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Resolution of quantitative resistance to clubroot into QTL-specific metabolic modules

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Abstract

Plant disease resistance is often under quantitative genetic control. Thus, in a given interaction, plant cellular responses to infection are influenced by resistance or susceptibility alleles at different loci. In this study, a genetic linkage analysis was used to address the complexity of the metabolic responses of Brassica napus roots to infection by Plasmodiophora brassicae. Metabolome profiling and pathogen quantification in a segregating progeny allowed a comparative mapping of quantitative trait loci (QTLs) involved in resistance and in metabolic adjustments. Distinct metabolic modules were associated with each resistance QTL, suggesting the involvement of different underlying cellular mechanisms. This approach highlighted the possible role of gluconasturtiin and two unknown metabolites in the resistance conferred by two QTLs on chromosomes C03 and C09, respectively. Only two susceptibility biomarkers (glycine and glutathione) were simultaneously linked to the three main resistance QTLs, suggesting the central role of these compounds in the interaction. By contrast, several genotype-specific metabolic responses to infection were genetically unconnected to resistance or susceptibility. Likewise, variations of root sugar profiles, which might have influenced pathogen nutrition, were not found to be related to resistance QTLs. This work illustrates how genetic metabolomics can help to understand plant stress responses and their possible links with disease.

Keywords: Brassica napus, clubroot, dwarfing gene bzh, metabolomics, metabolic quantitative trait loci, oilseed rape, Plasmodiophora brassicae, quantitative resistance.

Introduction

Plant cellular responses to pathogens include constitutive and induced metabolic adjustments and the biosynthesis of a vast diversity of specialized small compounds (Dixon, 2001; Piasecka et al., 2015), whose toxicity can be challenged by tolerance mechanisms in pathogens (Osbourn et al., 1995; Enkerli et al., 1998, Coleman et al., 2011; Kettle et al., 2015; Pedras and Abdoli, 2017). The development of metabolomic approaches (Fiehn et al., 2000) led to renewed investigation of how pathogenic processes reprogram plant primary metabolism (Schultz et al., 2013; Rojas et al., 2014) and has provided opportunities to broaden our view about the roles of specialized metabolites possibly involved in plant defence (Heuberger, 2014;
Tenenboim and Brotman, 2016). To take advantage of the large accessibility to metabolic phenotypes provided by these approaches, strategies are required to identify biochemical traits that are positively or negatively linked to pathogen and symptom development.

Metabolic profiling has often been used to compare pairs of plant genotypes showing contrasting resistance or susceptibility to a given pathogen strain, thus allowing the identification of candidate compounds putatively involved in plant defence or susceptibility (Ludwig-Müller et al., 1997; Lopez-Gresa, 2010; Sade et al., 2015; Yogendra et al., 2014). There can, however, be possible drawbacks to such comparative approaches, the first being that metabolic diversity among plant genotypes is controlled by a wealth of loci (Rowe et al., 2008; Chen et al., 2014), many of which might be unrelated to the resistance/susceptibility to a given pathogen strain. This problem can be solved by using near-isogenic lines, which allow the metabolic processes putatively connected to one given resistance locus to be specifically addressed (Gunniah, 2012). However, the metabolic comparison of resistant versus susceptible genotypes may be more complicated when examining quantitative resistances, controlled by the combined influence of sets of quantitative trait loci (QTLs). Quantitative resistance is widespread among plant–pathogen interactions (St Clair, 2010); most of the time, both in genotypes harbouring major resistance genetic factors and in apparently fully susceptible accessions, the level of symptoms is modulated by susceptibility or resistance alleles at low-effect QTLs, where each allele can provide a particular contribution to the complexity of metabolic responses (Corwin and Kliebenstein, 2017).

In this regard, the combination of metabolomics and quantitative genetics has been of great value in dissecting the roles of plant secondary metabolites in the interactions between plants and insects (Kliebenstein et al., 2002; Meihls et al., 2013). However, such approaches have been implemented only rarely in the field of plant–pathogen interactions (Denby et al., 2004; Koutouan et al., 2018). In the present study, we focused our efforts on deciphering the metabolic regulations triggered during the interaction between oilseed rape (Brassica napus) and the agent of clubroot, Plasmopara brassicae. This species is a eukaryotic (Rhizaria), telluric, and obligate biotroph, with the agent of clubroot, Plasmodiophora brassicae (Bergendahl et al., 2014). During the interaction between oilseed rape (Brassica napus) and subsequent substantial agronomic losses in Brassica crops worldwide. Genetic studies on the resistance of Brassicaceae species to different isolates of P. brassicae have highlighted major genetic resistance factors (Piao et al., 2009; Ueno et al., 2012; Hatakeyama et al., 2013; Kato et al., 2013; Chu et al., 2014; Rahman et al., 2014; Zhang et al., 2014; Yu et al., 2016; Zhang et al., 2016), and specific and broad-range QTLs (Manzanares-Dauleux et al., 2000a, 2003; Rocherieux et al., 2004; Joubault et al., 2008a; Chen et al., 2013; Tomita et al., 2013; Lee et al., 2016). As far as the information is accessible, genetic resistance factors appear to exert their control on pathogen development only after the completion of the primary phase (Kroll et al., 1983; Hatakeyama et al., 2013). One usual metabolic hallmark of susceptibility to clubroot infection is the accumulation of pathogen-synthesized trehalose, mostly during the later stages of infection (Keen and Williams, 1969; Broman et al., 2003; Gravot et al., 2011). In a previous study, we also pointed out that an accumulation of amino acids together with glutathione and S-methylcysteine during early secondary infection is correlated with susceptibility to clubroot in B. napus accessions (Wagner et al., 2012). However, the functional significance of these metabolic deviations remain unexplained.

The objective of the present study was to relate quantitative genetics to both metabolomics and pathogen-resistance traits, to get new insights into the metabolic modules associated with allelic variations at each QTL involved in the control of partial resistance. This work was focused on the progeny derived from two parental genotypes: Darmor-bzh, which displays a high level of partial resistance to the P. brassicae isolate eH, mainly controlled by the locus PbBr1, and Yudal, which shows an intermediate level of resistance to eH, controlled by several QTLs (Manzanares-Dauleux et al., 2003; Laperche et al., 2017; Aigu et al., 2018). We started the study with a histological and PCR-based evaluation of pathogen development over the duration of infection, to identify the earliest time point at which post-invasion partial resistance was distinguishable between the two parental genotypes. Then, at the earliest time point, metabolomic traits were investigated in roots of a segregating progeny derived from a Darmor-bzh × Yudal cross. Interlaced relationships between metabolites and the resistance-associated QTLs involved were then analysed using Cytoscape-based network representations.

Materials and methods

Plant and pathogen materials

Darmor-bzh is a winter dwarf line of B. napus, nearly isogenic to the genotype Darmor (Fousset et al., 1995), with an available genome sequence (Chalhoub et al., 2014). Yudal is a spring inbred line of B. napus selected from a Korean population. The segregating doubled haploid (DH) population used in this study was derived from an initial cross between the genotypes Darmor-bzh and Yudal (Fousset et al., 1996). The progeny, hereafter called the DY population, was first genotyped by Delourme et al. (2006). Based on these data, a subset of 130 DH lines was selected from this population, using the MapPop software (Brown and Vision, 2000) to optimize recombination events over the whole genome.

eH is a selection isolate of P. brassicae (Filhing et al., 2003). This isolate belongs to the most virulent pathotype, P1, according to Somé et al. (1996). This isolate was multiplied on the highly susceptible B. napus spp. pekinensis cv. Granaz (genotype ECDS) and its pathogenicity profile was verified in parallel in each experiment, using the host differential set described by Somé et al. (1996).

Pathogenic assays

Most clubroot assays in this study (except the study of primary infection; see below) were performed using a previously reported procedure (Wagner et al., 2012). In brief, plants were sown in plates filled with a mixture of two-thirds compost and one-third vermiculite, then cultivated under controlled conditions (16 h light at 22°C/8 h dark at 18°C). Plantlets were
inoculated 7 days after germination, using the eH isolate of *P. brassicae* \((10^7\text{ spores mL}^{-1})\) according to Manzanares-Dauleux *et al.* (2000b), and non-inoculated plants were treated with distilled water (control treatment).

**Evaluation of infection kinetics in genotypes Darmor-bzh and Yudal**

To study primary infection in root hairs (Fig. 1A), seedlings of genotypes Darmor-bzh and Yudal were germinated on moistened paper in petri dishes, and then cultivated hydroponically: the system consisted of a plastic pipette tip box and each plantlet was grown in one hole in the box. The root system of the plantlets was completely immersed in 400 ml of tap water (for non-inoculated plants) or 400 ml of an inoculum suspension at a concentration of \(10^7\) spores ml\(^{-1}\) (for inoculated plants). Plantlets were sampled at 10 days post-inoculation (dpi) and their roots were directly stained with 1% toluidine blue (Sigma Aldrich, St. Louis, MO, USA) for microscopy.

Secondary infection kinetics was followed using a clubroot assay organized in a randomized complete block design with four replications. In each experiment, plants of Darmor-bzh and Yudal were inoculated with isolate eH and grown using previously reported procedures (Wagner *et al.*, 2012). Root samples were collected at 14, 21, 28, 35, and 42 dpi. In each block, each sample consisted of a bulk of 27 plants, except for the non-inoculated 'Yudal' sample.

**Fig. 1.** Time course of compatible interactions in the two genotypes of *B. napus*, Darmor-bzh and Yudal, inoculated with the eH isolate of *P. brassicae*. (A) Histopathological characterization of clubroot infection in the root hairs at 10 days post-infection (dpi) and from transverse sections of the roots at 35 dpi of inoculated and non-inoculated plants. Staining with 1% toluidine blue allows visualization of *P. brassicae* inside root hairs during the primary phase, and inside cortical cells during the secondary phase. Bars=25 μm. (B) Typical root symptoms at 49 dpi. Bars=1 cm. The white ellipse on the central picture indicates the presence of a small gall. (C) Dynamics of pathogen root invasion in Darmor-bzh and Yudal followed by quantitative PCR using DNA extracted from whole infected roots. The internal transcribed spacer region of *P. brassicae* (**PbITS**) was amplified and compared with the *Cruciferin A* gene of *B. napus* (**BnCruA**), and the ratio of pathogen to plant DNA (**Pb**) was calculated. Samples were analysed at 14, 21, 28, 35, and 42 dpi. Data are expressed as the mean ±SE of four replicates. Asterisks indicate significant differences in Pb values between Darmor-bzh and Yudal at the same sampling time (\(P<0.05\), Wilcoxon-Mann-Whitney test). (This figure is available in colour at JXB online.)
first sampling point at 14 dpi, where 54 plants were collected to obtain enough plant material for further analyses. Six additional plants per block and per genotype were included in the experimental design to evaluate symptom severity at 49 dpi (Fig. 1B). At each sampling date, total roots were harvested and immediately frozen in liquid nitrogen and stored at −80°C before further analyses.

Additional plants were included in this assay for histological evaluation of *P. brassicae* development during secondary infection. Roots of inoculated and non-inoculated plants of genotype Darmor-bzh were collected weekly from 14 to 35 dpi and immediately immersed in 2% glutaraldehyde in phosphate buffer (0.1 mM, pH 7.2) before subsequent histological investigations. Root samples previously fixed in glutaraldehyde were dehydrated according to Wagner et al. (2012) and embedded in Technovit 7100 resin ( Heraeus Kulzer, Wehrheim, Germany) according to the supplier’s instructions. Thick sections (4 µm) were cut with a rotary microtome (Microm Microtech, Francheville, France) and stained with 1% toluidine blue O (Sigma Aldrich).

**Clubroot assays with doubled haploid progeny**

A clubroot assay was performed using isolate eH, the parental lines Darmor-bzh and Yudal, and 130 DH progeny lines. The experiment consisted of two independent biological replicates. For each biological replicate, 10 blocks of 18 non-inoculated plants. DH lines were represented and Y udal, and 130 DH progeny lines. The experiment con-
The 19 linkage groups, named A01 to A10 for the A genome and C01 to C09 for the C genome, covered 2087.7 cM. QTL analyses were performed using the R/QTL package (Broman et al., 2003; Arends et al., 2010). Composite interval mapping was carried out for each of the 231 metabolites, as well as for Pb and the DI. The number of covariates was set according to a first run of simple interval mapping for the trait under investigation. Logarithm of the odds (LOD) thresholds were estimated using 1000 permutations (Churchill and Doerge, 1994), and QTL confidence intervals were defined according a LOD drop-off of one unit.

A meta-analysis was carried out using Biomercator V4.2 (Goffinet and Gerber, 2000, Arcade et al., 2004, Sosnowski et al., 2012). In this study, metaQTLs were defined as genetic regions where multiple QTLs were colocated, and were obtained by integrating information from individual QTL detections (metabolites, DI, and Pb) using the Meta-analysis tool. The number of metaQTLs per linkage group was assessed according to the Veyrieras algorithm (Veyrieras et al., 2007) and the Bayesian Information Criterion. Each individual QTL contributed to one or several metaQTLs.

**Cytoscape-based representation of QTL/trait networks**

The full list of QTLs/phenotypic traits was treated as a directed network. In this network, nodes represent loci or traits, and edges represent statistically significant links between loci and phenotypic traits (from the QTL analysis described above). Edge widths were used to represent $R^2$, that is, the percentage of variance explained by each locus for a given phenotypic trait. Edge colour was used to represent positive or negative allele effects on phenotypic traits. This network was represented graphically using Cytoscape software (Shannon et al., 2003).

**Results and discussion**

**Partial resistance and metabolomic responses to clubroot infection in the Darmor-bzh and Yudal parental lines of B. napus**

A first series of experiments was conducted to monitor pathogen development in the *B. napus* parental genotypes Darmor-bzh and Yudal. Primary plasmodia were observed in root hairs of both genotypes at 10 dpi (Fig. 1A). At 14 dpi, small secondary plasmodia were detected in the cortical cells of both Darmor-bzh and Yudal (data not shown). At 35 dpi, infected roots of Yudal displayed extensive hyperplasia and hypertrophied plant cells containing well-developed secondary plasmodia of *P. brassicae* (Fig. 1A). At the same sampling point in Darmor-bzh, secondary plasmodia were also observed, but they were much less developed than in Yudal, and the infected cells were of similar size or even smaller than uninfected cells in the non-inoculated plants. The DI at 49 dpi was 45 for Yudal; clubs were observed on the primary root system in the majority of plants (Fig. 1B) and most of the leaves were wilted. The DI for Darmor-bzh at the same time point was 28, and only a few small galls had developed on the secondary root system. Fully susceptible genotypes from the differential reference set (Somé et al., 1996) displayed DI values of 100. This confirmed that the Darmor-bzh and Yudal genotypes harbour partial resistance to eH.

*Fig. 2.* Selected primary metabolite and glucosinolate contents in inoculated (with *P. brassicae* eH isolate) and non-inoculated (control) roots of the parental lines at 21 dpi. Y, Yudal, D-bzh, Darmor-bzh. (A) Organic acids. (B) Soluble carbohydrates. (C) Free amino acids. (D) Glucosinolates (arbitrary units, based on peak surface area and sample DW). For each condition, analyses were performed using 10 replicates (non-inoculated) or 20 replicates (inoculated with the eH isolate), each replicate consisting of at least 12 plants that were pooled for metabolite extractions. Error bars indicate SE. Different letters indicate significant differences in mean values ($P<0.05$, Tukey method), except for sucrose ($\alpha=0.062$).
The Pb value was similar in both genotypes at 14 dpi, the beginning of the secondary phase of infection (Fig. 1C). In Yudal, Pb increased by approximately two orders of magnitude between 14 and 35 dpi. By contrast, in the infected roots of Darmor-bzh, Pb remained consistently low throughout the whole time-course. Pb values of the two genotypes were significantly different at 21 dpi, after the beginning of the secondary phase of infection (Fig. 1C). In infected roots of Yudal, Pb increased by approximately two orders of magnitude between 14 and 35 dpi. By contrast, in the infected roots of Darmor-bzh, Pb remained consistently low throughout the whole time-course. Pb values of the two genotypes were significantly different at 21 dpi, after the beginning of the secondary phase of infection (Fig. 1C).

Cluster 1: metabolites constitutively accumulated at a higher level in Darmor-bzh and Yudal; cluster 2: metabolites accumulated at a higher level in Yudal than in Darmor-bzh; cluster 3: metabolites accumulated at a higher level in inoculated roots of Yudal and Darmor-bzh; cluster 4: metabolites accumulated at a higher level in inoculated roots of Yudal compared with non-inoculated roots and compared with inoculated roots of Darmor-bzh; cluster 5: metabolites that accumulated at a higher level in inoculated roots of Yudal compared with non-inoculated roots and compared with inoculated roots of Darmor-bzh.

**Fig. 3.** Constitutive and clubroot-induced root metabolic differences between inoculated (control) and non-inoculated (with *P. brassicae* eh) roots of Yudal and Darmor-bzh. Cluster 1: metabolites constitutively accumulated at a higher level in Darmor-bzh than in Yudal; cluster 2: metabolites constitutively accumulated at a higher level in Yudal than in Darmor-bzh; cluster 3: metabolites accumulated at a higher level in inoculated roots of Darmor-bzh compared with non-inoculated roots and compared with inoculated roots of Yudal; cluster 4: metabolites accumulated at a higher level in inoculated roots of Yudal compared with non-inoculated roots and compared with inoculated roots of Darmor-bzh. Metabolomic data are presented as mean-centred values from 10 replicates (non-inoculated) or 20 replicates (inoculated with the eh isolate), each replicate sample consisting of pooled roots from 12 plants. Red indicates higher levels of accumulation and blue indicates lower levels. XCMS-generated mass-tags were named based on their *m/z* (M) and retention time (T). Statistically significant differences (P<0.05) were inferred from Student's t-test.
(neoglucobrassicin) and 4-methoxy-3-indolyl glucosinolate (4-methoxyglucobrassicin), and to the aromatic glucosinolate glucoraphanin (2-phenylethyl glucosinolate). In addition, a series of low-intensity mass signals were found associated with root aliphatic glucosinolate content (glucoerucin, glucoraphanin, glucobrassicanapin, and glucosinate; and 27 unknown analytes, including M655T655, 2012); several additional compounds, such as a salicylic glucoside; and 27 unknown analytes, including M655T655 (with a mass signal ~10-fold higher in inoculated roots of Yudal compared with Darmor- bzh), Cluster 3 mostly included amino acids (threonine, phenylalanine, tryptophan, isoleucine, and valine) that were specifically triggered by *P. brassicae* inoculation in the more resistant accession Darmor- bzh.

A further analysis of this extended metabolite dataset allowed us to identify four categories of analytes showing a contrast between the two genotypes in inoculated roots (Fig. 3): metabolites that are constitutively more accumulated in one of the two genotypes (cluster 1, Darmor- bzh; cluster 2, Yudal), and metabolites whose accumulation is triggered by *P. brassicae* inoculation in only one of the two genotypes (cluster 3, Darmor- bzh; cluster 4, Yudal). Cluster 4 included a large number of compounds that accumulated specifically in inoculated roots of the more susceptible accession, Yudal. This large set of metabolites included trehalose and S-methylcysteine, two analytes that we have previously reported as clubroot susceptibility biomarkers (*Wagner et al.*, 2012); several additional compounds, such as a salicylic glucoside; and 27 unknown analytes, including M655T655 (with a mass signal ~10-fold higher in inoculated roots of Yudal compared with Darmor- bzh), Cluster 3 mostly included amino acids (threonine, phenylalanine, tryptophan, isoleucine, and valine) that were specifically triggered by *P. brassicae* inoculation in the more resistant accession Darmor- bzh.

The biological role of the above-described metabolic features at 21 dpi in the expression of clubroot resistance (or susceptibility) in Darmor- bzh and Yudal is difficult to decipher. Indeed, any compound within these four clusters can a priori promote or impede the infection process; for instance, glucobrassicin and other indole glucosinolates may support the biosynthesis of auxin, thus possibly favouring gall development (*Ludwig-Müller, 2009*), or alternatively might support the synthesis of defensive phytoalexins (*Clay et al.*, 2009; *Klein et al.*, 2017). Moreover, Darmor- bzh and Yudal displayed compatible interactions with

\[ \text{(5.2)} \text{than in Yudal bzh cose was markedly higher in Darmor-} \]

In Yudal, inoculation also led to an important decrease in glucose content from 39 to 21 µmol g DW\(^{-1}\), and to a statistically significant (*P*=0.019) increase in trehalose content from 0.6 to 1 µmol g DW\(^{-1}\) (*Supplementary Table S1*), consistent with the previously reported induction of fermentation processes and hypoxia responses during clubroot infection in a susceptible genotype of *Arabidopsis thaliana* (*Gravot et al.*, 2016).

The increasing accumulation of trehalose (presumably synthesized by *P. brassicae*) is usually observed during the secondary phase of clubroot infection, finally reaching 10–50 µmol g DW\(^{-1}\) in fully developed clubs (5–6 weeks post-inoculation in rapeseed) (*Keen and Williams, 1969; Broman et al., 2003; Gravot et al., 2011; Wagner et al., 2012*). In the present study, the slight increase in trehalose content in Yudal at 21 dpi is indicative of an already active secondary-phase development of *P. brassicae*.

Concatenation of datasets from targeted and untargeted metabolic profiling, resulting in a full dataset of 231 analytes (*Supplementary Table S1*), broadened the view on possible metabolic contrasts between infected and non-infected roots of the two genotypes. PCA of this dataset (*Supplementary Fig. S2*) indicated that ‘genotype’ was the major factor influencing metabolic contrasts between samples (axis 1), followed by ‘inoculation’, which led to a clear metabolic shift in roots of Yudal (axis 2) and, to a lesser extent, in Darmor- bzh (axis 3).

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The biological role of the above-described metabolic features at 21 dpi in the expression of clubroot resistance (or susceptibility) in Darmor- bzh and Yudal is difficult to decipher. Indeed, any compound within these four clusters can a priori promote or impede the infection process; for instance, glucobrassicin and other indole glucosinolates may support the biosynthesis of auxin, thus possibly favouring gall development (*Ludwig-Müller, 2009*), or alternatively might support the synthesis of defensive phytoalexins (*Clay et al.*, 2009; *Klein et al.*, 2017). Moreover, Darmor- bzh and Yudal displayed compatible interactions with
isolate eH, and accordingly it is expected that both genotypes would show some metabolic responses associated with susceptibility. Conversely, previous studies (Laperche et al., 2017; Aigu et al., 2018) have highlighted that both Darmor-bzh and Yudal harbour resistance alleles at different QTLs, and consequently both genotypes were expected to exhibit metabolic features associated with resistance (the mechanisms of which could be either similar or different in the two accessions). To extend the understanding of the relationship between partial resistance and metabolic traits, it was necessary to compare the mapping of QTLs associated with the control of disease traits and metabolic profiles in the two genotypes. To this end, a genetic analysis of Pb, DI, and metabolite patterns was conducted on a 130 DH progeny derived from the cross Darmor-bzh × Yudal.

QTLs controlling pathogen infection and the root metabolome

PCR-based quantification of pathogen relative growth (expressed here as Pb) was performed at 21 dpi in the progeny, where this variable showed a continuous pattern of distribution (Fig. 4A), suggesting polygenic control of pathogen growth. The distribution of DI at 49 dpi in the progeny showed a continuous but near bimodal shape, suggesting the involvement of a major-gene resistance in the control of the club phenotype (Fig. 4B). The values of DI were plotted against the values of Pb for each DH line (Fig. 4C), revealing that DI and Pb were not linearly correlated. This may reflect a saturation in the evaluation of symptoms with the DI scale. The DI takes account of the proportion of galls relative to asymptomatic roots, but it does not take account of the size of fully developed clubs, and thus does not distinguish nuances between highly susceptible genotypes. This argument may, however, not explain everything, as a low Pb value at 21 dpi in a given recombinant line did not allow the prediction of the final DI at 49 dpi, which may also suggest the existence of genetic factors exerting their effects after 21 dpi.

Genetic analysis by composite interval mapping of Pb led to the identification (Supplementary Table S2) of one moderate-effect QTL ($R^2=27.9\%$) on chromosome C09.
(resistance allele from Darmor-bzh) and two weak-effect QTLs on chromosomes C03 and C07 (resistance allele from Yudal). DI was found to be under the control of one major QTL (R²=78.8%) on C09, consistent with the phenotype distribution curve (Fig. 4B). The moderate-effect QTL C09_Pb and the major-effect QTL C09_DI, both detected at 59 cM, colocalized with the major resistance locus, first described as Pb-Bn2 by Manzanares-Dauleux et al. (2003) and further reported by Laperche et al. (2017) and Aigu et al. (2018). Four additional minor QTLs were found to be involved in the control of DI; these were located on chromosomes A01, A10, C02, and C03. The latter two QTLs, on C02 (position=130.8 cM) and C03 (position=35.2 cM), colocalized with QTLs involved in resistance to isolate eH reported by Laperche et al. (2017) and Aigu et al. (2018). In those previous studies, the genetic effect of these QTLs was found to be enhanced by decreasing the nitrogen supply (Aigu et al. 2018). Plants were cultivated under a non-limited nitrogen supply in the present study, and consequently the two QTLs on chromosomes C02 and C03 had only a low effect on resistance. Finally, the genetic architectures of Pb and DI were overlapping only for the QTL on C09 (Pb-Bn2). Other loci controlling Pb (C03, position=2.5 cM, and C07, position=67.7 cM) were also found to be involved in delaying the progression of P. brassicae infection during the early phase, without affecting the final development of clubroot symptoms. By contrast, for example, the resistance QTL on C02 appeared to control final symptom development but without any effect on the early progression of P. brassicae infection (Supplementary Table S2). These results can be compared with other reports of age-related or ontogenetically conditional resistance loci in quantitative resistance to pathogens (examples are given in Corwin and Kliewen, 2017). The fact that Pb and DI do not measure equivalent traits (the first is indicative of pathogen development, whereas the second is indicative of infection-triggered cellular proliferation) may also explain why the genetic architecture of club development overlaps only partially with that of P. brassicae multiplication. Similar phenomena have been previously reported for clubroot resistance in B. napus by Aigu et al. (2018).

Targeted and untargeted metabolite profiling was therefore performed on inoculated root samples of the 130 DH lines at 21 dpi together with the parental lines (detailed results are presented in Supplementary Table S1). QTL analysis was performed for the 231 analytes. No QTL was detected for 55 analytes (including sucrose; the other 54 analytes were unidentified compounds). Nevertheless, 374 metabolite QTLs (mQTLs) were detected (details are given in Supplementary Table S2). One to seven QTLs (most often one or two QTLs) were detected for each analyte. The mean R² per QTL was 12% (range 1.2–66.4%). To simplify further analysis of colocalizations, both mQTLs and resistance QTLs (controlling either Pb or DI) were processed by a QTL meta-analysis, allowing their aggregation on a set of 70 consensus genetic intervals (metaQTLs) (Fig. 5, Supplementary Table S3, Supplementary Figs S3 and S4). When necessary, some mQTLs were simultaneously attributed to two adjacent metaQTLs (details of weighting factors are given in Supplementary Table S3). Such cases were taken into account in further analysis when weighting factors exceeded 20%. Among the 70 metaQTLs identified, 18 contained only one QTL. This included the metaQTL A01_60.54, which exerted a small effect on DI (R²=1.4%) but was not associated with any mQTL, and the QTL A06_11.5, which exerted strong control (R²=40%) on the content of the unidentified root compound M501T921.

By contrast, four genomic regions were involved in the control of more than 20 mQTLs (Fig. 5). The metaQTL C09_58.88 was a major ‘hot-spot’ containing 78 mQTLs, including 35 mQTLs with R²>15%. This mQTL was colocalized with a major resistance QTL controlling DI and Pb (R²=78.8% and 27.9%, respectively). The metaQTL C07_79.74 was associated with 27 mQTLs, and colocalized with a resistance QTL contributing to the control of Pb (resistance allele from Yudal, R²=6.9%). This locus included the major mQTL controlling gluconasturtiin (R²=30.2%), with the Yudal allele being associated with a higher gluconasturtiin content.

Three adjacent metaQTLs (A06_100.49, A06_108.63, and A06_115.03) were found on chromosome A06 along a 15 cM genomic region. Altogether, the metaQTLs from this region were involved in the control of 59 metabolites. No clubroot resistance QTL was detected in this region in our study. A06_115.03 corresponded to the bzh genomic region introduced into the oilseed rape genotype Darmor-bzh from a mutant line obtained by chemical mutagenesis of the genotype Primor (Foisset et al., 1995). Some of the metabolic features observed in the present study were likely caused by this dwarfism allele (bzh), which corresponds to the DELLalingene BnaA06g34810D (genetic position 114.3 cM, physical position 23008797–23010993), which is involved in the repression of gibberellin responses (Zhao et al., 2017). The metaQTL C03_19.48 contained 21 mQTLs. The confidence intervals calculated from the meta-analysis suggested that this metabolic hot-spot did not include the two resistance QTLs detected nearby on the same chromosome: one resistance QTL exerting a moderate effect on Pb (R²=8.9%) and one resistance QTL exerting a low effect on DI (R²=1.5%) (Fig. 5).

The fact that two of the four hot-spot metabolic metaQTLs were genetically unrelated to partial resistance can be viewed as supporting the hypothesis that not every single metabolic contrast observed between Yudal and Darmor-bzh was necessarily connected to the fate (i.e. resistance or susceptibility) of the interaction with P. brassicae. Furthermore, from this dataset it also appeared that several compounds linked to resistance-related metaQTLs were also under the genetic control of other resistance-unrelated metaQTLs, thus questioning the putative functional link of those compounds with resistance. A network-based approach was thus necessary to shed more light on the complexity of locus/metabolite/resistance relationships.

Network-based analysis of links between metabolite fingerprints and resistance QTLs

Similarities and differences between metabolic patterns controlled by each of the four hot-spots were compared
by a Cytoscape-drawn directed network (represented in Supplementary Fig. S5). This representation highlighted not only how each hot-spot or metaQTL displayed its own pattern of metabolites, but also allowed an effective visualization of metabolites that are under the simultaneous control of two or even three of the four metaQTLs.

A first example is a series of amino acids (represented in green in the lower left of Supplementary Fig. S5), which were controlled by several mQTLs clustered under the metaQTL C07_79.74 (which is also associated with clubroot resistance) and other mQTLs clustered under the metaQTL A06_115.03 associated with the mutation bzh. Then, the rise of the amino acid contents could be independently triggered by two independent processes: an enhanced root metabolic sink caused by P. brassicae when a susceptible allele is present at C07_79.74 (Keen and Williams, 1969; Li et al., 2018), or an influence of the dwarfism gene bzh on shoot-to-root metabolite allocation (in line with Elias et al. 2012).

Fig. 6 illustrates the similarities and differences between metabolic patterns observed at 21 dpi associated with each of the three QTLs controlling pathogen development at 21 dpi (expressed as Pb). The amounts of glutathione and glycine are lower (indicated by blue arrows in Fig. 6) in the presence of alleles that decrease Pb at the three metaQTLs C03_1.01, C07_79.74, and C09_58.88. The apparently pivotal significance of these two nitrogen-containing compounds is consistent...
with previous results highlighting these compounds as susceptibility biomarkers (Wagner et al., 2012). In this previous study, Wagner et al. (2012) also identified additional susceptibility-related metabolites, including S-methylcysteine, trehalose, and alanine, which in the present study have been found to be regulated by the QTLs C07_79.74 and C09_58.88. As already mentioned, a large number of specific metabolic features were found to be associated with the metaQTL C09_58.88 (comprising one major-effect resistance QTL and 78 mQTLs). For a large number of these compounds, their increased accumulation was controlled by the resistance allele (indicated by red arrows in Fig. 6), suggesting that some of these unidentified compounds might be good candidates for causal factors involved in defence (detailed below). The metaQTL C03_1.01 included a major effect mQTL ($R^2=30.2\%$) controlling gluconasturtiin content, and a set of 13 mQTLs. The increased accumulation of gluconasturtiin was controlled by the resistance allele, which may suggest a positive role exerted by this glucosinolate on partial resistance. The resistance allele at QTL C07_79.74 was clearly involved in repressing the content of a series of amino acids, some of which were also repressed by C09_58.88, whereas others were specifically controlled by QTL C07_79.74.

A similar network connecting mQTLs at 21 dpi to QTL involved in the control of DI at 49 dpi (Supplementary Fig. S6), indicated that at 21 dpi only the major QTL, Pb-Bn2 (C09_58.88), was associated with a rich metabolic network. By contrast, minor QTLs involved in the control of DI displayed only faint metabolic fingerprints. This result was consistent with the fact that those minor QTLs did not colocalize with any QTLs controlling pathogen development (i.e. Pb) at 21 dpi. Together, these results indicate that for the QTLs with

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**Fig. 7.** Sub-networks representing interconnected genetic architectures of clubroot resistance traits with (A) organic acids, (B) non-structural carbohydrates, and (C) glucosinolates. Nodes represent metabolites (circles), resistance traits (squares), or loci defined from the QTL meta-analysis (diamonds). The width of the edges indicates the quantitative effect of each QTL ($R^2$) for a given QTL. When metaQTLs are associated with the control of Pb or DI, red and blue directed edges indicate positive or negative effects, respectively, on trait values (metabolite, Pb, or DI), exerted by Darmor or Yudal alleles at the QTL (as indicated in the diamonds).
a low effect on DI, underlying resistance processes are set up after 21 dpi.

In the first part of this study, comparative targeted analysis from parental lines had highlighted constitutively higher contents of organic acids, glucose, and glucosinolates in the accession Yudal compared with Darmor-bzh (Fig. 2), raising the question of the role of these compounds during *P. brassicae* infection. Citric acid, which accumulated at constitutively higher levels in the roots of Yudal, was found to be under the control of four QTLs, three of which were involved in the

**Fig. 8.** Sub-networks representing interconnected genetic architectures of the pathogen–plant genomic DNA ratio (Pb) at 21 dpi with clusters of compounds highlighted in Fig. 3. (A) Metabolites accumulated during early clubroot infection only in Darmor-bzh and not in Yudal. (B) Metabolites accumulated at constitutive higher levels in the roots of Darmor-bzh (compared with Yudal). The width of the edges is indicative of the quantitative effect of each QTL (\( R^2 \)) for a given QTL. Red and blue edges indicate positive and negative effects, respectively, on trait values (metabolite or Pb), controlled by the Darmor allele at the metaQTL C09_58.88, or the Yudal allele at the metaQTLs C07_79.74 and C03_1.01.
control of clubroot resistance traits (Fig. 7A). For each of these three QTLs, the resistance allele was associated with a decrease of citric acid content, thus suggesting that constitutively high citric acid levels might contribute to susceptibility. Indeed, with levels of more than 20 µmol g DW−1 in roots, this compound might be a primary source of ready-to-use organic carbon for B. brassicae. By contrast, glucose content was under the control of seven QTLs (Fig. 7B), of which only one (R² = 7.6%) aggre-gated with the metaQTL A10_68.42, which colocalized with a QTL involved in the control of DI. At this locus, the resistance allele was from Darmor-bzh and was associated with an increase of glucose content, which is inconsistent with the initial hypothesis that higher glucose contents in Yudal may favour pathogen development.

The constitutively high level of gluconasturtiin in Yudal was mostly controlled by the Yudal allele at the metaQTL C03_1.01, which also contributed to the reduction of Pb at 21 dpi (Fig. 7C). To a minor extent, the Darmor resistance allele at the metaQTL C09_58.88 also contributed to increase the content of gluconasturtiin. This genetic architecture thus suggests that high root gluconasturtiin contents might contribute to partial resistance. By contrast, neoglucobrassicin (the most abundant indole glucosinolate in roots) was under the control of only two QTLs with low R² values (36.5% and 23.7%, respectively), indicating that the constitutively high levels of those two compounds in Darmor-bzh was related to the presence of the Darmor-bzh resistance allele at the metaQTL C09_58.88. This ‘coincidence’ consistently suggested a possible role for these two unidentified molecules as candidate key players in the partial resistance driven by C09_58.88. Additional high-resolution MS/MS approaches allowed more precise m/z measurements of parental ions, and further investigation of their corresponding ion fragments (Supplementary Fig. S7). These fingerprints did not, however, match any referenced molecule available in public databases. Additional semiprep-scale purification of these compounds would be required for further investigations of their chemical structures by nuclear magnetic resonance spectroscopy.

Conclusions

Allelic variations affecting plant metabolism can potentially affect the plant/pathogen trophic relationship, regulate plant defence responses, or modulate the negative consequences of pathogen development. However, identifying connections between given metabolic features and disease resistance can be a difficult task when dealing with quantitative resistances controlled by series of resistance QTLs. In this work, metabolomic analysis of the B. napus genotypes Darmor-bzh and Yudal, which express contrasting degrees of clubroot resistance/susceptibility, highlighted differentiated constitutive metabolic features in non-inoculated plants, and dissimilar inducible responses to infection. A detailed analysis of metabolic and pathogen responses in the progeny of those two accessions allowed comparison of the QTL architectures governing clubroot resistance traits and metabolic profiles.

Conclusions from the present work should be nuanced by possible inaccuracies in the estimation of QTL colocalizations. In addition, some QTLs may have been missed in the analysis, due to the limited number of recombinant lines in this study (130 DH lines). However, from a broad perspective, our approach allowed us to build an interesting contrasted map of metabolic features associated with different metaQTLs. To the best of our knowledge, this work is the first report of network (Cytoscape)-based representation of multi-trait QTL interactions for the investigation of plant responses to a pathogen. As shown here, this approach can help to test hypotheses, and to get a clear view of the complexity of genetic networks influencing metabolic responses and disease resistance. Our main conclusions are that (i) contrasting metabolic features are associated with metaQTLs controlling pathogen development, suggesting that those resistance loci may mobilize different physiological responses; (ii) a significant part of the genotype-specific metabolic responses to infection is not genetically connected to either resistance or susceptibility; and (iii) several constitutively accumulated metabolites may be linked to partial resistance.

From the seminal works of Keen and Williams (1969) to the recent study of Wałerski et al. (2018), it has been clearly shown that modulation of the carbohydrate economy plays a
major role in the development of clubroot. It would thus have been conceivable that allelic variations affecting the amount of soluble carbohydrates could affect the course of infection. However, despite differences in the accumulation of glucose, fructose, and sucrose between the roots of Yudal and Darmor-bzh, these accumulations were found not to be related to the genetic factors controlling partial resistance. By contrast, our work underlined that partial resistance to clubroot is genetically related to the control of constitutive accumulation of citric acid (which may represent a consistent source of organic carbon for *P. brassicae*), glucostaurin, and the two unknown analytes M320T2161 and M368T3033.

Supplementary data

Supplementary data are available at *JXB* online.

Table S1. Full dataset of metabolomic analyses of the parental lines and in the Darmor-bzh × Yudal progeny.

Table S2. Detailed results from the QTL analysis.

Table S3. Detailed results from the metaQTL analysis.

Table S4. Formatted network data for Cytoscape analysis.

Table S5. Node attributes for Cytoscape analysis.

Fig. S1. Overview of the metabolomic workflow.

Fig. S2. PCA representation of metabolomic profiles.

Fig. S3. Detailed graphic view of mQTL colocalizations on the genome A.

Fig. S4. Detailed graphic view of mQTL colocalizations on the genome C.

Fig. S5. Sub-network representing mQTLs connected to the four genomic ‘hot-spots’ (>20 colocalized mQTLs).

Fig. S6. Sub-network representing mQTLs connected to the five QTL involved in the control of disease index.

Fig. S7. MS/MS high-resolution spectra (negative mode) of compounds corresponding to the two mass tags M320T2161 and M368T3033.

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