Genetic factors affecting Fusarium head blight resistance improvement from introgression of exotic Sumai 3 alleles (including \(Fhb1\), \(Fhb2\), and \(Fhb5\)) in hard red spring wheat

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**Abstract**

**Background:** Fusarium head blight resistance genes, \(Fhb1\) (for Type-II resistance), \(Fhb2\) (Type-II), and \(Fhb5\) (Type-I plus some Type-II), which originate from Sumai 3, are among the most important that confer resistance in hexaploid wheat. Near-isogenic lines (NILs), in the CDC Alsask (susceptible; \(n = 32\)) and CDC Go (moderately susceptible; \(n = 38\)) backgrounds, carrying these genes in all possible combinations were developed using flanking microsatellite markers and evaluated for their response to FHB and deoxynivalenol (DON) accumulation in eight environments. NILs were haplotyped with wheat 90 K iSelect assay to elucidate the genomic composition and confirm alleles’ presence. Other than evaluating the effects of three major genes in common genetic background, the study elucidated the epistatic gene interactions as they influence FHB measurements; identified loci other than \(Fhb1\), \(Fhb2\), and \(Fhb5\), in both recurrent and donor parents and examined annotated proteins in gene intervals.

**Results:** Genotyping using 81,857 single nucleotide polymorphism (SNP) markers revealed polymorphism on all chromosomes and that the NILs carried < 3% of alleles from the resistant donor. Significant improvement in field resistance (Type-I + Type-II) resulted only among the CDC Alsask NILs, not the CDC Go NILs. The phenotypic response of NILs carrying combinations of Sumai 3 derived genes suggested non-additive responses and \(Fhb5\) was as good as \(Fhb1\) in conferring field resistance in both populations. In addition to \(Fhb1\), \(Fhb2\), and \(Fhb5\), four to five resistance improving alleles in both populations were identified and three of five in CDC Go were contributed by the susceptible parent. The introgressed chromosome regions carried genes encoding disease resistance proteins, protein kinases, nucleotide-binding and leucine rich repeats’ domains. Complex epistatic gene-gene interactions among marker loci (including \(Fhb1\), \(Fhb2\), \(Fhb5\)) explained > 20% of the phenotypic variation in FHB measurements.

**Conclusions:** Immediate Sumai 3 derivatives carry a number of resistance improving minor effect alleles, other than \(Fhb1\), \(Fhb2\), \(Fhb5\). Results verified that marker-assisted selection is possible for the introgression of exotic FHB resistance genes, however, the genetic background of the recipient line and epistatic interactions can have a strong influence on expression and penetrance of any given gene.

**Keywords:** Cereals, Epistatic interactions, Fusarium head blight, \(Fhb1\), \(Fhb2\), \(Fhb5\), Scab, Sumai 3, Wheat
Background

Wheat (*Triticum* spp. L.) is one of the most important field crops worldwide as it serves as staple food for a large proportion of the global population. Wheat production is challenged by several constraints and *Fusarium* head blight is one of the major biotic limitations. There are several *Fusarium* spp. that cause head blight or scab; *Fusarium graminearum* Schwabe (syn. Gibberella zeae Schw. [Petch]) is the main culprit in North America, and it is also hosted by maize (*Zea mays* L.) and barley (*Hordeum vulgare* L.) [1, 2]. Direct yield loss from the disease is due to shrivelled grain with lower test weight or even failure of seed formation. Loss in marketability from mycotoxins accumulation is an even bigger concern from an international trade perspective. The accumulation of harmful mycotoxins, particularly deoxynivalenol (DON) and its acetylated forms (3-ADON and 15-ADON), may render the grain unsuitable for food or feed. The majority of wheat growing countries have defined certain threshold limits for the presence of DON in the grain to be able to export or import across international boundaries and many beverage and food industries have self-imposed even greater restrictions [1].

An integrated approach for FHB management is imperative and genetic resistance is an integral part of such disease management approach. Resistance to FHB in wheat is inherited quantitatively and strongly influenced by the environment [3]. Genetic studies in wheat have identified many useful loci for improvement in complex traits, such as FHB; unfortunately, many of them remain un- or under-utilized in plant breeding programs mainly because of the complex nature of resistance [3]. In spite of a tremendous amount of FHB resistance breeding efforts, genetic gain has been moderate [4]. Efficient introgression of QTL associated with FHB resistance into elite germplasm requires the use of linked genetic markers to facilitate marker-assisted selection (MAS); however, the linkage phase between the marker(s) and the QTL cannot always be inferred among genetic back-grounds, unless strong linkage disequilibrium exists [5]. Of more than 100 QTL identified for resistance to FHB, only seven have been formally designated as Mendelized genes: *Fhb1* derived from Sumai 3 [6], *Fhb2* from Sumai 3 [7], *Fhb3* from *Leymus racemosus* [8], *Fhb4* from Wangshuibai [9], *Fhb5* from Wangshuibai and Sumai 3 [10], *Fhb6* from *Elymus tsukushiensis* [11], and *Fhb7* from *Thinopyrum ponticum* [12]. Based on host re-ponse, the expression of resistance is classified into five different types: Type-I (resistance to initial pathogen infection), Type-II (resistance to fungal spread in the spike), Type-III (resistance to toxin accumulation or the ability to degrade the mycotoxins), Type-IV (resistance to kernel infection), and Type-V (tolerance to yield loss) [13–15]. Type-I and Type-II resistance are more widely exploited and Type-III resistance has gained importance as it is important to maintain grain end-use quality. All types of resistance are generally moderately to well correlated.

The discovery of promising QTL is a preliminary step in a MAS program, but validation of such loci in multiple genetic backgrounds and environments is equally important [16]. The actual effect of the QTL, usually identified from bi-parental populations such as recombinant inbred lines or double haploid lines, is dependent on the alleles and allelic frequencies present at the locus, as well as epistatic interactions among QTL and other genes, which are usually over-estimated in the original mapping population [5]. Near-isogenic lines (NILs) are advantageous for studying phenotypic effects attributable to a specific QTL or gene as the genetic background is fixed, which in turn maintains morphological and phenological traits of the plants that might influence the trait under study [17]. NILs are particularly attractive to breeders for traits that are introduced from exotic parents or wide crosses as they allow confirmation of allelic effects on traits of interest. Additionally, by fixing the genetic background, NILs serve as an ideal source for fine-mapping, gene expression profiling, and hypothesis-driven biological experimentation [18].

Canadian wheat growers have witnessed several FHB epidemics in the last two decades, particularly in eastern Canada and the province of Manitoba in western Canada [2, 19]. However, in last 10–15 years, FHB epidemics in Saskatchewan are not uncommon; attributable to the increasing proportion of the more aggressive 3-ADON chemotype in the *F. graminearum* population [19, 20]. These epidemics spurred research to improve genetic resistance and management options for FHB. The majority of the resistance genes currently available originate from Asian or Brazilian wheats; however, breeders in North America are reluctant to use exotic sources in their programs due to linkage drag (for example, shattering and susceptibility of Sumai 3 to other pathogens). In an effort to introgress resistance into Canadian hard red spring wheat, the bread wheat breeding program at the Crop Development Centre (CDC) utilized 04GC0139 (pedigree: ND2710/RL4851/BW278; carrying *Fhb1, Fhb2,* and *Fhb5*), a derivative of Sumai 3, to cross with CDC wheat cultivars. The current project used NILs in CDC Alsask and CDC Go backgrounds to study the effects of these three major genes and their combinations: *Fhb1, Fhb2,* and *Fhb5*. The effects of introgressing *Fhb1* and *Fhb5* on disease resistance are reported in previously published studies from North America [18, 21, 22], Europe [23, 24], and China [25]. The majority of these studies used RILs/NILs from *F2* derived inbreds through enforced inbreeding or doubled haploid lines and only Salameh et al. [24] and Xue et al. [25] used BC2 or BC3.
of the polymorphic SNPs among the NILs, most of the markers were located on Chromosomes 3B (452), 5A (444), 5B (341), and 6B (510) in CDC Alsask, and on 1A (127), 2A (167), 3B (83), 5A (226), and 7A (188) in CDC Go. The chromosomes 3B, 5A, 6B carrying Fhb1, Fhb5, and Fhb2 were anchored with 38% of the total polymorphic SNPs in CDC Alsask and 26% in CDC Go.

With the help of previously published studies or the use of a consensus map and physical location of the SNPs, the markers on Chromosomes 3B, 5A, and 6B from the 90K assay were identified and used to define haplotype segments carrying Fhb1, Fhb2, and Fhb5 in both populations (Fig. 1) [27–29]; Ron Knox, unpublished data. The classification of the NILs into gene classes using microsatellite markers was in agreement with SNP markers for all three genes in both populations with the exception of Fhb1 in two NIL entries (Go2 and Go6) in the CDC Go background. The two inconsistent NIL entries for presence/absence of Fhb1 were classified using the functional gene-specific, pore-forming toxin (PFT) protein marker [30]. The CDC Alsask NILs carried more alleles (difference in recombination rate, a function of genetic background) from the donor parent as compared to the CDC Go NILs (Fig. 1; Additional file 1 Table S1, Figures S1-S3). The genomes of the CDC Alsask NILs carried up to 2.7% of the resistant donor’s alleles and the CDC Go NILs 0.9% (Additional file 1 Table S1). On Chromosomes 3B (carrying Fhb1), 5A (carrying Fhb5), and 6B (carrying Fhb2), the CDC Alsask NILs had up to 9.9, 11.6, and 12.6% of the resistant donor parent alleles, respectively, while the CDC Go NILs had 2.2, 6.7, and 1.7%.

FHB evaluations and heritability estimates
In all environments, FHB inoculations were successful and there was sufficient disease pressure in all environments to discriminate NIL entries as indicated by longer environmental vectors in biplots which were proportional to the standard deviation in the phenotypic data (Additional file 1: Figure S4). Also, there was positive and significant correlations (P = 0.05) among most of the environments in both CDC Go and CDC Alsask populations, indicated by acute angles between environmental vectors. Two axes of the biplots explained ~ 56% of the phenotypic variation in both populations. For GH evaluations, F-values were significant for variation among gene classes in both NIL populations for GH14, GH21, and GH_AUDPC (Table 1). The F-values for entry nested within gene class were significant for GH14, GH21, and GH_AUDPC in the CDC Go population, but only for GH_AUDPC in the CDC Alsask population. The two F. graminearum chemotypes differed for GH14 and GH_AUDPC in both populations, whereas the interaction of chemotype by gene was significant only for
GH_AUDPC in the CDC Alsask population. For field evaluations, F-values were significant only for gene classes for all FHB parameters in both populations. Insignificant F-values for entry nested within gene class for INC, SEV, IND, and DON indicated that the NIL entries within any given gene class behaved similarly. Broad-sense heritability ($H^2$) estimates were high for GH14, GH21, GH_AUDPC, moderate for INC, SEV, IND and weak for DON. In both GH and field evaluations, the random effect of environment (site-year) was significant ($P = 0.05$; data not presented).

In the CDC Alsask population, for GH14, NILs carrying $Fhb1$, $Fhb5$, $Fhb2 + Fhb5$ and $Fhb1 + Fhb2 + Fhb5$ reduced FHB severity (SEV) as compared to the recurrent parent and the susceptible checks (Table 2). Other genes or gene combinations tended to lower the disease as compared to the recurrent parent and the susceptible checks, but differences were not statistically significant. For GH21 and GH_AUDPC, all genes and their combinations reduced disease compared to the recurrent parent and the susceptible check. For the CDC Go population, except for $Fhb2$, all other genes or their combinations reduced disease severity in GH14. For GH21, only NILs carrying all three genes reduced FHB severity and for GH_AUDPC, all genes classes except $Fhb1$ and $Fhb2$ had lower disease severity than the recurrent parent or the susceptible check. Pyramiding $Fhb1 + Fhb2 + Fhb5$ in the CDC Alsask background reduced severity (relative to the recurrent parent) by 37% for GH14, 27% for GH21, and 39% for GH_AUDPC, and in the CDC Go background by 62, 33, and 49%, respectively. Although NILs carrying all three genes performed better than the intermediate/moderately resistant (MR) check, i.e. AC Barrie, the improvement was not comparable to the resistant check or the resistant donor parent. It is important to mention that the NILs that did not carry $Fhb1$, $Fhb2$, or $Fhb5$ also reduced disease by 1–13% in both populations, and the combination of any two of these genes in the CDC Alsask population did not improve resistance as compared to NILs carrying the genes singly, which indicated significant gene interactions. The 3-ADON chemotype resulted in higher FHB severity, for GH14 and GH_AUDPC, as compared to the 15-ADON chemotype in both populations (Fig. 2).

In the field evaluation of the CDC Alsask population, only gene combinations of $Fhb2 + Fhb5$ and $Fhb1 + Fhb2 + Fhb5$ reduced FHB incidence and all three genes

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**Table 1** Analysis of variance (F-values; across all environments) for near-isogenic lines (NILs) in CDC Go and CDC Alsask backgrounds, carrying all combinations of three Fusarium head blight (FHB) resistance genes: $Fhb1$, $Fhb2$, and $Fhb5$. Fixed effects of FHB resistance genes and entry (nested within gene) are provided for FHB severity assessed in the greenhouse at 14 days post inoculation (GH14), 21 days post inoculation (GH21), area under disease progress curve (GH_AUDPC), field incidence (FLD_INC), field severity (FLD_SEV), field FHB index (FLD_IND), deoxynivalenol (FLD_DON) accumulation, and broad-sense heritability ($H^2$). For greenhouse data, the effect of chemotype (3ADON or 15ADON), and chemotype by gene interaction is also presented.

| Effect               | dF | GH14       | GH21       | GH_AUDPC   | FLD_INC   | FLD_SEV   | FLD_IND   | FLD_DON   |
|---------------------|----|------------|------------|------------|-----------|-----------|-----------|-----------|
| **CDC Alsask**      |    |            |            |            |           |           |           |           |
| Gene (G)            | 12 | 82.47***   | 364.28*    | 178.70***  | 9.60***   | 16.82***  | 13.25***  | 5.20***   |
| Gene (entry)        | 29 | 2.29*      | 3.42*      | 3.90***    | 1.86*     | 1.21*     | 0.98*     | 1.04*     |
| Chemotype (C)       | 1  | 11.38**    | 1.41*      | 19.80***   | –         | –         | –         | –         |
| C*G                 | 12 | 0.99*      | 1.01*      | 2.17*      | –         | –         | –         | –         |
| $H^2$ (%)           | –  | 87.5       | 92.4       | 90.6       | 77.3      | 52.3      | 59.6      | 21.4      |
| **CDC Go**          |    |            |            |            |           |           |           |           |
| Gene (G)            | 12 | 19.71***   | 25.19***   | 25.60***   | 11.47***  | 9.74***   | 10.92***  | 7.39***   |
| Gene (entry)        | 30 | 2.03**     | 1.65*      | 2.04**     | 1.07**    | 1.03**    | 0.95**    | 1.13**    |
| Chemotype (C)       | 1  | 8.03*      | 3.02*      | 7.38*      | –         | –         | –         | –         |
| C*G                 | 12 | 0.99*      | 1.15*      | 1.12*      | –         | –         | –         | –         |
| $H^2$ (%)           | –  | 80.9       | 90.4       | 86.3       | 58.0      | 56.3      | 45.0      | 15.4      |

Note: *, **, ***: significant at $P < 0.05$, $P < 0.001$, $P < 0.0001$, respectively; ns – not significant

*aDegree of freedom*
or their combinations reduced severity and FHB index compared to the recurrent parent (Table 3). The DON toxin was reduced only by \( \text{Fhb1} + \text{Fhb5} \), \( \text{Fhb2} + \text{Fhb5} \), and \( \text{Fhb1} + \text{Fhb2} + \text{Fhb5} \). For CDC Go, only the combination of all three genes reduced incidence and index, whereas all other gene classes were comparable to CDC Go. The DON accumulation was reduced by \( \text{Fhb5} \) and combinations of \( \text{Fhb5} \) with \( \text{Fhb1} \) and \( \text{Fhb2} \), and \( \text{Fhb1} + \text{Fhb2} + \text{Fhb5} \). The combination of these three genes reduced FLD_INC, FLD_SEV, FLD_IND, and FLD_DON by 9, 32, 37, and 49% in the CDC Alsask background, and by 14, 20, 26, 40% in the CDC Go background, respectively.

### Marker main effects and epistatic interactions

Epistatic interaction analyses were carried out between marker pairs for all marker loci and multiple genome-wide interactions were identified that influenced FHB parameters (Table 4). Statistically significant epistatic interactions (\( P < 0.001 \)) among other marker loci and \( \text{Fhb1} \), \( \text{Fhb2} \), and \( \text{Fhb5} \) were identified; interactions with \( \text{Fhb5} \) were the most common among the three genes, however, the nature of the epistatic interactions (additive/additive-dominant) could not be defined. In the CDC Alsask population, epistatic interactions explained up to 18.1% of the variation for FLD_INC, 24.4% for FLD_SEV, 25.9% for FLD_IND, and 16.4% for FLD_DON.

### Table 2

Means and standard errors for FHB severity assessed in the greenhouse at 14 days post inoculation (GH14), 21 days post inoculation (GH21), and area under disease progress curve (GH_AUDPC) in gene classes and check lines for CDC Alsask and CDC Go near-isogenic lines. Means within each column for each population followed by the same letter are not statistically significantly different according to Fisher’s least significant difference (LSD) at \( P = 0.05 \).

| Gene/genotype | GH14 (%) | SEM | PDR | GH21 (%) | SEM | PDR | GH_AUDPC | Mean | SEM | PDR |
|---------------|----------|-----|-----|----------|-----|-----|----------|-----|-----|-----|
| **CDC Alsask** |          |     |     |          |     |     |          |     |     |     |
| CDC Teal (susceptible check) | 72.7 a | 2.3 | - | 98.6 a | 1.1 | - | 977.3 a | 26.9 | - |
| CDC Alsask (recurrent parent) | 59.1 bc | 1.8 | - | 97.2 a | 1.5 | - | 862.0 b | 21.6 | - |
| AC Barrie (moderately resistant/intermediate check) | 49.1 d| 3.3 | - | 83.3 b | 2.0 | - | 676.3 de | 32.1 | - |
| ND2710 (resistant check) | 7.4 i | 1.6 | - | 15.4 d | 4.1 | - | 124.2 g | 24.1 | - |
| 04GC0139 (resistance donor parent) | 6.1 i | 1.4 | 89.7 | 7.6 d | 1.3 | 92.2 | 87.9 g | 14.7 | 89.8 |
| Null (n = 4) | 55.3 bcd | 1.6 | 64 | 91.3 ab | 1.9 | 6.1 | 749.3 c | 18.1 | 13.1 |
| Fhb1 (n = 6) | 42.2 gh | 2.0 | 28.6 | 82.4 b | 2.5 | 15.2 | 607.2 ef | 22.8 | 29.6 |
| Fhb2 (n = 4) | 47.6 c-g | 2.0 | 19.5 | 88.0 b | 2.5 | 9.5 | 673.3 d | 22.1 | 21.9 |
| Fhb5 (n = 2) | 44.8 e-h | 3.0 | 24.2 | 80.1 bc | 3.9 | 17.6 | 613.9 def | 33.4 | 28.8 |
| Fhb1 + Fhb2 (n = 2) | 48.4 b-h | 4.5 | 18.1 | 86.6 bc | 4.1 | 10.9 | 685.5 cde | 50.1 | 20.5 |
| Fhb1 + Fhb5 (n = 4) | 50.0 b-e | 2.4 | 15.4 | 82.1 bc | 2.0 | 15.5 | 666.9 de | 22.9 | 22.6 |
| Fhb2 + Fhb5 (n = 6) | 47.2 d-f | 1.4 | 20.1 | 81.1 bc | 1.7 | 16.6 | 640.9 de | 14.1 | 25.6 |
| Fhb1 + Fhb2 + Fhb5 (n = 4) | 37.0 h | 3.5 | 37.4 | 70.9 c | 4.7 | 27.1 | 529.8 f | 39.2 | 38.5 |
| **CDC Go** |          |     |     |          |     |     |          |     |     |     |
| CDC Teal (susceptible check) | 72.7 a | 7.1 | - | 98.7 a-d | 7.2 | - | 977.6 a | 68.9 | - |
| CDC Go (recurrent parent) | 78.0 a | 7.1 | - | 100.0 a-d | 7.2 | - | 917.8 ab | 68.9 | - |
| AC Barrie (moderately resistant/intermediate check) | 49.1 bc | 7.1 | - | 83.3 bcd | 7.2 | - | 676.6 cd | 68.9 | - |
| ND2710 (resistant check) | 7.4 e | 7.1 | - | 15.5 f | 7.1 | - | 124.5 f | 68.9 | - |
| 04GC0139 (resistance donor parent) | 5.9 e | 7.1 | 92.4 | 7.9 f | 7.1 | 92.1 | 87.6 f | 68.9 | 90.5 |
| Null (n = 7) | 67.9 a | 2.9 | 12.9 | 99.0 a | 2.7 | 1.0 | 844.5 ab | 27.5 | 8.0 |
| Fhb1 (n = 6) | 63.4 b | 3.1 | 18.7 | 95.8 abc | 3.0 | 4.2 | 805.6 bc | 29.5 | 11.1 |
| Fhb2 (n = 4) | 68.2 a | 3.7 | 12.6 | 97.5 ab | 3.6 | 2.5 | 848.6 ab | 35.4 | 7.5 |
| Fhb5 (n = 2) | 53.1 bc | 5.1 | 31.9 | 83.2 d | 5.1 | 16.8 | 686.6 d | 49.2 | 25.2 |
| Fhb1 + Fhb2 (n = 4) | 45.8 c | 3.7 | 41.3 | 88.6 bcd | 3.6 | 11.4 | 653.8 d | 35.5 | 28.8 |
| Fhb1 + Fhb5 (n = 6) | 47.2 c | 3.1 | 39.5 | 87.0 d | 3.0 | 13.0 | 657.4 d | 29.5 | 28.4 |
| Fhb2 + Fhb5 (n = 4) | 52.4 c | 3.8 | 32.8 | 87.1 cd | 3.7 | 12.9 | 693.9 d | 36.0 | 24.4 |
| Fhb1 + Fhb2 + Fhb5 (n = 5) | 29.8 d | 3.4 | 61.8 | 67.1 e | 3.2 | 32.9 | 464.8 e | 32.0 | 49.4 |

*Least squares mean; bStandard error of the mean; cPercent disease reduction compared to recurrent parent*
accumulation (Table 4). Similarly, for CDC Go, up to 16.6% of the phenotypic variation was explained for GH_AUDPC, 18.7% for FLD_INC, 10.7% for FLD_SEV, 18.9% for FLD_IND, and 22.2% for FLD_DON accumulation. In addition to \( Fhb1 \), \( Fhb2 \), and \( Fhb5 \), four stable alleles (stable alleles in this paper are those identified in multiple environments) in CDC Alsask NILs and five in CDC Go NILs were identified that conferred resistance against FHB (Tables 5 and 6). Of the resistance improving alleles, one of the four in CDC Alsask and three of the five in CDC Go were contributed by the susceptible recurrent parent. One of the four alleles identified in CDC Alsask on chromosome 6A overlapped with QTL \( Qfhb.ndwp-6A \) reported by Zhao et al. [29]; they mapped the QTL from ND2710, an advanced breeding line from North Dakota, USA and derivative of Sumai 3. The favourable alleles at 1DS, 6AS, \( Qfhb.ndwp-6A \), 7BS loci in CDC Alsask reduced FHB index up to 23.4, 15.7, 17.2, and 17.2% and and DON accumulation by as much as 26.0, 13.7, 37.8, and 14.8%, respectively (Table 5). In CDC Go, the favourable alleles at 1DL, 2AL, 2DL, 6DS, 7AL loci reduced FHB index up to 19.0, 23.2, 16.7, 22.3, and 5.6% and DON accumulation up to 24.5, 19.4, 15.1, 18.8, and 18.8%, respectively (Table 6).

### Physical mapping and functional annotation

To determine the physical location of all genes/loci associated with FHB resistance, the corresponding SNP marker sequences were used. As expected, \( Fhb1 \) was located on the distal end of the Chromosome 3B between 8 and 21 Mb; \( Fhb2 \) between 159 and 234 Mb; \( Fhb5 \) in region between 46 and 111 Mb (Fig. 1). Other than FHB major genes, all other regions were physically located in a narrow interval with the exception of 6DS and 7AL in the CDC Go population (Table 6). Although expressed genes were retrieved for all FHB resistance governing regions, we have reported only the annotated genes for regions other than \( Fhb1 \), \( Fhb2 \), and \( Fhb5 \) (Table 7; Additional file 2). For functional annotation of expressed genes in \( Fhb1 \), \( Fhb2 \), and \( Fhb5 \) regions, readers are directed to studies by Rawat et al. [30], Dhokane et al. [31], Samad-Zamini et al. [32], and Schweiger et al. [33, 34]. Of the 33 expressed genes in the \( Fhb1 \) region, reported by Schweiger et al. [34], only six were found in the Chinese Spring database through POTAGE; whereas all six candidates for \( Fhb2 \) reported by Dhokane et al. [31] were retrieved from POTAGE (data not presented). Using POTAGE and POSEQ recombination bin carrying SNP markers associated with the Chromosome regions 1DS, \( Qfhb.ndwp-6A \), 1DL, 2AL, 2DL, 6DS, and 7AL, a total of 70, 147, 3, 85, 515, 416, and 161 expressed genes were retrieved (Additional file 2); however, in Table 7 we present only those expressed genes that were directly associated with the 90 K SNP marker sequences. The most important annotated genes, which could be the potential candidates for FHB resistance governing regions were disease resistance proteins, protein kinases (including mitogen-activated protein kinases), disease resistance nucleotide binding sites and leucine rich repeats (NBS-LRR), glutathione synthetase, dihydroflavonol 4-reductase (involved in flavonoid biosynthesis), glycosyl transferases, NAC domain containing protein and F-box domain containing proteins (Table 7; Additional file 2) as these have been reported to play a role in disease resistance (FHB and diseases of other crops) [31, 33–36].

### Discussion

In this study, we successfully introgressed Sumai 3 derived \( Fhb1 \), \( Fhb2 \), and \( Fhb5 \) genes in two elite hard red
spring wheat cultivars (CDC Go and CDC Alsask) using microsatellite markers. Although studies by Pumphrey et al. [18], McCartney et al. [21], Miedaner et al. [23], Salameh et al. [24], and Xue et al. [25] have also reported successful introgression and evaluation of Sumai 3 derived genes in elite wheat cultivars; our study has several advantages. Firstly, many of these studies did not perform repeated backcrossing and rather derived recombinant inbred lines involving multiple parents, which are expected to carry relatively larger proportion of the resistant donor, whereas we performed repeated backcrossing with implementation of markers at each BC cycle. Secondly, all the studies cited evaluated only Fhb1 and Fhb5 and ignored Fhb2, another well-characterized gene for FHB Type-II resistance [6, 31]. Lastly, we genotyped our NILs with a large number of SNP markers in addition to microsatellite markers and were able to evaluate polymorphism on all chromosomes and the marker-marker interactions based on phenotypic assessment in 8–9 environments. By repeated backcrossing, we were able to reduce the proportion of donor parent alleles to a large extent, which was even lower than the theoretically expected value of 6.25%. Similar results for introgression of four FHB resistance QTL were reported in Xue et al. [25] although their results could be biased as they used only 150 microsatellite markers. The results

| Gene/genotype                                      | FLD_INC (%) | FLD_SEV (%) | FLD_IND (%) | FLD_DON (ppm) |
|---------------------------------------------------|-------------|-------------|-------------|---------------|
|                                                   | Mean*       | SEMb        | PDRc        | Mean          | SEM          | PDR          | Mean          | SEM          | PDR          |
| CDC Alsask                                        |             |             |             |               |              |              |               |              |              |
| CDC Teal (susceptible check)                      | 78.7 a      | 4.3         | –           | 61.7 a        | 8.4          | –           | 51.1 a        | 6.9          | –           |
| CDC Alsask (recurrent parent)                     | 76.0 ab     | 4.4         | –           | 62.7 a        | 8.4          | –           | 48.8 a        | 7.0          | –           |
| AC Barrie (moderately resistant/intermediate check)| 58.6 d      | 4.7         | –           | 37.1 f        | 8.4          | –           | 22.1 d        | 7.0          | –           |
| ND2710 (resistant check)                          | 50.2 de     | 4.7         | –           | 18.8 g        | 7.0          | –           | 10.0 e        | 4.0          | –           |
| 04GC0139 (resistance donor parent)                | 48.4 e      | 4.3         | 36.3        | 25.7 g        | 7.1          | 59.0        | 12.2 e        | 4.1          | 75.0        |
| Null (n = 4)                                      | 74.0 abc    | 3.7         | 2.6         | 49.8 bc       | 7.9          | 206         | 37.8 b        | 6.8          | 225         |
| Fhb1 (n = 6)                                      | 75.6 ab     | 3.6         | 0.5         | 47.0 cd       | 7.8          | 25.0        | 36.5 bc       | 6.7          | 25.2        |
| Fhb2 (n = 4)                                      | 74.2 abc    | 3.7         | 2.4         | 51.3 b        | 7.9          | 18.2        | 39.8 b        | 6.8          | 18.4        |
| Fhb5 (n = 2)                                      | 74.8 abc    | 3.8         | 1.6         | 48.3 bc       | 7.9          | 23.0        | 36.7 bc       | 6.8          | 24.8        |
| Fhb1 + Fhb2 (n = 2)                               | 74.6 abc    | 3.9         | 1.8         | 46.1 cde      | 8.0          | 26.5        | 35.2 bc       | 6.8          | 26.4 bc     |
| Fhb1 + Fhb5 (n = 4)                               | 72.7 abc    | 3.7         | 4.3         | 47.4 bc       | 7.9          | 24.4        | 36.2 bc       | 6.8          | 25.8        |
| Fhb2 + Fhb5 (n = 4)                               | 68.3 c      | 3.6         | 10.1        | 44.6 de       | 7.8          | 28.9        | 31.3 c        | 6.8          | 35.9        |
| Fhb1 + Fhb2 + Fhb5 (n = 4)                        | 69.1 c      | 3.6         | 9.1         | 42.8 ef       | 7.9          | 31.7        | 30.6 cd       | 6.7          | 37.3        |
| CDC Go                                            |             |             |             |               |              |              |               |              |              |
| CDC Teal (susceptible check)                      | 76.7 a      | 8.0         | –           | 67.7 a        | 7.4          | –           | 51.7 a        | 7.9          | –           |
| CDC Go (recurrent parent)                         | 75.6 ab     | 8.0         | –           | 46.6 b        | 7.4          | –           | 37.2 b        | 7.9          | –           |
| AC Barrie (moderately resistant/intermediate check)| 55.1 d      | 8.1         | –           | 35.7 b        | 7.6          | –           | 21.1 d        | 8.0          | –           |
| ND2710 (resistant check)                          | 39.1 e      | 8.1         | –           | 19.4 c        | 7.6          | –           | 9.7 e         | 8.0          | –           |
| 04GC0139 (resistance donor parent)                | 37.5 e      | 8.0         | 50.4        | 19.1 c        | 7.4          | 59.0        | 7.8 e         | 7.9          | 79.0        |
| Null (n = 7)                                      | 72.9 abc    | 7.7         | 3.6         | 45.4 b        | 7.1          | 26.2        | 35.9 b        | 7.7          | 3.5         |
| Fhb1 (n = 6)                                      | 70.8 abc    | 7.7         | 6.4         | 43.6 b        | 7.1          | 64.2        | 33.9 bc       | 7.7          | 8.9         |
| Fhb2 (n = 4)                                      | 71.2 abc    | 7.7         | 5.8         | 43.0 b        | 7.2          | 7.7         | 33.4 bc       | 7.7          | 10.2        |
| Fhb5 (n = 2)                                      | 67.1 abc    | 7.8         | 11.2        | 43.4 b        | 7.3          | 7.3         | 32.8 bc       | 7.8          | 11.8        |
| Fhb1 + Fhb2 (n = 4)                               | 70.3 abc    | 7.7         | 7.0         | 40.7 b        | 7.2          | 12.7        | 31.4 bc       | 7.7          | 15.6        |
| Fhb1 + Fhb5 (n = 6)                               | 66.5 bc     | 7.7         | 12.0        | 40.8 b        | 7.1          | 12.4        | 30.6 bc       | 7.7          | 17.7        |
| Fhb2 + Fhb5 (n = 4)                               | 69.6 abc    | 7.7         | 7.9         | 42.3 b        | 7.2          | 7.9         | 32.8 bc       | 7.7          | 11.8        |
| Fhb1 + Fhb2 + Fhb5 (n = 5)                        | 65.0 cd     | 7.7         | 14.0        | 37.4 b        | 7.2          | 19.7        | 27.4 cd       | 7.7          | 26.3        |

*Least squares mean; bStandard error of the mean; cPercent disease reduction compared to recurrent parent

| Table 3 Means and standard errors for field incidence (FLD_INC), field severity (FLD_SEV), field FHB index (FLD_IND), and deoxynivalenol (FLD_DON) accumulation in gene classes and check lines for CDC Alsask and CDC Go near-isogenic lines. The data is combined over 6 environments. Means within each column for each population followed by same letter are not statistically significantly different according to Fisher’s least significant difference (LSD) at \( P = 0.05 \)
Table 4 Significant ($P = 0.001$) epistatic marker-marker interactions and percent phenotypic variation explained ($R^2$) by the interaction in CDC Alsask and CDC Go near-isogenic lines (NILs). Here AA and BB alleles are from recurrent parents CDC Go/CDC Alsask and resistance donor parent 04GC0139, respectively.

| Trait | Chromosome-Marker/loci and alleles (in parentheses) involved | $R^2$ |
|-------|----------------------------------------------------------|------|
| **CDC Alsask** | | |
| INC | 6A-Ku_c1976_663 (AA), 5B-wsnp_Ku_c12464_20125626 (AA) | 18.1 |
| | 6A-Excubiar_c1833_175 (AA), 5D-IACX6288 (AA) | 16.6 |
| | 6A-Ku_c1976_663 (AA), 5B-Excubiar_c29304_176 (AA), 5B-tplbo0027f13_1493 (AA) | 10.6 |
| | 5B-wsnp_Ku_c12464_20125626 (AA), 5B-BobWhite_c13340_412 (AA) | 5.8 |
| SEV | 5D-Excubiar_c34793_1260 (AA), 6A-RAC875_c13610_822 (BB), 6A-B500071571_51 (AA) | 24.4 |
| | Fhb1-3B-CAP7_c1576_371 (AA), 2D-Excubiar_c73791_215 (AA), 2D-IAAV8570 (AA), 2D-RAC875_c319_1776 (AA) | 17.4 |
| | 3D-Kuki_rep_c06809_457 (AA), 2D-Excubiar_c73791_215 (AA), 2D-IAAV8570 (AA), 2D-RAC875_c319_1776 (AA) | 6.8 |
| **IND** | | |
| | Fhb5-SA-B500077990_51 (BB), Fhb5-SA-Tdurum_contig10128_593 (BB), Fhb5-SA-B50007087_51 (BB), Fhb5-SA-B500065284_51 (BB), Fhb5-SA-B500075872_51 (BB), 5B-Ra_c322_1259 (BB), 5A-wsnp_Exc_c11300_18272248 (BB) | 24.8–25.9 |
| | 4B-B500022582_51 (BB), 4B-B500022582_51 (BB), 1D-RAC875_c10387_685 (AA), 1D-Kuki_c26168_713 (AA), 1D-BobWhite_c1715_887 (AA), 1D-Excubiar_c16569_532 (AA) | 20.7 |
| | 5B-Ex_c5594_2630 (AA), Fhb5-SA-B500077990_51 (BB), Fhb5-SA-Tdurum_contig10128_593 (BB), Fhb5-SA-B50007087_51 (BB), Fhb5-SA-B500065284_51 (BB), Fhb5-SA-B500075872_51 (BB), 5A-Ra_c322_1259 (BB), 5A-wsnp_Exc_c11300_18272248 (BB) | 7.8–19.7 |
| | 4A-wsnp_Exc_c14342_7887348 (BB), 2B-wsnp_Exc_c17576_26303707 (BB), 7B-CAP11_c0877_69 (AA) | 6.8–12.6 |
| | 6B-Tdurum_contig18111_179 (BB), 1A-Kuki_c23985_166 (BB), 1A-Excubiar_c75270_566 (BB), 1A-Tdurum_contig43646_147 (BB) | 12.4 |
| | 5B-Ex_c5594_2630 (AA), 1B-Tdurum_contig899_53 (BB), 1A-Tdurum_contig5560_193 (AA), 1B-Tdurum_contig42558_134 (BB) | 3.3–16.4 |
| | 5A-B500075872_51 (AB), 6B-Tdurum_contig8191_179 (AA) | 15.5 |
| | Fhb5-SA-B500077990_51 (AA), Fhb5-SA-B500077990_51 (AA), Fhb5-SA-B50007087_51 (AA), Fhb5-SA-B500065284_51 (AA), Fhb5-SA-B500065284_51 (AA), 5B-tplbo0027f13_127 (AA), 2A-wsnp_Exc_c54952_269 (BB), 3B-B500001335_51 (AA), 6D-Excubiar_c17241_388 (BB), 3A-B500036089_51 (AA), 3D-Kuki_rep_c11139_338 (BB) | 5.7–8.4 |
| | 7B-B500063852_51 (BB), 1D-B500015317_53 (BB), 1D-RAC875_c10387_685 (BB), 1A-Excubiar_c16569_532 (BB), 1D-RAC875_c10387_685 (BB) | 6.5 |
| | Fhb5-SA-B500077990_51 (AA), Fhb5-SA-Tdurum_contig10128_593 (AA), Fhb5-SA-B50007087_51 (AA), Fhb5-SA-B500065284_51 (AA), Fhb1-3B-B500063445_51 (AA), Un-B500064204_51 (AA), 1A-Kuki_c29150_143 (AA), 1D-Kuki_c26168_713 (AA), 1D-BobWhite_c1715_887 (AA), 1D-RAC875_c10387_685 (AA) | 3.2–6.2 |
| | Fhb2-6B-wsnp_Exc_c5058_8981554 (BB), 6D-Excubiar_c17241_388 (BB), 6B-Tdurum_contig42203_3670 (AA) | 6.1 |
| **CDC Go** | | |
| GH_AUDPC | Fhb5-SA-wsnp_Ra_rep_c69221_66574148 (AA), 2A-Tdurum_contig21786_270 (AA), 2D-IACX6802 (AA), 7D-RAC875_c11969_384 (AA), 4D-wsnp_ID_rep_c51623_35119179 (AA), 7A-RAC875_c22592_2255 (AA), 7D-tpblop041e14_1096 (AA) | 8.8–16.6 |
| | Fhb5-SA-wsnp_Ra_rep_c69221_66574148 (BB), Fhb5-SA-B500045284_51 (BB), 7D-RAC875_c11969_384 (AA), 4D-wsnp_ID_rep_c51623_35119179 (AA), 7D-tpblop041e14_1096 (AA) | 16.6 |
| INC | Fhb5-SA-B500045284_51 (BB), Fhb5-SA-barc1186 (BB), Fhb5-SA-B500041219_51 (BB), Fhb5-SA-barc117 (BB), Fhb5-5A-wsnp_Ra_rep_c69221_66574148 (BB), 2A-wsnp_Exc_c36481_44425685 (AA), 2D-Excubiar_c65796_394 (AA) | 11.0–18.7 |
| | Fhb5-SA-B500045284_51 (AA), Fhb5-SA-barc1186 (AA), Fhb5-SA-wmrc705 (AA), Fhb5-SA-barc117 (AA), Fhb5-5A-wsnp_Ra_rep_c69221_66574148 (AA), Fhb1-3B-RAC875_c4389_1412 (BB), 7A-RAC875_c61603_1052 (AA) | 11.9–16.6 |
| | Fhb5-SA-B500041219_51 (AA), Fhb5-SA-barc117 (AA) | 12.6 |
| | Fhb5-SA-B500041219_51 (BB), Fhb5-SA-wsnp_Ra_rep_c69221_66574148 (BB), 1A-IAAV4238 (AA), 1B-Excubiar_c35289_64 (AA), 1D-Excubiar_c26495_84 (AA), 2D-Kuki_c6176_1400 (AA) | 9.2–12.7 |
| | Fhb2-6B-Ra_c3381_1027 (AA), 2A-wsnp_Exc_c36481_44425685 (BB), 6D-B500010365_51 (AA) | 9.4 |
from our study and Xue et al. [25] indicated that the MAS is not only helpful in foreground selection of resistance genes or QTL, but to retain a major portion of the recurrent parent's genome, we can reliably quantify the allelic effects of Fhb1, Fhb2, and Fhb5 in our populations.

Theoretically, the variation in allelic composition of NILs is expected only for the chromosome carrying the gene of interest, but that is practically impossible, especially when microsatellites are used for selection that targets multiple sites in the genome of allopolyploids such as wheat. Therefore, allelic variation on all chromosomes for given SNP markers was expected. The SNP markers from the wheat 90K assay provided very useful information as they represented polymorphisms on all chromosomes and were uniformly distributed over all chromosomes [26]. A number of SNPs on 3BS (carrying Fhb1), 5AS (Fhb2), and 6BS (Fhb5), including those mapped in gene/QTL intervals were located together (physically) on the chromosome arms and inherited together as a haplotype block (Fig. 1), which could be attributed to strong linkage disequilibrium among markers. In particular, Fhb1 and Fhb5 were relatively large haplotype blocks with suppressed recombination; Fhb1 is a diverse haplotype from susceptible spring wheat lines including Chinese Spring [28, 34]. The Fhb5 gene was fine-mapped to the low recombination pericentromeric region of chromosome 5A and the SNPs in the gene interval were all mapped to the same region in our populations; polymorphism was absent for most of the chromosome region validating results of successful introgression of Fhb5. Low recombination frequencies in Fhb1 and Fhb5 regions could be another reason for the relatively large physical segments carrying exactly the same marker haplotypes on 3BS and 5AS.

Unlike most other studies where NIL/entry nested within gene/QTL class had significant variance estimates, our study indicated insignificant variation among NILs within the same QTL class (Table 1). Alternatively, all NILs carrying the same QTL behaved similarly in our populations. These results indicated that there was no or negligible recombination between the markers used for foreground selection and the gene under selection. Loss of target QTL/genes on successful backcrossing is quite possible (because of double crossover events), however, all three genes were recovered in both backgrounds, possibly by using multiple microsatellites flanking the genes at each BC cycle. Also, repeated backcrossing and a very small proportion of the resistant donor could have resulted in less confounding effects from other alleles inherited along with the three major genes under selection. Moderate to high heritability estimates for all FHB parameters suggested that a large proportion of the differences observed among the NILs has a genetic basis. Heritability estimates in GH evaluations were particularly strong, which was not surprising as the environmental variation was minimal in these cases. As expected, the 3-ADON chemotype of Fg resulted in higher FHB severity as compared to the 15-ADON chemotype in GH evaluations in both populations because the 3-ADON chemotype is known to be more aggressive
and produces more DON than the 15-ADON chemo-
type [19, 20, 37]. Despite the fact that the 3-ADON
chemotype resulted in higher disease severity, the differ-
ence between 3-ADON and 15-ADON chemotypes was
not significant 21 days post inoculation because resist-
ance to FHB in wheat was not complete and the Sumai
3 genes only slow fungal progression. With time (by 21
days after inoculation), both resistant and susceptible
spikes will exhibit FHB symptoms, particularly under con-
ductive conditions coupled with artificial inoculations.

Despite the tendency towards reduced FHB symptoms
(incidence, severity, and/or index) and DON accumula-
tion in NILs carrying Sumai 3 derived genes, it was not
significant for most of the gene classes in the CDC Go
population. This may have been due to the relatively
higher level of resistance in recurrent parent CDC Go as
compared to CDC Alsask [38]. Some level of resistance in
CDC Go compared to CDC Alsask was also evident
from the fact that CDC Go has three resistance improv-
ing alleles, whereas CDC Alsask has only one (Tables 5
and 6). Similar results on insignificant improvement in
FHB resistance in winter wheat cultivar ‘Apache’ (MR)
were reported on introgression of Fhb1 and Fhb5 [24]
and with introgression of Fhb1 in recipient lines carrying
good Type-I resistance [18]. Also, Pumphrey et al. [18]
did not detect significant differences for FHB disease se-
verity or the proportion of FDK in half of the families
contrasting for Fhb1. In practice, it is hard to combine
all favourable alleles in one genetic background, particu-
larly when both parents carry favourable alleles; there
was no NIL entry in either the CDC Go or the CDC
Alsask populations that carried all favourable alleles
from each parent. Similar to the results reported by Sal-
ameh et al. [24], NILs carrying none of three major FHB
genes (classified as ‘null’) in our study tended to improve
resistance compared to the recurrent parents and the
differences were actually significant in the CDC Alsask
NILs. The improved resistance of such NILs could be at-
tributed to some other minor favourable loci derived
from either parent (Tables 5 and 6). Our study and all
studies cited in our paper, report that even after pyra-
miding Fhb1, Fhb2, and Fhb5 in the same background,

Table 5: Single nucleotide polymorphism (SNP) markers (other than Fhb1, Fhb2, and Fhb5) associated (P < 0.05) with Fusarium head
blight index (FLD_IND), severity (FLD_SEV), incidence (FLD_INC), deoxynivalenol accumulation (FLD_DON), and area under disease
progress curve from greenhouse evaluation (GH_AUDPC) in CDC Alsask near-isogenic lines. The numbers not in parentheses
represent differences (in units for the traits) in LSmeans and the number in parentheses indicate percent disease reduction relative
to susceptible allele.

| Chr./Locus (allele)* | Physical interval (Mb) | 2011  | 2012  | 2013  | 2015  | 2016  | 2016S | Average | GH_AUDPC |
|---------------------|------------------------|-------|-------|-------|-------|-------|-------|---------|----------|
| FLD_IND             |                        |       |       |       |       |       |       |         |          |
| 1DS (BB)            | 10.39–26.18            | –     | 3.2 (7.2) | 6.9 (23.4) | –     | 3.6 (23.2) | –     | 2.0 (5.6) | 105.1 (15.9) |
| 6AS (BB)            | 3.35                   | 5.3 (8.0) | 3.8 (11.2) | 4.0 (13.6) | 7.8 (15.7) | –     | –     | 3.3 (9.1) | –        |
| Qfhb.ndwp-6A (BB)   | 602.5–611.8            | –     | –     | 3.7 (11.8) | 3.0 (17.2) | –     | –     | –        | –        |
| 7BS (AA)            | 170.52                 | 4.3 (6.4) | 3.5 (11.6) | –     | 9.1 (17.2) | –     | 3.2 (12.2) | 3.5 (9.3) | 68.9 (10.0) |
| FLD_SEV             |                        |       |       |       |       |       |       |         |          |
| 1DS (BB)            | 10.39–26.18            | –     | 2.9 (6.7) | 7.9 (17.6) | 2.4 (3.7) | 5.7 (23.3) | –     | 3.2 (6.7) | –        |
| 6AS (BB)            | 3.35                   | –     | 9.3 (20.1) | –     | 4.4 (6.8) | –     | –     | –        | –        |
| Qfhb.ndwp-6A (BB)   | 602.5–611.8            | –     | –     | 9.4 (18.3) | –     | –     | 3.3 (12.5) | –        | –        |
| 7BS (AA)            | 170.52                 | –     | 4.9 (10.5) | –     | 4.9 (7.3) | –     | 2.8 (8.4) | 2.6 (5.4) | –        |
| FLD_INC             |                        |       |       |       |       |       |       |         |          |
| 1DS (BB)            | 10.39–26.18            | –     | 3.8 (5.0) | 3.5 (5.5) | –     | –     | –     | –        | –        |
| 6AS (BB)            | 3.35                   | 5.2 (6.1) | –     | 8.3 (12.6) | 7.7 (10.2) | –     | –     | –        | –        |
| Qfhb.ndwp-6A (BB)   | 602.5–611.8            | –     | 5.9 (7.4) | –     | –     | –     | –     | –        | –        |
| 7BS (AA)            | 170.52                 | 4.1 (4.7) | –     | 7.9 (11.6) | 8.8 (11.2) | –     | –     | –        | –        |
| FLD_DON             |                        |       |       |       |       |       |       |         |          |
| 1DS (BB)            | 10.39–26.18            | 12.8 (23.8) | 9.8 (24.3) | 5.1 (23.6) | 2.6 (9.9) | –     | 1.3 (26.0) | 4.5 (18.1) | –        |
| 6AS (BB)            | 3.35                   | –     | 5.5 (13.7) | –     | 3.7 (13.7) | –     | –     | –        | –        |
| Qfhb.ndwp-6A (BB)   | 602.5–611.8            | 10.4 (17.3) | 11.4 (23.8) | 2.3 (10.1) | –     | –     | 1.4 (37.8) | –        | –        |
| 7BS (AA)            | 170.52                 | –     | 4.2 (10.1) | 3.4 (14.8) | 2.8 (10.1) | –     | –     | –        | –        |

*aHere AA and BB in parentheses indicates the CDC Alsask (recurrent susceptible parent) or 04GC0139 (resistance donor parent) alleles, respectively, that contribute resistance
*Refer to Zhao et al. (2018)
the improvement did not lead to development of any NIL or RIL as resistant as the donor parent or the resistant check. This indicates that Sumai 3 and its immediate derivatives include multiple other loci conferring FHB resistance. In our study, we identified 2–3 additional loci, derived from the resistant donor parent, but none of the loci were overlapping in both populations, which in part could explain the additional resistance in the NILs. In addition to the Sumai 3-derived chromosome regions/loci identified in our study, Anderson et al. [39] reported QTL (in addition to Fhb1, Fhb2, Fhb5) on 3AL and 6AS, and Zhou et al. [40] on 2B and 7A.

The genes expressed in the chromosome regions associated with resistance include a wide variety of proteins including disease resistance proteins, protein kinases and nucleotide-binding and leucine-rich repeat type proteins, which are most commonly associated with resistance to plant pathogens (Table 7; Additional file 2; [35]). The prediction of disease resistance proteins and kinases (highly expressed in spikes) in resistance conferring regions further validated our results and indicated their potential involvement in FHB suppression. Although genes listed in Table 7 are mostly expressed in spikes and/or grain, and are directly associated with marker sequences, these should be considered in future studies with caution because there were many other genes predicted in the regions listed in Additional file 2. The absence of genes that were predicted in the Fhb1 region were also absent from our POTAGE analyses, which could be attributed to the fact that this region was very diverse in susceptible lines in terms of gene content and size [34].

Table 6 Single nucleotide polymorphism (SNP) markers (other than Fhb1, Fhb2, and Fhb5) significantly associated (P < 0.05) with Fusarium head blight index (FLD_IND), severity (FLD_SEV), incidence (FLD_INC), deoxynivalenol accumulation (FLD_DON), and area under disease progress curve from greenhouse evaluation (GH_AUDPC) in CDC Go near-isogenic lines. The numbers not in parentheses represent differences (in units for the traits) in LSmeans and the number in parentheses indicates percent disease reduction relative to susceptible allele.

| Chr./Locus (allele) | Position (Mb) | 2010^a | 2011 | 2012 | 2013 | 2015 | 2016 | 2016S | Average | GH_AUDPC |
|---------------------|--------------|--------|------|------|------|------|------|------|---------|----------|
| FLD_IND             |              |        |      |      |      |      |      |      |         |          |
| 1DL (AA)            | 492.17–495.11| 3.3    | 4.1  | 3.6  | 3.1  | 80.1 |      |      |         |          |
| 2AL (AA)            | 717.8–718.73 | 2.8    | 4.5  | 3.9  | 2.1  | 2.2  | 100.4|      |         |          |
| 6DS (BB)            | 72.01–136.12 | –      | 6.8  | 4.6  | 4.0  | 2.6  |      |      |         | 116.7    |
| 7AL (BB)            | 519.96–619.15| –      | 3.4  | 3.6  | 2.1  |      |      |      |         |          |
| FLD_SEV             |              |        |      |      |      |      |      |      |         |          |
| 1DL (AA)            | 492.17–495.11| 4.4    | 4.2  | 5.1  | 4.5  | 2.6  |      |      |         |          |
| 2AL (AA)            | 717.8–718.73 | 4.2    | 6.7  | 5.4  | 4.8  | 2.4  |      |      |         |          |
| 6DS (BB)            | 72.01–136.12 | –      | 4.6  | 4.4  | 4.0  | 3.0  |      |      |         |          |
| 7AL (BB)            | 519.96–619.15| 5.0    | –    | –    |      |      |      |      |         |          |
| FLD_INC             |              |        |      |      |      |      |      |      |         |          |
| 1DL (AA)            | 492.17–495.11| 4.4    | 4.2  | 5.1  | 4.5  | 2.6  |      |      |         |          |
| 2AL (AA)            | 717.8–718.73 | 4.6    | 6.7  | 5.4  | 4.8  | 2.4  |      |      |         |          |
| 6DS (BB)            | 72.01–136.12 | –      | 4.6  | 4.4  | 4.0  | 3.0  |      |      |         |          |
| 7AL (BB)            | 519.96–619.15| 5.0    | –    | –    |      |      |      |      |         |          |
| FLD_DON             |              |        |      |      |      |      |      |      |         |          |
| 1DL (AA)            | 492.17–495.11| 4.4    | 4.2  | 5.1  | 4.5  | 2.6  |      |      |         |          |
| 2AL (AA)            | 717.8–718.73 | 4.6    | 6.7  | 5.4  | 4.8  | 2.4  |      |      |         |          |
| 6DS (BB)            | 72.01–136.12 | –      | 4.6  | 4.4  | 4.0  | 3.0  |      |      |         |          |
| 7AL (BB)            | 519.96–619.15| 5.0    | –    | –    |      |      |      |      |         |          |

^aHere AA and BB in parentheses indicates the CDC Go (recurrent susceptible parent) or 04GC0139 (resistance donor parent) alleles, respectively, that contribute resistance.
Table 7 List of genes (gene ID and name) annotated for single nucleotide polymorphism (SNP) loci conferring resistance to Fusarium head blight (FHB). For each annotated gene, Munich Information Center for Protein Sequences (MIPS) annotation hit ID is provided.

| Locus associated with FHB resistance | Gene ID       | Gene name                                      | MIPS annotation hit ID  | Comments                                      |
|-------------------------------------|---------------|-----------------------------------------------|-------------------------|-----------------------------------------------|
| 1DS 6AS                            | Traes_1DS_BDACE1560 | Disease resistance protein sp|Q9T048|DRL27_ARATH | Expressed in all plant parts                   |
|                                    | Traes_1DS_F3F17A72C  | Protein kinase superfamily protein AT5G28080.2 | High expression in stem and spike (Z32, Z39, Z65) |
|                                    | Traes_1DS_205D3AC88 | Disease resistance protein CC-NBS-LRR class family AT5G63020.1 | Highly expressed in spike (Z39) |
|                                    | Traes_1DS_4E3A925B9  | Leucine-rich repeat receptor-like protein kinase family protein AT4G08850.1 | Expressed in all plant parts |
|                                    | Traes_6AS_318DA417A | Protein kinase AT3G25490.1 | Only expressed in leaf (Z23, Z71), stem (Z65), and spike (Z65) |
|                                    | Qfhb.ndwp-6Aa  | F-box/RNI-like superfamily protein AT3G2692.2 | Highly expressed in spike (all stages) |
|                                    | Traes_6AL_8ASE06C77  | LRR receptor-like serine/threonine-protein kinase UniRef90_M8CZR7 | Expressed in all plant parts except root |
|                                    | Traes_6AL_1B43FE620  | Lysine-specific histone demethylase 1 homolog 3 sp|Q9CAE3|LDL3_ARATH | Highest expression in spike (Z32 and Z65) |
|                                    | Traes_6AL_8BA1FF8B2 | NAC domain containing protein 2 AT5G04410.1 | Expressed in all plant parts |
|                                    | Traes_6AL_F759812CF  | Acyl-CoA-binding domain-containing protein 4 sp|Q9MA55|ACBP4_ARATH | Expressed in all plant parts |
| 7BS                                | –             | –                                             | –                       | –                                             |
| 1DL                                | Traes_1DL_6305C4C8E | Histone deacetylase AT5G22650.1 | Highly expressed in root, stem, and spike (Z65) |
| 2AL                                | Traes_2AL_57CC28FD8  | dihydroflavonol 4-reductase AT5G42800.1 | Only expressed in grain (Z71) |
|                                    | Traes_2AL_3341560A9  | ATP binding protein UniRef90_UP0002BC9F6D | Highly expressed in grain (Z85) |
|                                    | Traes_2AL_D4EED56CE  | histone-lysine N-methyltransferase AT3G21820.1 | Highly expressed in grain (Z85) |
|                                    | Traes_2AL_B01F4C113  | polymerase delta 4 AT1G09815.1 | Highly expressed in grain (Z75) |
| 2DL                                | Traes_2DL_2249CE82  | Regulator of chromosome condensation RCC1 family protein AT3G63800.1 | Highest expression in spike (Z65) |
| 6DS                                | Traes_6DS_352313CDF | glutathione synthetase 2 AT5G27380.1 | Expressed in all parts |
|                                    | Traes_6DS_A9E719CC8  | Pentatricopeptide repeat-containing protein sp|Q9SVH0|PP329_ARATH | Highly expressed in spike (Z32) |
|                                    | Traes_6DS_E0FD61378  | Unknown sp|Q9SSB5|PRS7A_ARATH | Only expressed in spike (Z65) |
|                                    | Traes_6DS_C9398DB9C  | Synaptotagmin-5 sp|Q9MA55|METK4_HORVU | Highly expressed in grain (Z85) |
|                                    | Traes_6DS_78871A7EA  | 26S protease regulatory subunit 7 homolog A sp|Q9TW32|PPIB_DICDI | Expressed in all parts |
|                                    | Traes_6DS_762984823  | S-adenosylmethionine synthase 4 sp|P17573|LIP1_GEOCN | Highly expressed in grain (Z85) |
| 7AL                                | Traes_7AL_13DE4F55  | Lipase 1 sp|P17573|LIP1_GEOCN | Highly expressed in leaf and spike (Z65) |
|                                    | Traes_7AL_677F23CE  | F-box domain containing protein sp|Q9T048|DRL27_ARATH | Highly expressed in leaf and spike (all stages) |
|                                    | Traes_7AL_F60FF74CA  | Rel1 family protein AT4G39220.1 | Expressed in all plant parts |
|                                    | Traes_7AL_89E0BA362  | Peptidyl-prolyl cis-trans isomerase B sp|Q9TW32|PPIB_DICDI | Expressed in all plant parts |
|                                    | Traes_7AL_C501CCF17  | Similar to RCD (ribose catalytic domain) one 1 AT2G35510.1 | Highly expressed in spike and grain (all stages) |
|                                    | Traes_7AL_S30CAE15B  | mitogen-activated protein kinase ATSG19010.1 | Expressed in all plant parts |
|                                    | Traes_7AL_99483DCCC  | Mitochondrial substrate carrier AT2G46320.1 | Highly expressed in grain (Z85) |
observation from our results was that Sumai 3 genes did not show additive responses for field resistance, particularly in the CDC Go population (Table 3). The expression of Fhb5, which is considered to confer mainly Type-I resistance, was as strong as Fhb1 (Type-II resistance) in both populations, indicating that Fhb5 may also confers some level of Type-II resistance. The non-additive response of Sumai 3 derived genes or non-significant reduction even upon introgression of major genes such as Fhb1 or Fhb5 suggests epistatic or gene-gene interactions, which are often speculated, but overlooked in such studies. With the given marker density and good sample size in both populations, we were able to underpin the markers/genes involved in significant epistatic interactions in both populations that explained > 20% of the phenotypic variation of all FHB parameters. Epistatic marker-marker interactions were previously reported for some other diseases of wheat particularly for stem rust Ug99 resistance [41, 42]; however, it is worth mentioning that interactions reported by Yu et al. [41] explained less than 9% of the phenotypic variation which could be attributed to the nature of resistance in rusts (vertical/qualitative) vs FHB (horizontal/quantitative). Additionally, the role of environment in epistatic interactions and complex traits such as FHB was also significant, which is why epistatic interactions in our study accounted for a relatively large part of the total phenotypic variation [3, 16]. Frequent involvement of Sumai 3 derived genes, particularly Fhb5, in epistatic interactions also suggests their critical role in FHB resistance. Although the nature of epistasis could not be determined in our study, the significant involvement of Fhb1, Fhb2, and Fhb5 in interactions along with other loci (from both the recurrent and the donor parent) could explain the non-additive phenotypic expression in our populations and possibly other studies.

**Conclusions**

The present study has elucidated the effects of Sumai 3 introgressions on FHB disease resistance and resistance to DON accumulation. As next goal, in another study, we also utilized the set of NILs to evaluate the effect (linkage-drag) of introgressed major genes as well as minor loci on agronomic and end-use quality traits [43]. Before breeders can utilize any identified/mapped QTL or gene in their breeding program, validation using MAS is usually warranted because the effect is not always similar in all genetic backgrounds. NILs with improved resistance and phenological similarity to more advanced elite lines can easily be used for MAS in wheat breeding programs. However, the allelic effect on FHB resistance could differ depending on genetic background.
and complex epistatic interactions, thus affecting expression and penetrance of the genes in the recipient lines. Although our study suggested that improved resistance in lines carrying so-called ‘native’ resistance may not be as much as in S or MS lines, rare transgressive segregants can also be obtained from such cultivars/lines, which in turn again depends on their genetic background. In fact, Sumai 3 itself was a transgressive segregant from its parents [4]. The importance of ‘native’ resistance in local elite cultivars should not be ignored while breeding for FHB resistance in wheat.

Methods
NIL development using marker-assisted background selection
F4 populations were developed from two backcross populations CDC Go*4/04GC0139 and CDC Alsask*4/04GC0139. Line 04GC0139 (Triticum aestivum L.) was derived from Sumai 3 and has a high level of resistance to FHB, kindly provided by Dr. Julian Thomas (retired) of the Cereal Research Centre, Agriculture and Agri-Food Canada, Winnipeg, Manitoba. Line 04GC0139 has ND2710 and BW278 (pedigree: AC Domain*2/Sumai 3) in its pedigree which are both derivatives of Sumai3. The line 04GC0139 carries three well-characterized genes for resistance to FHB on chromosomes 3BS (Fhb1), 6BS (Fhb2) and 5AS (Fhb5) (G.S. Brar, unpublished data). The hard red spring wheat cultivar CDC Go (pedigree: Grandin/SD3055) is moderately susceptible (MS) to FHB, and CDC Alsask (pedigree: AC Elsa/AC Cora) is susceptible (S) (Anonymous 2015). The NILs were developed by backcrossing F1 plants to the recurrent parents (CDC Go or CDC Alsask) and the F1 at each BC cycle was screened for Fhb1, Fhb2 and Fhb5. Approximately 2100 BC2F1 and 1300 BC3F1 hybrid seeds were generated by hand crossing and 90% of the seeds were germinated and haplotyped. During the selfing process, 123 BC3F1-derived families were advanced and approximately 7000 F2 seedlings were grown, which were used to generate seed for F3 plots grown in the field in 2009. Two hundred spikes were harvested from each plot and haplotyped in the F4 generation. A total of 70 lines (38 from CDC Go and 32 from CDC Alsask) representing all eight possible combinations of FHB genes were recovered.

Microsatellite and SNP genotyping
Genomic DNA was extracted from grain and/or leaf tissue with the DNeasy 96 Plant Kit (Qiagen, Mississauga, ON). Quantification of DNA was done by fluorometry using Hoechst 33258 stain. During population development, a total of seven simple sequence repeats markers associated with Fhb1, Fhb2, or Fhb5 were screened on genomic DNA. Markers umn10 for Fhb1 on chromosome 3BS [44], gwm133 and gwm644 for Fhb2 on chromosome 6BS [7], and gwm304, barc117, wmc705 and gwm293 for Fhb5 on chromosome 5AS were used [10]. Each SSR primer pair was modified by addition of the M13 sequence to the 5’ end of the forward primer during synthesis. Fluorescent dye (either HEX, FAM or NED) was used to label the universal M13 primer. The PCR reactions consisted of 1.5 μl 10× PCR buffer, 1.5 (or 0) mM of MgCl2, 0.2 mM of each dNTP, 0.04 μM of M13 sequence-modified forward primer, 0.16 μM of reverse primer, 0.152 μM of universal dye-labelled M13 primer, 1 U of Taq DNA polymerase, and 50 ng of genomic DNA. The total PCR volume was 15 μL. Temperature cycling included 94 °C for 30 s, 56 °C (or 62 °C) for 50 s, 72 °C for 55 s, 94 °C for 30 s, 54 °C (or 60 °C) for 50 s, 72 °C for 55 s, 94 °C for 30 s, 52 °C (or 58 °C) for 50 s, 72 °C for 55 s, 94 °C for 30 s, 50 °C (or 56 °C) for 50 s, 72 °C for 55 s, then 25 cycles of 94 °C for 30 s, 51 °C for 50 s, 72 °C for 55 s, then 1 cycle of 72 °C for 10 min. Primers were first assessed for polymorphism on 2% (w/v) agarose gel stained with 0.5 μg/ml ethidium bromide, then further tested for polymorphism by capillary electrophoresis (CE) using an AB13100 Genetic Analyzer (Applied Biosystems). For CE, 1 μl of diluted PCR product (diluted 1/5, 1/10 or 1/20 in deionized water, depending on band intensity visualized on agarose gel) was combined with 9.0 μl HiDi formamide (ABI, Foster City, CA, USA) and 0.09 μl of 500 ROX size standard. Samples were run on a 36 cm capillary array, processed with Applied Biosystems Data Collection Software version 2.0, and genotyped using GeneMapper version 3.0. The presence of Fhb1 in NILs was also confirmed with the KASP assay [44, 45].

The NILs were genotyped with seven microsatellite markers while in the developmental phase in 2008–2011. To confirm the genotype of the NILs, some additional microsatellite markers (from fine-mapping studies reporting a narrow QTL interval) were used in 2017–2018 i.e. gwm493, gwm533, and functional marker for pore-forming toxin (PFT) protein for Fhb1 [6, 30], Fhb2-CAPS3 for Fhb2 [29], barc180 and barc186 for Fhb5 [28]. Additionally, the NILs were genotyped along with the parents using the wheat 90,000 iSelect assay comprised of 81,587 SNPs [26] to better understand the genomic composition and haplotype structure of the NILs. The SNP alleles were called using GenomeStudio (Illumina Inc., San Diego, CA, USA) and filtered based on polymorphisms between parents.

Greenhouse FHB evaluations
The 38 CDC Go and 32 CDC Alsask NILs, the parents and a number of check cultivars [CDC Teal as S check, AC Barrie as intermediate (l)/moderately resistant (MR)]
check, and ND2710 as resistant (R) check] were assessed for FHB symptoms in 2010 in a greenhouse (GH) equipped with incandescent lamps, 16 h photoperiod and 22/16 °C day/night temperatures. Two isolates of F. graminearum, M09–07-1, a 3-ADON chemotype (NRRL 52068) and M1–07-2, a 15-ADON chemotype (NRRL 47847), were used for inoculations [19]. At 50% anthesis, a main stem spike (two florets leaving lower two-third of the spike) on each plant was inoculated with a 10 μl macroconidial spore suspension (50,000 spores/ml) containing 0.02% Tween 20. The inoculations were performed as described in [6]. Single floret inoculation was performed to evaluate Type-II resistance. A total of three plants per replication were inoculated and there were three replications in total. The FHB severity was rated as the percentage of infected spikelets per spike at 7 (GH7), 14 (GH14) and 21 days (GH21) post inoculation. Area under disease progress curve (GH_AUDPC), used as a measure of FHB severity over time, was calculated according to [46].

Field FHB evaluations
The same NILs, parents and check cultivars evaluated in the greenhouse were also assessed in the field nursery at Carman, MB from 2010 to 2013 and 2016, Saskatoon, SK in 2016, and at Morden, MB in 2015. In 2010, the CDC Alsask population was not evaluated in the field because the seed under multiplication in the greenhouse was not ready for field planting. The field trial was set up as a randomized complete block design with two replicates in Carman and four replicates in Morden and Saskatoon. Plots at Morden and Carman consisted of single 1.5 m and 1 m rows, respectively, and in the Saskatoon nursery in hills. Sowing density was approximately 80 seeds per row and 30 seeds per hill. At Carman, every plot in the nursery was artificially inoculated with a suspension of F. graminearum macro-conidia prepared with the isolates M9–07-1 (3-ADON), M7–07-1 (3-ADON), M1–07-1 (15-ADON) and M3–07-2 (15-ADON). The isolates used were originally provided by Dr. Jeannie Gilbert at the CRC-AAFC. Isolates were cultured in Spezieller Nährstoffarmer Agar (SNA) for seven days and then incubated in Carboxymethyl Cellulose (CMC) media for another seven to ten days. The number of spores was counted to calculate their concentration. Prior to field application, the suspension of the four isolates was mixed in equal proportions (based on macro-conidia concentration) to provide a total concentration of 50,000 macro-conidia spores/ml. The field application was achieved using a CO2 backpack sprayer and directed to the wheat spikes at flowering (anthesis) stage. A second application was performed to the same rows three days later. After each inoculation, plots were mist irrigated overnight. Visual assessments of disease incidence (% of infected spikes in the plot) and severity (% of spikelets infected on the infected spikes) were made on each plot 18–21 days after the first inoculation. Fusarium head blight index for each plot was calculated as follows: (disease incidence x disease severity)/100. At Morden, MB and Saskatoon, SK, irrigated nurseries were inoculated with air-dried corn spawn (colonized by F. graminearum) at 50% anthesis. Each plot was assessed using an FHB index (%incidence x %severity/100) (FLD_IND) based on disease incidence (%) (FLD_INC) and severity (%) (FLD_SEV) at 21 to 23 dpi. Cultivars CDC Teal (S), AC Barrie (moderately resistant, MR), and ND2710 (resistant, R) were included as checks in Morden and Saskatoon. Cultivars AC Vista, and CDC Teal were used as S check, AC Cora as I check, 5602HR as MR, and FHB37 as R check in Carman nursery. Up to 50 spikes of each NIL were harvested by hand and retained for DON quantification (FLD_DON).

DON quantification
The spikes of each NIL were harvested from two replicates at the fully ripe stage (BBCH 92; [47]) and dried to minimal water content. Approximately 50–100 g samples of each NIL were ground to a fine powder with a laboratory mill and stored at −20 °C until further processing. Analysis of DON was carried out using ELISA based assays [48] and a Neogen commercial kit. Measurements were performed in two technical replications of each biological replication. Detailed information on the Neogen ELISA assay are available in Additional file 1 (Protocol #1).

Physical mapping and functional annotation
All SNP markers from the wheat 90 K assay were physically positioned on the Chinese Spring wheat reference genome sequence. The SNP-bearing sequences were probed to the entire bread wheat NRGene wheat genome assembly RefSeq ver. 1.0 (International Wheat Genome Sequencing Consortium, https://wheat-urgi.versailles.inra.fr/Seq-Repository/Assemblies) using an in-house BLAST portal. The best hits, based on sequence similarity and cumulative alignment length percentage of matches, were considered. For annotation, the wheat genome scaffolds carrying the marker were retrieved from the BLAST searches and used to find genes expressed on the scaffolds using POTAGE (PopSeq Ordered Triticum aestivum Gene Expression) [49]. POTAGE integrates map location with gene expression, infers functional annotation and visualizes these data through a web browser interface. The map location (implemented in POTAGE) were based on the wheat POPSEQ map of the 90 double haploid individuals of the synthetic W7984 X Opata M85 population, where SNP markers are anchored to contigs in linear order [50].
**SNP data and marker-marker epistatic interaction analyses**

For haplotype analyses and to assign each NIL entry to a QTL class, SNP markers tightly linked to SSR markers or mapped to Fhb1, Fhb2, and Fhb5 regions were considered [28,29; Ron Knox, unpublished data]. The SNP markers flanking the Fhb1 region of Carberry were provided by Dr. Ron Knox (AAFC, Swift Current) and were also mapped in ND2710 by Zhao et al. [29]. The introgressed haplotypes from the resistant donor parent were visualized using Graphical Genotypes software ver. 2.0 [51]. To analyze the phenotypic data as influenced by all polymorphic markers, genotypic and phenotypic data were used to test for epistatic interactions. Epistatic interactions between markers with significant main effects (Fhb1, Fhb2, and Fhb5) were tested as well as all other markers regardless of significance. A linear regression model was used to calculate P-values for pairwise as well as multiple marker-marker interactions using an in-house designed script in the R environment [52]. A false discovery rate of 0.05 was used as a threshold for significant interactions. Epistatic interactions were analyzed according to [53] and modeled as follows:

\[
y = \mu + Z_1y_1 + (Z_i \times Z_j)y_{ij} + e
\]

Where: \( y \) is the \( n \times 1 \) vector for phenotypic observation, \( \mu \) is the population mean, \( Z_1 \) is a vector (\( Z_1, ..., Z_m \))\(^T\), for the genotype indicators of locus \( l \), \( Z_{ij} \) takes one of two values (\(-1, +1\)) depending on which parental allele was passed on to line \( i \), for locus \( l \), \( y_1 \) is the additive (main) effect of locus \( l \), \( y_{ij} \) is the epistatic effect between loci \( l \) and \( j \), and \( e \) is the residual error vector.

From each chromosome, one marker from each group of redundant/co-segregating markers was chosen for the epistatic interaction analyses. Any marker-marker interaction for a given phenotype was declared significant at \( P = 0.001 \).

**Statistical and phenotypic data analyses**

The phenotypic data collected from field and greenhouse evaluations was subjected to correlation and analysis of variance (ANOVA). Before conducting ANOVA, assumptions of independence, normal distribution and homogeneity of residuals for all class variables were verified using Shapiro-Wilk and Levene’s tests implemented in procedure UNIVARIATE in SAS (Statistical Analytical Software) ver. 9.4 (SAS Institute, Inc., Cary, NC). Heterogeneous variances, if any, were modeled using the ‘repeated/group = effect’ statement in procedure MIXED [54]. Variance component estimates and corresponding \( F \)-values were calculated using the procedure MIXED in SAS ver. 9.4 with the ‘ddfm = kenwardroger’ option to approximate degrees of freedom. Mean separation was conducted using the least significant difference (LSD) test (Fisher’s least significant difference). All tests used a nominal alpha level of 0.05. Broad-sense heritability (\( H^2 \)) was calculated as described in [55]. Pearson’s correlation coefficients among various parameters were calculated using procedure CORR in SAS. Associations among environments, genotypes, and the genotype by environment interaction were analyzed and visualized using biplot analyses [56] in the R environment using the GGEbiplotGUI package [57]. For biplot analyses, the following settings were used: singular value portioning, environment-metric preserving; and genotype by environment scaling, according to the standard deviation; centered by environment (\( G + G*E \)).

**Additional files**

**Additional file 1:** Protocol #1: Procedure for Neogen enzyme linked immune-sorbent assay (ELISA) for deoxynivalenol (DON) quantification in Fusarium head blight infected grains. **Table S1.** Proportions of the recurrent parent (RP) and donor parent (DP) genomes in the near-isogenic lines for CDC Go and CDC Alsask streams based on 81,587 SNP markers from 90K Select assay. Here: A, B, H, U represent recurrent parent, donor parent, heterozygous, and unknown alleles, respectively. **Figure S1.** Polymorphism in CDC Go and CDC Alsask near-isogenic lines (NILs). **Figure S2.** Polymorphism in CDC Alsask near-isogenic lines on Chromosomes other than 3B, 5A, 6B. **Figure S3.** Polymorphism in CDC Go near-isogenic lines on Chromosomes other than 3B, 5A, 6B. **Figure S4.** GGE Biplots for CDC Go and CDC Alsask near-isogenic lines (NILs).

**Additional file 2:** List of annotated genes in the intervals of FHB resistance loci (XLSX 429 kb)

**Abbreviations**

AUDPC: Area under disease progress curve; CDC: Crop Development Centre; DON: Deoxynivalenol; ELISA: Enzyme linked immunosorbant assay; FG: Fusarium graminearum; FHB: Fusarium head blight; GH: Greenhouse; INC: Incidence; IND: Index; MAS: Marker-assisted selection; MR: Moderately resistant; MS: Moderately susceptible; NIL: Near-isogenic line; POTAGE: PopSeq Ordered Triticum aestivum Gene Expression; QTL: Quantitative trait loci; S: Susceptible; SEV: Severity; SNP: Single nucleotide polymorphism; SSR: Simple sequence repeat

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**Availability of data and materials**

All data pertinent to this publication is available within article or online as Additional file.

**Authors’ contributions**

PJH, CJP, and GSB conceived the research project. PJH, CJP, and HRK supervised and coordinated the research. YR performed greenhouse experiments and marker-assisted selection of NILs. ABB phenotyped NILs from 2010 to 2013, and in 2016 at Carman, MB. GSB performed phenotyping in 2016 at Saskatoon, SK. MAH phenotyped the material in 2015 at Morden,
MB. GSB collected data on DON accumulation, analyzed data and wrote manuscript. All authors read and approved the manuscript.

Ethics approval and consent to participate
The experiments described in the manuscript were conducted according to ethical standards of Canada.

Consent for publication
Not applicable.

Competing interests
The authors declare that they do not have any competing interests.

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References
1. McMullen M, Bergstrom G, de Wolf E, Dill-Macky R, Hershmann D, Shaner G, van Sanford D. A unified effort to fight an enemy of wheat and barley: Fusarium head blight. Plant Dis. 2012;96:1712–28.
2. Gilbert J, Haber S. Overview of some recent research developments in Fusarium head blight of wheat. Can J Plant Pathol. 2013;35:149–74.
3. Buertmayr H, Ban T, Anderson JA. QTL mapping and marker-assisted selection for Fusarium head blight resistance in wheat: a review. Plant Breed. 2009;128:1–26.
4. Bai G, Su Z, Cai J. Wheat resistance to Fusarium head blight. Can J Plant Pathol. 2018; [Published online: https://doi.org/10.1007/s00035-018-01476-11].
5. Beavis WD. QTL analyses: power, precision, and accuracy. In: Paterson AH, editor. Molecular dissection of complex traits. CRC press. Boca Raton; 1998. p. 145–62.
6. Cuthbert PA, Somers DJ, Thomas J, Cloutier S, Brûlé-Babel A. Fine mapping Fhb1, a major gene controlling Fusarium head blight resistance in bread wheat (Triticum aestivum L.). Theor Appl Genet. 2006;112:1465–72.
7. Cuthbert PA, Somers DJ, Brûlé-Babel A. Mapping of Fhb2 on chromosome 6BS: a gene controlling Fusarium head blight field resistance in bread wheat (Triticum aestivum L.). Theor Appl Genet. 2007;114:429–39.
8. Qi L, Pumphrey M, Friebie B, Chen P, Gill B. Molecular cytogenetic characterization of alien introgressions with gene Fhb3 for resistance to Fusarium head blight disease of wheat. Theor Appl Genet. 2008;117:155–66.
9. Xue S, Li G, Jia H, Xu F, Lin F, Tang M, Wang Y, An X, Xu H, Zhang L, Kong Z, Ma Z. Fine mapping Fhb4, a major QTL conditioning resistance to Fusarium infection in bread wheat (Triticum aestivum L.). Theor Appl Genet. 2010;121:147–56.
10. Xue S, Xu F, Tang M, Zhou Y, Li G, An X, Lin F, Xu H, Jia H, Zhang L, Kong Z, Ma Z. Precise mapping Fhb5, a major QTL conditioning resistance to Fusarium infection in bread wheat (Triticum aestivum L.). Theor Appl Genet. 2011;123:1055–63.
11. Cao J, Bockus JW, Feng Y, Chen P, Qi L, Sehgal SK, Danilova TV, Koo DH, Friebie B, Gill BS. Chromosome engineering, mapping, and transferring of resistance to Fusarium head blight disease from Elymus trichodes to wheat (Triticum aestivum L.). Theor Appl Genet. 2015;128:1019–27.
12. Guo J, Zhang X, Hou Y, Cai J, Shen X, Zhou T, Xu H, Ohm HW, Wang L, Li A, Han F, Wang H, Kong L. High-density mapping of the major Fhb8 resistance gene Fhb8 derived from Thinopyrum ponticum and its pyramiding with Fhb1 by marker-assisted selection. Theor Appl Genet. 2015;128:2301–16.
13. Schroeder H, Christensen J. Factors affecting resistance of wheat to scab caused by Gibberella zeae. Phytopathology. 1963;53:831–8.
14. Miller JD, Young JC, Sampson RD. Deoxynivalenol and Fusarium head blight resistance in spring cereals. Phytopathology. 1985;113:359–67.
15. Mesterhazy A, Bartok T, Mirocha CG, Komorocy AR. Nature of wheat resistance to Fusarium head blight and the role of deoxynivalenol for breeding. Plant Breed. 1999;118:97–110.
16. Bernardi R. Molecular markers and selection for complex traits in plants: learning from the last 20 years. Crop Sci. 2008;48:1649–64.
17. Kaeppeler SM, Phillips RL, Kim TS. Used of near-isogenic lines derived by backcrossing or selfing to map qualitative traits. Theor Appl Genet. 1993;87:233–7.
18. Pumphrey MO, Bernardi R, Anderson JA. Validating the Fhb1 QTL for Fusarium head blight resistance in near-isogenic wheat lines developed from breeding populations. Crop Sci. 2007;47:200–6.
19. Gilbert J, Brûlé-Babel A, Guerrieri AT, Clear RM, Patrick S, Slusarenko K, Wolfe C. Ratio of 3-ADON and 15-ADON isolates of Fusarium graminearum recovered from wheat kernels in Manitoba from 2008 to 2012. Can J Plant Pathol. 2014;36:54–63.
20. Ward TJ, Clear RM, Rooney AP, D’Ollenn K, Gaba D, Patrick S, Starksy DE, Gilbert J, Geiser DM, Nowicki TW. An adaptive evolutionary shift in Fusarium head blight pathogen populations is driving the rapid spread of more toxigenic Fusarium graminearum in North America. Fungal Genet Biol. 2008;45:473–84.
21. McCartney CA, Somers DJ, Fedak G, DePauw RW, Thomas J, Fox SL, Humphreys DG, Lukow O, Savard ME, McCallum BD, Cao W. The evaluation of FHB resistance QTLs introgressed into elite Canadian spring wheat germplasm. Mol Breeding. 2007;20:209–21.
22. Balut AL, Clark AJ, Brown-Guedria G, Souza E, Van Sanford DA. Validation of Fhb1 and Qfhs.nau-2DL in several soft red winter wheat populations. Crop Sci. 2013;53:934–45.
23. Miedaner T, Wilde F, Steiner B, Buerstmayr H, Korzun V, Ebmeyer E. Stacking quantitative trait loci (QTL) for Fusarium head blight resistance from non-adapted sources in an European elite spring wheat background and assessing their effects on deoxynivalenol (DON) content and disease severity. Theor Appl Genet. 2006;112:562–9.
24. Salameh A, Buerstmayr M, Steiner B, Neumayer A, Lemmens M, Buerstmayr H. Effects of introgression of two QTL for Fusarium head blight resistance from Asian spring wheat by marker-assisted backcrossing into European winter wheat on fusarium head blight resistance, yield and quality traits. Mol Breeding. 2011;28:485–94.
25. Xue S, Li G, Jia H, Lin F, Cao Y, Xu F, Tang M, Wang Y, Wu X, Zhang Z, Zhang L, Kong Z, Ma Z. Marker-assisted development and evaluation of near-isogenic lines for scab resistance QTLs of wheat. Mol Breeding. 2010;25:397–405.
26. Wang S, Wong D, Forrest K, et al. Characterization of polyplid wheat genome diversity using a high-density 90 000 single nucleotide polymorphism array. Plant Biotechnol J. 2014;12:787–96.
27. Macferri M, Ricci A, Cattiveli L, Buerstmayr H. High-resolution mapping of the pericentromeric region on wheat chromosome arm 5AS harbouring the Fusarium head blight resistance QTL Qfhs.ifa-5A. Plant Biotechnol J. 2013;11:648–63.
28. Buerstmayr M, Steiner B, Wagner C, Schwarz P, Brugger K, Barabash D, Volante A, Vale G, Cattivelli L, Buerstmayr H. High-resolution mapping of the FHB resistance QTL on chromosome 5AS harbouring the Fusarium head blight resistance QTL Qfhs.ifa-5A. Plant Biotechnol J. 2011;13:1046–56.
29. Zhao M, Wang G, Leng Y, Wang Hui X, Pi X, Grosz M, Mengouo M, Zhong S. Molecular Mapping of Fusarium head blight resistance in the spring wheat variety ND2710. Phytopathology. 2018;108:972–9.
30. Rawat N, Pumphrey MO, Liu S, Zhang X, Tiwari VK, Ando K, TrickHN, Bockus WW, Akhunov E, Anderson JA, Gill BS. Wheat Fhb1 encodes a chimeric lectin with agglutinin domains and a pore-forming toxin-like domain conferring resistance to Fusarium head blight. Nat Genet. 2016;48:1576–80.
31. Dhokane D, Karre S, Kushalappa AC, McCartney C. Integrated metabolon-transcriptomics reveals Fusarium head blight candidate resistance genes in wheat QTL-Fhb2. PLoS One. 2016;11:e0155881.
32. Samad-Zaamin M, Schweiger W, Nussbaum T, Mayer KFX, Buerstmayr H. Time-course expression QTL-atlas of the global transcriptional response of wheat to Fusarium graminearum infection. Plant Biotechnol J. 2017;15:1455–64.
33. Schweiger W, Steiner B, Ametz C, Siegwart G, Wiesenberger G, Berthiller F, Lemmens M, Jia H, Adam G, Muehlbauer G, Kreil DP, Buerstmayr H. Transcriptomic characterization of two major Fusarium resistance quantitative trait loci (QTLs), Fhb1 and Qfhs.nau-6A, identifies novel candidate genes. Mol Plant Pathol. 2013;14:772–85.
34. Schweiger W, Steiner B, Vautrin S, Nussbaumer T, Siegwart G, Zamini M, et al. Suppressed recombination and unique candidate genes in the divergent haplotype encoding Fhb1, a major Fusarium head blight resistance locus in wheat. Theor Appl Genet. 2016;129:1607–23.
35. Bent AF, Mackey D. Elicitors, effectors, and R genes: the new paradigm and a lifetime supply of questions. Annu Rev Phytopathol. 2007;45:399–436.
36. Deokar A, Sagi M, Daba K, Tar'an B. QTL sequencing strategy to map genomic regions associated with resistance to ascochyta blight in chickpea. Plant Biotechnol J. 2018;1–14. https://doi.org/10.1111/pbi.12964.
37. Gilbert J, Clear RM, Ward TJ, Gaba D, Tekauz A, Turkington TK, Woods SM, Novicki T, O'Donnell K. Relative aggressiveness and production of 3- or 15-acetyl deoxynivalenol and deoxynivalenol by Fusarium graminearum in spring wheat. Can J Plant Pathol. 2010;32:146–52.
38. Anonymous. Varieties of grain crops 2015. In: 2015 Guide. Regina: Saskatchewan Seed Growers Association; 2015.
39. Anderson JA, Stack RW, Liu S, Waldron BL, Fjeld AD, Coyne C, Moreno-Sevilla B, Fetch JM, Song QJ, Cregan PB, Frohberg RC. DNA markers for Fusarium head blight resistance QTLs in two wheat populations. Theor Appl Genet. 2001;102:164–8.
40. Zou WC, Kolb FL, Bai GH, Dornier LL, Yao JB. Effect of individual Sumai 3 chromosomes on resistance to scab spread within spikes and deoxynivalenol accumulation within kernels in wheat. Hereditas. 2002;137:81–9.
41. Yu L-X, Lorenz A, Rutkoski J, Singh RP, Bhavani S, Huerta-Espino J, Sorrells ME. Association mapping and gene-gene interaction for stem rust resistance in CIMMYT spring wheat germplasm. Theor Appl Genet. 2011;123:1257–68.
42. Yu L-X, Morgounov A, Wanyera R, Kees M, Singh SK, Sorrells M. Identification of Ug99 stem rust resistance loci in winter wheat germplasm using genome-wide association analysis. Theor Appl Genet. 2012;125:749–58.
43. Brar GS, Pozniak CJ, Kitcher HR, Hulc PJ. Evaluation of Fusarium head blight resistance genes Fhb1, Fhb2, and Fhb5 introgressed into elite Canadian hard red spring wheats: effect on agronomic and end-use quality traits and implications for breeding. Mol Breeding. 2019;39:44. https://doi.org/10.1007/s11032-019-0957-8.
44. Liu S, Pumphrey MO, Gill BS, Trick H, Zhang JX, Chalhoub B, Anderson JA. Towards positional cloning of Fhb1, a major QTL for Fusarium head blight resistance in wheat. Cereal Res Comm. 2008;36:195–201.
45. Rasheed A, Wen WE, Gao FM, Zhai SN, Jin H, Liu JD, Guo Q, Zhang YJ, Dressigacker S, Xia XC, He ZH. Development and validation of KASP assays for genes underpinning key economic traits in bread wheat. Theor Appl Genet. 2016;129:1843–60.
46. Buerstmayr H, Steiner B, Lemmens M, Ruckenbauer P. Resistance to Fusarium head blight in winter wheat: heritability and trait associations. Crop Sci. 2000;40:1012–8.
47. Lancashire PD, Bleiholder H, Langeluddecke P, Stauss R, van den Boom T, Weber E, Witzenberger A. A uniform decimal code for growth stages of crops and weeds. Ann Appl Biol. 1991;119:561–601.
48. Sinha RC, Savard ME, Lau R. Production of monoclonal antibodies for the specific detection of deoxynivalenol and 15-acetyldeoxynivalenol by ELISA. J Agric Food Chem. 1995;43:1740–4.
49. Suchecki R, Watson-Haigh NS, Baumann U. POTAGE: a visualization tool for speeding up gene discovery in wheat. Sci Rep. 2017;7:14315.
50. Chapman JA, Mascher M, Bulac A, et al. A whole-genome shotgun approach for assembling and anchoring the hexaploid bread wheat genome. Genome Biol. 2015;16:26.
51. Van Berloo R. GGT 2.0: versatile software for visualization and analysis of genetic data. J Heredity. 2008;99:232–6.
52. Chapman JA, Mascher M, Bulac A, et al. A whole-genome shotgun approach for assembling and anchoring the hexaploid bread wheat genome. Genome Biol. 2015;16:26.
53. Xu S, Jia Z. Genomewide analysis of epistatic effects for quantitative traits in barley. Genetics. 2007;175:1955–63.
54. Friend CL, Milliken GA, Stoup WW, Wolflinger R, Schabenberger O. SAS for mixed models. 2nd ed. Cary: SAS Institute Inc.; 2006.
55. Brar GS, Fuentes-Davilla G, He X, Sansaloni CP, Singh RP, Singh PK. Genetic mapping of resistance in hexaploid wheat for a quarantine disease: Karnal bunt. Front Plant Sci. 2019;10:897. https://doi.org/10.3389/fpls.2018.01497.
56. Yan W, Tinker NA. Biplot analysis of multi-environment trial data: principles and applications. Can J Plant Sci. 2006;86:623–45.
57. Frutos E, Pascual Galindo M, Leiva V. An interactive biplot implementation in R for modeling genotype-by-environment interaction. Stoch Environ Res Risk Assess. 2014;28:1629–41.