Detection of the quantitative trait loci for \( \alpha \)-amylase activity on a high-density genetic map of rye and comparison of their localization to loci controlling preharvest sprouting and earliness

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Abstract The objectives of the research were to determine the position of quantitative trait loci (QTL) for \( \alpha \)-amylase activity on the genetic map of a rye recombinant inbred line population—S120 × S76—and to compare them to known QTL for preharvest sprouting and heading earliness. Fourteen QTL for \( \alpha \)-amylase activity on all seven chromosomes were identified. The detected QTL were responsible for 6.09–23.32% of \( \alpha \)-amylase activity variation. The lowest LOD value (2.22) was achieved by locus QAa4R-M3 and the highest (7.79) by locus QAa7R-M1. Some QTL intervals for features of interest overlapped partially or completely. There were six overlapping QTL for \( \alpha \)-amylase activity and preharvest sprouting (on 1R, 3R, 4R, 6R, 7R) and the same number for preharvest sprouting and heading earliness (on 1R, 2R, 6R, 7R). Furthermore, there was one interval partially common to all three traits, mapped on the long arm of chromosome 1R. Testing of lines originating from hybrid breeding programs, such as S120 and S76, may provide important information about the most significant genes and markers for selection in commercial breeding. Among the statistically significant markers selected in the Kruskal–Wallis test (\( P < 0.005 \)), there were 55 common ones for preharvest sprouting and heading earliness (1R, 2R, 6R), 30 markers coinciding between \( \alpha \)-amylase activity and preharvest sprouting (5R, 7R) and one marker for \( \alpha \)-amylase activity and heading earliness (6R).

Keywords \( \alpha \)-amylase • DArT • Heading earliness • High density genetic map • Preharvest sprouting • QTL • RIL • Rye

Introduction

Recent advances in molecular marker technology and the development of high-density molecular marker linkage maps have provided powerful tools for elucidating the genetic basis of quantitatively inherited traits (Kunpu et al. 2009). The first high-resolution map of rye constructed with the use of
Diversity Arrays Technology (DArT) (Jaccoud et al. 2001) was created in 2009 (Bolibok-Bragoszewska et al. 2009). The next high-density linkage map of rye, based on DArT and PCR markers, was recently used for detecting quantitative trait loci (QTL) for preharvest sprouting (PHS) and heading earliness (Myśkowski 2011). These agriculturally important traits are connected with two crucial events in the plant’s life cycle. First is the end of dormancy and seedling emergence, because until then there is no photosynthesis and plants cannot gain mass. The second event is anthesis which is important because it determines the time of grain maturity during the season (Jamieson et al. 1998). The appropriate timing of flowering is a critical adaptive trait for the propagation and survival of a plant species. It should be late enough to avoid frequent exposure of the sensitive reproductive organs to freezing temperatures early in the season, but not so late as to expose the crop to damaging drought conditions and high temperatures during anthesis and grain filling (Chen et al. 2009). Premature breaking of dormancy can result in PHS of spikes in wet weather conditions. PHS is a major cause of excess α-amylase activity (AA), which impairs grain quality since enzymatic hydrolysis of starch during food manufacture can lead to processing problems and unsatisfactory end products (Graybosch et al. 2000; Lunn et al. 2001; Mares et al. 2002; Tjin Wong Joe et al. 2005). Much research on wheat grains has proved that there are a number of additional causes for the deterioration of meal properties, which reflects low falling number. These include late maturity α-amylase (LMA) (Mrva and Mares 1996a, b, 1999, 2002, 2008), also known as prematurity α-amylase (PMAA) in the UK (Lunn et al. 2001; Tjin Wong Joe et al. 2005), and retained pericarp α-amylase (RPAA) (Lunn et al. 2001).

It is well established that the genetic mechanism of AA, PHS and heading/flowering in crops is complex. Earliness is controlled by photoperiodic response, chilling requirement and narrow sense earliness. Each of these components are under the control of multiple loci, localized on most of the wheat and barley (bibliography in: Cockram et al. 2007; Kunpu et al. 2009) and all rye chromosomes (Plaschke et al. 1993; Masoń and Milczarski 1999; Börner et al. 2000; Korzun et al. 2001; Stojałowski and Łapiński 2002; Myśkowski 2011). All chromosomes of Triticeae are also involved in determining PHS and AA (Masoń and Milczarski 2005, 2009, and bibliography therein).

Extensive research involving crop species has been conducted on the relationship between flowering time (heading date), plant adaptation and yield (Laurie 1997). Several data suggest that loci controlling photoperiod response, vernalization requirement and earliness per se may exert pleiotropic effects on yield and yield-related traits (Worland et al. 1998; Buck-Sorlin and Börner 2001; Lewis et al. 2008). Studies of QTL controlling PHS and AA in rye demonstrated that these two systems coincide in great part (Masoń and Milczarski 2005, 2009; Masoń et al. 2011). Partial overlap between QTL for heading earliness (HE) and PHS in rye has been found by Myśkowski (2011). However, a combined relationship between AA, PHS and earliness has not been defined yet. The objectives of our research were to determine the position of QTL for AA on the genetic map of a rye recombinant inbred line (RIL) population—S120 × S76—and to compare them to known QTL for PHS and HE.

Materials and methods

The mapping population of S120 × S76 consisted of 143 genotypes of the RIL-F8 generation. The pedigree of parental inbred lines is described in Myśkowski et al. (2001). S120 and S76 differ in values of AA, PHS and time of ear emergence.

The genetic map of the S120 × S76 RIL population has been constructed by Myśkowski (2011) with the use of JoinMap 3.0 software (Van Ooijen and Voorrips 2001). All seven linkage groups comprised 1285 DArT loci and 62 PCR-based loci. Individual chromosomes included from 123 (5R) to 261 (6R) loci, and spanned distances of 76 cM (3R) to 233 cM (1R). The whole map length was 962 cM and the average density varied from 0.9 to 2.4 markers/cM (average distance 0.4–1.1 cM).

AA was analyzed during the period 2008–2010, for self-pollinated plants of the RIL-F8 to RIL-F10 generations. Plant material was grown in an experimental field at the West Pomeranian University of Technology in Szczecin, Poland. Each RIL was represented by 1–8 spikes (4–5 for most genotypes) from different plants. Equal amounts of flour (250 mg) obtained from all seeds with no visible
signs of sprouting were used for AA determination. The method of estimating the level of AA was described by Masojc and Larsson-Rażnikiewicz (1991). Additionally, PHS was measured in the year 2010 as a percentage of growing seeds per total seeds in the ear, after watering of ten mature, harvested spikes, according to the method described by Masojc et al. (2007).

Statistical analysis was carried out using the STATISTICA version 9.0 package (http://www.statsoft.com). Broad-sense heritability ($h^2$) was calculated using the formula $h^2 = VG/V_F$, where $VG$ is the genetic variance estimated as a difference between $V_F$ and $V_E$; $V_F$ is the RIL population variance, $V_E$ is the mean variance of parental lines S120 and S76, estimated with using at least ten plants of each one.

The significance of difference between parental lines trait values was established by employing the Cochran and Cox test.

The relationship between segregations of single marker and trait was analyzed with the Kruskal–Wallis test using MapQTL 5.0 package (Van Ooijen 2004).

Linkage analysis was performed using the composite interval mapping (CIM) method (Zeng 1994) with Windows QTL Cartographer version 2.51 (http://www.statgen.ncsu.edu/qtlcart/ WQTLCart.htm, Wang et al. 2007). The step size chosen for all QTL was 2 cM. Significant thresholds for declaring the presence of a QTL were estimated from 1,000 permutations of the data (Doerge and Churchill 1996).

Results of composite interval mapping performed on the data of AA and new data of PHS were compared with localization of QTL for PHS and HE obtained previously by Myśków (2011) with the use of the same genetic map of the S120 × S76 RIL population.

Results

Phenotypic variation and correlation analysis

Parental phenotypic variation and the distribution among RILs for AA, PHS and HE in different years are shown in Table 1. Parental lines differed significantly ($P = 0.001$) with respect to PHS and HE, but not significantly in the case of AA values. The measured target traits varied in the RIL population following a continuous distribution representing a normal phenotypic segregation for QTL mapping. All trait value ranges were higher in the RIL population than in both parents, suggesting that transgression was observed. The estimated heritability of AA, HE and PHS was 24.7, 34.5 and 46.7%, respectively.

Pairwise correlation coefficients between the traits are given in Table 2. Significant correlations were observed between result data obtained in all years for PHS, and for almost all variants of HE and for AA assessed in 2008 and 2009. There was no correlation found between different traits, except one—between AA and PHS during the 2009 season.

### Table 1 Characteristics of α-amylase activity (AA), preharvest sprouting (PHS) and heading earliness (HE), in mapping population S120 × S76

| Trait | Mean value ± standard error for parental line in population | Year | Mean value in population | Variation range in population |
|-------|-------------------------------------------------------------|------|--------------------------|--------------------------------|
|       | S120 | S76 |       |                          |                                |
| AA (mm) | 9.3 ± 0.7 | 10.6 ± 0.8 | 2008 | 10.1 | 6–16.5 |
|       |       |       | 2009 | 11.4 | 6–16  |
|       |       |       | 2010 | 10.2 | 6–19  |
| PHS (%) | 68.1 ± 1.3 | 25.6 ± 1.3 | 2007 | 28.1 | 0–88.9 |
|       |       |       | 2008 | 37.9 | 0–100 |
|       |       |       | 2009 | 41.4 | 0–100 |
|       |       |       | 2010 | 48.8 | 0–100 |
| HE (°) | 7.4 ± 0.2 | 3.1 ± 0.2 | 2007 | 6.4  | 1–9   |
|       |       |       | 2008 | 4.7  | 1–9   |
|       |       |       | 2009 | 5.0  | 1–9   |
|       |       |       | 2010 | 5.8  | 1–9   |
Table 2  Phenotypic correlation between α-amylase activity (AA), preharvest sprouting (PHS) and heading earliness (HE) in rye RIL population of S120 × S76 cross

|       | AA09 | AA10 | AA  | HE07 | HE08 | HE09 | HE10 | HE  | PHS07 | PHS08 | PHS09 | PHS10 | PHS  |
|-------|------|------|-----|------|------|------|------|-----|-------|-------|-------|-------|------|
| AA08  | 0.45***| 0.17 | 0.77***| 0.01 | −0.16 | −0.14 | −0.04 | −0.11 | −0.07 | 0.22  | 0.19  | 0.06  | 0.18 |
| AA09  | −0.05 | 0.59***| −0.12 | −0.05 | 0.03  | 0.05  | 0.01  | 0.25  | 0.39***| 0.37***| 0.10  | 0.34***|
| AA10  | 0.69***| −0.17 | −0.02 | −0.03 | −0.07 | −0.05 | −0.07 | −0.03 | −0.04 | −0.03  | −0.03  | −0.05  |
| AA  | −0.13 | −0.10 | −0.06 | −0.03 | −0.07 | 0.03  | 0.21  | 0.19  | 0.04  | 0.15  |
| HE07  | 0.29* | 0.33**| 0.23  | 0.41***| 0.14  | 0.06  | 0.02  | 0.09  | 0.05  |
| HE08  | 0.63***| 0.67***| 0.85***| 0.00  | 0.04  | −0.02 | −0.02 | −0.02 | 0.01  |
| HE09  | 0.75***| 0.88***| −0.08 | 0.07  | 0.01  | 0.02  | 0.02  | 0.01  |
| HE10  | 0.91***| −0.05 | 0.09  | 0.02  | −0.04 | 0.02  | 0.02  |
| HE  | −0.04 | 0.08  | 0.02  | −0.01 | 0.01  | 0.02  | 0.02  |
| PHS07 | 0.48***| 0.51***| 0.35* | 0.55***|
| PHS08 | 0.72***| 0.53***| 0.86***|
| PHS09 | 0.61***| 0.88***|
| PHS10 | 0.83***|

*, **, *** Significance at $P < 0.01$, 0.001, 0.0001, respectively

a Mean values
QTL analysis

Putative QTL for AA detected in each year (2008–2010) and as a result of average values of all seasons are listed in Table 3. This table contains information about QTL for PHS detected in 2010 and those based on the mean values of the trait (2007–2010). Fourteen QTL for AA on all seven chromosomes and seven QTL involving chromosomes 3R, 5R and 6R for PHS were identified. Detected QTL were responsible for 6.09–23.32% and 7.51–17.08% of AA and PHS variation, respectively. The lowest LOD value of 2.22 was achieved by locus QAa4R-M3 and the highest—7.79—by locus QAa7R-M1. The range of LODs of QTL peaks for PHS varied from 2.35 to 4.32. Most of the alleles increasing AA were contributed by line S120. Four of the seven alleles enhancing PHS originated from parent S76. Three of them were localized on the 6RL chromosome close to each other (Fig. 1).

None of the QTL controlling AA were identified in 2 years. However, three of them were confirmed by the detection on the basis of mean values (Table 3). Three of the QTL for PHS detected in 2010 were mapped in an earlier study (Myśkó 2011) and the fourth was found additionally using the mean score over all years (Table 3).

Eighteen of 20 segregations of markers most strongly linked to the QTL were significantly associated with segregations of AA or PHS, as revealed by the Kruskal–Wallis test; six of them were significant at \( P < 0.001 \) (Table 3).

The locations of all QTL for AA, PHS and HE in the RIL population of S120 × S76 are shown in Fig. 1 and in Electronic Supplementary Material 1. If any locus was found in different years, it was presented only as a single rectangle and counted as a single QTL. There were 14 QTL detected for AA, 33 for PHS and 17 for HE. Considering QTL revealed twice (in 2 years or both in 1 year and as a result obtained on the basis of the average data), there were 4, 16 and 9 loci, for AA, PHS and HE, respectively.

Co-localization of QTL and common markers

Some QTL intervals for measured features overlapped partially or completely (Fig. 1, Supplement 1). There were six overlapping QTL for AA and PHS (on 1R, 3R, 4R, 6R, 7R) and the same number for PHS and HE (on 1R, 2R, 6R, 7R). Furthermore, there was one interval partially common to all three traits, mapped on the long arm of chromosome 1R.

Among the statistically significant markers selected using the Kruskal–Wallis test (\( P < 0.005 \)), there were 55 common ones for PHS and heading earliness: 29 located on chromosome 1R, 10 from linkage group 2R and 16 from chromosome 6R. There were 30 coincident markers identified between AA and PHS (one from linkage group 5R, the rest from 7R). There was one significant marker for AA and HE (6R). Markers common to two traits of interests are listed in Supplement 2.

Discussion

In rye, QTL mapping for AA and PHS has been an area of intense research, which has led to the identification of many loci and to the discovery of the phenomenon of overlapping intervals of QTL for these two traits (Masojć and Milczarski 2005, 2009). A large number of QTL showed relatively small effects and only a few major QTL with large phenotypic effects were previously identified. This presents a serious challenge when using single QTL with small effects for marker-assisted selection (MAS), and therefore there is a need to pay more attention to exploring those stable QTL detected in different environments/seasons and different genetic backgrounds and those revealing pleiotropic effects (Wang et al. 2009). The present research is an extension of earlier studies on AA and PHS and a summary of results from QTL analysis of AA, PHS and HE. This is the first attempt to use a high-density genetic map of rye for QTL analysis of these three important traits. It provides more opportunities for the identification of markers tightly linked to the target features. The study highlights the importance of some co-localized QTL, which could show the direction of studies on pleiotropic effects. The first case of application of an advanced RIL population in research on rye allowed for verification and phenotypic assessment in different environments.

AA QTL

Since in our study seeds were harvested at the stage of full ripeness, the majority of the α-amylase developed
Table 3  Characteristics of putative QTL and markers for α-amylase activity and pre-harvest sprouting in rye RIL population of S120 × S76 cross

| QTL   | Chromosome | Probably present on the other maps | Year                  | Mapping interval (cM) | LOD  | a (%) | R² (%) | Marker linked to a QTL | Highest significance in Kruskal–Wallis test | Mean value of genotype a | Mean value of genotype b |
|-------|------------|-----------------------------------|-----------------------|-----------------------|------|-------|--------|------------------------|---------------------------------------------|--------------------------|--------------------------|
| QAa1R-M1 | 1R        | +                                 | 2009, 2008–2010       | 8–23                  | 2.42 | 0.86  | 8.66   | Xpr47L1000             | ****                                         | 10.9                     | 8.8                      |
| QAa1R-M2 | 1R        | +                                 | 2009                  | 38–39                 | 2.40 | -0.89 | 7.63   | XrPt66654             | **                                          | 11.1                     | 10.3                     |
| QAa1R-M3 | 1R        | +                                 | 2008                  | 200–223               | 3.31 | -1.13 | 12.48  | XrPt509418             | ****                                         | 9.3                      | 10.8                     |
| QAa2R-M1 | 2R        | –                                 | 2010                  | 0–2                   | 2.57 | -1.30 | 8.20   | XrPt509653             | –                                           | 9.5                      | 11.0                     |
| QAa3R-M1 | 3R        | +                                 | 2010                  | 14–15                 | 2.77 | 1.37  | 9.40   | XrPt505544             | ****                                         | 11.3                     | 10.3                     |
| QAa3R-M2 | 3R        | +                                 | 2009                  | 58–60                 | 2.75 | 0.65  | 7.04   | XrPt390268             | ****                                         | 12.1                     | 11.0                     |
| QAa4R-M1 | 4R        | +                                 | 2009, 2008–2010       | 3–10                  | 2.71 | -0.70 | 7.70   | XrPt506669             | ******                                        | 11.1                     | 12.4                     |
| QAa4R-M2 | 4R        | –                                 | 2010                  | 36–38                 | 2.24 | 1.22  | 7.72   | XrPt506689             | ****                                         | 11.1                     | 9.6                      |
| QAa4R-M3 | 4R        | –                                 | 2008–2010             | 65–67                 | 2.22 | 0.64  | 6.09   | XrPt399485             | **                                          | 10.9                     | 9.7                      |
| QAa5R-M1 | 5R        | +                                 | 2009, 2008–2010       | 80–88                 | 2.79 | 0.75  | 10.18  | XrPt505693             | ***                                         | 12.2                     | 11.2                     |
| QAa6R-M1 | 6R        | –                                 | 2008                  | 35–36                 | 2.32 | -0.90 | 7.38   | XrPt389418             | **                                          | 9.4                      | 10.6                     |
| QAa6R-M2 | 6R        | –                                 | 2009                  | 42–43                 | 2.84 | 0.79  | 8.01   | XrPt401305             | **                                          | 12.1                     | 11.1                     |
| QAa6R-M3 | 6R        | +                                 | 2008                  | 95–99                 | 2.80 | 0.94  | 8.48   | XrPt506969             | **                                          | 11.5                     | 10.2                     |
| QAa7R-M1 | 7R        | +                                 | 2009                  | 89–104                | 7.79 | 1.31  | 23.32  | XrPt402657             | ******                                        | 12.4                     | 10.1                     |
| QPhs3R-M3 | 3R        | –                                 | 2009, 2010             | 31–33                 | 2.61 | -7.07 | 7.96   | XprL43L1100             | ****                                         | 40.8                     | 52.6                     |
| QPhs3R-M4 | 3R        | +                                 | 2009, 2010             | 42–43                 | 2.50 | 7.58  | 9.13   | XrPt411037             | **                                          | 35.7                     | 25.0                     |
| QPhs5R-M3 | 5R        | +                                 | 2010                  | 0–3                  | 2.35 | 6.77  | 8.75   | XrPt398706             | ******                                        | 50.5                     | 35.0                     |
| QPhs6R-M5 | 6R        | –                                 | 2010                  | 50–53                 | 2.42 | -6.53 | 7.51   | XrPt507720             | –                                           | 26.3                     | 31.0                     |
| QPhs6R-M6 | 6R        | –                                 | 2010                  | 58–63                 | 3.88 | -9.21 | 17.08  | XrPt509579             | ****                                         | 39.9                     | 53.7                     |
| QPhs6R-M4 | 6R        | –                                 | 2008, 2010             | 64–77                 | 3.39 | -7.07 | 8.96   | XrPt400777             | ****                                         | 39.9                     | 53.7                     |
| QPhs6R-M7 | 6R        | +                                 | 2010, 2007–2010       | 89–98                 | 4.32 | 8.42  | 11.65  | XrPt506969             | –                                           | 44.5                     | 38.2                     |

***, ****, ***** Significance at P < 0.05, 0.01, 0.005, 0.001, 0.0005, 0.0001, respectively

1 Values in (mm) for AA and in (%) for PHS
should represent LMA, which was shown to be the \(\alpha\)-AMY1 group of isozymes (Mrva and Mares 1999, 2002; Mares and Mrva 2008).

In order to find out whether the AA QTL reported here correspond to any of the known QTL, the map used in the present study was aligned to the other published genetic maps used in studies of QTL for AA and PHS. Nine out of 14 AA QTL reported here were probably mapped in the regions previously indicated in studies performed in two other rye populations (Mastojí and Milczarski 2005, 2009). QAa7R-M1 seems to be the most important due to its highest LOD of 7.79 and because it explained as much as 23% of AA. The DArT marker XrPt402657 linked to this QTL was statistically significantly related to the trait at \(P < 0.0001\) and it separates the genotypes into two groups with differing values of AA.

The most recent report on loci linked to AA detected using bidirectional selective genotyping (BSG) method coupled with molecular mapping (Mastojí et al. 2011) pointed to seven markers distributed on the long arm of chromosome 7R. QAa7R-M1 may contain one or a few of them. However, the loci revealed therein were considered to be the ones with minor value for AA synthesis; the approach presented by that team did not allow for estimated parameters obtained by QTL analysis. BSG gives no information about the relative magnitude of the QTL effects acting in favour of the valuable trait.

![Localization of \(\alpha\)-amylase activity QTL (white rectangles) and QTL for preharvest sprouting (grey rectangles) revealed in 2010 on the linkage map of S120 × S76 RIL population and their alignment with QTL for preharvest sprouting (grey rectangles) and heading earliness (black rectangles) detected previously (Myšków 2011). QTL detected both in 1 year and on the base of mean values are indicated by one asterisk. QTL detected in two seasons are indicated by two asterisks. The ruler on the left side shows map distances in cM. Markers situated on the right of each linkage group are those most strongly linked to the newly detected QTL. All markers from the map are listed in Supplement 1](image-url)
(Masojć et al. 2011). In a recently published report on localization of genes affecting AA (Tenhola-Roininen et al. 2011), only one QTL was detected in breeding materials analyzed, on the long arm of chromosome 5R. Interestingly, the coinciding locus detected by Masojć et al. (2011) was considered to belong to the hypostatic, less important class of loci. This leads to the question whether research performed only on genotypes very different with respect to AA and PHS, such as lines 541 and Ot1-3 used by Masojć et al. (2011), properly reflects the genetic diversity of actual breeding materials. Testing of populations developed from crosses between cultivars like Amilo and Voima (Tenhola-Roininen et al. 2011) or lines originating from hybrid breeding program, such as S120 and S76, may provide important information about the most significant genes and markers for selection in commercial breeding.

In the past, it has been suggested that the QTL identified in more than one environment or those identified using data pooled over environments are useful from the point of view of MAS. Furthermore, for use in MAS, it is desirable to have one or a few QTL, each with a major effect on the trait (Kulwal et al. 2005). In the light of this observation, the QTL from chromosome 7R with DArT marker XrPt402657 and the next eight QTL detected in the present study and reported previously should prove valuable in MAS aimed at improving the grain quality of rye in terms of decrease in AA.

Interestingly, QTL analysis performed on the F2-F3 populations of S120 × S76 (Myśków et al. 2010) allowed for the detection of only one QTL for AA, not corresponding to any of the QTL for AA revealed in the present study.

The comparable or even higher number of QTL for AA on the map presented compared to the results in other rye maps suggested that there is no need to detect significant differences between values of the trait of parental lines in QTL analysis. Both high- and low-activity QTL alleles were found in each parental line, which explains the similarity of the mean values of both lines and the appearance of transgressive recombinants in the segregating population. The lack of repeatable QTL through the years reflects the low heritability of AA. However, it does not diminish the importance of the QTL, since they are confirmed in different genetic backgrounds.

PHS QTL

Out of the seven QTL for PHS detected in 2010, three were homeologous to the loci from the same map identified before (Myśków 2011). One of these three loci, QPhs3R-M4, was probably detected either in the F2 population of S120 × S76, as a Phs1 (Myśków et al. 2010). However, QPhs5R-M3 was revealed in one season only; it probably coincided with Phs2 from the F2 population and with QTL for PHS or AA mapped on chromosome 5R, near the structural locus of α-amylase in other populations (Masojć et al. 2007, Masojć and Milczarski 2009). A marker linked to this locus, XrPt398706, was statistically significant at P < 0.0005. The next QTL apparently important was QPhs6R-M7 from group 6R, due to its identification using pooled data, its expression in different genetic backgrounds and because its interval overlapped with the region containing a locus controlling AA—QAa6R-M3.

In total 33 QTL for PHS were detected on the present map, 17 of them identified in at least two seasons. Similar to the case of QTL for AA analysis, this number exceeds the quantity of loci known from the earlier studies using the CIM method in a S120 × S76-F2 population (Myśków et al. 2010) and two others (Masojć and Milczarski 2009). This could partially be a result of different threshold values of LOD obtained with the permutation test in the present study and threshold values used in previously published reports (LOD = 3.0). This could also be the consequence of using a high-density map and more abundant RIL population. Even if the new QTL are of less importance, they add important information to the growing knowledge concerning PHS resistance and AA in rye.

HE QTL

Exploiting the high-density map of the S120 × S76 population resulted in detection of 17 QTL for HE falling on all chromosomes except 3R (Myśków 2011). Ten of them were identified in two seasons at least. This was the first report on QTL controlling earliness mapped on chromosome 1R in rye. There were therefore at least five new loci detected. QHe5R-M1 from chromosome 5R was probably homeologous to locus Vrn-R1, previously known as Sp1 (Plashke
et al. 1993), responsible for plant reaction for vernalization.

Relationship between AA, PHS and HE

Correlation between pairs of analyzed traits showed no significant relationship. AA and PHS also showed no correlation within two other mapping populations of rye (Twardowska et al. 2005), suggesting their independent genetic basis. However, it was found that QTL systems controlling PHS and AA partially overlap. Nine common QTL were detected, among 16 QTL for AA and 13 QTL for PHS (Masojc´ and Milczarski 2009). In the present study, six coinciding QTL for AA and PHS were found, which confirms the previous observations showing partial coincidence of the genomic regions controlling these two traits. Despite the higher density of our map, it is still not resolvable if loci for both traits are only linked or have pleiotropic effects. However, three perfectly overlapping QTL from chromosomes 1R, 3R and 6R, with the same markers linked both to AA and PHS QTL, make the second explanation probable. The fourth coinciding QTL from linkage group 6R had only a slight shift and the same marker linked, too. Such ideal or almost ideal matches of short intervals seem not to be accidental, taking into account that there is often no or weak correlation between QTL for one trait detected in different seasons.

There were no reports on the relationship between potentially related traits of PHS and HE in rye, except for the publication by Myśków (2011). The connection of PHS and HE in other crops is not well documented. PHS resistance was associated with later heading date (HD) in white wheat population. The mean HD and PHS scores were significantly negatively correlated ($r = -0.39$), probably due to a major HD QTL found to be tightly linked to the PHS QTL on chromosome 2B (Munkvold et al. 2009). Despite the lack of correlation between PHS and HE, implying their independent genetic basis, there were as many as six coinciding QTL for both traits found. It shows that the relationship between PHS and both AA and HE is on the congruent level. However, such an ideal overlapping as between some QTL for AA and PHS was not observed, which suggests linkage rather than a pleiotropic effect of the revealed loci.

Additionally, one interval partially overlapping for all three studied traits was detected, which was the first report of this kind. The reason for co-localization of the QTL for two or all three target features remains unknown and requires developing studies. Future research involving fine mapping of these coinciding QTL may resolve whether the co-localized QTL represent a single locus with pleiotropic effect or whether there are two linked loci.

The present results show that there is no rule that plants prone to PHS reveal high activity of LMA. Similarly, earlier heading plants are not always early sprouting, and are then not susceptible to PHS in wet weather conditions. In any case, focusing on overlapping QTL should allow the choice of markers beneficial in the selection of plant materials with favorable values of PHS, AA and HE. This work reveals 55 markers linked to PHS and HE simultaneously, 30 to PHS and AA and one to AA and HE. Knowledge of DArT marker sequences enables their conversion to PCR markers, which are easy to use in MAS.

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