Genome-wide association study and genomic prediction of white rust resistance in USDA GRIN spinach germplasm

Ainong Shi1,*, Gehendra Bhattarai1, Haizheng Xiong1, Carlos A. Avila2,*, Chunda Feng3, Bo Liu3, Vijay Joshi4, Larry Stein4,*, Beiquan Mou5,*, Lindsey J. du Toit6, and James C. Correll3,*

1Department of Horticulture, University of Arkansas, Fayetteville, AR 72701, USA
2Department of Horticultural Sciences, Texas A&M AgriLife Research and Extension Center, Weslaco, TX 78596, USA
3Department of Plant Pathology, University of Arkansas, Fayetteville, AR 72701, USA
4Texas A&M AgriLife Research and Extension Center, Uvalde, TX 78801, USA
5Crop Improvement and Protection Research Unit, USDA-ARS, Salinas, CA 93905, USA
6Washington State University, Mount Vernon, WA 98273, USA
*Corresponding authors. E-mail: ashi@uark.edu; Carlos.Avila@ag.tamu.edu; larrystein@tamu.edu; beiquan.mou@usda.gov; dutoit@wsu.edu, jcorrell@uark.edu

Abstract
White rust, caused by Albugo occidentalis, is one of the major yield-limiting diseases of spinach (Spinacia oleracea) in some major commercial production areas, particularly in southern Texas in the United States. The use of host resistance is the most economical and environment-friendly approach to managing white rust in spinach production. The objectives of this study were to conduct a genome-wide associating study (GWAS), to identify single nucleotide polymorphism (SNP) markers associated with white rust resistance in spinach, and to perform genomic prediction (GP) to estimate the prediction accuracy (PA). A GWAS panel of 346 USDA (US Dept. of Agriculture) germplasm accessions was phenotyped for white rust resistance under field conditions and GWAS was performed using 13,235 whole-genome resequencing (WGR) generated SNPs. Nine SNPs, chr2_53049132, chr3_58479501, chr3_95114909, chr4_9176069, chr4_17807168, chr4_83938338, chr4_87601768, chr6_1877096, and chr6_31287118, located on chromosomes 2, 3, 4, and 6 were associated with white rust resistance in this GWAS panel. Four scenarios were tested for PA using Pearson’s correlation coefficient (r) between the genomic estimation breeding value (GEBV) and the observed values: (1) different ratios between the training set and testing set (fold), (2) different GP models, (3) different SNP numbers in three different SNP sets, and (4) the use of GWAS-derived significant SNP markers. The results indicated that a 2- to 10-fold difference in the various GP models had similar, although not identical, averaged r values in each SNP set; using GWAS-derived significant SNP markers would increase PA with a high r-value up to 0.84. The SNP markers and the high PA can provide valuable information for breeders to improve spinach by marker-assisted selection (MAS) and genomic selection (GS).

Introduction
Spinach (Spinacia oleracea L.) is one of the important vegetable crops in the world economically, which was estimated to average $490 million (fresh and for processing) per year during 2018–20 in the United States (US), with 97% of the value for the fresh market [1]. Spinach is considered a “super food” due to a high concentration of phytonutrients and other health-promoting compounds, including vitamin A and C, carotenoid, lutein, folate, calcium, iron, and antioxidants [2, 3]. In the US, spinach has become very popular during the past two decades as healthy-conscious consumers have increased the consumption of leafy vegetables. To meet the greater demand for spinach, commercial production has evolved into high density (up to 10 million/ha) planting, year-round production cycles, overhead sprinkler irrigation systems, high fertilizer application, and expanded production areas; all of them create an environment conducive for the development of diseases. Several diseases reduce yield and quality individually or in combination, thus posing serious challenges to commercial spinach production. Spinach suffers from many diseases while downy mildew, white rust, Fusarium wilt, Stemphylium leaf spot, and anthracnose leaf spot are the most devastating and economically important diseases of spinach.

White rust of spinach is caused by Albugo occidentalis, an obligate oomycete that can reduce yield and quality [3–8]. White rust has been an endemic throughout the central and eastern US for many years but has also been reported in other parts of the world, including Greece [9], Mexico [10], and Turkey [11]. The persistent appearance...
of this disease and its expansion into wider geographic areas pose a significant challenge to the spinach industry in the US and world. If this pathogen is introduced into major US production areas in California and Arizona that produce over 90% of the fresh market product, it would be devastating to the US spinach industry as the vast majority of the cultivars adapted to these areas have little to no resistance incorporated. Resistance to white rust has been previously found in USDA spinach germplasm and breeding lines [4, 7, 12–14]. High levels of resistance to white rust have been reported in the spinach varieties developed at the spinach breeding program of University of Arkansas as this disease was a primary breeding goal [3]; thus, the Arkansas germplasm has been used as a source of resistance to transfer white rust resistance into several commercial cultivars. However, even this resistance material can suffer severe infection when conditions are highly conducive for the disease development [7, 12–14]. Yet, there is little information regarding the genetics of white rust resistance in spinach. Thus it is necessary to routinely evaluate and identify new spinach resources for white rust resistance for the development of cultivars with improved resistance. So far, only quantitative resistance has been found and utilized as no major genes have been reported for white rust resistance [5].

The publication of reference genomes and other new assemblies [15–19] has made genome-wide variant discovery in the germplasm panel and genome-wide association studies (GWAS) possible in spinach. With the decreasing genotyping cost in recent years and advanced statistical methods, GWAS and genomic selection (GS) approaches are commonly utilized to improve complex genetic traits in crops. GWAS, based on the genotyping and phenotyping of a natural germplasm population and high-density markers, has been employed to map simple to complex traits and identify candidate genes in many crops [20–22]. GWAS has been used in spinach for many traits, including surface texture, edge shape, and petiole color [23]; bolting, tallness, and erectness [24]; leafminer resistance [25]; oxalate concentration [26]; Verticillium wilt resistance [27], Stemphylium leaf spot resistance [28]; mineral nutrient contents [29]; white rust resistance [30]; growth habit [31]; anthracnose resistance [32]; and downy mildew resistance [33–36]. The identification of single nucleotide polymorphism (SNP) markers for the traits has provided valuable molecular tools for breeders to develop spinach cultivars more efficiently. It has been demonstrated that quantitative trait of white rust resistance in spinach can be screened under field conditions and is correlated with quantitative resistance to downy mildew [4, 37]. Using a panel of 267 spinach accessions with 6111 SNPs, Awika et al. (2019) [30] conducted GWAS analysis for white rust resistance and identified 448 minor alleles (SNPs) associated with white rust disease severity, which maybe be utilized in the selection for resistant plants.

Genomic prediction (GP) is emerging as a promising tool to improve the efficiency and speed of plant breeding. So far, GP has been reported in several crops [38–48] for various traits. Genomic estimation breeding value (GEBV) is a key step in GS. Several approaches have been proposed to determine GEBV, such as Bayesian methods (Bayes A, Bayes B, Bayes LASSO, and Bayes ridge regression) and BLUP methods (RR-BLUP, gBLUP, and cBLUP). The GS approaches have been adopted for variety of traits in various crops [49–55]. All articles reported the prediction accuracy (PA) estimates using the Pearson’s correlation coefficient (r) between the observed phenotypic values and the predicted GEBV for each trait in a validation set using several different models.

Currently, USDA-GRIN (Germplasm Resources Information Network) has approximately 400 spinach accessions, which were originally collected from 33 countries and represent a diverse germplasm collection. The overall objectives of this study were to evaluate USDA spinach germplasm for white rust resistance under the field conditions and to identify resistance-associated SNP markers through GWAS to conduct marker-assisted selection and genomic selection for white rust resistance.

Materials and methods
Plant materials (genome-wide association panel)
Three hundred forty-six spinach accessions of the USDA-GRIN spinach germplasm collection were phenotyped for white rust disease and genotyped by using whole-genome resequencing in this study (Table S1 where S signifies supplementary). The accessions in this GWAS panel were originally collected from 32 countries, with a majority (81.5%) from ten countries: Turkey (n = 107), United States (US) (n = 55), China (n = 25), North Macedonia (n = 22), Afghanistan (n = 20), Iran (n = 15), Belgium (n = 10), India (n = 10), Syria (n = 10), and Hungary (n = 8) (Table S1).

White rust phenotyping
The 346 spinach accessions were evaluated for white rust resistance in the Del Monte White Rust Nursery, Crystal City, Texas, during the three winter seasons of 2015–16, 2016–17, and 2017–18. The nursery is known to have high white rust disease pressure over the three decades. The field experiments were performed in a randomized complete block (RCB) design with two replications. In each block, each accession was planted in a 10-feet long row, three feet between rows, and 4-inch between plants within the row. Thus, there were about 30 plants in each row with 60 plants of each accession for evaluation each year. White rust disease was evaluated under natural disease pressure without introducing external inoculum. A susceptible cultivar, Viroflay, was planted as a spreader row on both sides of the tested genotypes.

White rust disease was rated using a scale of 0 to 10 whereby a 0 = no disease, 1 = <1% of the total leaf area covered with white rust infection, 2 = 1–10%, 3 = 11–20%, 4 = 21–30%, 5 = 31–40%, 6 = 41–50%, 7 = 51–60%, 8 = 61–70%, 9 = 71–80%, and 10 = > 80% infected leaf area. After...
65–70 days from planting, around ten plants of each genotype were scored for disease severity by estimating the proportion of total leaf area canopy with symptoms (chlorotic and necrotic lesions) and signs (sporulation and pustules). The white rust disease severity was recorded 2–3 times each season.

The 2017–2018 winter season trial had high white rust disease pressure among the three years of evaluations, while the disease severity was lower in the other two years (data not shown), thus only disease severity data from the 2017–18 winter season were reported in this study. The white rust response data of the 346 spinach accessions were analyzed for the analysis of variance (ANOVA) with the general linear models (GLM) in JMP Genomics 9 (SAS Institute, Cary, NC). Multiple comparisons among individual accessions were performed using the student T-test at \( \alpha = 0.05 \), and mean, range, standard deviation (SD), standard error (SE), and coefficient of variation (CV) of disease severity were computed. Distribution of white rust disease across accessions was drawn and the mean disease rating of each accession was used as the phenotypic data for GWAS.

**Genotyping**

DNA was extracted from fresh leaves bulked from 5–10 plants for each genotype. Qualified DNA for each sample was sheared randomly into 350-bp fragments by Covaris Ultrasonic Processor before sequencing. The construction of the DNA libraries followed the process of end repairing, adding A tails, purification, PCR amplification and library qualification [56]. The DNA libraries were pair-end sequenced by whole-genome resequencing (WGR) technology at 10x spinach genome size coverage generating about 10 Gb sequence data for each sample using Illumina NovaSeq Sequencer machine at BGI (https://www.bgi.com/). The spinach genome of Sp75 [18, 57] available at SpinachBase was used as a reference to map the WGR data of the 346 spinach genotypes using Burrows-Wheeler aligner software (BWA v0.7.8-r455 [58]). SAMtools (v 0.1.19-44428 cd) [58] were utilized to sort the bam files and remove duplication reads. The program Picard (v 1.111) [58] was used to merge the bam files from the same sample, and the GATK software (v 3.5) [59] was chosen to detect and filter SNPs and Indels.

Around 16 million raw SNPs were identified in the 346 spinach genotypes. Filtering and keeping the SNPs with minor allele frequency (MAF) \( >2\% \), missing allele \( <30\% \), and heterogeneous rate \( <50\% \), retained 2357260 SNPs distributed on six chromosomes (chr) that were used in this study. There are 217,531 SNPs on chr 1; 239,902 SNPs on chr 2; 651,097 SNPs on chr 3; 629,147 SNPs on chr 4; 334,526 SNPs on chr 5; and 1080 SNPs on chr 6 (FigShare:https://doi.org/10.6084/m9.figshare.17283194). The selected set of SNPs was included in the principal component analysis (PCA) and genetic diversity analysis. PCA and genetic diversity were analyzed with GAPIT 3 (Genomic Association and Prediction Integrated Tool version 3) [54, 60] by setting PCA = 2 to 10 and NJ tree = 2 to 10. Phylogenetic trees were drawn by using neighbor-joining (NJ) method.

**Association analysis**

GWAS was performed in a two step process. In the first step, 2357260 SNPs were used to perform GWAS implementing single marker regression (SMR), GLM (PCA), and MLM (PCA + K) methods in TASSEL 5 [61]. However, we only used the 8399 randomly selected SNPs to estimate PCA and Kinship matrices. PCA matrix was estimated with the PCA tool in TASSEL 5, setting covariance (alternative = correction) and the number of components = 2. Kinship (K) was estimated in TASSEL 5 by using Scald_IBS method. Based on GWAS analysis in TASSEL 5, there were 4836 SNPs with the logarithm of odds (LOD) \([-\text{Log}_{10}(P\text{-value})]\) > 4.0 either in SMR, GLM, or MLM (We use LOD instead of \(-\text{Log}_{10}(P\text{-value})\) in this article).

In the second step, 13235 SNPs (4836 associated SNPs in the first step plus the randomly selected 8399 SNPs used for PCA and Kinship analysis) (FigShare: https://doi.org/10.6084/m9.figshare.17283194) distributed on the six spinach chromosomes (Fig. 51) were used to perform GWAS using the SMR, GLM, and MLM models in TASSEL 5 and several models in GAPIT 3 [54, 60] program. In GAPIT3, GWAS was performed using the general linear model (GLM), mixed linear model (MLM), compressed MLM (cMLM) [62], Settlement of MLM Under Progressively Exclusive Relationship (SUPER) [63], multiple-locus MLM (MLMM), fixed and random model circulating probability unification (FarmCPU) [64] and bayesian-information and linkage-disequilibrium iteratively nested keyway (BLINK) [65] models. In addition, a t-test was conducted for all 13235 SNPs by using visual basic codes in Microsoft Excel 2016.

Multiple TASSEL and GAPIT models were used to find reliable and stable white rust resistance-associated SNP markers and candidate genes and QTL regions in spinach. The significant threshold of associations was calculated using Bonferroni correction of P-value with an \( \alpha = 0.05 \) \((0.05 / SNP \text{ number})\), and LOD value of 5.42 was used as significance threshold based on the 13235 SNPs in this study.

**Candidate gene identification/detection**

Genes were searched within 50 Kb on either side of significant SNPs of the spinach Sp75 genome annotation at the SpinachBase site (http://www.spinachbase.org/). Our emphasis was to find analogs of disease resistance genes near the significantly associated SNP markers.
Genomic prediction for genomic selection of white rust resistance

The ridge regression best linear unbiased prediction (rrBLUP) method was used to perform GP using the rrBLUP package [66] in R Version 4.0.5. In addition, GP was conducted with gBLUP and cBLUP implemented in GAPIT package [54]; Bayesian models including Bayes A, Bayes B, Bayes LASSO (BL), and Bayes ridge regression (BRR) implemented in BGLR package [67]; and random forest (RF) model implemented in Random Forest R package [68] and support vector machines (SVM) [68] implemented in kernlab packages. GP using these packages has been reported in previous studies [49–53].

GP for white rust resistance was performed in 346 spinach accessions based on ratios of training / testing sets, number of SNPs, and GP models. (1) GP was performed using nine different ratios of training / testing sets, 2 fold (1:1), 3-, 4-, 5-, 6-, 7-, 8-, 9-, and 10-fold (9:1) across three sets of SNPs: (i) all 13,235 SNPs, (ii) 40 SNP markers, and (iii) 9 SNP markers (GWAS-derived SNP markers). (2) Eight different SNP number sets from 9 SNPs to 4846 SNPs were used to estimate GP by BL for three SNP sets: (i) Set.13235SNP, (ii) Set.4836SNP_select, and (iii) Set.8399SNP.random. (3) GP was estimated with nine GP models, BA, BB, BL, BRR, SVM, RF, rrBLUP, gBLUP, and cBLUP, in cross-prediction for white rust resistance among seven SNP sets (all, 40 m, 9 m, 40r, 9r, 40rr, and 9rr). These seven SNP sets were selected based on results obtained and information provided in the result section. In addition, GWAS-derived SNP markers for GP were analyzed and discussed.

The PA for the tested models in this study was estimated by calculating the average Pearson’s correlation coefficient (r) between the GEBVs estimates from the training set and white rust phenotypic values in the validation set or testing set [40, 53, 69]. The training and testing sets were randomly generated 100 times; the average r-value was estimated; and distribution charts (boxplots) were drawn using the ggplot2 R package.

Results
Evaluation of white rust resistance

The white rust disease showed signs and symptoms on leaves, and the disease severity was recorded using the 0 to 10 disease scale (Fig. 1). The scale (0–10) in the 346 spinach accessions did not show a normal distribution but skewed toward a higher disease severity due to most material being highly susceptible (Fig. 2) in the association panel. The mean disease severity ranged from 1.0 to 6.5, averaged 4.8 with a standard deviation of 0.911 and the CV was 17.2%. The data showed an extensive range and variation of the white rust disease scale in the 346 accessions, confirming the suitability of the association panel for GWAS. The lines NSL 6098, PI 175311, PI 220686, PI 224959, PI 226671, PI 227045, and PI 648958 showed the highest white rust resistant levels with a score of 2.0 or less (Table 1 & S1), indicating their suitability as parents in breeding programs to develop white rust-resistant hybrids and cultivars.

Genetic diversity among white rust-resistant lines

Among the 346 spinach accessions, 23 showed white rust resistance with a rate 3 or below (Table 1), indicating that the 23 spinach accessions can be used as parents to develop new spinach cultivars or lines for white rust resistance in breeding. Five of the 23 accessions were originally collected from Afghanistan; two from China; two from India; five from Iran; three from Turkey; and six from United States (Table 1), indicating that the white rust resistance was mainly distributed among Asia and U.S. accessions in this study.

The genetic diversity analysis among the 23 accessions showed that (1) the accessions from the same country were located at neighbor each other with less genetic distance in the phylogenetic tree in most cases; (2) two clusters were grouped: the five accessions, PI 227045, PI 165994, PI 175311, PI 433210 and PI 648949 from Iran, China and India as one group, and other 18 accessions as another group; and (3) In group I, the four accessions, PI 207518, PI 220686, PI 211632, and PI 212120 had different genetic base from others as I-outlier (Fig. 3, Table 1). The phylogenetic analysis will provide information on how to utilize these white rust-resistant accessions.

PCA and phylogenetic analysis

Based on PCA and phylogenetic analysis when PCA = 2 to 10 by GAPIT 3 in the 346 spinach accessions with 8399 randomly selected SNPs distributed on six chromosomes, two sub-populations (clusters) were the most clearly divided in the GWAS panel of the 346 accessions (Fig. S2-1, S2-2, S2-3, S2-4, and Fig. 4). The GWAS panel can also be divided into three sub-populations (clusters or groups) but not for other sub-populations from PCA = 4 to 10 (Fig. 4, Fig. S2-1, S2-2, S2-3). Each of the 346 accessions was arranged into its position in a phylogenetic tree of two sub-populations by the neighbor-joining (NJ) method drawn by GAPIT 3 (Fig. S2-5). The NJ phylogenetic trees of two sub-populations and three sub-populations and the 3D graphical plot of PCA in two sub-populations and three sub-populations were shown in Fig. 4 and listed in Table S1.

Based on 2-cluster, Q1 and Q2 consisted of 301 (87.0%) and 45 (13.0%) accessions, respectively, nevertheless, based on 3-cluster (G1 to G3), there were 301 (87.0%) G1, 26 (7.5%) G2, and 19 (5.5%) G3 accessions, respectively (Table S2). Combining 2- and 3-cluster, all 301 accessions in Q1 stayed at the same cluster G1; but the 45 accessions in Q2 were divided into two groups G2 and G3 with 26 and 19 accessions, respectively (Table S2). The accessions from India, Japan, and Mongolia were grouped into Q2, G2, where the accessions were grouped from cluster Q1 based on two clusters and G2 based on three clusters in the panel; the majority of accessions from China plus all accessions from South Korea and Thailand (but only
one accession each of the two countries) to Q2.G3; and the accessions from other countries to Q1.G1 (Table S2).

**Association study**

Based on the six models in GAPIT 3 and three models in TASSEL 5 when PCA = 2, 40 SNPs, located on chrs 1, 2, 3, 4, 5, and 6, were associated with the white rust resistance (Table S3). The observed vs expected LOD $\{-\log_{10}(p)\}$ distributions in QQ-plots showed a large divergence from the expected distribution (Fig. S3 B), indicating that there were SNPs associated with the white rust resistance in the association panel. The Manhattan plot showed that a dozen SNPs with LOD value greater than 5.42 (significant threshold) across the six GWAS models from GAPIT 3 (Fig. S3 A & C), were associated with white rust resistance.

BLINK had SNPs with LOD greater than 5.42 on chrs 1, 2, 3, 4, and 6 and FarmCPU had SNPs with LOD > 5.42 on chr 2, 3, 4, 5, and 6 (Fig. 5, Table S3), indicating that there are SNP markers associated with white rust resistance. Gapit.SUPER, Gapit.GLM, Tassel.GLM and Tassel.SMR showed a peak at the region on chr 4, where a dozen SNPs had LOD > 5.42 and only this region had SNPs with LOD > 10 (Fig. 6, Fig. S4, Table S3), indicating there is a major QTL in the region of chromosome 4 for white rust resistance. However, the Tassel.MLM (Fig. S4), Gapit.MLM, and Gapit.MLMM (Fig. S5) don’t have any SNP with LOD > 5.42, but have dozen of SNPs with LOD > 3.0 (Fig. S4 and S5) and six SNPs had LOD score > 4.0 or 3.0 in the three models, indicating that there were small-effect QTLs for white rust resistance (Table S3).

After combining, nine SNPs, chr2_53049132, chr3_58479501, chr3_95114909, chr4_9176069, chr4_17807168, chr4_83938338, chr4_87601768, chr6_1877096, and chr6_31287118, located on chrs 2, 3, 4, and 6, respectively were selected as the associated SNP markers for white rust resistance.
Table 1. Top 23 spinach white rust resistant lines

| Line ID             | ACCESSION | NAME         | ORIGIN          | Country | 2-cluster | 3-cluster | Group | WR Scale |
|---------------------|-----------|--------------|-----------------|---------|-----------|-----------|-------|----------|
| PI222270_Iran.Q1.G1.3 | PI 222270 | Esfenaj      | Iran            | Iran    | Q1        | G1        | I     | 3.0      |
| PI222283_Iran.Q1.G1.3 | PI 222838 | Esfenaj      | Iran            | Iran    | Q1        | G1        | I     | 3.0      |
| PI224959_Iran.Q1.G1.2 | PI 224959 | Cornell ID #4| Iran            | Iran    | Q1        | G1        | I     | 2.0      |
| PI226671_Iran.Q1.G1.1 | PI 226671 | Cornell ID #10| Iran            | Iran    | Q1        | G1        | I     | 1.0      |
| PI171858_Turkey.Q1.G1.3 | PI 171858 | Harlan 6652  | Kastamonu, Turkey| Turkey | Q1        | G1        | I     | 3.0      |
| PI173131_Turkey.Q1.G1.3 | PI 173131 | Cornell ID #87| Malatya, Turkey | Turkey | Q1        | G1        | I     | 3.0      |
| PI171859_Turkey.Q1.G1.3 | PI 171859 | Harlan 6725  | Samsun, Turkey  | Turkey | Q1        | G1        | I     | 3.0      |
| PI648951_UnitedStates.Q1.G1.3 | PI 648951 | Cornell ID #275| Maryland, United States| United States | Q1        | G1        | I     | 3.0      |
| PI648957_UnitedStates.Q1.G1.3 | PI 648957 | 76 X 71      | Maryland, United States| United States | Q1        | G1        | I     | 3.0      |
| PI648958_UnitedStates.Q1.G1.1.5 | PI 648958 | Cornell ID #286| Maryland, United States| United States | Q1        | G1        | I     | 1.5      |
| PI648960_UnitedStates.Q1.G1.3 | PI 648960 | Cornell ID #288| Maryland, United States| United States | Q1        | G1        | I     | 3.0      |
| PI648961_UnitedStates.Q1.G1.3 | PI 648961 | 224 X 223    | Maryland, United States| United States | Q1        | G1        | I     | 3.0      |
| NSL6098_UnitedStates.Q1.G1.2 | NSL 6098  | Norfolk Savoy/Bloomsdale| Virginia, United States| United States | Q1        | G1        | I     | 2.0      |
| PI212119_Afghanistan.Q1.G1.3 | PI 212119 | Cornell ID #5| Afghanistan      | Afghanistan | Q1        | G1        | I     | 3.0      |
| PI207518_Afghanistan.Q1.G1.2.5 | PI 207518 | Cornell ID #30| Afghanistan      | Afghanistan | Q1        | G1        | I     | 2.5      |
| PI220686_Afghanistan.Q1.G1.2 | PI 220686 | Palek        | Afghanistan      | Afghanistan | Q1        | G1        | I     | 2.0      |
| PI211632_Afghanistan.Q1.G1.2.5 | PI 211632 | Cornell ID #35| Afghanistan      | Afghanistan | Q1        | G1        | I     | 2.5      |
| PI212120_Afghanistan.Q1.G1.2.5 | PI 212120 | Cornell ID #6| Afghanistan      | Afghanistan | Q1        | G1        | I     | 2.5      |
| PI648949_China.Q2.G3.3 | PI 648949 | II9A0323      | Beijing, China  | China    | Q2        | G3        | II    | 3.0      |
| PI433210_China.Q3.G3.2.5 | PI 433210 | 498          | China           | China    | Q2        | G3        | II    | 2.5      |
| PI165994_India.Q2.G2.3 | PI 165994 | Palak        | India           | India    | Q2        | G2        | II    | 2.0      |
| PI175311_India.Q2.G2.2 | PI 175311 | Palak        | India           | India    | Q2        | G2        | II    | 2.0      |
| PI227045_India.Q2.G2.2 | PI 227045 | Cornell ID #201| Iran            | Iran     | Q2        | G2        | II    | 2.0      |

The SNP, chr2_53049132, located at 53049132 bp on chr 2 had a significantly high LOD value at BLINK, SUPER, Gapit.GLMM, and Tassel.SMR with LOD values of 10.08, 5.96, 5.61, and 8.65, respectively (>5.42 threshold); high LOD value of 4.87 in FarmCPU, and a LOD value from 2.5 – 5.42 in other eight models except for Tassel.MLM with 1.87 (Table 2), indicating that chr3_58479501 was but not strongly associated with white rust resistance. Similar to SNP chr2_53049132, SNP, chr3_95114909 at 95114909 bp on chr 3 had significant LOD value >5.42 in BLINK and SUPER; a high LOD value (LOD >4.0) at Gapit.GLMM, Tassel.GLMM; but a low value (LOD <2.0) at FarmCPU, Gapit.MLM, MLMM, and Tassel.MLM (Table 2), indicating that chr3_95114909 was associated with white rust resistance but was not stable across all tested models. SNPs, chr4_9176069 at 9176069 bp on chr 4 had a high and significant LOD value >5.42 in BLINK, FarmCPU, SUPER, Gapit.GLMM, Tassel.GLMM, and SMR, but a low value (LOD <2.0) at Gapit.MLM, MLMM, and Tassel.MLM (Table 2), indicating...
Table 2. Nine SNP markers associated with white rust resistance in 346 USDA spinach germplasm accessions

| SNP     | Chr | Position | LOD (−log(P)) value using GAPIT 3<sup>a</sup> | LOD (−Log(P)) value in Tassela<sup>b</sup> | MAF(%) | LOD (−Log(P)) > 5.42 in GAPIT 3 R Package, and MLM, GLM, and SMR in TASSEL 5 | Dominance/recessive for white rust resistance |
|---------|-----|----------|-----------------------------------------------|--------------------------------------------|--------|-------------------------------------------------------------------------|-----------------------------------------------|
| chr2_53049132 | 2   | 53049132 | 2.48                                          | 1.32                                       | 5.96   | 5.61                                                                     | Blink,Super,Glm                                |
| chr3_58479501 | 3   | 58479501 | 2.14                                          | 2.43                                       | 2.47   | 4.07                                                                     | FarmCPU                                       |
| chr3_95114909 | 3   | 95114909 | 6.61                                          | 1.54                                       | 1.55   | 5.52                                                                     | Blink,Super                                    |
| chr4_9176069  | 4   | 9176069  | 5.86                                          | 1.94                                       | 1.97   | 13.54                                                                    | Blink,FarmCPU,Super,Glm                       |
| chr4_17807168 | 4   | 17807168 | 0.81                                          | 0.71                                       | 0.71   | 7.38                                                                     | Super,Glm                                     |
| chr4_83938338 | 4   | 83938338 | 5.76                                          | 1.03                                       | 1.04   | 5.07                                                                     | Blink,Glm                                     |
| chr5_87601768 | 5   | 87601768 | 2.68                                          | 4.02                                       | 4.17   | 2.39                                                                     | Blink,FarmCPU,Super,Glm                       |
| chr6_1877096  | 6   | 1877096  | 1.08                                          | 6.71                                       | 3.27   | 3.33                                                                     | FarmCPU,plus(MLM + MMLM>3)                    |
| chr6_31287118 | 6   | 31287118 | 2.98                                          | 3.85                                       | 3.98   | 3.64                                                                     | Blink,plus (mlm + mmlm>3)                     |

<sup>a</sup>SNP name defined as SNP on the chromosome plus its position on chromosome.<sup>b</sup>Lod (−Log(P)) value, where P is the P value from the six models, BLINK, FarmCPU, MLM, MMLM, SUPER, and GLM in GAPIT 3 R Package, and MLM, GLM, and SMR in TASSEL 5. A: Beneficial allele for white rust resistance, B: unbeneficial allele for susceptible.
that chr4_9176069 was a comparatively good marker for white rust resistance. The SNP, chr4_17807168 at 17807168 bp on chr 4 had a very high and significant LOD value >5.42 in SUPER, Gapit.GLMM, Tassel.GLMM and SMR but a very low value at other models with LOD <1.0 (Table 2), indicating that chr4_17807168 was associated with white rust resistance but did not show stability across all tested models. SNP, chr4_839383838 at 839383838 bp on chr 4 had a significant LOD value >5.42 in BLINK and Gapit.GLMM; high value (LOD = 5.01) at SUPER; and a LOD value >3.5 at both Tassel.GLMM and SMR; but a low value (LOD <1.2) at FarmCPU, Gapit.MLM, MLMM, and Tassel.MLM (Table 2), indicating that chr4_839383838 was associated with white rust resistance but did not show stability across all tested models. chr4_87601768 at 87601768 bp on chr 4 did not have a significant LOD value >5.42 but had value >2.0 across all tested nine models and the highest values across three MLM Models, Gapit.MLM, MLMM, and Tassel.MLM with LOD value >4.0 (Table 2), indicating that the SNP chr4_87601768 showed stability across nine models, although the LOD value was not high but significant at P = 0.01. chr6_1877096 at 1877096 bp on chr 6 had significant LOD value >5.42 at FarmCPU; high LOD value >3.0 at Gapit.MLM, MLMM, SUPER, Gapit.GLMM, and Tassel.MLM; >2.30 at Tassel.GLM and SMR; but low value with 1.08 at BLINK (Table 2), indicating that chr6_1877096 was associated with white rust resistance but did not show stability across all tested models. chr6_31287118 at 31287118 bp on chr 6 showed the best SNP markers with LOD value >2.4 at all nine models; 12.18 at BLINK; >3.5 at Gapit.MLM, MLMM, SUPER, Gapit.GLMM, and Tassel.MLM; >2.9 at FarmCPU and Tassel.GLMM; and 2.49 at SMR (Table 2), indicating that the SNP, chr6_31287118 was a very stable marker associated with white rust resistance.}

**Figure 4.** Population genetic diversity analysis in the association panel consisted of 346 USDA spinach germplasm accessions. Phylogenetic trees drawn by neighbor-joining (NJ) method in (A) two sub-population and (B) three sub-population, and 3D graphical plot of the principal component analysis (PCA) in (C) two sub-population and (D) three sub-population drawn by GAPIT 3. A large phylogenetic tree of the 3B can be visible for each of the 346 spinach accessions is shown in Supplementary Figure S2–5.

**T-test for association**

t-test showed dominance or recessive in each of the selected 40 SNP markers with LOD >2.0 at P = 0.01 level significance for white rust resistance (Table S4). Seven of the 40 SNPs, chr2_50382388, chr2_53049132, chr3_78126596, chr4_17691593, chr4_17807168, chr5_25899209, and chr5_51760073, did not have both homozygous genotypes (SNP homozygosity in the panel of 346 spinach accessions) but have heterogeneous genotypes and showed dominance for white rust resistance. In addition, 14 SNPs had only one spinach accession; four SNPs had two accessions; two SNPs had
three accessions; and three SNPs had four accessions with homozygosity in one of the SNP alleles, showing dominance or over-dominance (Table S4). Eight SNPs, chr3_58479501, chr4_9155049, chr4_9156552, chr4_9163612, chr4_83938338, chr4_86732255, chr5_28794483, and chr6_41345783 showed the significant differences between two homozygous SNP alleles at $P=0.01$ level (LOD $>2.0$) (Table S4). Five of the eight SNPs had only two or three accessions with homozygosity in one of the SNP alleles (Table S4).
Among the nine SNP markers selected (Table S4, Table 2), chr2_53049132 showed dominance with allele “T” as a beneficial allele for white rust resistance and “C” as an un-beneficial allele for susceptibility; chr3_95114909, chr4_9176069, chr4_17807168 and chr4_83938338 also showed dominance; chr3_58479501, chr6_1877096, and chr6_31287118 showed partial-recessive.

Candidate genes for white rust resistance
A total of 121 genes were listed in Supplementary Table S5 and they were located within 50 Kb distance from the 40 associated SNP markers in Table S3. All Leucine-Rich Repeat (LRR) genes plus those with less than 1 Kb distance from the associated SNP markers were listed in Table 3, where 13 genes were located at 12 associated SNP regions. Six SNPs were inside six genes and three SNPs were with a distance less than 1 Kb from a gene, respectively. Six SNPs were located less than 50 Kb from five disease resistance gene analogues encoding LRR domains (Table 3).

The six gene models, Spo01590, Spo23694, Spo12071, Spo12072, Spo14612, and Spo08236 contain a SNP marker, chr2_50382388, chr2_53049132, chr4_9152378, chr4_9170963, chr4_20532790, and chr4_86732255 respectively, on chrs 2 and 4 (Table 3). Whether these six gene models are related to white rust resistance needs further study. The Leucine-Rich Repeat (LRR) gene model, Spo01686 located at 50399687 – 50401105 bp on chr 2, is based on Sp75 spinach genome reference located near SNP marker chr2_50382388 (distance of 17.299 Kbp). Spo12068, located at 9097039 – 9101229 bp on chr 4, is located near SNP marker chr4_9077455 (distance of 19.548 Kbp). Both LRR models, Spo20901 and Spo20900, located at 17644108 –17646417 bp and 17674521 – 17684091 bp on chr 4, located near the SNP marker chr4_17691593 with distance 45.176 Kb and 7.502 Kbp, respectively. Spo04510, located at 31326235 – 31330354 bp in chr 6 is near the SNP marker chr6_31287112 (39.123,kbp) (Table 3). Further studies are needed to evaluate whether the five LRR genes are related to white rust resistance.

Genomic prediction with different SNP numbers
GP was performed with eight different SNP number sets (9, 40, 100, 200, 500, 1000, 2000, and 4836 SNPs) selected from three different SNP groups in cross-predictions for white rust resistance using BL model in three SNP sets: Set.13235SNP, Set.4836SNP.select, and Set.8839SNP.random (Datasets available at FigShare: https://doi.org/10.6084/m9.figshare.17283194). There were 24 combinations for GP analysis, consisting of eight SNP number sets selected from three SNP groups. Each GP analysis was run for 100 times to calculate GP statistical parameters and r values. The average r-value (r_y100) and SE of 100 runs for each GP combination are presented in the Table S7 and Fig. S7.

The results showed that the average r value (r_y100) decreased when decreasing the SNP number set (Table S7 and Fig. S7). From the Set.13235SNP, the average r-value (r_y100) was 0.79 when 4836 SNPs were used and decreased to 0.25 when 9 SNPs were used. In the Set.4836SNP.select, the average r-value (r_y100) was 0.82 when 4836 SNPs were used and decreased to 0.31 when 9 SNPs were used. In the two SNP groups, the r-value was higher than 0.50 when 100 SNPs or more were used. But the average r-value (r_y100) was very low in the Set.8839SNP.random of the SNP group; ranged from 0.19 to 0.07; and the highest was only 0.19 when 4836 SNPs were used (Table S7, Fig. S7), indicating that random SNP group without associated markers included can’t be used for white rust resistance in GS, but using associated SNP marker will increase the selection efficiency; and when >=100 SNPs with associated markers can be used as a set to select white rust resistance in GS.

Genomic prediction using different models
Nine GP models (BA, BB, BL, BRR, SVM, RF, rrBLUP, gBLUP, and cBLUP) were used to conduct GP for white rust resistance among seven SNP sets: all, 40 m, 9 m, 40r, 9r, 40rr, and 9rr, where all signifies all 13235 SNP set; 40 m is the 40 SNP markers in the Table S3; 9 m is the 9 SNP markers listed in Table 2; 40r is a SNP set consisted of 40 SNPs randomly selected from the 13235 SNP set; 9r is a SNP set of 9 SNPs randomly selected from the 13 235 SNP set; 40rr
Table 3. List of 13 genes located at 12 associated SNP regions including 6 SNPs on the 6 genes and 3 SNPs with a distance less than 1 Kb with a gene, respectively, and 6 SNPs with a distance less than 50 Kb with 5 disease resistance gene analogue LRR domain

| Gene ID   | Chr | Gene Start | Gene End | Gene Description                                                                 | Associated SNP | Chr SNP Position | Distance from the start position of the gene | Distance from the end position of the gene | Distance (Kb) |
|-----------|-----|------------|----------|-----------------------------------------------------------------------------------|----------------|-----------------|---------------------------------------------|--------------------------------------------|---------------|
| Spo15817  | 2   | 28461 449  | 28471 213| ABH1                                                                              | chr2_28471392 | 2 28471 392     | 9943                                        | 179                                        | <1 kb         |
| Spo01590  | 2   | 50372 467  | 50388 526| Ornithine amine dehydrogenase, beta chain-like / RIC1-like guanyl-nucleotide exchange factor | chr2_50382388 | 2 50382 388     | 9921                                        | −6138                                      | on gene       |
| Spo01686  | 2   | 50399 687  | 50401 105| Receptor-like kinase, Leucine-rich repeat (LRR)                                   | chr2_50349132 | 2 53049 132     | −17299                                      | −18717                                      | <20Kb         |
| Spo23694  | 2   | 53041 260  | 53050 511| Serine decarboxylase family protein                                               | chr4_9152378  | 4 9152 378      | 1246                                        | −2402                                      | on gene       |
| Spo12068  | 4   | 9097 039   | 9101 229 | Receptor-like protein kinase, Leucine-rich repeat (LRR)                           | chr4_9170963  | 4 9170 963      | 524                                         | −7089                                      | on gene       |
| Spo12071  | 4   | 9151 132   | 9154 780 | 60S ribosomal protein L7a, putative                                               | chr4_17691593 | 4 17691 593     | 47458                                       | 45760                                      | <50Kb         |
| Spo12072  | 4   | 9170 439   | 9178 052| Lecithin:cholesterol acyltransferase family protein                               | chr4_17644108 | 4 17644 108     | 47485                                       | 47485                                      | <50Kb         |
| Spo20901  | 4   | 9170 439   | 9178 052| Lecithin:cholesterol acyltransferase family protein                               | chr4_17644108 | 4 17644 108     | 47485                                       | 47485                                      | <50Kb         |
| Spo20900  | 4   | 17674 521  | 17684 091| Receptor-like kinase, Leucine-rich repeat (LRR)                                   | chr4_17674 521| 4 17674 521     | 17072                                       | 17072                                      | <10Kb         |
| Spo14612  | 4   | 20529 360  | 20535 771| Calmodulin binding protein                                                        | chr4_20532 790| 4 20532 790     | 3430                                        | −2981                                      | on gene       |
| Spo08236  | 4   | 86730 376  | 86739 358| Nuclear transport factor 2B                                                       | chr4_86732 255| 4 86732 255     | 1879                                        | −7103                                      | on gene       |
| Spo04510  | 6   | 31326 235  | 31330 354| Receptor-like protein kinase 2, Leucine-rich repeat (LRR)                          | chr6_31326 235| 4 31326 235     | 1879                                        | −7103                                      | on gene       |
| Spo25888  | 6   | 41342 127  | 41345 503| Dihydroflavonol 4-reductase                                                       | chr6_41345 783| 6 41345 783     | 3656                                        | 280                                        | <1 kb         |
is a set consisting of 40 SNPs randomly selected from the random 8839 SNP set; and 9r is a set of 9 SNPs randomly selected from the random 8839 SNP set (Table S8, Fig. 7). The six GP models (BA, BB, BL, BRR, gBLUP, and cBLUP) had similar high average r-value ($r_{\bar{Y}100}$) $> 0.82$ in all 13235-SNP set; $> 0.73$ in the 40-SNP marker set; and $> 0.59$ in the 9-SNP marker set. The other three models, SVM, RF, and rrBLUP, still had a high average r-value ($r_{\bar{Y}100}$) $> 5.2$ in the all.

SNP set. SVM and RF have r-value $> 0.63$ in the 40-SNP marker, and $> 0.47$ in 9-SNP marker set, but rrBLUP had a low value with 0.38 and 0.25, respectively (Table S8, Fig. 7), indicating that the six GP models (BA, BB, BL, BRR, gBLUP, and cBLUP) are good GP models to be utilized in GS for selecting white rust resistance in spinach. All nine models had low $r_{\bar{Y}100}$ values in the four random sets (40r, 9r, 40rr, and 9rr), suggesting that we can’t use a small SNP number randomly selected from a million SNP set in GS for white rust resistance.

**Genomic prediction using GWAS-derived SNP markers**

A higher r-value ($r_{\bar{Y}100}$) were observed when using the GWAS-derived SNP marker sets: $> 0.73$ in 40-SNP marker set (m40) and $> 0.59$ in 9-SNP marker set (m9) across the six GP models (BA, BB, BL, BRR, gBLUP, and cBLUP) (Table S8, Fig. 7). An averaged 0.75 of $r_{\bar{Y}100}$ value in 40 m set and 0.61 in 9 m set were calculated across nine GP models, which were much higher than those from the four randomly selected SNP sets: either 40r, 9r, 40rr, or 9rr across all tested nine GP models (Table S8, Fig. 7), indicating that the GWAS-derived SNP marker sets had high PA and suggesting that we can use the GWAS-derived SNP markers in GS for selecting white rust resistance.

**Discussion**

**Evaluation of white rust**

White rust is a non-culturable oomycete pathogen making it difficult to screen and select spinach genotypes for resistance. Currently, no efficient method has been developed to evaluate white rust resistance in greenhouse or growth chamber conditions as disease severity among known resistant and susceptible genotypes is difficult to discriminate in a single disease cycle [70]. Therefore, field evaluations, whereby spinach genotypes can be evaluated over multiple secondary infection cycles over a longer period of time, can be used to evaluate and select white rust-resistant lines in spinach [4]. For example, some spinach genotypes that are known to be highly resistant to white rust get infected, but the lesions develop somewhat slower and the rust pustules may not even break through the epidermis. As a result, fewer infections occur on a given plant and the difference between a highly susceptible and highly resistant line in a single infection cycle may not be noticeable, but the differences become more pronounced over a
longer period of time under field conditions. However, even in a white rust disease nursery, as was used in this study, disease severity is still highly dependent on the environment from year to year, resulting in a low and unpredictable selection efficiency. Establishing a uniform spread of consistent white rust nurseries under field setting is difficult to accomplish, requiring multiple years of no-rotation spinach crop to build enough inoculum in the soil. In this study, the 346 USDA spinach germplasm accessions were evaluated for white rust resistance in the Del Monte White Rust Nursery in Crystal City, Texas, for three years during the three winter seasons of 2015–16, 2016–17, and 2017–18. The nursery was used for spinach white rust evaluation for commercial spinach hybrids, germplasm, and breeding lines where heavy disease pressure had consistently been observed for 30 years when environmental conditions were favorable for the disease. Because the disease severity was relatively low in the winter seasons 2015–2016 and 2016–2017 including the known susceptible control lines, the white rust disease severity ratings were not robust (i.e. false low ratings due to escape from disease). However, disease severity was high in the winter 2017–2018 season due to the favorable environment. Therefore, this 2017–2018 disease severity report was only used to conduct GWAS and identify SNP markers associated with white rust. Based on the white rust phenotypic data, 23 of 346 spinach accessions showed relatively high resistance to white rust. The accessions showing higher resistance with a disease severity scale of 2.0 or less (Table 1 & S1) can be used as parents in breeding programs to develop white rust-resistant lines and cultivars.

Genome-wide association study and SNP marker identification for white rust resistance

In this study, GWAS was performed in two steps. In the first step, 4836 SNP markers associated with white rust resistance were identified from 2357260 SNPs using TASSEL 5. In the second step, 13235 SNPs (the 4836 SNPs from the first step plus the randomly selected 8399 SNPs) were used to conduct GWAS by implementing multiple models, including six models in GAPIT 3, three models in TASSEL 5, and t-test. Forty SNP markers were associated with white rust resistance with LOD >5.42 in one of the six tested MLM models (Gapit.MLM, MLMM, SUPER, FarmCPU, BLINK, or Tassel.MLM). After combining analysis of the six models in Gapit 3 and three models in TASSEL 5, nine SNPs located on chrs 2, 3, 4, and 6 were relatively consistent across the models and were selected as the associated SNP markers for white rust resistance in this study (Table 2). Awika et al. (2019) [30] conducted GWAS analysis for white rust resistance in a panel of 267 spinach accessions with 6111 SNPs and reported a total of 448 minor alleles (SNPs) associated with white rust severity. None of the 448 SNPs reported by Awika et al. (2019) [30] was validated in this study since their approach targeted factors associated with susceptibility. Therefore, it is possible to combine resistance and susceptible associated SNPs found in this study to improve spinach resistance. As expected, we observed differences in the number of identified associated SNPs when using TASSEL 5 vs GAPIT 3 GWAS tools or different models such as BLINK, FarmCPU, GAPIT.MLM and GAPIT.GLM. The same differences have been widely reported and discussed in several publications [30–36, 53]. In this study, we selected SNPs as the markers associated with white rust resistance by multiple models combined, including six models in GAPIT 3 and three models in TASSEL 5 if the SNP had a significant LOD value across multiple models.

Candidate genes for white rust resistance

In this study, a total of 121 genes were identified to be located within 50 Kb distance from the 40 associated SNP markers (Table S3). Thirteen genes located at 12 associated SNP regions were selected as candidates for white rust resistance, among them, five were disease resistance gene analogue with LRR domain (Table 3). However, further studies are needed to confirm whether the five LRR genes are related to white rust resistance.

Despite the success of GWAS in identifying genetic loci associated with several agronomic and disease resistance-related traits, it will be challenging to pinpoint the causal gene in each of these loci. A successful GWAS only identifies probable genomic regions but requires subsequent characterization for validating the actual identification of causal relationship with disease using proteomics/ transcriptional profiling. Given the complexity and nature of the white rust, most genes identified in our study, whether a given gene is likely to be involved in determining a resistant phenotype alone would need cloning individual genes in the appropriate genetic background. Most verified plant disease resistance genes isolated to date contain a nucleotide-binding site and leucine-rich repeat (NBS-LRR) domains similar to the one we identified in this study. Activated NB-LRRs represent a tip of the signaling cascade that triggers defense responses and not the causal genes defining the resistance alone. It will be impulsive to correlate expression patterns of the identified candidate genes and disease reaction. The percent variation (R-square%) explained by individual SNP (Table 3) shows the strength of these variants at the evolutionary and population levels in defining resistance. Hence, the genes identified in this study open new avenues to design white rust resistance through systematic integration of selected accessions in the breeding program of spinach and other closely related species. Detailed characterization of these genes, although intriguing, is beyond the scope of this paper. However, we will continue pursuing the relationship of these genes in resistance mechanisms as a future work through additional studies.

Genomic prediction

In this study, the nine-fold sets from 2 fold (1:1) to 10-fold (9:1) had similar, although not identical, averaged
r values ($r_{100}$) in each SNP set using the same model, either BL or rrBLUP (Table S6, Fig. S6). When increasing the fold number, the SE was also increased, suggesting that a larger training set and the smaller testing set would increase the error. The 2-fold set has a smaller r-value and showed that a smaller training set would have less PA with a smaller r-value. Shi et al. (2021) [53] also reported that different training/testing ratio sets showed similar trends in GP analysis for soybean cyst nematode resistance in common bean. Ravelombola et al. (2021) [49] reported similar results for growth habit, flowering time, and grain yield in the cowpea population evaluated under drought conditions. Keller et al. (2020) [71] reported time, and grain yield in the cowpea population evaluated resistance in common bean. Ravelombola et al. (2021) [49] reported that different training/testing ratio sets showed ever, the average r-value ($r_{100}$) when 4836 SNPs were used and decreased to 0.31 when 9 SNP were used. In the Set.8839SNP.random, which was randomly selected set: Set.8839SNP.select, the average r-value ($r_{100}$) was 0.79 when 4836 SNPs were used and decreased to 0.25 when 9 SNPs were used. In the Set.4836SNP.select, the average r-value ($r_{100}$) was 0.82 when 4836 SNPs were used and decreased to 0.31 when 9 SNPs were used. In the two SNP groups, the r-value was higher than 0.50 when 100 SNPs or more were used. However, the average r-value ($r_{100}$) was very low in randomly selected set: Set.8839SNP.random, which was randomly selected from 217531 SNPs (4.06%); ranged from 0.07 to 0.19 (Table S7, Fig. S7), indicating that random SNP group without associated markers included cannot be used for white rust resistance in GS, but using associated SNP marker will increase the selection efficiency. From this study, 100 or more SNPs with associated markers can be used as a set to select white rust resistance in GS. In most reports, the smaller the number of SNPs resulted in lowering the PA [40, 53, 71, 72]. Zhang et al. (2016) [40] estimated PA (r-value) of seed size of 309 soybean accessions and reported $r = 0.85$ using 2000 SNPs or 31045 SNPs; and lowered to 0.8 when 1000 SNPs or 500 SNPs were used.

In this study, GP was performed using GWAS-derived SNP markers, either 4836 selected SNPs, 40-SNP marker set, or 9-SNP marker set had higher PA (r-value) than the randomly selected SNP sets in all of the tested GP models (Table S6 and S8). The results suggest the advantage of using the GWAS-derived SNP markers in GS for white rust resistance. Zhang et al. (2016) [40] reported that the r values were 25% higher when using GWAS-derived SNP markers than using the same number of randomly selected SNPs for seed size in soybean. Qin et al. (2019) [72] also reported that the average r values were higher when using SNP markers for 15 amino acid contents in soybean seeds. Spindel et al. (2016) [73] developed a GS model that combines RR-BLUP with GWAS derived-markers and reported that this new model outperformed for a variety of traits in multiple environments. Thus, using GWAS-derived SNP markers to perform GS is an approach that combines MAS and GS and can be used in the real-world breeding programs. However, the predictive ability may be biased when GWAS-associated SNP markers are used to predict the GEBVs in the same GWAS panel. The GP will probably be lower if prediction performance is tested in other panels with different individuals. Similar approaches have been tested for many traits in several crops and found it practical to do genome breeding using GWAS-derived SNP markers [49–51, 53, 72]. Therefore, GP approach combining both MAS and GS through GEBVs using associated SNP markers would be valuable in molecular breeding for white rust resistance in spinach and for other quantitative traits in other plant species, and assessment of genomic prediction potential is ongoing on several important traits in spinach [74].

**Conclusion**

In this study, 346 USDA spinach germplasm accessions were phenotyped for white rust resistance under field conditions; 23 accessions showed white rust resistance or intermediate resistance with a disease rate 3.0 or less based on 0–10 scale; and the seven accessions, NSL 6098, PI 175311, PI 220686, PI 224959, PI 226671, PI 227045, and PI 648958 showed the highest white rust resistant levels with a score of 2.0 or less, indicating their suitability as parents in breeding programs to develop white rust-resistant hybrids and cultivars. Genome-wide association study (GWAS) was performed in the 346 accessions with 13235 SNPs and identified nine SNPs, chr2:53,049,132, chr3:58,479,501, chr3:95,114,909, chr4:9,176,069, chr4:17,807,168, chr4:83,938,338, chr4:87,601,768, chr6:1,877,096, and chr6:31,287,118, located on chromosomes 2, 3, 4, and 6 associated with white
rust resistance. Genomic prediction (GP) was tested for prediction accuracy (PA) using Pearson’s correlation coefficient (r) between the genomic estimation breeding value (GEBV) and the observed values. High averaged r values were observed in each SNP set using different GP models and up to 0.84 when using GWAS-derived significant SNP markers. The SNP markers and the high PA can provide valuable information for breeders to improve spinach by marker-assisted selection (MAS) and genomic selection (GS).

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Author contributions
AS, JC, CA, BM, LdT, and LS were the principal investigator (PI) of the project. All authors, AS, GB, HX, CA, CF, BL, VJ, LS, BM, LdT, and JC, were involved in the phenotyping and performed white rust resistant evaluation. AS performed genotyping, genomic and statistical analysis, and wrote the draft of the manuscript. GB and HX assisted in genotyping and data analysis. All authors have edited, reviewed and approved the manuscript.

Data availability statement
The datasets presented in this study are available in Tables, Figures, Supplementary Tables, and Supplementary Figures. The SNP data are available in FigShare https://doi.org/10.6084/m9.figshare.17283194. The accession number(s) used in this study can be found in the article/Supplementary Material.

Conflict of interest
The authors declare that they have no conflict of interest.

Supplementary data
Supplementary data is available at Horticulture Research online.

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