). Moreover, the mechanism involved in this pathogenicity could
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40 472 be explained by the presence of transposable elements, which is a characteristic feature
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42 473 of *transposa* isolates. Thus, Baulcombe (2013) explained that transposon small RNA
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44 474 (sRNA) molecules are associated with the suppression of host defences, which may have
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46 475 important implications for the pathogen arms race. This idea is supported by Weiberg et
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48 476 al. (2013), who founded that transposon sRNA molecules derived from *B. cinerea* can act
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50 477 as effectors to suppress host immunity and play a positive role in pathogenicity. Thus,
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52 478 although we did not consider the high phenotypic variability in this study, it has been
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5 479 demonstrated that the two major sympatric transposon genotypes (*transposa* and *vacuma*)
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7 480 are present similarly in Chile as in France (Martinez et al. 2003, 2008). They also tend to
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9 481 have similar characteristics in both countries (Muñoz et al. 2002); consequently, this
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11 482 variability should not affect the results to a great extent.
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15 16 484 *Effect of grape maturity on disease susceptibility*

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18 485 The fruit maturity was identified as a major factor determining the cultivar susceptibility
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20 486 to *B. cinerea*. Several studies, often based on one selected model cultivar, have
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22 487 demonstrated that increasing sugar concentration with the phenological stage in maturing
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24 488 grape berries promotes infection and colonization by *B. cinerea*. Some of these studies
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26 489 also demonstrated that the presence of sugar in berry exudates stimulates the germination
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28 490 and mycelium growth of *B. cinerea* (Kosuge and Hewitt 1964, Blakeman 1975, Coley-
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30 491 Smith et al. 1980, Doneche 1986, Padgett and Morrison 1990, Vercesi et al. 1997,
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32 492 Deytieux et al. 2009). Despite several authors having demonstrated the relationship
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34 493 between sugar concentration and pathogen infection, few works have revealed a
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36 494 correlation between increasing maturity and progress of disease severity, and they mostly
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38 495 used a single cultivar (Fermaud et al 2011), not a set of different cultivars. Studies related
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40 496 to the infection by the pathogen and the solid soluble contents of grapes have been
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42 497 conducted, in particular by Mundy and Beresford (2007), who established clearly a
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44 498 significant and positive linear regression between berry sugar concentration and the
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46 499 percentage of rotted berries. Furthermore, regarding the maturity effect, the susceptibility
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48 500 of berries increased during ripening (Kretschmer et al. 2007), and, more precisely, a
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50 501 positive, close and sigmoid relationship between maturity variables and *B. cinerea*
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52 502 susceptibility was established by Deytieux-Belleau et al. (2009). This last study
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5 503 demonstrated that severity of *B. cinerea* increases regularly during berry maturity,
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7 504 reaching a maximum at the over-maturity stage: then, this relationship can be represented
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9 505 by a sigmoid curve. In our study, these relationships were exponential, showing that the
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11 506 most mature grapevine cultivars were the most susceptible to the pathogen. These
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13 507 cultivars were mostly white cultivars, in which the sugar content is, generally, higher than
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15 508 in black ones (Doneche 1986). If we had measured the disease severity of cultivars in a
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17 509 more advanced state of maturity, these results may have been similar. Moreover, the most
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19 510 mature cultivars correspond to the earliest cultivars. They could also have been more
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21 511 attacked because they were exposed, in a susceptible, mature stage, for a longer time
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23 512 under favourable conditions for infection and disease development.
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28 513 In addition to the maturity, other factors may account for the variability in susceptibility.
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30 514 For example, the less-susceptible cultivars, according to the disease incidence and
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32 515 severity, were in both countries black cultivars. In contrast, the most susceptible cultivars
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34 516 were white and pink ones. This relationship between susceptibility and berry colour was
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36 517 expected because it has been shown that the susceptibility of grapes may be affected by
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38 518 the concentration of phenolic compounds in grapes (Frankel et al. 1995, Goldberg et al.
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40 519 1995), and particularly, the tannin content within the berry skin (Deytieux-Belleau et al.
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42 520 2009). These results confirmed previous studies (Goetz et al. 1999, Xie and Dixon 2005)
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44 521 that demonstrated that black cultivars are less susceptible to *B. cinerea* than white or pink
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46 522 cultivars. In addition, the compactness of clusters has been shown to be an important
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48 523 morphological feature that affects the susceptibility to *B. cinerea* by affecting the
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50 524 microclimate and the thickness and wax content of the berry cuticle (Marois et al. 1986,
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52 525 Vail and Marois 1991, Percival et al. 1993, Fermaud et al. 2001). In this study, we
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54 526 observed a clear trend in the vineyard conditions that the cvs with more compact clusters
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4 527 were more severely attacked and more susceptible to the pathogen. In contrast, we noted
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7 528 that the less-attacked cvs presented looser clusters and were classified as less susceptible
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9 529 to *B. cinerea*. This corroborates a previous study that showed a positive correlation
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11 530 between BBR development and cluster compactness (Hed et al. 2009). Lastly, and in
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13
14 531 addition to the fruit maturity, berry skin colour and cluster compactness, which also may
15
16 532 affect the susceptibility to BBR, there are other predisposal factors, such as genetic
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18 533 (morphological, anatomical and chemical features of the berry skin), physical (wounds),
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20 534 environmental (climate and weather conditions) and agronomic (cultural practices)
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23 535 (Latorre et al. 2015). For agronomic factors, after the climate influence, vegetative growth
24
25 536 and canopy development are considered the second most important factors favouring *B.*
26
27 537 *cinerea* development (Valdés-Gómez et al. 2008). Then, some morphological factors
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29 538 related to cluster architecture, e.g., the bunch mass and berry number, also have an
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31 539 important influence on BBR epidemics (Vail and Marois 1991, Valdés-Gómez et al.
32
33 540 2008). The bunch mass has been positively and significantly correlated with the BBR
34
35 541 incidence and considered more relevant than the yield to account for disease development.
36
37 542 This factor contributes largely to cluster compactness; thus, it can be considered as a key
38
39 543 morphological feature that increases *B. cinerea* susceptibility (Valdés-Gómez et al.
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41 544 2008). Although in this work we did not consider any of these factors, they should be
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43 545 further studied in future works addressing cv susceptibility to the pathogen.
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51 ***Main findings and implications for IPM and climatic change adaptations***

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54 548 As previously reported, our results also confirmed that environmental conditions are a
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56 549 main factor in the disease epidemiological development (Savage and Sall 1984, Thomas
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58 550 et al. 1988, English et al. 1989, Fermaud et al. 2001, Valdés-Gómez et al. 2008, Ciliberti
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4 551 2015, 2016). The contrasting climatic conditions in the two regions studied led to different
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6
7 552 levels of disease infection, due principally very different amounts and distributions of
8
9 553 rainfall. Rainfall, which is predominantly at the origin of increased relative humidity and
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11 554 wetness duration in the vineyards, was found to be of primary importance in disease
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13 555 development (Ciliberti 2015, 2016). Thus, in France, all cultivars were attacked by *B.*
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15 556 *cinerea*, and they presented more advanced disease development than in Chile. Although
16
17 557 under Chilean conditions, no cultivars seemed to be very susceptible, considering the low
18
19 558 disease severity values, it was possible to classify them according to their susceptibility.
20
21 559 This classification was similar to that in France, thus demonstrating that climate does not
22
23 560 change the susceptibility of cultivars. However, when the climatic conditions are not
24
25 561 favourable to the pathogen development, it is difficult to differentiate resistant from
26
27 562 intermediate cultivars because the latter do not develop the disease at all. This situation
28
29 563 was observed, in particular, in grapes that were not sprayed with water in Chile (data no
30
31 564 shown). Thus, the decision to apply a fungicide to these cultivars based on their
32
33 565 susceptibility classification to BBR would be more difficult. Furthermore, it is interesting
34
35 566 to note that future climatic conditions in the Bordeaux region could be relatively similar
36
37 567 to the current climatic conditions characterizing the Chilean region considered in the
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39 568 present study (Pañitrur-De la Fuente et al. 2016). Under this context of climate change,
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41 569 strategies may be orientated by adapting the cultivar choice to future possible climatic
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43 570 scenarios, considering both the potential disease development and the associated cultivar
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45 571 susceptibility.
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47 572 Further investigation should be conducted to better understand the relationships between
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49 573 the classification of cultivars according to their susceptibility to *B. cinerea* and other
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4 574 variables (e.g., clone, vigour, and rootstock) to develop management and integrated pest
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6 575 management strategies.
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11 577 **Conclusions**

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14 578 The results of this study demonstrated that the classification of different wine cultivars
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16 579 according to their susceptibility to *B. cinerea* was generally similar in both countries,
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18 580 despite the contrasting climatic conditions and management practices. Sauvignon Blanc
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20 581 and Gewürztraminer were the most-susceptible cultivars, whereas Petit Verdot, Cabernet
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22 582 Sauvignon, Mourvèdre and Syrah were rather resistant or highly resistant. These results
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24 583 are in accordance to previous studies; however, for the other cvs that we evaluated, their
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26 584 ranking differed to some extent compared with data from the literature. This difference is
27
28 585 presumably caused by variations in the agronomic and/or environmental conditions under
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30 586 which the field experiments were performed. The interfering effects of various factors,
31
32 587 such as clone, rootstock, and cluster compactness related to flower abortion are discussed
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34 588 in detail and should be considered in further studies aiming to compare cultivar
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36 589 susceptibility to the pathogen.
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42 590 The maturity of cultivars seems to be a major determining factor in the susceptibility to
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44 591 *B. cinerea*. In our study, the relationship between fruit maturity and susceptibility to the
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46 592 pathogen was positive and exponential, indicating that the most mature grapevine
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48 593 cultivars were the most susceptible. This could be explained by the increasing sugar
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50 594 concentrations in ripening berries, which promote fungal colonization, and by the longer
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52 595 time during which later grapevine cultivars are exposed to favourable conditions for
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54 596 disease development.
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4 597 The cultivar is a principal and permanent factor affecting the susceptibility to *B. cinerea*,
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6 598 which could be modified by climate and agronomic management, which are considered
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8 599 as variable factors. Thus, the cultivar remains a key parameter in decision support
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10 600 systems, and the fruit maturity could be used to support this. Further investigation should
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12 601 be conducted to better understand the relationship between susceptibility to *B. cinerea*
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14 602 and other variables (e.g., clone, vigour, and rootstock) to develop management and
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16 603 integrated pest management strategies.
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860 **Tables**

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864 **Table 1.** Susceptibility to *B. cinerea* of 13 grapevine cultivars according to different
865 literature sources.

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Cultivar	a	b	c	d	e	f	g	h
Grenache Noir	4	3	-	-	4	-	3	4
Cabernet Franc	3	-	-	-	-	-	4	1
Petit Verdot	0-1	-	-	-	-	-	1	1
Cabernet Sauvignon	2	-	0	1	1	0	1	1
Mourvèdre	-	-	-	-	-	-	1	-
Merlot	3	-	-	-	-	-	3	3
Syrah	2	-	1	3	3	-	-	2
Cot	3	-	-	-	-	-	3	3
Roussanne	4	-	-	-	-	-	-	4
Chardonnay	4	-	2	2	3	-	3	3
Pinot Noir	3	4	2	3	4	-	-	3
Gewürztraminer	4	-	-	-	-	-	1	4
Sauvignon Blanc	4	-	4	3	4	-	1	4

867 a = Dubos (2002), b = Dry and Gregory (1990), c = Orffer (1979), d = Jackson and
868 Schuster (1987), e = Robinson (1986), f = Marois et al. (1992), g = Galet (1988), h =
869 ACTA (1980); 0 = highly resistant, 1 = resistant, 2 = intermediate, 3 = susceptible, 4 =
870 highly susceptible.

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Table 2. Cultivars evaluated at each experimental site in France and in Chile

Tour Blanche (France)	Grande Ferrade (France)	Panguilemo (Chile)	Common cultivars (France and Chile)
Alicante Bouschet	Cabernet Franc	Cabernet Franc	Cabernet Franc
Cabernet Franc	Cabernet Sauvignon	Cabernet Sauvignon	Cabernet Sauvignon
Cabernet Sauvignon	Carignan	Carménère	Chardonnay
Carignan	Chardonnay	Chardonnay	Cot
Chardonnay	Chenin	Cot	Gewürztraminer
Chenin	Cot	Gewürztraminer	Grenache Noir
Cinsault	Gamay	Grenache Noir	Merlot
Colombard	Grenache Noir	Marsanne	Mourvèdre
Cot	Marselan	Merlot	Petit Verdot
Folle Blanche	Merlot	Mourvèdre	Pinot Noir
Gamay	Mourvèdre	Petit Verdot	Roussanne

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5	Gewürztraminer	Muscadelle	Pinot Gris	Sauvignon Blanc
6	Grenache Blanc	Petit Verdot	Pinot Noir	Syrah
7	Grenache Noir	Pinot Noir	Roussanne	
8	Gros Manseng	Riesling	Sangiovese	
9	Melon	Roussanne	Sauvignon Blanc	
10	Merlot	Sauvignon Blanc	Sauvignon Gris	
11	Mourvèdre	Semillon	Syrah	
12	Muscadelle	Tempranillo	Tempranillo	
13	Muscat Petit Grain	Touriga Nacional		
14	Negrette	Ugni Blanc		
15	Petit Manseng	Vioigner		
16	Petit Verdot			
17	Pinot Noir			
18	Riesling			
19	Rolle			
20	Roussanne			
21	Sauvignon Blanc			
22	Semillon			
23	Syrah			
24	Tannat			
25	Ugni Blanc			
26	Vioigner			
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Table 3. Field characteristics of the experimental fields.

Property	France		Chile
	Tour Blanche	Grande Ferrade	Panguilemo
Experimental Period	2011, 2012, 2014	2011	2013-14, 2014-15
Vineyard planting year	1995	2009	2006
Rootstock	3309	SO4	Own-rooted
Location (WGS84)	44°32' N, 0°21' W	44°47'N, 0°34' W	35°22' S, 71°36' W
Spacing (m x m)	1.8 x 0.9	1.8 x 1.0	2.0 x 1.0
Trellis/Pruning system		VSPSystem ^a / Two-bilateral	
Irrigation system	Non-irrigated	Non-irrigated	Drip irrigation (one dropper per plant with a flow rate of 4 L / h)

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891 **Table 4.** Mean disease incidence and severity values (%) for each cultivar under field
 892 conditions in the “Tour Blanche” site (France) over three seasons.
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Cultivar	Disease incidence (%)			Disease severity (%)		
	2011	2012	2014	2011	2012	2014
Alicante Bouchet	30.7cdef	35.8bcd	33.4bcdefg	3.3ab	3.1abc	1.2a
Cabernet Franc	13.2abc	16.4ab	7.1a	0.4a	0.4a	0.2a
Cabernet Sauv.	27.6bcde	10.7ab	26.6abcde	0.9a	0.3a	0.5a
Carignan	37.9def	10.5ab	25.8abcde	1.9ab	0.3a	0.6a
Chardonnay	79.5klmn	93.1j	51.9fghijk	11.5abcde	26.4fgh	10.2defg
Chenin	94.5mn	49.4cdef	37.0cdefgh	39.0i	7.4abcd	1.7ab
Cinsault	54.9fghijk	29.8abc	55.6ghijk	5.9abcd	2.3ab	3.2abc
Colombard	18.8abcd	29.2abc	36.4cdefgh	0.7a	1.0a	1.3ab
Cot	41.2defg	46.3cde	40.5defghi	4.7abc	2.5ab	2.0abc
Folle Blanche	89.2lmn	92.8j	81.8lm	29.3ghi	32.6h	29.7i
Gamay	51.0efghi	25.6abc	51.7fghijk	13.7bcdef	3.9abc	11.1efg
Gewürztraminer	64.8ghijkl	63.5efghi	68.4jklm	19.3efg	23.3efgh	11.7fg
Grenache Blanc	65.8hijkl	86.0ij	33.4bcdefg	17.1def	17.1def	2.9abc
Grenache Noir	34.9cdef	5.6a	11.8ab	4.0abc	0.2a	0.2a
Gros Manseng	0a	15.1ab	12.4ab	0a	0.3a	0.1a
Melon	42.9defgh	73.7fghij	67.6jklm	4.5abc	10.3abcd	14.5g
Merlot	33.1cdef	15.6ab	51.3fghijk	1.2a	1.2a	3.3abcd
Mourvèdre	21.8abcd	22.5abc	26.4abcde	1.8ab	0.9a	0.7a
Muscadelle	75.9jklmn	88.2ij	51.4fghijk	17.7def	14.4bcdef	5.3abcdef
Muscat petit grain	97.2n	86.7ij	46.9efghij	29.8ghi	12.0abcde	4.4abcde
Negrette	83.8lmn	57.3defg	58.1hijk	24.4fgh	8.6abcd	7.0abcdef
Petit Manseng	12.6abc	18.1ab	13.6abc	0.3a	0.3a	0.2a
Petit Verdot	3.3ab	13.4ab	22.3abcd	1.3a	0.2a	0.4a
Pinot Noir	77.8jklmn	70.2efghij	74.0klm	32.7hi	15.6cdef	21.7h
Riesling	97.7n	91.2j	61.5ijkl	65.7j	47.1i	5.1abcdef
Rolle	48.5efghi	24.3abc	31.0bcdef	3.3ab	0.9a	2.7abc
Roussanne	88.6lmn	63.2efghi	43.1defghi	31.2ghi	7.3abcd	2.1abc
Sauvignon Blanc	71.3ijklm	96.2j	61.8ijkl	15.3cdef	30.6gh	8.3bcdefg
Semillon	96.2n	84.6hij	86.7m	39.2i	19.3defg	11.6fg
Syrah	37.0cdef	58.0defgh	11.5ab	2.6ab	11.8abcde	1.4ab
Tannat	22.4abcd	10.5ab	24.2abcde	1.1a	0.9a	0.4a
Ugni Blanc	43.3defgh	32.5abcd	56.1ghijk	2.8ab	1.1a	1.9abc
Viogner	53.5fghij	80.4ghij	53.6fghijk	8.4abcde	13.2abcde	8.7cdefg

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906 **Table 5.** Mean disease incidence and severity values (%) for each cultivar under field
 907 conditions in the “Grande Ferrade” site (France) in the 2011 season.
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Cultivar	Disease incidence (%)	Disease severity (%)
Cabernet Franc	100.0e	36.8efg
Cabernet Sauvignon	83.3bc	15.8abc
Carignan	96.3de	25.9bcde
Chardonnay	92.4cde	39.5efg
Chenin	96.4de	33.9def
Cot	100.0e	37.1efg
Gamay	93.7cde	28.8cde
Grenache Noir	91.7cde	10.1ab
Marselan	71.3ab	7.3a
Merlot	97.7de	28.6cde
Mourvèdre	65.0a	5.1a
Muscadelle	100.0e	47.7fg
Petit Verdot	98.8e	34.6def
Pinot Noir	85.4cd	18.9abcd
Riesling	95.9cde	26.0bcde
Roussanne	98.6e	51.2g
Sauvignon Blanc	98.8e	40.5efg
Semillon	100.0e	30.3cde
Tempranillo	100.0e	48.0fg
Touriga Nacional	98.8e	33.8def
Ugni Blanc	93.8cde	14.8abc
Viogner	97.5de	42.1efg

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Table 6. Mean disease incidence and severity values (%) for each cultivar under field
 conditions in Chile over two seasons.

Cultivar	Disease incidence (%)		Disease severity (%)	
	2013-14	2014-15	2013-14	2014-15
Cabernet Franc	0a	0a	0a	0a
Cabernet Sauvignon	0a	0a	0a	0a
Carménère	0a	0.3a*	0a	0a
Chardonnay	1.07a	2.7ab	0.05ab	0.30a
Cot	0a	0a	0a	0a
Gewürztraminer	8.11c	12.0cd	0.24d	2.25ab
Grenache Noir	0a	0.25a*	0a	0a
Marsanne	0.01a	0.18a*	0.01ab	0a
Merlot	0a	0a	0a	0a
Mourvedre	0a	0a	0a	0a
Petit Verdot	0a	0a	0a	0a
Pinot Gris	2.33ab	9.75bcd	0.12bc	0.78a
Pinot Noir	0.72a	3.93ab	0.06ab	0.30a

Roussanne	0.47a	0.98a	0.03ab	0.23a
Sangiovese	0a	6.05abc	0a	0.8a
Sauvignon Blanc	4.72bc	16.88d	0.19cd	3.85b
Sauvignon Gris	1.28a	37.7e	0.048ab	9.80c
Syrah	0a	0.25a	0a	0.03a
Tempranillo	0a	2.53ab	0a	0.10a

*When there is a value for the incidence but the severity is 0, it is because the severity value is less than 0.001.

Table 7. Comparison of the susceptibility to *B. cinerea* of 13 grapevine cultivars according sources and our results

Cultivar	Mean lit.	Sd lit.	Our res.	Sd res.
Grenache Noir	4	0.5	0	1.0
Cabernet Franc	3	1.5	0	1.6
Petit Verdot	1	0.3	0	1.6
Cabernet Sauvignon	1	0.7	0	1.2
Mourvèdre	1	-	0	0.5
Merlot	3	0	1	1.5
Syrah	2	0.8	1	1.2
Cot	3	0	1	1.5
Roussanne	4	0	2	1.2
Chardonnay	3	0.8	3	1.2
Pinot Noir	3	0.8	3	1.3
Gewürztraminer	3	1.7	4	0
Sauvignon Blanc	3	1.2	4	0.5

0 = highly resistant, 1 = resistant, 2 = intermediate, 3 = susceptible, 4 = highly susceptible; Mean lit = Mean of literature source, Our res = Results of our study; Sdlit = standard deviation of literature sources, Sd res = standard deviation of our results.

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5 944 **Figure legends**
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7 946 **Figure 1.** Monthly mean rainfall (mm) in France (a) and Chile (b) and mean air
8 947 temperature (°C) in France (c) and Chile (d) during all seasons. The horizontal dotted
9 948 lines in (c) and (d) represent the mean air temperature (°C) in each season. Bud =
10 949 Budbreak; Flo = Flowering; Ver = Veraison; Har = Harvest.
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12 951 **Figure 2.** Cluster classification of cultivars in France in the sites “Tour Blanche” (a) and
13 952 both “Grande Ferrade and “Tour Blanche” (b) according to their severity values.
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15 954 **Figure 3.** Cluster classification of cultivars in Chile according to their severity values.
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17 956 **Figure 4.** Box plot of cultivars according to the susceptibility index. HR = Highly
18 957 Resistant; R = Resistant; I = Intermediate; S = Susceptible; HS = Highly Susceptible. The
19 958 vertical line in each box and the cross represent the median and mean value of the SI,
20 959 respectively.
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22 961 **Figure 5.** Relationship between the maturity of cultivars (F Mat) and susceptibility to
23 962 BBR (SI), assessed at different dates, in France and Chile.
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25 964 **Figure 6.** Relationship between the maturity of cultivars (F Mat_adj) and susceptibility
26 965 to BBR (SI) at both sites, France (a) and Chile (b), during all study seasons.
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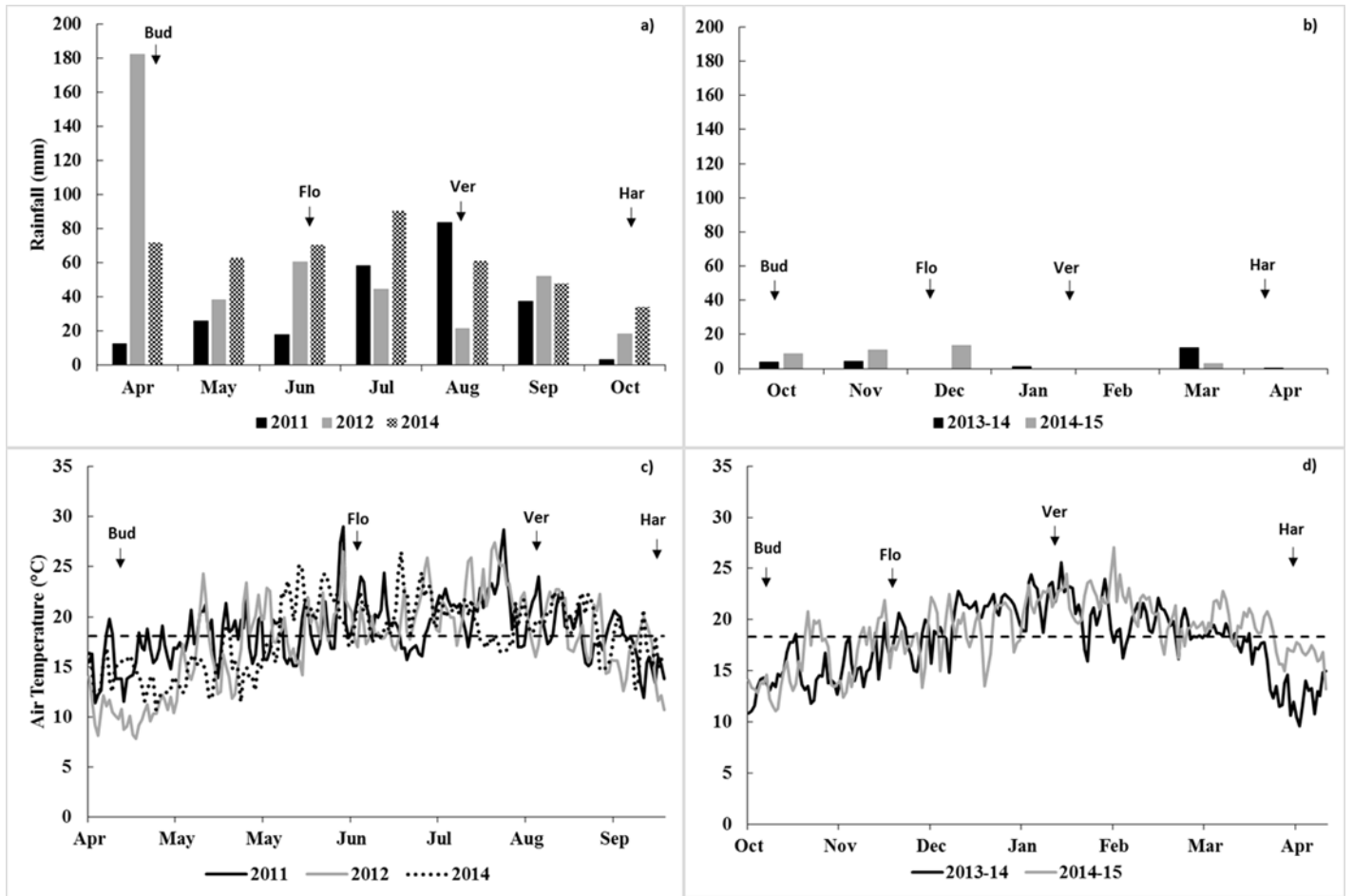


Figure 1. Monthly mean rainfall (mm) in France (a) and Chile (b) and mean air temperature (°C) in France (c) and Chile (d) during all seasons. The horizontal dotted lines in (c) and (d) represent the mean air temperature (°C) in each season. Bud = Budbreak; Flo = Flowering; Ver = Veraison; Har = Harvest.

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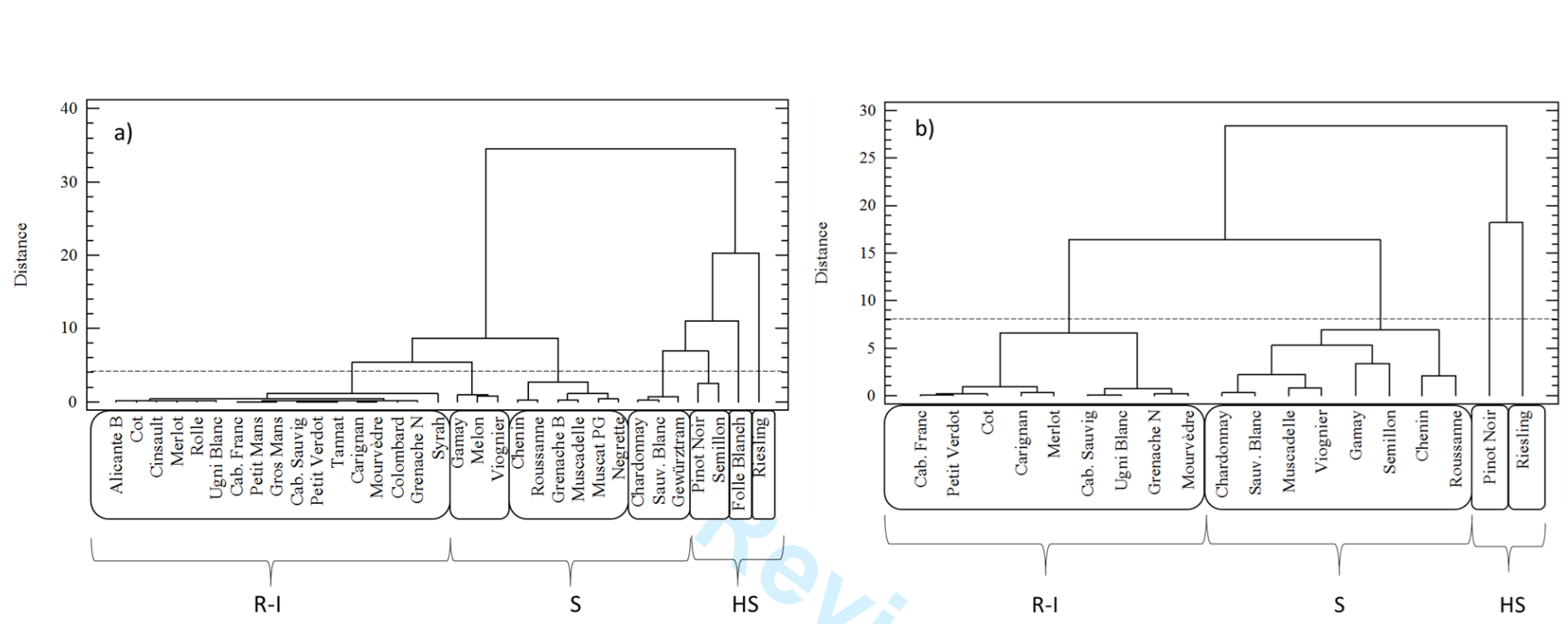


Figure 2. Cluster classification of cultivars in France in the sites “Tour Blanche” (a) and both “Grande Ferrade and “Tour Blanche” (b) according to their severity values.

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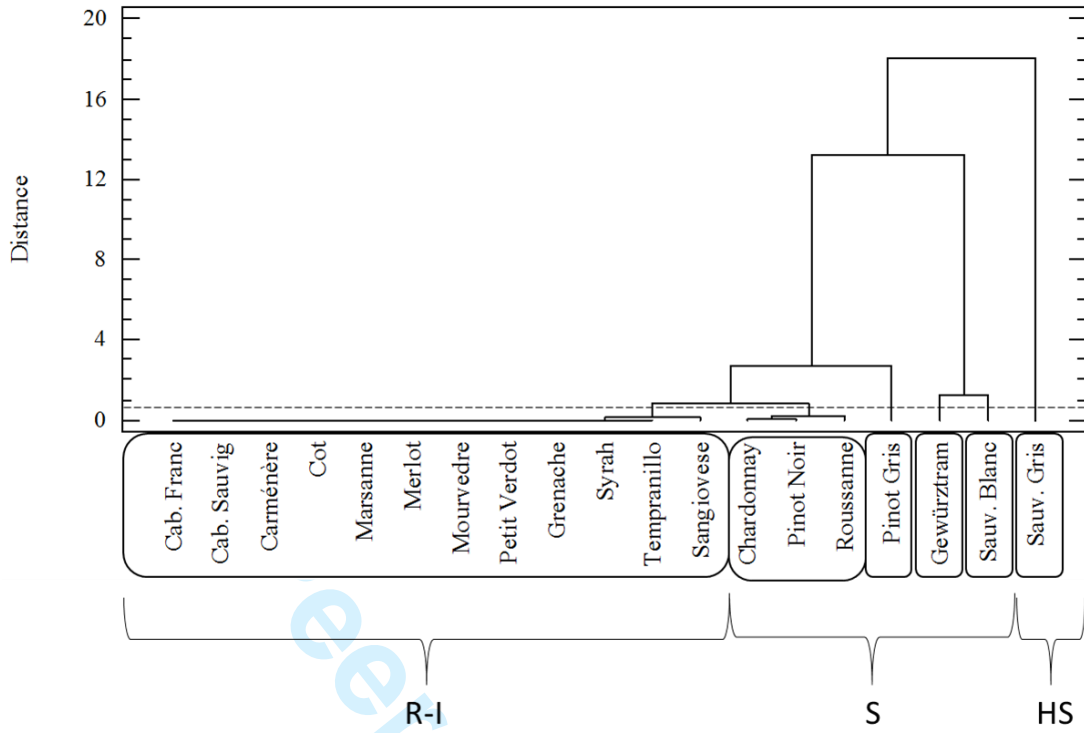


Figure 3. Cluster classification of cultivars in Chile according to their severity values.

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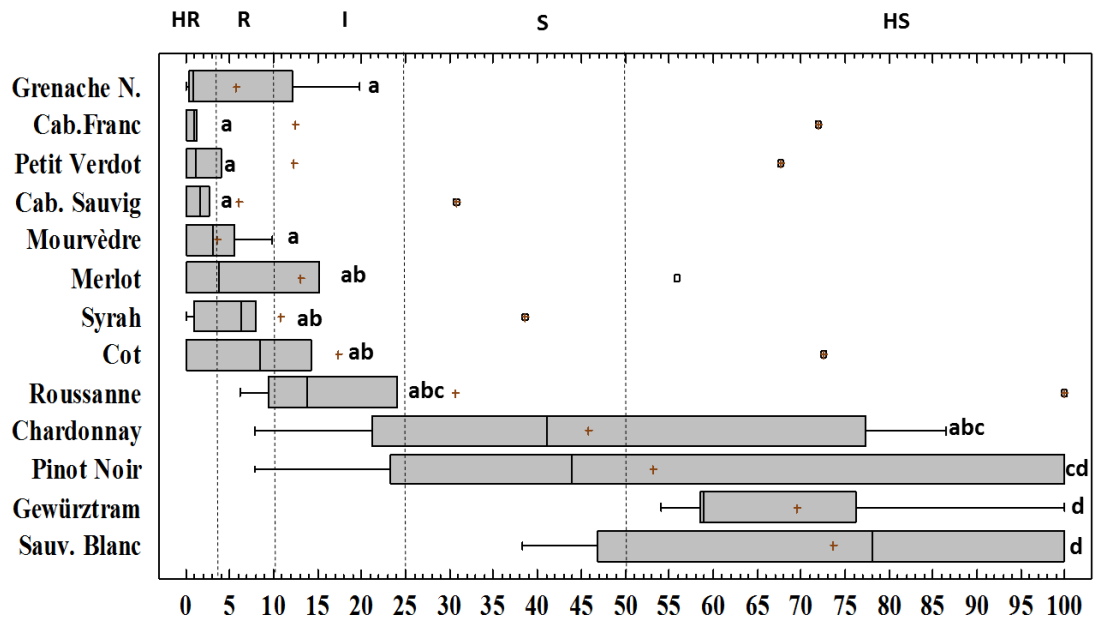


Figure 4. Box plot of cultivars according to the susceptibility index. HR = Highly Resistant; R = Resistant; I = Intermediate; S = Susceptible; HS = Highly Susceptible. The vertical line in each box and the cross represent the median and mean value of the SI, respectively.

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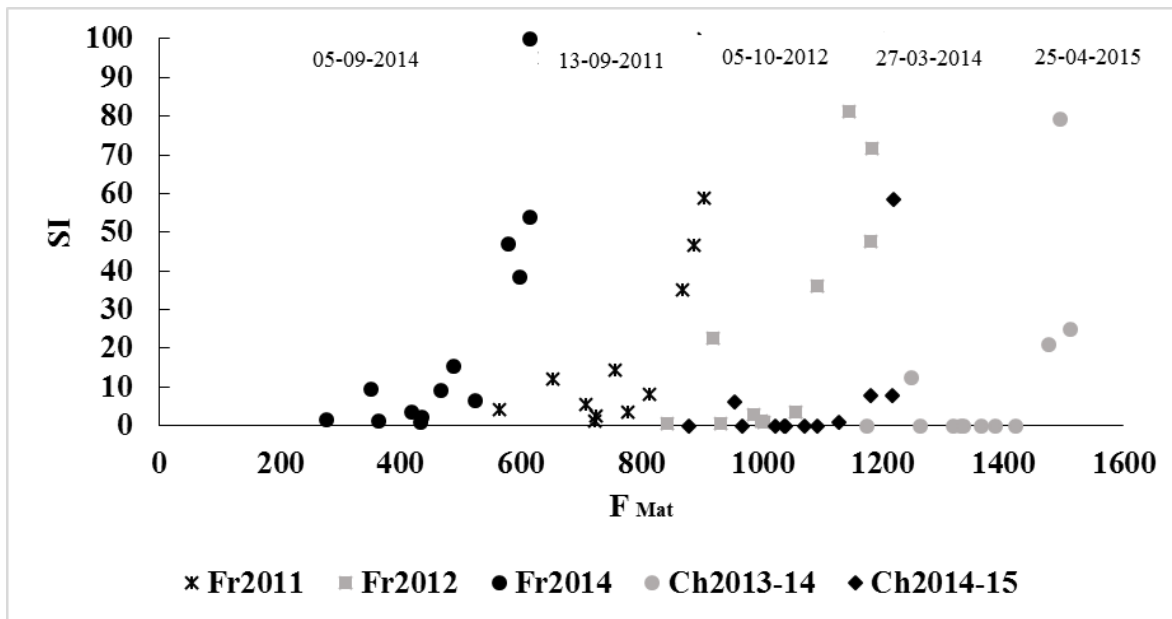


Figure 5. Relationship between the maturity of cultivars (F Mat) and susceptibility to BBR (SI), assessed at different dates, in France and Chile.

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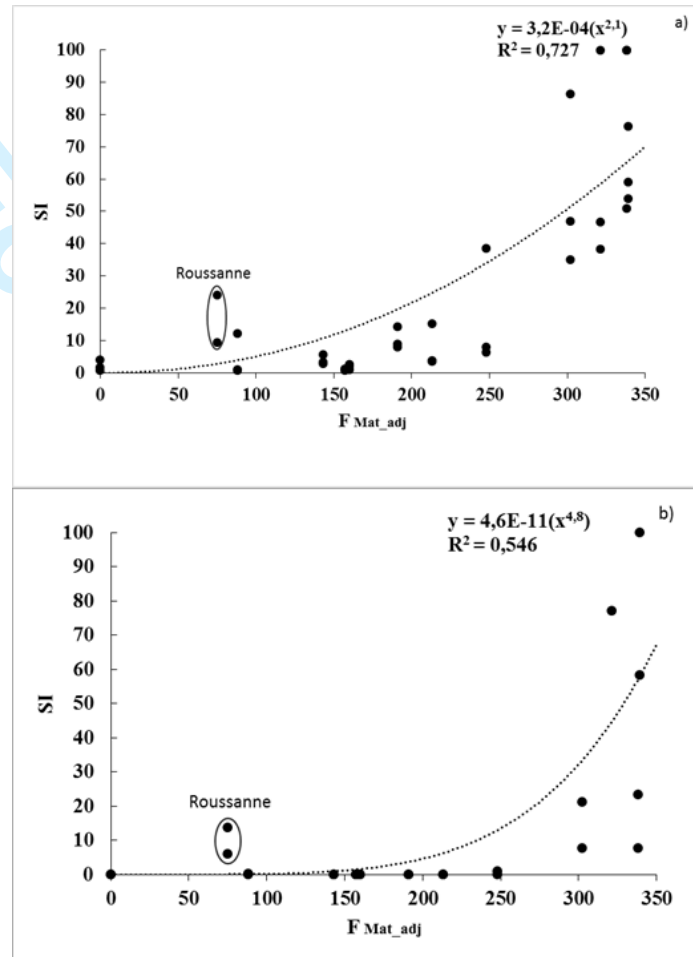


Figure 6. Relationship between the maturity of cultivars (F Mat_adj) and susceptibility to BBR (SI) at both sites, France (a) and Chile (b), during all study seasons.