A new three-locus model for rootstock-induced dwarfing in apple revealed by genetic mapping of root bark percentage

Article

Published Version

Creative Commons: Attribution 3.0 (CC-BY)

Open access


It is advisable to refer to the publisher's version if you intend to cite from the work. See Guidance on citing.

To link to this article DOI: http://dx.doi.org/10.1093/jxb/erw001

Publisher: Oxford University Press

All outputs in CentAUR are protected by Intellectual Property Rights law, including copyright law. Copyright and IPR is retained by the creators or other copyright holders. Terms and conditions for use of this material are defined in the End User Agreement.

www.reading.ac.uk/centaur
RESEARCH PAPER

A new three-locus model for rootstock-induced dwarfing in apple revealed by genetic mapping of root bark percentage

Nicola Harrison1,2*, Richard J. Harrison1,2, Nuria Barber-Perez1, Emma Cascant-Lopez1, Magdalena Cobo-Medina1, Marzena Lipska1, Rebeca Conde-Ruíz1, Philip Brain1, Peter J. Gregory1,2 and Felicidad Fernández-Fernández1

1 East Malling Research, New Road, East Malling, Kent ME19 6BJ, UK
2 Centre for Food Security, School of Agriculture, Policy and Development, University of Reading, Whiteknights, PO Box 237, Reading RG6 6AR, UK

* Correspondence: nicola.harrison@emr.ac.uk

Received 17 July 2015; Accepted 9 December 2015

Abstract

Rootstock-induced dwarfing of apple scions revolutionized global apple production during the twentieth century, leading to the development of modern intensive orchards. A high root bark percentage (the percentage of the whole root area constituted by root cortex) has previously been associated with rootstock-induced dwarfing in apple. In this study, the root bark percentage was measured in a full-sib family of ungrafted apple rootstocks and found to be under the control of three loci. Two quantitative trait loci (QTLs) for root bark percentage were found to co-localize to the same genomic regions on chromosome 5 and chromosome 11 previously identified as controlling dwarfing, Dw1 and Dw2, respectively. A third QTL was identified on chromosome 13 in a region that has not been previously associated with dwarfing. The development of closely linked sequence-tagged site markers improved the resolution of allelic classes, thereby allowing the detection of dominance and epistatic interactions between loci, with high root bark percentage only occurring in specific allelic combinations. In addition, we report a significant negative correlation between root bark percentage and stem diameter (an indicator of tree vigour), measured on a clonally propagated grafted subset of the mapping population. The demonstrated link between root bark percentage and rootstock-induced dwarfing of the scion leads us to propose a three-locus model that is able to explain levels of dwarfing from the dwarf ‘M.27’ to the semi-invigorating rootstock ‘M.116’. Moreover, we suggest that the QTL on chromosome 13 (Rb3) might be analogous to a third dwarfing QTL, Dw3, which has not previously been identified.

Key words: Apple, grafting, Malus × domestica, root anatomy, root bark percentage, root cortex, rootstock, rootstock-induced dwarfing, scion.

Introduction

The technique of grafting has been practised for hundreds of years in order to maintain, crop and propagate many different fruits (Feree and Carlson, 1987). Grafted trees are typically composed of two parts, the rootstock and the scion. The rootstock contributes the root system and the base of the stem while the scion is grafted on top of the rootstock and is
the part of the tree that produces the fruit. The point of grafting between the two plant parts is known as the graft union (Mudge et al., 2009). All temperate tree fruit crops grown commercially throughout the developed world are grafted onto rootstocks selected for specific traits such as growth control and early fruit production. Growth-controlling rootstocks have enabled the transition of low-yielding traditional orchards to high-density modern fruiting orchards, mainly through the vigour control they impart to the scion to produce compact trees (Mudge et al., 2009). This vigour control is gradual and cumulative, and interactions between rootstock, environment and scion can introduce variation into the growth of the composite tree. Rootstocks can also influence precocity and yield efficiency in addition to controlling some diseases (Rom and Carlson, 1987). Between 1913 and 1915, researchers at the Wye College Fruit Experiment Station at East Malling collected a range of clonally propagated apple rootstocks used by nurseries in the UK and continental Europe and embarked on a process of detailed phenotypic characterization. This led to the description of the first nine ‘Malling’ series of apple rootstocks (‘M.1’ to ‘M.9’), which included both dwarfing (‘M.8’ and ‘M.9’) and non-dwarfing types (‘M.1’ and ‘M.2’; Hatton, 1917; Hatton, 1918). Breeding programmes in Europe, North America and New Zealand have since used the ‘Malling’ series to develop novel apple rootstocks with improved pest and disease resistance that are better adapted to their local conditions (Cummins and Aldwinckle, 1974; Webster et al., 2000). Most commercially available dwarfing apple rootstocks can be traced to just one or two original sources, namely ‘M.9’ and ‘M.8’ (Russo et al., 2007), which have been found to be closely related to Malus × domestica using molecular analysis (Oraguzie et al., 2005; Fazio et al., 2011). A practical problem when using invigorating genotypes as a source of pest or disease resistance in breeding programmes is that the dwarfing effect of one parent is frequently lost in the first generation. As a result of this, dwarfing genotypes only reappear when F1 genotypes (derived from a dwarfing and a non-dwarfing parent) are crossed again to another dwarfing genotype. This is a serious limitation for breeding programmes because of the long generation cycles required to confidently evaluate rootstock performance and its effect on the scion. Robust molecular markers strongly linked to rootstock-induced dwarfing are a much-needed tool to hasten rootstock breeding.

Assessment of vigour control by a rootstock is not easy and earlier work to determine the genetic basis of vigour control in apple rootstocks has focused on above-ground traits such as trunk cross-sectional area, or visual assessment of vigour. This determines the ability of the rootstock to control in apple rootstocks has focused on above-ground traits such as growth control and early fruit production. Growth-controlling rootstocks have enabled the transition of low-yielding traditional orchards to high-density modern fruiting orchards, mainly through the vigour control they impart to the scion to produce compact trees (Mudge et al., 2009). This vigour control is gradual and cumulative, and interactions between rootstock, environment and scion can introduce variation into the growth of the composite tree. Rootstocks can also influence precocity and yield efficiency in addition to controlling some diseases (Rom and Carlson, 1987). Between 1913 and 1915, researchers at the Wye College Fruit Experiment Station at East Malling collected a range of clonally propagated apple rootstocks used by nurseries in the UK and continental Europe and embarked on a process of detailed phenotypic characterization. This led to the description of the first nine ‘Malling’ series of apple rootstocks (‘M.1’ to ‘M.9’), which included both dwarfing (‘M.8’ and ‘M.9’) and non-dwarfing types (‘M.1’ and ‘M.2’; Hatton, 1917; Hatton, 1918). Breeding programmes in Europe, North America and New Zealand have since used the ‘Malling’ series to develop novel apple rootstocks with improved pest and disease resistance that are better adapted to their local conditions (Cummins and Aldwinckle, 1974; Webster et al., 2000). Most commercially available dwarfing apple rootstocks can be traced to just one or two original sources, namely ‘M.9’ and ‘M.8’ (Russo et al., 2007), which have been found to be closely related to Malus × domestica using molecular analysis (Oraguzie et al., 2005; Fazio et al., 2011). A practical problem when using invigorating genotypes as a source of pest or disease resistance in breeding programmes is that the dwarfing effect of one parent is frequently lost in the first generation. As a result of this, dwarfing genotypes only reappear when F1 genotypes (derived from a dwarfing and a non-dwarfing parent) are crossed again to another dwarfing genotype. This is a serious limitation for breeding programmes because of the long generation cycles required to confidently evaluate rootstock performance and its effect on the scion. Robust molecular markers strongly linked to rootstock-induced dwarfing are a much-needed tool to hasten rootstock breeding.

Assessment of vigour control by a rootstock is not easy and earlier work to determine the genetic basis of vigour control in apple rootstocks has focused on above-ground traits such as trunk cross-sectional area, or visual assessment of growth habit and internode length (Seleznyova et al., 2003; Rusholme-Pilcher et al., 2008).

Previous studies by Rusholme-Pilcher et al. (2008) and Fazio et al. (2014) identified two quantitative trait loci (QTLs) associated with rootstock-induced dwarfing: Dwarfing 1 (Dw1), at the top of linkage group/chromosome 5 (LG05) between markers CH03a09 and NZraAM18-700, and Dwarfing 2 (Dw2), on linkage group/chromosome 11 (LG11) between markers CH02d08 and C13243 (Rusholme-Pilcher et al., 2008; Fazio et al., 2014). The second QTL, Dw2, was initially identified assuming an additive model with Dw1, but was proposed to have non-additive effects after further analysis. The genetic model put forward by Fazio et al. (2014) was more predictive where either of the dwarfing loci was homozygous, and gave less accurate predictions of phenotype where either or both loci were heterozygous (Fazio et al., 2012). A recent study by Foster et al. (2015) identified Dw2 as an additive QTL, and they suggested that ‘Dw1’ has a stronger effect on rootstock-induced dwarfing than Dw2, and that Dw2 may act as an enhancer of ‘Dw1’ (Foster et al., 2015). Foster et al. found similar results to Fazio et al., with their model only able to account for 68.6% of the observed variation. Both studies reported cases where Dw1 was present yet the expected phenotype did not behave as predicted.

While many previous studies on dwarfing apple rootstocks have concentrated upon measuring a secondary conferred trait (i.e. the manifestation of vigour control in the scion), this study focuses on a primary rootstock trait known as root bark percentage. A high proportion of root cortical cells (hereafter termed root bark percentage) in the root of an apple rootstock has previously been associated with the ability of the rootstock to reduce the vigour of a grafted scion (Beakbane and Thompson, 1947). The aim of the present study was to identify QTLs for root bark percentage and to confirm whether there was an association with rootstock-induced dwarfing in the M432 mapping population. We used the M432 apple rootstock mapping population together with its associated genetic map (Fernández-Fernández et al., 2012; Antanavičiute et al., 2012) to identify QTLs associated with root bark percentage. We also present a model of rootstock-induced dwarfing, describing the nature of allelic variation and how this determines the ability of the rootstock to control vigour.

Materials and methods

Plant material

M432 field population

The M432 apple rootstock mapping progeny used in this study has been previously reported (Fernández-Fernández et al., 2012). It is a subset of a larger seedling population raised in 2004 from a cross between the dwarfing rootstock ‘M.27’ (‘M.13’ × ‘M.9’) and a semi-vigorous rootstock ‘M.116’ (‘M.106’ × ‘M.27’). The M432 family originally comprised 257 individuals of which 120 (population A) were maintained as low hedges (ungrafted and cut hard to encourage shoot production) for 10 growing seasons. The remaining individuals were budded with a columnar scion (East Malling Research (EMR) selection number SA544-28) in 2005 (population B). From the latter population, 68 (population B1) were cut back below the graft union in December 2007 and treated as population A until they were all grubbed (removal of trees from a field planting) in December 2014. The rest of the seedlings (population B2) were evaluated as part of the breeding programme for their effect on the scion and lifted in 2015 for propagation. The mapping progeny comprised a total of 140 seedlings (89 and 51 from populations A and B1, respectively). DNA from which was used to develop a combined simple sequence repeat (SSR) linkage map (Fernández-Fernández et al., 2012) and single nucleotide polymorphism (SNP)-based maps of the parental genomes (Fernández-Fernández et al., 2012; Antanavičiute et al., 2012; Troggio et al., 2013).
Reference rootstock cultivars in the field
Three replicates each of four commercially available rootstock cultivars (‘M.9’, ‘M.26’ ‘M.27’ and ‘M.106’) were interspaced in the planting of the M432 progeny and were treated like population B1.

Grafted subset of M432 mapping population
In December 2010, four or more replicates of each of the seedlings in the M432 mapping population (populations A and B1) were propagated through hardwood cuttings. Successfully rooted cuttings were grafted with ‘Royal Gala’ in January 2012 and grown in pots outdoors for three growing seasons. Replicates of the parental genotypes (‘M.27’ and ‘M.116’) were also grafted and grown in pots alongside the seedlings. In January 2015, trees from 37 seedlings (for which five or more replicates grafted with ‘Royal Gala’ were available) as well as the parents were measured for rootstock-induced dwarfing traits (e.g. stem diameter).

Determination of root bark percentage
M432 field population
In 2014, the 122 seedlings of the M432 mapping population that survived the 10 growing seasons were lifted from the field. Twelve root segments (2–8 mm in diameter, 100–120 mm in length) were excised from each root system using secatores, placed into a labelled polythene bag with moist tissue to prevent desiccation of the roots and stored at 4 °C before analysis. The roots were then carefully washed using a nailbrush to remove all the soil. For each root segment, a scalpel or a utility knife was used to remove a ring of bark (cortex) approximately 2–3 mm in length, leaving behind the stele of the root. Digital calipers were used to make pairs of measurements of the root with and without the bark. The cross sectional area of the root was calculated for each sample, assuming that the root section was a perfect cylinder. The percentage of root bark at a standard root diameter of 7.5 mm was then inferred using regression analysis. For the grafted subset of the M432 mapping population (which assumes an additive model of genetic effects) was not used (Lark et al., 1995). The K-W QTL analysis was carried out in the MapQTL5 software package (Kyazma, Wageningen, The Netherlands), using the map published by Antanaviciute et al. (2012).

Marker development
The highest scoring SNP marker for each QTL was used to select the chromosomal regions for marker development. Microsatellite markers were manually designed, using the draft rootstock whole genome alignment (N. Harrison, unpublished data), in regions where significant SNPs from the QTL mapping were located. Using the rootstock whole genome alignment for chromosomes 5, 11 and 13, three primer pairs were designed around simple sequence repeats and indel features, and screened on the M432 mapping population for segregation. The primer sequences are given as follows with primer name followed by the 5’–3’ DNA sequence: EM_Rb1_F ggtgtaaggagcttgatag; EM_Rb1_R acatatatcatctccgact; EM_Rb2_F pagaataggagcccattgag; EM_Rb2_R gcagacctggcccagac; EM_Rb3_F gaggaatataataatgagaa; EM_Rb3_R gcagacactgcttggtcaca.

Sequence-tagged site marker analysis
Primer pairs were labelled on the forward primer with 6-FAM fluorescent dye (IDT, Belgium) using an M13-tailed primer in a two-step reaction as described by Schuelle (2000). PCR reactions for sequence-tagged site (STS) markers were performed using the ‘Type-it’ PCR mastermix (Qiagen) following the manufacturer’s recommendations, in a final volume of 12.5 µl. PCR reactions were carried out using the following PCR cycles: an initial denaturation
step of 94 °C for 5 min was followed by 10 cycles of 94 °C for 30 s, an annealing temperature of 55 °C decreasing by 0.5 °C per cycle until 50 °C for 45 s and 72 °C for 60 s, followed by 25 cycles of 94 °C for 30 s, 50 °C for 45 s and 72 °C for 60 s with a final extension step at 72 °C for 15 min. PCR products were fractionated by capillary electrophoresis through a 3130 Genetic Analyzer (Applied Biosystems). Data generated were collected and analysed using the GENESCAN and GENOTYPER (Applied Biosystems) software.

Statistical analysis

Statistical analyses were performed using the statistical software package ‘R’ version 3.1.0 using the lm function (R Core Development Team, 2008) and GenStat for Windows 14th Edition (VSN International). The models presented were developed by sequentially adding fixed effects to the regression analysis, beginning with single additive factors (chr 5, chr 11, chr 13), followed by interaction terms. Final models were selected, based upon significant terms in the model selection.

Results

Root bark percentage

Root bark percentage was examined in the field-grown, ungrafted M432 rootstock population (populations A and B1) and found to vary in a genotype-dependent manner. Field controls included ‘M.27’, ‘M.9’, ‘M.26’ and ‘M. M.106’ (Table 1); unfortunately, liners—young rootstocks, typically 1 or 2 years old—of ‘M.116’ were not available for planting when the field plot was established and it is not, therefore, included in Table 1. (Liners can be produced by hardwood cuttings or micro-propagation or lifted from a stool bed as a single stem rooted trunk; they can be bench grafted in the winter and then planted out in the field or pots or ‘lined’ in a nursery field for summer budding.) Table 1 shows that dwarfing rootstocks had a higher percentage of root bark than semi-invigorating rootstocks. It is noteworthy that ‘M.27’, which has been characterized as more dwarfing than ‘M.9’, has a slightly lower root bark percentage than its semi-invigorating parent ‘M. M.106’ (52.5%, Table 1), it was still significantly different from its dwarfing parent ‘M.27’ (Fig. 1 and Table 1). These preliminary data agree with expectations derived from much earlier studies (Beakbane and Thompson, 1947).

Variation in root bark percentage

The log-transformed distribution of root bark percentages for the 122 individuals of the M432 mapping population had a unimodal distribution pattern with clear segregation (Fig. 2). Log-transformed values were normally distributed and were used for subsequent QTL mapping of loci involved in modulating root bark percentage.

Identification of QTLs

QTL mapping was carried out using K-W testing and revealed three loci involved in the determination of root bark percentage, located on chromosomes 5, 11 and 13 (Table 2 and Fig. 3A), labelled Rb1(a), Rb2(b) and Rb3(c), respectively. Rb1(a) was present only in ‘M.27’, while Rb2(b) and Rb3(c) were detected in both parental genotypes. All QTLs were highly significant (Table 2) with Rb1(a) and Rb2(b) showing clear maxima on their respective linkage groups (Fig. 3B).

Plots of the progeny grouped by single SNP markers (STS markers in the case of QTL Rb2 in order to resolve all parental haplotypes), reveal that, on average, each marker alone appears to have only a moderate effect on root bark percentage depending upon the allelic state (Fig. 4A–C). However, pairwise marker analysis revealed that there were substantial non-additive effects between Rb1(a) and Rb2(b) (Fig. 4D and Supplementary Fig. S1 at JXB online), but not Rb1(a) and Rb3(c) (Fig. 4E). These observations suggest a linear mixed model. A model that included an interaction between Rb1(a) and Rb2(b) was significant, but inclusion of interaction terms between Rb1(a) and Rb3(c), or Rb2(b) and Rb3(c) did not significantly improve this model (Supplementary Table S1). Specifically, the activity of Rb1(a) was dependent on the presence of Rb2(b) and vice versa (Fig. 4D and Supplementary Fig. S2). This is substantiated by the observation that the classes containing Rb1(a)

Table 1. Root bark percentage in common field-grown apple rootstocks, measured after 10 growing seasons

<table>
<thead>
<tr>
<th>Rootstock</th>
<th>Average root bark percentage (at 7.5mm)</th>
<th>SEM</th>
<th>Vigour</th>
</tr>
</thead>
<tbody>
<tr>
<td>M27</td>
<td>72.92</td>
<td>2.00</td>
<td>Strongly dwarving</td>
</tr>
<tr>
<td>M9</td>
<td>76.89</td>
<td>0.76</td>
<td>Dwarfing</td>
</tr>
<tr>
<td>M26</td>
<td>64.05</td>
<td>2.13</td>
<td>Semi-dwarfing</td>
</tr>
<tr>
<td>MM106</td>
<td>52.56</td>
<td>4.31</td>
<td>Semi-invigorating</td>
</tr>
</tbody>
</table>

SEM: standard error of the mean.

Fig. 1. Root bark microscopy images depicting the root bark percentages for the two parents of the M432 mapping population, ‘M.27’ and ‘M.116’.
but not Rb2(b), or Rb2(b) in the absence of Rb1(a) display phenotypes that are, on average, similar to the class lacking both Rb1(a) and Rb2(b) (Fig. 4D). The class of individuals with the highest root bark percentage in this pairwise comparison is the class containing Rb1(a) and two copies of Rb2(b) (one allele from ‘M.27’ and one from ‘M.116’). Alleles inherited from ‘M.27’ and ‘M.116’ in repulsion to the Rb2(b) allele, i.e alleles Rb2(BB′) in Fig. 4, appear to have different effects upon the ‘penetrance’ or expression of the root bark phenotype when present with the Rb2(b) allele (Fig. 4D). The Rb2(B) allele from ‘M.27’ appears to suppress the effects of Rb2(b) (though not to the level of the Rb2-lacking class), while the Rb2(B′) allele from ‘M.116’ increases the root bark percentage to levels comparable to the homozygous Rb2(bb) class, though with greater variance than the homozygous class.

Again, as seen with Rb1(a) when Rb2(b) was present (Fig. 4D), the expression of Rb3(c) appeared to be dependent on the combination of alleles at Rb2, with the Rb2(B′) allele apparently compensating for the homozygous state of Rb3(cc). In this pairwise analysis, it was clear that the allelic status of Rb1 confounded the analysis of Rb2 and Rb3 due to small sample sizes of some genotypic classes. If there were no effect of Rb2 in the absence of Rb3, the genotypic class ‘Rb2(bb) Rb3(CC)’ would predict a low root bark percentage. However, on examination of the allelic status of Rb1 in this class, which comprised only two individuals, it was found that both individuals contained Rb1(a) and therefore had a high root bark percentage due to the interaction between Rb1(a) and Rb2(b).

To test whether the effect of Rb3(c) occurs regardless of the allelic status of Rb1 (Fig. 4E) or Rb2 (Fig. 4F), a three-way analysis of Rb loci was carried out. With Rb2(BB′), expression of Rb3(c) appeared to show only a pronounced difference in the homozygous state (see genotypes CCBB′, CcBB′ and ccbB′ in Fig. 4F). Comparisons of homozygous (CC) and heterozygous (Cc) classes of Rb3, grouped by Rb1 and Rb2 allelic status, revealed no significant differences in a regression analysis (data not shown) and were therefore pooled, to create a total of 16 genotypic classes, reducing the number of missing classes in the data. Regression analysis of these data revealed that there was a significant three-way interaction between Rb1(a), Rb2(b) and Rb3(c) (Supplementary Table S1).

When data were grouped by Rb1, Rb2 and Rb3 (Supplementary Fig. S2) considering chromosome 13 in all three classes, Rb3(CC) homozygotes (Supplementary Fig. S2A), Rb3(Cc) heterozygotes (Supplementary Fig. S2B) and Rb3(cc) homozygotes (Supplementary Fig. S2C), much of the variation in root bark percentage in Rb3(cc) homozygotes in the absence of Rb2(b) was explained by the presence or absence of Rb1(a), with Rb1(a) giving the higher root bark percentage. This indicates that there is an interaction between Rb1(a) and Rb3(c) in the absence of Rb2(b). It is also noteworthy that regardless of the allelic status of Rb1 in the Rb3(cc) homozygous class, mean levels of root bark percentage were almost the same when Rb2(Bb) heterozygotes were considered, indicating that the functional status of Rb2 also affects Rb3(cc) expression (Supplementary Fig. S2).

A full three-locus analysis, taking into account the full allelic status of Rb1, Rb2 and Rb3 was not feasible due to the small sample size obtained once the population was further split into its representative genotypes. For this population there are up to 64 possible genotypic classes and the dataset does not contain even the 24 possible classes described in this paper; for example, the AAbbcc genotypic class is missing from the dataset. However, there were six individuals with Rb1(a) present, and both Rb2(b) and Rb3(c) in the homozygous states, i.e. Aabbcc. The average root bark percentage in this class was 81%, which was the highest of any genotypic class with a sample size of >2. There were three individuals where the homozygous state at all QTL loci was AABB′CC, and the average root bark percentage in this class was 58% (the genotype and root bark percentage

![Histogram of phenotypic distribution of log-transformed root bark percentage data for the M432 mapping population.](image)

Fig. 2. Histogram of phenotypic distribution of log-transformed root bark percentage data for the M432 mapping population.

Table 2. Most significant SNP markers associated with the QTLs Rb1, Rb2 and Rb3

<table>
<thead>
<tr>
<th>Chr</th>
<th>Position (cM)</th>
<th>SNP markers associated with the QTLs Rb1, Rb2 and Rb3</th>
<th>K statistic</th>
<th>Significance</th>
<th>Marker origin</th>
</tr>
</thead>
<tbody>
<tr>
<td>5</td>
<td>20.47</td>
<td>RosBREEDSNP_SNP_AC_2429897_Lg5_00179_MAF40_1681882_exon1</td>
<td>32.1</td>
<td>*</td>
<td>M27</td>
</tr>
<tr>
<td>11</td>
<td>23.637</td>
<td>RosBREEDSNP_SNP_CA_8702100_Lg11_00735_MAF40_1677605_exon4</td>
<td>13.409</td>
<td>*</td>
<td>M27 and M116</td>
</tr>
<tr>
<td>13</td>
<td>5.6</td>
<td>RosBREEDSNP_SNP_GT_2194655_Lg13_00098_MAF30_</td>
<td>12.648</td>
<td>*</td>
<td>M27 and M116</td>
</tr>
</tbody>
</table>

Chr: chromosome.
Significance levels: ****: 0.0005; ******: 0.0001.
of each individual in the M432 population are presented in Supplementary Table S3).

Root bark percentage and rootstock-induced dwarfing

To test for an association between root bark percentage and rootstock-induced dwarfing in the M432 population, a small subset (n=38) of the M432 population that had been clonally propagated and grafted was used. Stem diameter was measured in this subset and found to be significantly negatively correlated (r = –0.54, P < 0.0004) with root bark percentage (Fig. 5). Using a linear mixed model, the effects of Rb3(c) on stem diameter were examined. Model selection was carried out using log-transformed data as previously described, by sequentially adding terms. Presented below are two genetic models: model (a) postulated by Fazio et al. (2014) using their Dw notation, and model (b), the result of our model selection process using Rb notation (Supplementary Table S2):

Model (a): Stem diameter ~ Dw1 + Dw2 + Dw1 × Dw2
Model (b): Stem diameter ~ Rb1 + Rb2 + Rb1 × Rb2 + Rb2 × Rb3

Model (b) was a significantly better fit than model (a), with interactions detected between Rb1(a) and Rb2(b), and Rb2(b) and Rb3(c) (note that it was not possible to fit a three-way interaction model due to multiple missing genotypic classes in this subset of the population). Model (b) explained 84% of the variance observed in the grafted subset of the population (Supplementary Table S2), compared with 53% for model (a). Adding Rb2 × Rb3 to the model significantly increased the percentage of the variance accounted for by 31% (P = 0.001), demonstrating that Rb3(c) has a non-additive effect on stem dwarfing in the presence of Rb2(b). This is consistent with the results obtained for root bark percentage and can therefore be considered to be analogous to a new putative QTL for rootstock-induced dwarfing (Dw3). Although the dataset is limited, it can be seen that the effects of Rb3(c)/Dw3 (Supplementary Fig. S3A–C) are largely consistent with the root bark percentage data, with Rb1(A), Rb2(B) and Rb3(C) plants displaying the largest stem diameter (Supplementary Fig. S3A), and heterozygous Rb3Cc plants displaying a dwarfing phenotype dependent upon Rb1 and Rb2 (Supplementary Fig. S3B), which is enhanced in Rb3Cc homozygotes (Supplementary Fig. S3C). In this case, significant differences were observed between classes heterozygous at Rb3Cc and homozygous at Rb3cc, indicating that Rb3 may have more profound effects upon stem diameter than root bark percentage.

QTL in other common rootstocks

Three STS markers were developed that are closely linked to the root bark QTL and screened on a selection of apple...
A three-locus model for rootstock-induced dwarfing

1877

rootstock material (Table 3). Interestingly, only ‘M.26’, ‘M.27’ and ‘M.9’ contained all Rb QTL, while other rootstocks contained either Rb1(a) and/or Rb3(c) (‘M.1’, ‘M.6’, ‘M.7’ and ‘M.16’, Table 3). The rootstocks ‘Polish 22’ and ‘Mac 9’ are both dwarfing and yet lack both Rb1 and Rb3, only containing markers for Rb2/Dw2. This raises the possibility that there are other alleles or loci interacting with Rb2/Dw2 that are absent from other ‘M.9’-derived material and that either they are unmapped in the M432 cross or recombination has occurred between markers and QTL in these accessions. This information reinforces our finding that Rb2 is a crucial locus for highly dwarfing rootstocks.

Discussion

Measurements of rootstock-induced dwarfing

Rootstock-induced dwarfing presents a ‘difficult-to-measure’ trait in rootstock studies that changes with time and can be highly influenced by many factors including soil type, climate and the interaction between rootstock and scion genotypes. Typical measurements of tree vigour are performed over several years to fully characterize the influence that a rootstock confers to a scion (Seleznyova et al., 2003). This study explored a previously reported association between root bark percentage and rootstock-induced dwarfing in apple
rootstocks (Beakbane and Thompson, 1947). The measurement of a primary rootstock trait confers a large advantage when phenotyping rootstocks for dwarfing ability, and root bark percentage is relatively easy to measure in comparison with other standard measurements of tree vigour. By measuring root bark percentage in the M432 mapping population, we were able to confirm a link between root bark percentage and rootstock-induced dwarfing, with a higher root bark percentage in the ungrafted rootstock correlating with a decrease in stem diameter when the same rootstock was grafted with a scion. Though the association between root bark percentage and rootstock-induced dwarfing was made in this population, further assessment is needed in a wider selection of germplasm to determine whether root bark percentage is a robust trait that is linked to stem dwarfing in other populations.

Genetic interaction and model selection

This study identified QTLs for root bark percentage in the M432 mapping population, identifying three QTLs in this field grown rootstock population. Two strong QTLs, \( Rb1 \) and \( Rb2 \), were identified and found to co-localize with previously determined major dwarfing loci, \( Dw1 \) and \( Dw2 \), as well as QTLs for early bearing, flower density and fruit yield (Rusholme-Pilcher et al., 2008; Fazio et al., 2014), indicating that these chromosomal regions contain valuable rootstock traits. The positions of \( Rb1 \) and \( Rb2 \) concur broadly with the previously reported locations of two QTLs for stem dwarfing, \( Dw1 \) and \( Dw2 \). The first of these has been reported as approximately 14 cM down linkage group 5, strongly linked to the marker CH03a09 (Rusholme-Pilcher et al., 2008). This microsatellite marker is placed approximately 7.6 Mb along chromosome 5 (as judged by BLASTn of sequence XM_008357860.1). From our analysis, the most significant SNP marker is approximately 14.5 Mb along the physical map, though the significant QTL region encompasses the region in which \( Dw1 \) is contained. The large interval of significance for \( Rb1 \) in our study is due to the lack of observable recombination events on this linkage group in M27, whereby 63 out of 140 offspring showed no discernable recombination event on this linkage group. The actual number of markers used for the study of \( Rb1 \) on linkage group 5 was 88 segregating markers. Subsequent development of a microsatellite marker to determine specific haplotypes was mapped to a position 2.4 Mb along chromosome 5.

\( Dw2 \) has been reported as lying at approximately 12 Mb along the physical map of chromosome 11 flanked by CH02d08 and C13243 (Fazio et al., 2014). The most significant SNP linked to \( Rb2 \) maps to 8.4 Mb along the physical map of chromosome 11, while the most significant SSR is CH03d08, which maps <1 cM away from CH02d08 in our map, confirming that \( Rb2 \) and \( Dw2 \) co-locate.

The third QTL, \( Rb3 \), was located on chromosome 13, a region that has not been previously identified in association with any important rootstock trait. \( Rb3 \) is positioned approximately 2.2 Mb along the physical map of chromosome 13 and does not co-localize with any known dwarfing locus. To our knowledge, this region on chromosome 13 has not

### Table 3. Analysis of the three Rb molecular markers developed in this study in a selection of apple rootstocks

<table>
<thead>
<tr>
<th>Apple rootstock</th>
<th>Pedigree</th>
<th>210</th>
<th>212</th>
<th>222</th>
<th>276</th>
<th>296</th>
<th>339</th>
<th>243</th>
<th>252</th>
<th>200</th>
<th>Vigour prediction</th>
<th>Actual vigour</th>
</tr>
</thead>
<tbody>
<tr>
<td>M.1</td>
<td>Unknown</td>
<td>x</td>
<td>x</td>
<td>×</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>V</td>
<td>v</td>
</tr>
<tr>
<td>M.2</td>
<td>Unknown</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>V</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>M.6</td>
<td>Unknown</td>
<td>x</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>V</td>
<td>v</td>
</tr>
<tr>
<td>M.7</td>
<td>Unknown</td>
<td>x</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>V</td>
<td>v</td>
</tr>
<tr>
<td>M.9</td>
<td>Unknown</td>
<td>x</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>SD</td>
<td>SD</td>
</tr>
<tr>
<td>M.10</td>
<td>Unknown</td>
<td>x</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>SD</td>
<td>SD</td>
</tr>
<tr>
<td>M.11</td>
<td>Unknown</td>
<td>x</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>V</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>M.12</td>
<td>Unknown</td>
<td>x</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>V</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>M.13</td>
<td>Unknown</td>
<td>x</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>V</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>M.16</td>
<td>Unknown</td>
<td>x</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>V</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>M.26</td>
<td>M.16 × M.9</td>
<td>x</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>D</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>M.27</td>
<td>M.13 × M.9</td>
<td>x</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>D</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>M.116</td>
<td>M.27 × M.106</td>
<td>x</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>V</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>M.106</td>
<td>N. Spy × M.1</td>
<td>x</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>V</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mac 9</td>
<td>M.9 open pollination</td>
<td>x</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>V</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Northern Spy</td>
<td>Unknown</td>
<td>x</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>V</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ottawa 3</td>
<td>Robin Crab × M.9</td>
<td>x</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>D</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Polish 22</td>
<td>M.9 × Common</td>
<td>x</td>
<td>x</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>V</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

V: vigorous; SI: semi-invigorating; SD: semi-dwarfing; D: dwarfing; DD: highly dwarfing.
been associated with any scion tree architecture phenotype. Through the use of a clonally propagated subset of the mapping population, a link between the effects of all three root bark loci and that of scion dwarfing was established.

The identification of eight genotypic classes (under a two-locus model) using a combination of SNP and microsatellite markers led to the clear observation that $Rb2(b)$ on LG11 was absolutely required for the effect of $Rb1(a)$ on LG05 to be expressed, demonstrating that there are non-additive effects of allelic combination upon the phenotype and that there is negative epistasis acting between the two loci. Furthermore, depending upon the allelic combination and allelic dosage at $Rb2$, the effect of $Rb1(a)$ varied, indicating that some $Rb2$ alleles were better able to compensate for the effect of $Rb1(a)$ than others. The effect of $Rb3(c)$ on root bark percentage appears in most situations to require $Rb1(a)$ or $Rb2(b)$ to be present to have a large effect (Fig. 4F and Supplementary Fig. S2). Furthermore, when $Rb3(c)$ was analysed in relation to stem dwarfing, it was found to interact significantly with $Rb2(b)$ to alter stem diameter (Supplementary Table S2 and Supplementary Fig. S3)—note that a full three-way analysis could not be performed in this experiment. In this study, the model of rootstock-induced dwarfing presented by Fazio et al. (2014), could not distinguish genotypically between the dwarfing ‘M27’ rootstock and the semi-invigorating ‘M.116’ rootstock (Fazio et al., 2014). The new three-locus model developed in this study provided resolution between the two closely related genotypes and, in addition, increased the percentage variance accounted for to 84%, significantly improving the total variance accounted for by 31% ($P=0.001$).

A central role for $Rb2/Dw2$

Previous studies have emphasized the dominant role of $Dw1(Rb1)$ in rootstock-induced dwarfing (Rusholme-Pilcher et al., 2008; Fazio et al., 2014; Foster et al., 2015), although these same studies reported cases where $Dw1$ was present yet the expected phenotype did not behave as predicted. In addition, Foster and co-workers state: ‘$Dw1$ has a stronger effect than $Dw2$’ and that ‘$Dw2$ alone cannot induce dwarfing’, and furthermore, Fazio et al reported peculiarities in that ‘homozygous individuals for non-dwarfing alleles at either $Dw2$ or $Dw1$ invalidate the effect of the dwarfing ability of the other dwarfing locus’. Our approach in this study has enabled us to identify epistatic interactions between all three $Rb/Dw$ loci, and furthermore, determine that $Rb2/Dw2$ plays a significant role in rootstock-induced dwarfing. Previous reports that several rootstocks contain $Dw1$ yet have vigorous phenotypes, can now be explained in the light of our findings that $Rb2/Dw2$ or $Rb3/Dw3$ would need to be present for the dwarfing phenotype to manifest itself. Through a marker screen of selected rootstocks, we found the dwarfing rootstocks ‘Polish 22’ and ‘Mac 9’ only contain markers for $Rb2/Dw2$. This provides support for a central role for $Dw2/Rb2$, and as previously mentioned, also raises the possibility of other interacting loci that are yet to be determined. In addition, the dwarfing rootstock ‘Ottawa 3’ does not have any of the markers linked to the three $Rb$ QTLs, suggesting the $Rb$ markers are unlinked to the QTLs in this genotype or that the dwarfing QTLs, previously identified by Fazio et al. (2014), have arisen independently, though this is unlikely with the dwarfing ‘M.9’ as a parent.

Root bark percentage and rootstock-induced dwarfing

It cannot be stated that the amount of root bark directly affects dwarfing, as this could be the result of a pleiotropic effect of an as-yet-unidentified molecular process. However, vigour control over the scion is still observed with dwarfing rootstocks when they are used as an interstock (a stem piece of dwarfing rootstock grafted in between the rootstock and the scion to create a three-part composite tree), with the degree of dwarfing related to the length of the interstock such that a longer interstock confers a greater degree of dwarfing (Feree and Carlson, 1987). In addition, bark grafts have also been shown to reduce tree vigour, whereby a ring of bark from a tree is removed and replaced with the bark of a dwarfing rootstock (Lockard and Schneider, 1981). In both cases, the bark is implicated in the process of vigour control to the scion; in the latter case, the causative agent is reduced to a far simpler component. However, if constituents within the bark are responsible for rootstock-induced dwarfing, there is currently little information to suggest the identification of the genes responsible or the primary mechanisms underlying this phenomenon, although key hormones such as auxin, abscisic acid and cytokins, as well as chemical compounds including phenolic acids and flavonoids, have all been implicated (Beakbane, 1956; Yadava and Lockard, 1977; Kamboj et al., 1999; Van Hooijdonk et al., 2010).

In woody perennial systems, there is a paucity of information on the development of the secondary vascular cambium and the cellular processes involved in the determination of cambial cell fate. The vascular cambium of trees undergoes asymmetric cell division to differentiate into secondary phloem and xylem cells. It is the cambium that is responsible for the secondary growth leading to the radial thickening of trees (Esau, 1965). The developmental and regulatory networks of vascular cambium differentiation are little understood, yet vascular cellular organization and ontogeny are central to all plant functions, playing a vital role in plant growth and development. Fundamental studies into root bark development and the genetic mechanisms underlying the determination of cambial cell fate in the developing stem and root system would provide important insights into many aspects of woody perennial development. Further research is needed to understand the underlying causes of high root bark percentage in apple rootstocks and how these cellular mechanisms may interact with the scion, including the correlation between root bark percentage and rootstock-induced dwarfing. An alternative explanation for the tight linkage between $Rb$ and $Dw$ loci might be that they are a consequence of linkage drag (high root bark percentage being unintentionally selected alongside rootstock-induced dwarfing) in breeding programmes. However, it is unlikely that QTLs for root bark variation and scion dwarfing would be genetically linked...
to all three rootstock dwarfing loci without a mechanistic explanation.

Conclusions

This study has demonstrated a link between root bark percentage and rootstock-induced dwarfing in a rootstock mapping population. Through the identification of three QTLs for root bark percentage, a three-locus model for predicting rootstock-induced dwarfing to the scion has been developed. The rootstocks ‘M.27’ and ‘M.116’ confer different levels of vigour control towards the scion, and with the identification of 24–64 genotypic classes, we are now able to differentiate genotypically between ‘M.27’ and ‘M.116’ allowing resolution of dominance effects. In addition, we can use the information provided by the development of a three-locus model to further our understanding of rootstock-induced dwarfing by developing new mapping populations that will enable fine mapping of each QTL to identify the genes underlying it and to study the effects of individual QTL. Furthermore, this model can now be incorporated into rootstock breeding programmes to aid marker-assisted breeding strategies. This study is an important step towards the understanding of the genetic mechanism(s) underlying an important rootstock trait and has provided useful molecular markers for marker-assisted breeding.

Supplementary data

Supplementary data are available at JXB online.

Figure S1. Plot of means for chromosome 5 and chromosome 11 based on the model selection procedure.

Figure S2. Plot of three-locus analysis of means for log-transformed root bark percentage.

Figure S3. Plot of three-locus analysis of means for log-transformed stem diameter.

Table S1. Model selection for root bark percentage.

Table S2. Model selection for stem diameter.

Table S3. The genotype and root bark percentage of each individual in the M432 population.

Acknowledgements

This research was supported by grants awarded to NH from the Department for Environment Food & Rural Affairs (Defra-CH0108) and the Horticultural Development Company (HDC-TF212). NH was also supported by funding from the BBsRC, the East Malling Trust and The University of Reading awarded to PJG. We would like to acknowledge financial support through the Leonardo Scholarship Fund for scholarships granted to NBP, ECL and RCR. RCR’s internship was also supported by the commercial rootstock breeding programme at EMR (EMRC). We thank the farm and glasshouse staff of East Malling Research for their care and maintenance of the plant material throughout this study.

References


Foster TM, Kelton J-M, Chagné D,ustin DS, Gardiner SE. 2015. Two quantitative trait loci, Dw1 and Dw2, are primarily responsible for rootstock-induced dwarfing in apple. Horticulture Research 2, 15001.


Lark KG, Chase K, Adler F, Mansur LM, Orf JH. 1995. Interactions between quantitative trait loci in soybean in which trait variation at one locus is conditional upon a specific allele at another. Proceedings of the National Academy of Sciences, USA 92, 4656–4660.


