Blossom blight resistance in peach: heritability and segregation in progenies from reciprocal crosses

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ABSTRACT

Blossom blight and brown rot caused by Monilinia fructicola is the most important peach disease in Brazil. Genetic resistance is a control strategy that is gaining importance in breeding programs worldwide. This study aimed to identify genotypes with higher levels of blossom blight resistance to estimate the heritability of this character; study the frequency distribution in populations; and test the possibility of maternal effect. Blossom blight susceptibility was tested in reciprocal hybridizations seedlings, as well as their parents. The detached flower technique was used in a randomized complete block design, considering each genotype as a treatment. Flower inoculation was made by spraying a M. fructicola suspension and evaluations were carried out after 72 and 120 hours using a scale of five severity levels. The studied populations presented low phenotypic variability regarding the flower resistance/susceptibility to M. fructicola, being most of them susceptible or very susceptible. Among the tested genotypes, the cultivars Maciel and Cerrito showed less blossom blight susceptibility, transmitting this character to their progenies. Heritability estimates of the blossom blight resistance were medium to low. The low heritability and its distribution in the progenies suggest that the character has additive inheritance, without detecting deviations associated with maternal effects.

Keywords: Prunus persica (L.) Batsch; Monilinia fructicola (Winter) Honey; genetic resistance.

INTRODUCTION

The fungus Monilinia fructicola (Winter) Honey, is the most important pathogen of the peach culture in Brazil, as well as in other producing regions of the world due to the large losses it may cause, when its control is incorrectly done (Adaskaveg et al., 2008; Agrios, 1998; Fortes & Martins, 1998; May-de Mio et al., 2008, 2014; Ogawa et al., 1995). This pathogen may attack the peach during the whole cycle, but especially during flowering and fruit ripening, which are the most susceptible phases (Bleicher, 1997; May-de Mio et al., 2014). The disease cycle begins during flowering, causing blossom blight. Normally necrotic flowers remain attached to the branch, which may be infected by the fungus, resulting in cankers and twig blight (May-de Mio et al., 2008, 2014; Mondino et al., 2010). The blossom blight is the primary infection of the disease and has great epidemiological importance, since it is an inoculum source for secondary infections in the fruits, directly by the production of conidia or in form of latent infections on the fruit in formation, or developing only during the ripening stage of the fruit (Garcia-Benitez et al., 2016, 2017; May-de Mio et al., 2014; Mondino et al., 2010; Thomidis, 2017). The first symptoms observed in fruits (brown rot) are small and circular brown spots. At the maturation stage, infected fruits develop a firm rot, brown lesion that advances rapidly, covering the entire fruit. On the lesion, the fungus sporulation appears powdery and grayish color (May-de Mio et al., 2014; Mondino et al., 2010).

Under mild temperature, humid and rainy weather conditions, this disease can cause total crop loss and, in an attempt to reduce these losses, growers may weekly
apply fungicides. Nowadays, with the increasing concern about the environment and the health of growers and consumers (Baró-Montel et al., 2019; Elshafie et al., 2015), as well as the occurrence of fungus strains resistant to the main used fungicide molecules (Luo et al., 2010; Hily et al., 2011; Zhu et al., 2012; Chen et al., 2017; Fu et al., 2017) the importance of other control strategies is emphasized. Among them, genetic resistance is the most efficient way to control the disease, reducing production costs and environmental impact.

The selection of resistant genotypes is still very limited due to the lack of knowledge of high resistance or immunity sources (Raseira & Franzon, 2014). The resistance to *M. fructicola* is a quantitative and polygenic trait in peach, considered as a character of difficult transmission from the parents to the progenies and highly influenced by the environment (Wagner Júnior et al., 2005; Raseira & Franzon, 2014). However, there are significant differences in susceptibility among the available genotypes (Adaskaveg et al., 2008; Santos & Ueno, 2014).

There are evidences that there is no correlation between flower and fruit resistance (Fabiane, 2011; Santos et al., 2012; Wagner Júnior et al., 2005). In cv. Bolinha for example, that has been widely studied as a standard of brown rot resistance, there was low level of resistance in flowers, unlike to the reaction in fruits (Santos et al., 2012). Therefore, the selection of resistant genotypes must be done independently, for the blossom blight and brown rot in fruits (Raseira & Franzon, 2014; Wagner Júnior et al., 2005), however there are few studies focused on resistance in flowers. Thus, the aims of this work were: to identify genotypes with higher levels of resistance in flowers; to estimate the heritability; to study their distribution in populations; and to test the possibility of maternal effect.

**MATERIAL AND METHODS**

**Plant material**

The study was performed at Embrapa Clima Temperado, in Pelotas, RS, Brazil (31°40’S, 52°26’W, 57 masl), in the years 2015 and 2016. The susceptibility to blossom blight in peaches was tested in reciprocal hybridizations seedlings (F₁ progenies), as well as on their parents. The reciprocal F₁ progenies used were: 2008.159 (Conserva 1526 × ‘Cerrito’) and 2009.38 (‘Cerrito’ × Conserva 1526); 2012.26 (Cascata 1055 × ‘Chimarrita’) and 2012.43 (‘Chimarrita’ × Cascata 1055); 2012.49 (Conserva 672 × Conserva 1526) and 2012.61 (Conserva 1526 × ‘Cerrito’); 2012.52 (Conserva 947 × Conserva 1600) and 2012.66 (Conserva 1600 × Conserva 947); 2012.68 (Conserva 1662 × ‘Maciel’) and 2012.88 (‘Maciel’ × Conserva 1662). The progenies had a minimum of seven and a maximum of 25 evaluated seedlings.

It should be noted that the phytosanitary management of peach seedlings, in the Embrapa Clima Temperado breeding program, is restricted to fungicide applications in the winter period, except in years of extreme weather conditions (such as excess rain). During spring and summer, only insecticides are applied to pest management, aiming to select less susceptible seedlings to the main pathogens, as *M. fructicola*.

**Experimental design and treatments**

For testing the reaction to blossom blight, the technique of detached flowers cited by Fabiane (2011) was used as the most efficient technique to this purpose. The experiment was arranged in a randomized complete block design, considering each genotype as one treatment (seedlings and parents). For the seedling 12 flowers were inoculated, divided into three replicates of four flowers. In the case of the parents, there were three plants obtained by budding (clones) in which three replicates of four flowers were evaluated per clone. Four more flowers per genotype (or clone) without inoculation (control), were also observed to estimate the proportion of latent inoculum coming from the field. The selections Conserva 655 and Cascata 1055 were used as standard of high and low susceptibility to blossom blight, respectively (Fabiane, 2011). The cultivar Bolinha, standard of resistance to *M. fructicola* in fruits, was also included.

**Pathogen culture, conidia productMion, and inoculation**

The fungus isolate was obtained from mummified fruits, infected by *M. fructicola*, collected at four different sites of Embrapa Clima Temperado peach orchards (Pelotas, RS, Brazil). From these, fragments of approximately 5 mm were collected and transferred to Petri dishes containing Potato Dextrose Agar (PDA) culture medium and incubated in a growth chamber at 25 ± 2°C for seven to ten days, with 12 hours light. Contamination with other fungi or bacteria was eliminated by successive passages until the pure culture was obtained. The obtained fungus isolate was stored in test tubes with PDA culture medium in a cold chamber (4 ± 1°C). Whenever necessary, the fungus was cultured on ripe peach fruits and then, re-inoculate in Petri dishes with PDA.

The conidia were removed from the cultures of *M. fructicola* with seven to ten days of incubation, with a brush and 10 mL of distilled water. The suspension was then filtered and the concentration of conidia was determined using an optical microscope and a Neubauer chamber. The concentration was adjusted to 1 x 10⁵ conidia mL⁻¹ (Fabiane, 2011; Santos et al., 2012; Wagner Júnior et al., 2005).

Rev. Ceres, Viçosa, v. 68, n.6, p. 555-563, nov/dec, 2021
Productive branches containing flower buds at half inch green and pink stages, stages 3 and 4 according to Chapman & Catlin (1976), were collected from the individual plants to be tested. The branches were prepared by removing opened or damaged flowers and were kept in buckets with water, inside a cold room during 48 hours at 4 ± 1°C, in order to standardize flowering (Santos et al., 2012), and also to avoid or reduce the contamination with pathogens (Luo et al., 2010; May-de Mio et al., 2008). After 48 hours in the cold chamber, the branches were left for another 24 hours at room temperature for the opening of flowers (anthesis). Finished this period, 16 opened flowers, without disease symptoms, were selected from each branch group. Plastic boxes (50 × 35 × 10 cm) with phenolic foam with cells (2.5 × 2.5 × 3.8 cm) (Green-up®) previously washed in running water for 30 minutes were used. One flower with a small portion of the twig was fixed in each cell.

The inoculation was done by spraying, using a fine droplet spray, with approximately 0.8 mL of the M. fructicola conidial suspension, per box (Fabiane, 2011; Santos et al., 2012) containing between 140 to 200 flowers. Adjustment of the spray volume was done using water-sensitive cards, aiming the correct coverage of flowers.

**Blossom blight evaluation**

After inoculation, the boxes with the flowers, were covered with a plastic bag and placed in a growth chamber (Fitotron), with 23°C±1°C temperature, 75% humidity and 12h light. After 72 and 120 hours, the incidence and severity of the blossom blight were evaluated. Those flowers with petals with necrotic spots were considered infected (Fabiane, 2011; Santos et al., 2012; Wagner Júnior, 2003). Severity was assessed on a grading scale from 0 to 4 (Table 1 and Figure 1).

The grading scale used (Table 1) was elaborated aiming at its ease use during evaluation. The scale limits that determine the scores correspond to the percentage of the flower area with the presence of necrotic spots. The limits described in the grading scale were transformed into a scale of figures (Figure 1), with the ImageJ program and photographs of flowers evaluated 72 and 120 hours after inoculation (hai) with M. fructicola, under the same conditions of the experiment. Therefore, the standard of this scale was specific for artificially inoculated flowers and under the conditions of this experiment.

**Statistical and genetic analysis**

To evaluate the segregation of the progenies and to test for the possibility of maternal effect influence, relative frequency histograms of the severity data were constructed. The maternal effect comparing the progeny of one of the crosses with the reciprocal progeny was also tested using the Mann-Whitney test at 5% significance (Londero et al., 2009; Wu et al., 2012).

The variance observed between the three clones of each parent gave the environmental effect estimation, and the average of parents’ variances was used as the estimated environmental variance (§2). The observed variance among plants of the same progeny was used as the total estimated phenotypic variance (§2). The estimated genetic variance (§2) was calculated by subtracting the environmental variance from the phenotypic variance of each progeny. Broad-sense heritability (H) for the character of resistance to blossom blight was estimated dividing the genetic variance of each population by the total variance of the same population. The calculation was based on the data obtained in the two evaluation seasons, so was divided by two (number of environments) (Dirlewanger et al., 2012; Griffiths et al., 2015):

**RESULTS AND DISCUSSION**

A high variability in the incidence of blossom blight in non-inoculated flowers (field infestation) was detected. This fact was indicated by the high phenotypic variance observed, mainly, in the evaluation after 72 hours, among the evaluated individuals, which was 1015.07 and 610.53 for progenies, and 1144.94 and 729.66 for parents, in the years of 2015 and 2016, respectively (Table 2). This shows the presence of inoculum in the orchard during blossom.

The variability among the studied genotypes can be explained by the different levels of susceptibility to the disease associated with the genotype, by differences in blossom time (temporal) and geographic location within the Embrapa orchard (spatial) (Santos et al., 2012). The high incidence of blossom blight, even without a previous inoculation has already been reported (Keske et al., 2010; Santos et al., 2012) and is due to the high pressure of natural inoculum present in peach orchards in southern Brazil (Fortes & Martins, 1998; May-de Mio et al., 2008; 2014) and favorable climatic conditions for the disease at flowering season (Figure 2).

It was possible to evaluate the reaction to M. fructicola in flowers of 129 and 148 seedlings in 2015 and 2016, respectively. Nine parents of those seedlings progenies plus the genotypes Conserva 655 and ‘Bolina’ were evaluated in both years. The lack of data for all individuals available is due to the fact that many of them did not bloom (either by age, plant size, or adverse climatic factors) or the flowers were not in adequate condition during the experiment period. The vegetative cycle of 2015 and 2016 was characterized by the occurrence of higher temperatures during the flowering period (July to August), and high rainfall (Figure 2, conditions that favored the incidence of M. fructicola on flowers. There
were cases in which the samples had to be discarded due to the high level of infection in flower buds coming from the orchard.

The score range for the disease incidence was the maximum possible (0 to 100% incidence) in flowers without inoculation, in the evaluations performed in the 72 h and in the two years of evaluation, for both, progenies and parents. For the inoculated flowers the score, after 72 hours, also ranged from 0 to 100% in the year 2015, however in 2016, it was between 50 and 100% and 44.44 to100%, for the progenies and parents respectively (Table 2).

The phenotypic variability regarding resistance/susceptibility to *M. fructicola* in inoculated flowers was low, with most genotypes being classified as susceptible or very susceptible to the disease. These genotypes presented overall averages of incidence between 86.94 and 96.94% (72 hai), and between 91.71 and 99.28% (120 hai). The overall averages of severity degree were between 1.36 and 2.24 (72 hai), and between 2.54 and 3.48 (120 hai) (Tabela 2).

Analyzing the percentage of lesions incidence in the flowers of the selections and cultivars tested without inoculation, the genotypes with the lowest incidence were ‘Maciel’, Conserva 672, Conserva 1526, ‘Cerrito’ and Cascata 1055 with less than 65% when evaluated at 72 hours, and less than 80% when evaluated at 120 hours (Figure 3).

When inoculated, the genotypes that presented the lowest susceptibility were ‘Maciel’ and ‘Cerrito’, with 55.88 e 77.78% average incidence, respectively, in the evaluation performed at 72 hours, and less than 100% when evaluated at 120 hours (Figure 3).

In the evaluation performed at 120 hai, only the cultivars Maciel and Cerrito had an incidence lower than 90%, while most of the genotypes showed around 100% incidence. Thus the evaluation at 120 hai, under the tested conditions, did not differentiate the genotypes (Figure 3).

‘Cerrito’, Cascata 1055 and ‘Maciel’ presented the highest percentage of flowers (considering the two years of evaluation) within categories 0 and 1 (severity scale), with 64.89, 80.95 and 93.55%, respectively, in the evaluation performed 72 hai (Figure 4). When evaluated at 120 hai, only the cultivars ‘Cerrito’ and ‘Maciel’ presented a considerable percentage within these categories, 28.17 and 44.44%, respectively.

The progenies 2008.159, 2009.38, 2012.68 and 2012.88 presented the lowest incidence (<85%) and severity (Figure 4). This indicates that the less susceptible ‘Cerrito’ and ‘Maciel’, used as parents transmitted this trait to their

| Score | Description                                                                 |
|-------|------------------------------------------------------------------------------|
| 0     | Without infection                                                            |
| 1     | Necrotic spots on the petals covering > 1% d” 20% of the surface              |
| 2     | Necrotic spots on the petals > 20% ≤ 40%                                     |
| 3     | Necrotic spots on the petals > 40% ≤ 60%                                     |
| 4     | Necrotic spots on the petals > 60%                                          |

Figure 1: Severity scale used to evaluate the blossom blight in peach flowers artificially inoculated by spraying.
Blossom blight resistance in peach: heritability and segregation in progenies from reciprocal crosses

The severity scale of blossom blight was not fully efficient to quantify susceptibility to *M. fructicola*. This was evidenced by the lack of normal distribution in the histograms of relative frequencies (Figure 4). When evaluated 72 hai most of the genotypes remained in category 1 (necrotic spots covering between 1% and 20% of the petals) (Table 1). Similarly, when evaluations were performed 120 hai, most genotypes were located in the category 4 (necrotic spots covering more than 60% of the petals) (Table 1). This suggests that there should be more categories between these percentages to segregate more levels of severity to the disease. Although this has not great practical importance, it is interesting in the epidemiology of the disease since genotypes that develop infections and sporulations more rapidly, increase the rate of disease dissemination (May-de Mio *et al.*, 2014; Mondino *et al.*, 2010; Rios & Debona, 2018).

Observing the histograms, different behavior was not evident when a genitor was used as female or male parent, leading to the conclusion that there is no maternal effect. (Dini *et al.*, 2019). The hypothesis of maternal effect in the five reciprocal crosses studied was tested through the

Table 2: Descriptive statistics of the variables incidence and severity of blossom blight in 10 reciprocal peach progenies and their parents, evaluated after 72 and 120 hours, with and without artificial inoculation, in the years 2015 and 2016, Embrapa Clima Temperado, Pelotas, Rio Grande do Sul, Brazil

|        | 72 hours | 120 hours |        | 72 hours | 120 hours |        | 72 hours | 120 hours |
|--------|----------|-----------|--------|----------|-----------|--------|----------|-----------|
|        | Inc. (%) | Sev. (0 to 4) | Inc. (%) | Sev. (0 to 4) | Inc. (%) | Sev. (0 to 4) | Inc. (%) | Sev. (0 to 4) |
|        |          |           |        |          |           |        |          |           |
| Means  | 69.93    | 0.81      | 84.15  | 1.27     | 93.08     | 2.24   | 97.14    | 3.48      |
| Median | 75.00    | 0.75      | 100    | 1.00     | 100       | 2.46   | 100      | 4.00      |
| PV     | 1015.07  | 0.25      | 565.88 | 0.55     | 321.35    | 1.08   | 88.69    | 0.69      |
| SD     | 31.86    | 0.50      | 23.79  | 0.74     | 17.93     | 1.04   | 9.42     | 0.83      |
| CV (%) | 45.56    | 62.19     | 28.27  | 58.72    | 19.26     | 46.51  | 9.69     | 23.90     |
| Minimum| 0.00     | 0.00      | 0.00   | 0.00     | 0.00      | 0.00   | 41.67    | 0.75      |
| Maximum| 100      | 2.75      | 100    | 3.75     | 100       | 3.91   | 100      | 4.00      |

2015 Parents

|        | 72 hours | 120 hours |        | 72 hours | 120 hours |        | 72 hours | 120 hours |
|--------|----------|-----------|--------|----------|-----------|--------|----------|-----------|
|        | Inc. (%) | Sev. (0 to 4) | Inc. (%) | Sev. (0 to 4) | Inc. (%) | Sev. (0 to 4) | Inc. (%) | Sev. (0 to 4) |
| Means  | 57.41    | 0.66      | 72.22  | 1.15     | 86.94     | 1.95   | 91.71    | 3.27      |
| Median | 50.00    | 0.50      | 75.00  | 1.00     | 100       | 2.08   | 100      | 3.75      |
| PV     | 1144.94  | 0.24      | 1169.87| 0.73     | 798.29    | 1.04   | 380.88   | 1.29      |
| SD     | 33.84    | 0.49      | 34.20  | 0.86     | 28.25     | 1.02   | 19.52    | 1.14      |
| CV (%) | 58.94    | 74.69     | 47.36  | 74.52    | 32.50     | 52.33  | 21.28    | 34.77     |
| Minimum| 0.00     | 0.00      | 0.00   | 0.00     | 0.00      | 0.00   | 30.00    | 0.58      |
| Maximum| 100      | 2.00      | 100    | 3.25     | 100       | 3.83   | 100      | 4.00      |

2016 Parents

|        | 72 hours | 120 hours |        | 72 hours | 120 hours |        | 72 hours | 120 hours |
|--------|----------|-----------|--------|----------|-----------|--------|----------|-----------|
|        | Inc. (%) | Sev. (0 to 4) | Inc. (%) | Sev. (0 to 4) | Inc. (%) | Sev. (0 to 4) | Inc. (%) | Sev. (0 to 4) |
| Means  | 85.81    | 0.97      | 96.29  | 1.64     | 96.94     | 1.66   | 99.24    | 3.12      |
| Median | 100      | 1.00      | 100    | 1.50     | 100       | 1.50   | 100      | 3.25      |
| PV     | 610.63   | 0.16      | 138.68 | 0.49     | 55.13     | 0.48   | 13.70    | 0.67      |
| SD     | 24.71    | 0.40      | 11.78  | 0.70     | 7.42      | 0.69   | 3.70     | 0.82      |
| CV (%) | 28.80    | 40.57     | 12.23  | 42.57    | 7.66      | 41.56  | 3.73     | 26.16     |
| Minimum| 0.00     | 0.00      | 25.00  | 0.25     | 50.00     | 0.63   | 75.00    | 0.88      |
| Maximum| 100      | 2.50      | 100    | 3.50     | 100       | 4.00   | 100      | 4.00      |

Results of 129 e 148 different genotypes belonging to F₁ progenies, in the 2015 and 2016 seasons, respectively; 9 parental genotypes were evaluated (advanced selections and cultivars); Scale of 0 to 4 as being 0 petals without lesions and 4 petals with 60% or more of the area with lesions. PV = phenotypic variance; SD = standard deviation; CV = coefficient of variation.
Mann-Whitney test. The tested contrast, in all cases, was F1 progeny versus its reciprocal F1 progeny for the studied parameters (incidence and severity). The test was not significant ($p > 0.05$), for any of the progenies, that is, there were no significant differences between the reciprocal progenies, indicating that there is no maternal effect on the transmission of this trait (Dini et al. 2019; Londero et al.; 2009).

The estimated $H^2$ values were medium to low, varying between 11 and 43%, depending on the studied population, with an average of 24% for incidence and 17% for severity.

In the study of Wagner Júnior (2003) in which the $H^2$ for blossom blight was estimated through the incidence, the values ranged between 6 and 66%, depending on the studied population, with averages between 30 and 42%. This same author also concluded that the $H^2$ for this trait is low and highly variable among populations.

Further studies aiming at adjusting a phenotyping protocol for this character become necessary to be able to correctly differentiate genotypes with different levels of genetic resistance. This should include mainly, studies related to the concentration of conidia to be used, phenological state of the flower, as well as the elaboration of a scale of severity more efficient to differentiate the genotypes.

On the other hand, the high latent infection in non-inoculated flowers, leads to conclude that a good treatment and sanitary practices should be used in commercial orchards, whereas a source of high resistance level to blossom blight is not available. Moreover, for rapid screening of seedlings, the field conditions are sufficient to cause a high level of infection and select the least susceptible to be further evaluated with artificial inoculation, reducing significantly the number of individuals to be tested.

**Figure 2:** Average temperatures and daily rains in July and August of 2015 and 2016, Embrapa Clima Temperado, Pelotas, Rio Grande do Sul, Brazil. Source: AGROMET/CPACT/EMBRAPA (2018).

**Figure 3:** Blossom blight incidence in peach genotypes submitted or not to M. fructicola inoculation, evaluated after 72 and 120 hours. The columns correspond to the average of two harvest seasons (2015 and 2016) and the vertical bars in each column refer to the standard error. Embrapa Clima Temperado, Pelotas, Rio Grande do Sul, Brazil.
CONCLUSIONS

The studied populations presented low phenotypic variability regarding to resistance/susceptibility to *Monilinia fructicola*, with most genotypes being susceptible or very susceptible. The cultivars Maciel and Cerrito are less susceptible to blossom blight, transmitting this character to their progenies. The heritability of the resistance to blossom blight in peach is medium to low.

Figure 4: Severity of blossom blight in F1 progenies and their parents inoculated with *M. fructicola* and evaluated 72 and 120 hours after inoculation, years 2015 and 2016. The severity scale used (0 to 4) is detailed in Table 1 and in Figure 1. Embrapa Clima Temperado, Pelotas, Rio Grande do Sul, Brazil.
No maternal effect on the transmission of susceptibility to blossom blight was detected.

Future studies should aim at adjust a phenotype protocol for this character, mainly related with the conidia concentration to be inoculated, as well as the elaboration of a more detailed scale of disease severity, in order to accurately differentiate genotypes with different levels of susceptibility.

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Rev. Ceres, Viçosa, v. 68, n.6, p. 555-563, nov/dec, 2021
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