Machine Learning based Genome-Wide Association Studies for Uncovering QTL Underlying Soybean Yield and its Components

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Research Article

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Abstract

Genome-wide association study (GWAS) is currently one of the most recommended approaches for discovering quantitative trait loci (QTL) associated with complex traits in plant species. Insufficient statistical power is a limiting factor that current conventional GWAS methods are suffering from, especially in narrow genetic bases plants such as soybean. In this study, we evaluated the potential use of two machine learning (ML) algorithms, support vector machine (SVR) and random forest (RF), in GWAS and compared them with two conventional methods of mixed linear models (MLM) and fixed and random model circulating probability unification (FarmCPU), for identifying QTL associated with soybean yield and its components. In this study, important soybean yield component traits, including the number of reproductive nodes (RNP), non-reproductive nodes (NRNP), total nodes (NP), and total pods (PP) per plant along with yield and date to maturity were assessed using a panel of 227 soybean genotypes evaluated across four environments. Using the SVR-mediated GWAS method, we were able to discover a greater number and most relevant QTL associated with the target traits, supported by the functional annotation of candidate gene analyses. This study for the first time demonstrated the potential benefit of using sophisticated mathematical approaches, such as ML algorithms, in GWAS for identifying QTL that can improve the efficiency of genomic-based breeding programs.

Key Message

Implementing machine learning algorithms in GWAS can simultaneously consider a wide range of interconnected biological processes and mechanisms that shape the phenotype of complex traits such as yield and its components in soybean.

Introduction

Soybean (Glycine max [L.] Merr.) is known as one of the most important legume crops with substantial economic value (Rębilas et al. 2020). Soybean is widely used for food, feed, fiber, biodiesel, and green manure (Carter et al. 2018; Temesgen and Assefa 2020). Despite the importance of genetic improvement in soybean yield, the soybean germplasm has in general a narrow genetic basis, especially within the North American germplasm, which has resulted in limited historical enhancement of the genetic gain (Xavier and Rainey 2020). Therefore, there is a great need for analytical breeding to explore the optimum genetic potential for soybean yield (Mangena 2020; Suhre et al. 2014).

Analytical breeding strategy as an alternate breeding approach requires a better understanding of the factors, or individual traits, responsible for more complex characteristics and phenomenon such as plant growth, development, and yield (Richards 1982). This strategy considers highly correlated secondary traits with the trait of interest as the selection criteria that can make empirical selection more efficient for improving the genetic gain (Reynolds 2001; Richards 1982; Xavier and Rainey 2020). The application of the analytical approaches in plant breeding programs has been limited due mainly to limited resources for evaluating several secondary traits that are mostly time and labor-consuming (Richards 1982; Xavier
et al. 2018). Most of analytical breeding studies were conducted on small populations with limited number of genotypes and, therefore, the results with limited generalization ed in the limitation of the knowledge in the genome-to-phenome analysis process (Kahlon et al. 2011; Nico et al. 2019; Robinson et al. 2009).

Yield potential in soybean is mainly determined by its yield component traits that are the total number of pods, seeds, and nodes, which can be divided into reproductive and non-reproductive nodes, in plants as well as seed size (Pedersen and Lauer 2004; Reynolds 2001; Xavier et al. 2018; Xavier and Rainey 2020; Yoosefzadeh-Najafabadi et al. 2021b). Of these traits, the total number of nodes and pods play more important roles in seed yield production (Robinson et al. 2009; Yoosefzadeh-Najafabadi et al. 2021b).

Several studies reported a steady increase in the total number of nodes and the total number of pods in soybean cultivars from 1920 to 2010 (Kahlon et al. 2011; Suhre et al. 2014; Xavier and Rainey 2020). These findings may highlight the importance and potential use of the phenotypic and genotypic information on these traits, along with yield per se, as selection criteria in cultivar development programs (Ma et al. 2001).

Genetic information of soybean yield component traits can accelerate the efficiency of cultivar development programs through selecting genotypes in improved genetic gains (Xavier and Rainey 2020). Genome-Wide Association Studies (GWAS), as one of the most common genetic approaches, can be implemented on genetically diverse populations to detect the quantitative trait loci (QTL) associated with the soybean yield component traits (Kaler et al. 2020). Associated QTL can be used for screening large soybean populations in a more cost-effective and timely manner (Xavier et al. 2018). Up to date, several GWAS approaches such as mixed linear models (MLM), multiple loci linear mixed model (MLMM), and fixed and random model circulating probability unification (FarmCPU) have been developed for genetic studies of complex traits (Kaler et al. 2020). However, due to the narrow genetic base of some plant species, including soybean, these conventional approaches may not have sufficient statistical power to detect reliable QTL (Kaler et al. 2020; Mohammadi et al. 2020; Xavier and Rainey 2020). Therefore, the development of more sophisticated statistical methods can help to establish effective GWAS methods for plant species with narrow genetic bases.

Current GWAS methods are based on the conventional statistical methods that are useful for studying less complex traits in plant species with broader genetic bases (Lipka et al. 2015; Pasaniuc and Price 2017). Machine learning (ML) algorithms as powerful and reliable mathematical methods have been considered as an alternative to conventional statistical methods for dealing with large data set as well as conducting GWAS analyses (Xavier and Rainey 2020). Recently, the use of ML algorithms has been reported in different areas such as plant science (Hesami et al. 2020; Yoosefzadeh-Najafabadi et al. 2021a), animal science (Tulpan 2020), human science (Chen and Vergheese 2020), engineering (Kim et al. 2020), and computer science (Jordan and Mitchell 2015). The application of ML algorithms in GWAS was previously investigated in human science by Szymczak et al. (2009). In this study, the potential use of different ML algorithms such as artificial neural networks (ANN), Bayesian network analysis (BNA), and random forests (RF) are explained for using in GWAS studies focused on human disease studies.
One of the most common used ML algorithms is RF, developed by Breiman (2001), which generates a series of trees from the independent samples for a better prediction performance (Meinshausen 2006). The latter algorithm has been widely used in plant genomics (Ogutu et al. 2011), phenomics (Yoosefzadeh-Najafabadi et al. 2021a), proteomics (Jamil et al. 2020), and metabolomics (Sun et al. 2020).

The first and only use of the RF-mediated GWAS in soybean, for detecting the genomic regions association with yield component traits, was reported by Xavier and Rainey (2020). Support vector machine (SVM) is another common algorithm that can detect behavior and patterns of nonlinear relationships (Auria and Moro 2008; Hesami and Jones 2020; Su et al. 2017). Theoretically, SVM should have high performance due to the use of structural risk minimization instead of the empirical risk minimization inductive principles (Belayneh et al. 2014; Yoosefzadeh-Najafabadi et al. 2021a). There is a significant number of reports on the successful using of SVM in prediction problems (Denton and Salleb-Aouissi 2020; Duan et al. 2005; Hesami et al. 2020; Tulpan 2020; Yoosefzadeh-Najafabadi et al. 2021a). Support vector regression (SVR) is known as the regression version of SVM that commonly used for continuous variables. There are also reports on the successful use of SVR for addressing plant prediction problems (Awad and Khanna 2015). However, the potential use of SVR in GWAS is still unexplored remained to be discovered in the area of plant science.

In this study we aimed to: (1) gain a better understanding of the genetic relationships between soybean yield and its component traits, and (2) investigate the potential use of RF and SVM algorithms in GWAS for discovering QTL underlying soybean yield components in compared with the conventional GWAS methods of MLM and FarmCPU. The results of this study will help soybean breeders and geneticists to have a better perspective of exploiting ML algorithms in GWAS studies and may offer new genomic tools for screening high yielding genotypes with improved genetic gain in large breeding populations.

Materials And Methods

Population and experimental design

An GWAS panel of 250 soybean genotypes was grown at the University of Guelph, Ridgetown Campus in two locations, Palmyra (42°25'50.1"N 81°45'06.9"W, 195 m above sea level) and Ridgetown (42°27'14.8"N 81°52'48.0"W, 200m above sea level) in Ontario, Canada, in two consecutive years, 2018 and 2019. The panel used in this study consisted of the main germplasm of the soybean breeding program at the University of Guelph, Ridgetown Campus, that has been established over 35 years for cultivar development and genetic studies. The randomized complete block design (RCBD) with two replications was used for all four environments. In general, there were 500 and 1000 research plots per environment and year, respectively. Each plot consisted of five 4.2 m long rows with 57 seeds per m² seeding rate.

Phenotyping
In this experiment, soybean seed yield \( (t \text{ ha}^{-1} \text{ at } 13\% \text{ moisture}) \) for each plot was estimated by harvesting three middle rows and adjusted based on the maturity data. Soybean seed yield components, including the total number of reproductive nodes per plant (RNP), the total number of non-reproductive nodes per plant (NRNP), the total nodes per plant (NP), and the total number of pods per plant (PP), were measured using 10 randomly selected plants from each plot. The maturity was recorded as the number of days from planting to physiological maturity (R7, (Fehr and Caviness 1971) for each genotype.

**Genotyping**

Young trifoliate leaf tissue for each soybean genotype from the first replication of the trail at the Ridgetown in 2018, were collected and in a 2 mL screw-cap tube. The leaf samples were freeze-dried for 72 hours, using the Savant ModulyoD Thermoquest (Savant Instruments, Holbrook, NY). By using the DNA Extraction Kit (SIGMA®, Saint Louis, MO), DNA was extracted for soybean genotypes, and the quantity of DNAs was checked via Qubit® 2.0 fluorometer (Invitrogen, Carlsbad, CA). For genotyping-by-sequencing (GBS), DNA samples were sent to Genomic Analysis Platform at Université Laval (Laval, Quebec, Canada). The GWAS panel was genotyped via a GBS protocol based on the enzymatic digestion with ApeKI (Sonah et al. 2013). Single-nucleotide polymorphisms (SNPs) were called by the Fast GBS pipeline (Torkamaneh et al. 2020), using Gmax_275_v2 reference genome. Markov model was used to impute the missing loci, and SNPS with a minor allele frequency (MAF) less than 0.05 were removed below the threshold. In total, after checking the quality of reading sequence and removing SNPs with more than 50% heterozygosity, 23 genotypes were eliminated from the experiment and 17,958 high-quality SNPs from 227 soybean genotypes used for genetic analysis.

**Statistical analyses**

The best linear unbiased prediction (BLUP) as one of the common linear mixed models (Goldberger 1962) was used to estimate the genetic values of each soybean genotype. Also, R package lme4 (Bates et al. 2014) was used to analyze yield and yield components with ‘environment’ as a fixed effect and ‘genotype’ as a random effect. To control for the possible soil heterogeneity among the plots within a given block and reduce the associated experimental errors, nearest-neighbor analyses (NNA) was used as one of the common error control methods (Bowley 1999; Katsileros et al. 2015; Stroup and Mulitze 1991). Outliers were determined in the raw dataset based on the protocols proposed by Bowley (1999) and treated the same as missing data points in the analysis. Overall, the following statistical model (Eq. 1) was used in this study:

\[
Y_{ij} = \mu + f(s) + G_i + E_j + GE_{ij} + \epsilon_{ij}, \ i = 1, ..., k; \ j = 1, ..., n \quad (\text{Eq. 1})
\]

Where \( Y_{ij} \) stands for the trait of interest (soybean seed yield and yield component traits) as a function of an intercept \( \mu \), \( f(s) \) stands for the spatial covariate, \( G_i \) is the random genotype effect, \( E_j \) stands for the
fixed environment effect, GE\textsubscript{ij} is the genotype x environment interaction effect, and \(\varepsilon\textsubscript{ij}\) stands for the residual effect.

The heritability (Eq. 2) was calculated for soybean seed yield and yield components using \textit{lme4} open-source R package (Bates et al. 2007) based on the following equation:

\[ H^2 = \frac{\sigma_G^2}{\sigma_G^2 + \sigma_E^2} \]  

(Eq. 2)

where \(\sigma_G^2\) stands for the genotypic variance, and \(\sigma_E^2\) is the environmental variance.

**Analysis of population structure**

A total of 17,958 high-quality SNPs from 227 soybean genotypes were used to conduct population structure analysis using \textit{fastSTRUCTURE} (Raj et al. 2014). Five runs were conducted for \(K\) set from 1 and 15 to estimate the most appropriate number of subpopulations by using the \(K\) tool from the \textit{fastSTRUCTURE} software.

**Association studies**

Since different GWAS methods may capture different genomic regions (Yang et al. 2018). Therefore, MLM and FarmCPU (two most common GWAS methods) and RF and SVM (two most common machine learning algorithms) were used in this study. MLM and FarmCPU were implemented by using \textit{GAPIT} package (Lipka et al. 2012), and RF, as well as SVM, were conducted through the \textit{Caret} package (Kuhn et al. 2020) in R software version 3.6.1. A brief description of each of the GWAS methods is provided below:

**Mixed Linear Model (MLM)**

This GWAS is based on the likelihood ratio between the full model, consisting of the marker of interest, and the reduced model, which is known as the model without the marker of interest (Wen et al. 2018).

**Fixed and random model circulating probability unification (FarmCPU)**

This GWAS takes the advantages of using MLM as the random model, and stepwise regression as the fixed model iteratively (Liu et al. 2016). False discovery rate (FDR) is used for setting the threshold both in the FarmCPU and MLM models (Benjamini and Hochberg 1995).

**Random Forest (RF)**
This machine-learning algorithm was first implemented by Xavier and Rainey (2020) in a soybean GWAS study. This method is known as the powerful non-parametric regression approach that is derived from aggregating the bootstrapping in various decision trees (Breiman 2001). In this experiment, a 1000-set of decision trees constructed the forest, and the GWAS analysis was done by measuring the importance of each feature (Botta et al. 2014), which was an SNP in this study.

Support vector regression (SVR)

This machine learning algorithm is known as one of the common supervised learning methods in prediction problems (Cortes and Vapnik 1995). This algorithm is based on constructing a set of hyperplanes that can be useful in regression problems (Fletcher 2009). The association statistics in this algorithm can be achieved by estimating the feature importance that was previously proposed by Weston et al. (2001). In this experiment, SNP markers were selected as inputs, and the traits were selected as target variables for estimating the feature importance.

Variable Importance measurement

As one of the common indices for tree-based algorithms, the impurity index was chosen as the metric of the feature importance for the RF algorithm. Regarding the SVR algorithm, the variable importance method for SVR Weston et al. (2001) was implemented in this dataset. For both algorithms, the importance of each SNP was scaled based on 0 to 100 percent scale. Since there is no confirmed way of defining the significant threshold in the tested algorithms, the global empirical threshold that provides the empirical distribution of the null hypothesis (Churchill and Doerge 1994; Doerge and Churchill 1996) was used for establishing threshold in this study. The global empirical threshold was estimated based on fitting the ML algorithm, storing the highest variable importance, repeating 1000 times, and select the SNPs based on \( \alpha = 0.05 \).

Data-driven model processes

In order to estimate the feature importance in RF and SVR algorithms, a five-fold cross-validation strategy (Siegmann and Jarmer 2015) with ten repetitions was applied on the dataset. All of the tested machine learning algorithms were optimized for their parameters for this dataset accordingly.

Extracting candidate genes underlying detected QTLs

For each tested GWAS model, the flanking regions of each QTL was determined using LD decay distance (Fig.1), and then potential candidate genes were retrieved using the G. max cv. William 82 reference genome, gene models 2.0 in SoyBase (https://www.soybase.org). After listing potential candidate genes in defined windows around each significant SNP, at the peak of each QTL, Gene Ontology annotation, GO
visualization

All of the visualizations in this study were conducted using the ggplot2 package (Wickham 2011) in R version 3.6.1 software and Microsoft Excel software (2016).

results

phenotyping evaluations

The panel, consisted of 227 soybean genotypes, showed a different level of variations among the genotypes for seed yield, maturity, and yield component traits. The distribution of the phenotypic measures for the target traits across the four environments is presented in Fig. 2. The highest heritability was observed for maturity with an estimate value of 0.78 followed by NP, RNP, NRNP, and PP, with estimate values of 0.34, 0.33, 0.31, and 0.30, respectively (Fig. 2). The lowest heritability was estimated for yield with an estimate value of 0.24 (Fig. 2). Soybean seed yield and PP showed the highest variability across the environments (Fig. 2).

The linear correlations among all the measured traits were estimated using the coefficients of correlation (r). Based on the results (Fig. 3), all traits were positively correlated with each other, except NRNP that was negatively associated with yield, maturity, RNP, NP, and PP. NP showed the highest correlation with the RNP (r = 0.97) and NRNP (r = -0.63). RNP had the highest correlation (r =0.86) with yield among all the tested yield components (Fig. 3).

genotyping evaluations

For the tested GWAS panel, high-quality SNPs were obtained from 210M single-end Ion Torrent reads that were proceeded with Fast-GBS.v2. From 40,712 SNPs, 17,958 SNPs were polymorphic and mapped to 20 soybean chromosomes. The minimum number of 403 SNPs were mapped on 403 and 1780 on chromosomes 11 and the maximum number of 1780 were mapped on chromosome 18. Overall, the average number of SNPs across all the 20 chromosomes was 898, with the mean density of one SNP for every 0.12 cM across the whole genome.

population structure and kinship
The structure profile for the tested population is presented in Fig. 4. The result of genotypic evaluations suggested that the tested GWAS panel was composed of four to seven subpopulations. Therefore, we chose to conduct the structure analysis using K=7 as the appropriate K for the structure profile of the tested GWAS panel (Fig. 4). In order to reduce the confounding, the kinship was also estimated between genotypes of the GWAS panel.

**GWAS analysis**

The average value for soybean maturity in the tested GWAS panel was 106 days with a standard deviation of 5 days (Fig. 3). Association analysis by the MLM method identified nine associated SNP markers located on chromosomes 2 and 19 (Fig. 5a). Using FarmCPU, a total of nine associated SNP markers were located on chromosomes 2, 19, and 20 (Fig. 5a). By using the RF method, the total of three SNP markers on chromosomes 3, 7, and 16 were associated with the soybean maturity, whereas SVR-mediated GWAS detected 10 associated SNP markers located on chromosomes 2, 6, 10, 16, 19, and 20 (Fig. 5a). SVR-mediated GWAS detected five QTL directly related to the reproductive period and R8 full maturity (Table 1).

The average soybean seed yield in the GWAS panel was 3.5 t ha\(^{-1}\) with a standard deviation of 0.45 t ha\(^{-1}\) (Fig. 3). Using MLM, FarmCPU, RF, and SVR approach, we identified two, three, five, and 18 SNP markers associated with the yield, respectively (Fig. 5b). The SNP markers identified by MLM and FarmCPU were located on chromosomes 6 and 8. Using the RF-mediated GWAS method, associated SNP markers were located on chromosomes 4, 7, 12, and 17. By using the SVR-mediated GWAS method, the SNP markers were identified on chromosomes 3, 4, 6, 7, 15, 19, and 20 (Fig. 5b). In SVR-mediated GWAS, the identified QTL were co-localized with eight previously reported yield-related QTL such as seed yield, seed weight, and seed set (Table 2). However, other tested GWAS methods was not able to find QTLs that were co-localized with any previously reported QTL associated with seed yield (Table 2).

The average NP in the tested GWAS panel was 15.21 nodes with a standard deviation of 0.77 nodes (Fig. 3). By using the MLM and FarmCPU methods, one and two associated SNP markers were detected, respectively (Fig. 6a). Four and ten SNP markers associated with NP were detected using RF and SVR methods, respectively. SVR-mediated GWAS was the only method that identified three previously reported NP-related QTL (Table 3). The average NRNP was 3.33 nodes with a standard deviation of 0.28 nodes (Fig. 3). A total of two, three, five, and ten SNP markers were detected associated with NRNP using the MLM, FarmCPU, RF, and SVR methods, respectively (Fig. 6b). The detected SNP markers using the SVR method were located on chromosomes 4, 7, 18, 19, and 20, whereas SNP markers identified through RF were located on chromosomes 1, 4, 7, 18, and 19 (Fig. 6b). Chromosomes number 4, 8, and 15 were identified as carrying SNP markers with NRNP using FarmCPU and the MLM method identified SNP markers located on chromosomes 8 and 15. Most of the detected QTL using all GWAS methods co-localized with previously reported QTL related to seed weight, seed protein, water use efficiency, first flower, and soybean cyst nematode (Table 4).
The average RNP was 11.89 nodes with a standard deviation of 0.98 nodes (Fig. 3). Based on the results of MLM and FarmCPU methods, four associated SNP markers with RNP were located on chromosomes 8 and 19. Using the RF method, four associated SNP markers were identified on chromosomes 8, 9, 15, and 20. Using the SVR method, 11 SNP markers were detected associated with RNP located on chromosomes 4, 7, 8, 15, 18, 19, and 20 (Fig. 7a). Regardless of the type of GWAS methods used in this study, we found SNP markers associated with the trait on chromosome 8. The position of the associated SNP marker on chromosome 8 was identical using all GWAS methods (~ 450 Kbp). The list of detected QTLs for RNP is also presented in Table 5.

The average value for PP in the tested GWAS panel was 45.02 pods with a standard deviation of 8.54 pods. We did not detect any SNP marker associated with PP using the MLM or FarmCPU methods. However, by using the RF method, four SNP markers located on chromosomes 7, 10, 18, and 20 were found to be associated with PP (Fig. 7b). Twelve SNP markers were detected associated with PP using SVR. The markers were located on chromosomes 6, 9, 10, 11, 15, 18, and 19 (Fig. 7b). The associated SNP markers in chromosome 10 were found both in RF and SVR with 4.6 cM distance far from each other. In PP, MLM and FarmCPU did not detect any related QTL for this trait, while SVR-mediated GWAS was identified seven previously reported QTL directly related to the pod number (Table 6).

Extracting candidate genes undelaying detected QTLs

According to the flanking regions of each QTL, which was determined using LD decay distance, 150-kbp upstream and downstream of each SNP’s peak were considered to identify potential candidate genes (Fig. 1). Candidate genes were extracted within the flanking region of each QTL with high allelic effects using the gene annotation, enrichment tools and the information from previous studies (Table S1). For date to maturity, three peak SNPs (Chr2_695362, Chr2_720134, and Chr19_47513536) had the highest allelic effects than other detected peak SNPs (Fig. 8a). On the basis of the gene annotation and expression within the QTL, Glyma.02g006500 (GO:0015996) and Glyma.19g224200 (GO:0010201), which respectively encode chlorophyll catabolic process and phytochrome A (PHYA) related genes, were identified as the strong candidate genes for maturity. Glyma.02g006500 (GO:0015996) was exactly detected in the peak SNP position of Chr2_695362, whereas Glyma.19g224200 (GO:0010201) was 119 Kbp far from the detected peak SNP at Chr19_47513536. The yield-related QTL with the peak SNP positioned on Chr7_1032587 had the highest allelic effect in comparison with other detected peak SNPs (Fig. 8b). Within a 77 Kbp away from the detected peak SNP (Chr7_1032587), Glyma.07G014100 (GO:0010817), which encodes the regulation of hormone levels, was identified as the strongest candidate genes in yield. For the NP trait, two peak SNPs, Chr7_1032587 and Chr7_1092403, had the highest allelic effects among all the detected peak SNPs (Fig. 8c). In this study, the Chr7_1032587 SNP was found to be associated with yield, NP, and NRNP. The Glyma.07G025500 (GO:0009693) and Glyma.08G065300 (GO:0042546) genes, which encode UBP1-associated protein 2C and cell wall biogenesis, respectively, were candidate as plausible genes influencing both NP and NRNP. Both detected gene candidates were exactly collocated at the corresponding peak SNPs at Chr7_1032587.
and Chr8_5005929 (Fig. 8d). Regarding peak SNPs associated with RNP, the highest allelic effects were found in peak SNPs of Chr9_40285014 and Chr15_34958361 (Fig. 8e). The Glyma.15G214600 (GO:0009920) and Glyma.15G214700 (GO:0009910) genes, which encode cell plate formation involved in plant-type cell wall biogenesis and acetyl-CoA biosynthetic process, respectively, were nominated as strong candidate genes governing NRNP. Glyma.15G214600 (GO:0009920) and Glyma.15G214700 (GO:0009910) were 127 and 90 Kbp far from the peak SNP at Chr15_3495836, respectively. For the PP trait, the highest allelic effects were found in peak SNPs at Chr7_15331676, Chr11_5245870, and Chr18_55469601 (Fig. 8f). The Glyma.07G128100 (GO:0009909) gene, which encodes the regulation of flower development, was the strongest candidate genes that can affect PP. Glyma.07G128100 (GO:0009909) is located in the peak SNP position, Chr7_15331676.

**Discussion**

One of the objectives of this study was to attain a better understanding of the roles of soybean yield component traits in the production of total seed yield and how these traits can be used for facilitating the development of high-yielding soybeans with improved genetic gains. The genetic dissection of soybean yield components and establishing genetic and genomics toolkits that can be used for either designing crosses or screening large breeding populations for selecting genotypes with improved yield components will facilitate the improvement of yield genetic gains in new cultivars (Cooper et al. 2009; Hu et al. 2020; Xavier and Rainey 2020). For this aim, a wide range of analyses, including Pearson correlation, normality and distribution plots, GWAS both in combined and separate environments, and functional annotation of candidate genes and QTL, were performed in this study. The collective evaluation of the mentioned analyses contributed to building the wide perspectives of the genetic architecture of the soybean yield component traits. While high phenotypic variations were observed for yield and PP in the panel, date to maturity and NP had the lowest phenotypic variations across the tested environments. These findings are in line with the results of previous research studies on yield component traits (Kahlon and Board 2012; Xavier and Rainey 2020), in which they found higher variation for the total seed yield and total pods per plant. The heritability and correlation analyses showed that NP had the highest heritability and significant linear correlations with RNP and PP. Also, PP had the highest correlation with yield among all the tested soybean yield components. The number of nodes and pods in soybean are known as the two of the key soybean yield components that play important roles in determining the final soybean seed yield (Herbert and Litchfield 1982; Kahlon and Board 2012; Xavier and Rainey 2020). Previous studies reported low heritability rates for soybean yield components, especially NP and PP (KUSWANTORO 2017; Sulistyo and Sari 2018; Xavier et al. 2016a; Xavier and Rainey 2020). These traits are significantly affected by environmental factors (Price and Schluter 1991). Although for a given trait heritability indicates the strength of the relationship between phenotype and genetic variability, it does not necessarily indicate the value of the trait for genetic studies (Cassell 2009). Different low heritable traits are highly correlated with significant economic traits (Cassell 2009). In soybean, for example, yield can be considered as the most important economic trait that is highly determined by its component traits.
GWAS is known as one of the most important genetic toolkits for detecting QTL associated with quantitate traits (Kaler et al. 2020). There are several statistical methods implemented in GWAS for improving the detection of associated SNP markers with the trait of interest. While conventional GWAS are appropriate approaches for detecting SNP markers with large effects on complex traits, they are, however, underpowered for the simultaneous consideration of a wide range of interconnected biological processes and mechanisms that shape the phenotype of complex traits (Lee et al. 2020). Therefore, using variable importance values in ML algorithms for identifying SNP-trait associations may improve the power of GWAS for discovering variant-trait association with higher resolution (Szymczak et al. 2009). The variable importance methods based on linear and logistic regressions, support vector machines, and random forests are well established in the literature (Grömping 2009; Williamson et al. 2020; Wu and Liu 2009; Yoosefzadeh-Najafabadi et al. 2021a). Among all the tested GWAS methods in this study, SVR-mediated GWAS was the best method to detect SNP markers with high allelic effects associated with the tested traits. The advantage of SVR-mediated GWAS, over conventional GWAS models, can be explained by the presence of a nonlinear relationship between input and output variables, which is used to build an algorithm with accurate prediction ability (Kaneko 2020). Therefore, reliable genomic regions can be discovered by SVR-mediated GWAS because of its ability to consider the interaction effects between SNPs rather than \( p \)-values for individual SNP-trait GWAS tests.

In this study, none of the QTL identified using MLM, FarmCPU, and RF was reported to be associated directly with soybean maturity. However, using SVR-mediated GWAS, five QTL were detected on chromosomes 16 and 19 specifically related to the soybean maturity. All the detected QTL were previously reported by Sonah et al. (2015) and Zhang et al. (2015b) in separate studies as it presented in table 1. Also, the peak SNP position of Chr19_47513536 detected by SVR-mediated GWAS had the highest allelic effect among all the detected SNPs in soybean maturity, which is in line with Sonah et al. (2015). For soybean seed yield, five QTL discovered by SVR-mediated GWAS were reported previously (Copley et al. 2018; Hu et al. 2014), while none of the detected QTL from other tested GWAS methods was previously reported for this trait. There was no previous study on the genetic structure of NRNP and RNP; therefore, all the identified QTL in this study are reported for the first time. For PP, conventional GWAS methods were not able to detect any QTL. However, using SVR-mediated GWAS, a total of seven QTL were distinguished to be related to pod numbers aligned with previous studies (Zhang et al. 2015a). It is necessary to be mentioned that the average allelic effects of the QTL presented in this study, Fig. 8, was estimated using the equation develop by Pimentel et al. (2015) but not directly by the GWAS methods. The RF and SVR-mediated GWAS methods do not specifically provide an allele effect therefore, the aim of this study was mostly focused on detecting the associated genes and QTL underlying the soybean yield, maturity, and yield components.

The results of candidate gene identifications within identified QTL using SVR-mediated GWAS analyses revealed important information. For example, from all the detected genes using SVR-mediated GWAS for maturity, candidate gene \textit{Glyma.02g006500} (GO:0015996) is a protein ABC transporter 1, that is annotated as a chlorophyll catabolic process and located exactly in the peak SNP position at Chr02_695362. ATP-binding cassette (ABC) transporter genes play conspicuous roles in different plant
growth and developmental stages by transporting different phytochemicals across endoplasmic reticulum (ER) membranes (Hwang et al. 2016). Because of the central roles of ABC transporters in transporting biomolecules such as phytohormones, metabolites, and lipids, they play important roles in plant growth and development as well as maturity (Block and Jouhet 2015; Hwang et al. 2016). Moreover, recent studies revealed that ER uses fatty acid building blocks made in the chloroplast to synthesize Triacylglycerol (TAG). Therefore, ABC transporter genes are important for the normal accumulation of Triacylglycerol (TAG) during the seed-filling stage and maturity (Block and Jouhet 2015; Kim et al. 2013). Additionally, Glyma.19g224200 (GO:0010201) in E3 locus, which was previously discovered by Buzzell (1971) and molecularly characterized as a phytochrome A (PHYA) gene (Watanabe et al. 2009), was detected through the SVR-mediated GWAS. Phytochromes, through PHYTOCHROME INTERACTING FACTOR (PIF), regulate the expression of some specific genes encoding rate-limiting catalytic enzymes of different plant growth regulators (e.g., abscisic acid, gibberellins, auxin) and, therefore, play crucial roles in plant maturity (Legris et al. 2019). In addition, PHYB is inactivated after imbibition shade signals, which repress PHYA-dependent signaling in the embryo that results in the maturity of seeds by preventing germination (Casal 2013; De Wit et al. 2016). This is obtained by regulating the balance between abscisic acid and gibberellin. Subsequently, abscisic acid transports from the endosperm to the embryo by ABC transporter (De Wit et al. 2016).

Regarding NRNP, candidate gene Glyma.07G205500 (GO:0009693- UBP1-associated protein 2C) that annotated as ethylene biosynthetic process was located exactly at the peak SNP position on Chr7_37469678, was detected by SVR-mediated GWAS. An interaction screen with the heterogeneous nuclear ribonucleoprotein (hnRNP) results in the production of oligouridylatebinding protein 1 (UBP1)-associated protein (Lambermon et al. 2002). It has been well documented that this protein plays important roles in several physiological processes such as responses to abiotic stresses (Li et al. 2002), leaf senescence(Kim et al. 2008), floral development (Streitner et al. 2008), and chromatin modification (Liu et al. 2007). In addition, previous studies showed that the production of productive or non-reproductive nodes is completely accompanied by the upregulation or downregulation of this protein (Bäurle and Dean 2008; Na et al. 2015). In addition, Glyma.08G065300 (GO:0042546- MADS-box transcription factor) that is associated with cell wall biogenesis, was located in the SNP position of Chr8_5005929. The genes of the MADS-box family can be considered as the main regulators for cell differentiation and organ determination (Lee et al. 2013). The floral organ recognition MADS-box family has been categorized into A, B, C, D, and E classes. Among these classes, class E was shown to be associated with reproductive organ development (Hussin et al. 2021). Indeed, activation or repression of this transcription factor leads to the development of nodes to productive or non-productive nodes (Ditta et al. 2004; Gao et al. 2010; Liu et al. 2013).

Gene expression data provided by Severin et al. (2010) noted that 20 candidate genes for PP that were detected using the SVR-mediated GWAS were expressed in flowers, 1 cm pod (7 DAF), pod shell (10-13 DAF), pod shell (14-17 DAF) and seeds. In PP, most of the genes detected by SVR-mediated GWAS are associated with auxin influx carrier or auxin response factors (ARFs), gibberellin synthesis, and response to brassinosteroid (Lin et al. 2020; Yin et al. 2018). Song et al. (2020) and Li et al. (2018a) also reported
that some genes related to PP were associated with embryo development, stamen development, ovule development, cytokinin biosynthesis, and response gibberellin that we also identified in our study. Soybean seed yield significantly depends on seed number and seed size (Liu et al. 2010; Rotundo et al. 2009). These two factors are determined from fertilization to seed maturity. Therefore, soybean seed development can be divided into three stages or phases: pre-embryo or seed set, embryo growth or seed growth, and desiccation stages or seed maturation phases (Ruan et al. 2012; Weber et al. 2005). In Arabidopsis, a complex signaling pathway and regulatory networks, including sugar and hormonal signaling, transcription factors, and metabolic pathway, have been reported to be involved in seed development (Le et al. 2010; Orozco-Arroyo et al. 2015). Several key genes and transcription factors (e.g., LEAFY COTYLEDON 1 (LEC1), LEC2, FUSCA3 (FUS3), AGAMOUS-LIKE15 (AGL15), ABSCISIC ACID INSENSITIVE 3 (ABI3), YUCCA10 (YUC10), ARFs) have been determined to control several downstream plant growth regulators pathways to the seed development (Lepiniec et al. 2018; Pelletier et al. 2017; Sun et al. 2010). Indeed, a high ratio of abscisic acid to gibberellic acid can regulate seed development (Figueiredo and Köhler 2018; Wang et al. 2016). The downregulation of FUS3 obtains this through repressing GA3ox1 and GA3ox2 and activating ABA biosynthesis (Weber et al. 2005). In soybean, RNA seq analysis for the seed set, embryo growth, and early maturation stages of developing seeds in two soybeans with contrasting seed size showed cell division and growth genes, hormone regulation, transcription factors, and metabolic pathway are involved in seed size and numbers (Du et al. 2017).

**Conclusion**

A better understanding of the genetic architecture of the yield component traits in soybean may enable breeders to establish more efficient selection strategies for developing high-yielding cultivars with improved genetic gains. Major yield component traits such as maturity, NP, NRNP, RNP, and PP play important roles in determining the overall yield production in soybean. This study verified the importance of those traits, using correlation and distribution analyses, in determining the total soybean seed yield. Furthermore, by testing different conventional and ML-mediated GWAS methods, this study demonstrated the potential benefit of using ML-mediated methods in GWAS. SVR-mediated GWAS outperformed all the other methods tested in this study, and therefore, it is recommended as an alternative to conventional GWAS methods with a greater power for detecting genomic regions associated with complex traits such as yield and its components in soybean, and possibly other crop species. To the best of our knowledge, this study is the first endeavor in which SVR was used for GWAS analyses in plants. In order to verify the causal relationship between identified QTL and the target phenotypic traits, we identified candidate genes within each QTL region using gene annotation procedures and information. The results demonstrated the efficiency of SVR-mediated GWAS in detecting reliable QTL that can be used for marker-assisted selections. Nevertheless, further investigation is recommended to confirm the efficiency of SVR-mediated GWAS for discovering genomic regions associated with complex traits in other plant species.

**Declarations**
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Author Contribution

ME conceptualized, designed and directed the experiments. MY-N performed the experiments, modeled, summed up, and wrote the manuscript. ST participated in candidate gene analyses; DT, DTOR, ST, IR, and ME revised the manuscript and validated the results. All authors have read and approved the final manuscript.

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Conflict of Interest

The authors declare that they have no conflict of interest.

Data Availability Statement

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

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Tables
Table 1. The list of detected QTL for soybean maturity using different GWAS methods in the tested soybean population.

| GWAS Method | Chromosome | Peak SNP position | Co-located QTL | Environment | Reference |
|-------------|------------|-------------------|----------------|-------------|-----------|
| MLM         | 2          | 2212910           | Sclero 3-g31   | NA          | (Moellers et al. 2017) |
|             | 8233782    |                   | Seed Weight 6-g1 | NA          | (Sonah et al. 2015) |
| FarmCPU     | 2          | 2212910           | Sclero 3-g31   | NA          | (Moellers et al. 2017) |
|             | 8233766    |                   | Seed Weight 6-g1 | NA          | (Sonah et al. 2015) |
|             | 20         | 37765851          | WUE 2-g53      | NA          | (Kaler et al. 2017) |
| RF          | 3          | 2978272           | Leaflet area 1-g2.1 | NA          | (Fang et al. 2017) |
|             |            |                   | Leaflet width 1-g4.1 | NA          | (Fang et al. 2017) |
|             |            |                   | Leaflet area 1-g2.2 | NA          | (Fang et al. 2017) |
|             |            |                   | Leaflet width 1-g4.2 | NA          | (Fang et al. 2017) |
|             |            |                   | Salt tolerance 1-g12 | NA          | (Kan et al. 2015) |
|             | 16         | 5730281           | Plant height 6-g17 | NA          | (Zhang et al. 2015b) |
|             |            |                   | Plant height 1-g17 | NA          | (Zhang et al. 2015b) |
|             |            |                   | First flower 4-g63 | NA          | (Mao et al. 2017) |
|             | 17         | 34757372          | SDS root retention 1-g6 | NA          | (Bao et al. 2015) |
| SVR         | 2          | 695362            | Seed linolenic 2-g1 | NA          | (Leamy et al. 2017) |
|             |            |                   | Seed linolenic 2-g2 | NA          | (Leamy et al. 2017) |
|             |            | 720134            | SDS 1-g12.1      | 2           | (Wen et al. 2014) |
|             |            |                   | SDS 1-g12.2      | 2           | (Wen et al. 2014) |
| Sample Code | Description                     | Year | Reference               |
|-------------|---------------------------------|------|-------------------------|
| 827374      | SDS 1-g12.3                     | 2014 | (Wen et al. 2014)       |
| 1595239     | Shoot Cu 1-g8                   | 2018 | (Dhanapal et al. 2018)  |
| 1689395     | Seed oil 5-g3                   | 2018 | (Sonah et al. 2015)     |
| 2438652     | Reproductive period 4-g16       | 2018 | (Zhang et al. 2015b)    |
| 2460921     | Reproductive period 2-g16       | 2018 | (Zhang et al. 2015b)    |
| 47513536    | R8 full maturity 4-g1           | 2018 | (Sonah et al. 2015)     |
| 47513572    | First flower 4-g81              | 2018 | (Mao et al. 2017)       |

*a Detected in separate environments in addition to the combined environment. 1) 2018Ridgetown, 2) 2019Ridgetwon, 3) 2018Palmyra, 4) 2019Palmyra, NA) Not found in any separate environment.

MLM: Mixed Linear Model, FarmCPU: Fixed and random model circulating probability unification, RF: Random Forest, SVR: Support Vector Regression
Table 2. The list of detected QTL for soybean yield using different GWAS methods in the tested soybean population.

| GWAS Method | Chromosome | Peak SNP position | Co-located QTL       | Environment | Reference         |
|-------------|------------|-------------------|----------------------|-------------|-------------------|
| MLM         | 5          | 34391386          | Ureide content 1-g16.1 | NA          | (Ray et al. 2015) |
|             |            |                   | Ureide content 1-g16.2 | NA          | (Ray et al. 2015) |
| FarmCPU     | 5          | 34391386          | Ureide content 1-g16.1 | NA          | (Ray et al. 2015) |
|             |            |                   | Ureide content 1-g16.2 | NA          | (Ray et al. 2015) |
| RF          | 7          | 1032587           | WUE 2-g18             | NA          | (Kaler et al. 2017) |
| SVR         | 3          | 36309302          | First flower 4-g10    | NA          | (Mao et al. 2017) |
|             |            |                   | First flower 3-g2     | NA          | (Hu et al. 2014)  |
|             |            |                   | Seed weight 4-g3      | NA          | (Hu et al. 2014)  |
|             |            |                   | Seed yield 4-g2       | NA          | (Hu et al. 2014)  |
|             |            |                   | R8 full maturity 3-g3 | NA          | (Hu et al. 2014)  |
|             |            | 37617293          | Plant height 3-g17    | NA          | (Contreras-Soto et al. 2017) |
|             |            |                   | Leaflet shape 1-g1.1  | NA          | (Fang et al. 2017) |
|             |            |                   | Leaflet shape 1-g1.2  | NA          | (Fang et al. 2017) |
|             |            |                   | Leaflet shape 1-g1.3  | NA          | (Fang et al. 2017) |
|             |            |                   | Seed set 1-            | NA          | (Fang et al.       |
| g32.1 | 2017) |
|-------|-------|
| Seed set 1- | g32.2 | NA | (Fang et al. 2017) |
| Seed yield 4-g4 | NA | (Hu et al. 2014) |
| WUE 2-g18 | NA | (Kaler et al. 2017) |
| SCN 5-g35 | NA | (Li et al. 2016) |
| Seed weight 5- | g20 | NA | (Zhang et al. 2016) |
| Seed weight 4- | g18 | NA | (Hu et al. 2014) |
| Seed yield 4-g5 | NA | (Hu et al. 2014) |
| Shoot Zn 1- | g28.1 | NA | (Dhanapal et al. 2018) |
| Shoot Zn 1- | g28.2 | NA | (Dhanapal et al. 2018) |
| Shoot Zn 1- | g29.1 | NA | (Dhanapal et al. 2018) |
| Shoot Zn 1- | g29.2 | NA | (Dhanapal et al. 2018) |
| Shoot Zn 1- | g29.3 | NA | (Dhanapal et al. 2018) |

a Detected in separate environments in addition to the combined environment. 1) 2018Ridgetown, 2) 2019Ridgetwon, 3) 2018Palmyra, 4) 2019Palmyra, NA) Not found in any separate environment.

MLM: Mixed Linear Model, FarmCPU: Fixed and random model circulating probability unification, RF: Random Forest, SVR: Support Vector Regression
Table 3. The list of detected QTL for soybean total number of nodes per plant (NP) using different GWAS methods in the tested soybean population.

| GWAS Method | Chromosome | Peak SNP position | Co-located QTL | Environmenta | Reference                  |
|-------------|------------|-------------------|----------------|--------------|-----------------------------|
| FarmCPU     | 19         | 40131952          | Pubescence density 1-g17 | NA           | (Chang and Hartman 2017)    |
|             |            |                   | Seed weight 9-g5.1 | NA           | (Copley et al. 2018)        |
| RF          | 4          | 1205787           | Shoot Ca 1-g10    | NA           | (Dhanapal et al. 2018)      |
|             | 6          | 50570624          | Seed set 1-g51.1  | NA           | (Fang et al. 2017)          |
|             |            |                   | Seed set 1-g43.1  | NA           | (Fang et al. 2017)          |
|             |            |                   | Seed set 1-g25.1  | NA           | (Fang et al. 2017)          |
|             |            |                   | Seed set 1-g43.2  | NA           | (Fang et al. 2017)          |
|             |            |                   | Seed set 1-g25.2  | NA           | (Fang et al. 2017)          |
|             |            |                   | Seed set 1-g51.2  | NA           | (Fang et al. 2017)          |
|             | 6          | 50570473          | Seed set 1-g43.3  | NA           | (Fang et al. 2017)          |
|             |            |                   | Seed set 1-g51.3  | NA           | (Fang et al. 2017)          |
|             |            |                   | Seed set 1-g25.3  | NA           | (Fang et al. 2017)          |
|             |            |                   | Pod number 1-g3   | NA           | (Fang et al. 2017)          |
|             |            |                   | Seed palmitic 2-g2| NA           | (Fang et al. 2017)          |
|             |            |                   | Seed long-chain faty acid 1-g22 | NA | (Fang et al. 2017) |
| SVR         | 6          | 50570624          | Seed set 1-g51.1  | NA           | (Fang et al. 2017)          |
|             |            |                   | Seed set 1-g43.1  | NA           | (Fang et al. 2017)          |
|                      |          |         |                        |
|----------------------|----------|---------|------------------------|
| Seed set 1-g25.1     | NA       | (Fang et al. 2017) |
| Seed set 1-g43.2     | NA       | (Fang et al. 2017) |
| Seed set 1-g25.2     | NA       | (Fang et al. 2017) |
| Seed set 1-g51.2     | NA       | (Fang et al. 2017) |
| Seed set 1-g43.3     | NA       | (Fang et al. 2017) |
| Seed set 1-g51.3     | NA       | (Fang et al. 2017) |
| Seed set 1-g25.3     | NA       | (Fang et al. 2017) |
| Pod number 1-g3      | NA       | (Fang et al. 2017) |
| Seed palmitic 2-g2   | NA       | (Fang et al. 2017) |
| Seed long-chain fatty acid 1-g22 | NA | (Fang et al. 2017) |
| WUE 2-g18            | NA       | (Kaler et al. 2017) |
| WUE 2-g18            | NA       | (Kaler et al. 2017) |
| First flower 3-g4    | NA       | (Fang et al. 2017) |
| Leaflet shape 1-g4.1 | NA       | (Fang et al. 2017) |
| Leaflet shape 1-g4.2 | NA       | (Fang et al. 2017) |
| Leaflet shape 1-g4.3 | NA       | (Fang et al. 2017) |
| Seed stearic 4-g5    | NA       | (Li et al. 2015)   |
| Node number 1-g6.1   | NA       | (Fang et al. 2017) |
| Node number 1-g6.2   | NA       | (Fang et al. 2017) |
| Pod number 1-g1.1    | NA       | (Fang et al. 2017) |
| Trait                          | Value | Reference          |
|-------------------------------|-------|--------------------|
| Pod number 1-g1.2             | NA    | (Fang et al. 2017) |
| Pode number 1-g1.3            | NA    | (Fang et al. 2017) |
| WUE 3-g31                     | NA    | (Kaler et al. 2017) |
| Seed weight, SoyNAM 14-g28    | NA    | (Xavier et al. 2016b) |
| Lodging, SoyNAM 4-g15         | NA    | (Cook et al. 2014)  |
| Branching 1-g1.1              | NA    | (Fang et al. 2017)  |
| Plant height 5-g4.2           | NA    | (Fang et al. 2017)  |
| Plant height 5-g4.3           | NA    | (Fang et al. 2017)  |
| Shoot p 1-g30                 | NA    | (Dhanapal et al. 2018) |
| 19                            | 47350110 | (Fang et al. 2017)  |

* Detected in separate environments in addition to the combined environment. 1) 2018Ridgetown, 2) 2019Ridgetwon, 3) 2018Palmyra, 4) 2019Palmyra, NA) Not found in any separate environment.

MLM: Mixed Linear Model, FarmCPU: Fixed and random model circulating probability unification, RF: Random Forest, SVR: Support Vector Regression
Table 4. The list of detected QTL for soybean total number of non-reproductive nodes per plant (NRNP) using different GWAS methods in the tested soybean population.

| GWAS Method | Chromosome | Peak SNP position | Co-located QTL                | Environment   | Reference            |
|-------------|------------|-------------------|------------------------------|---------------|----------------------|
| MLM         | 15         | 10193796          | Seed protein 6-g2            | NA            | (Zhang et al. 2018)  |
|             |            |                   | Seed Arg 1-g4                | NA            | (Zhang et al. 2018)  |
|             |            |                   | Seed coat luster 1-g1.3       | NA            | (Fang et al. 2017)   |
| FarmCPU     | 15         | 10193796          | Seed protein 6-g2            | NA            | (Zhang et al. 2018)  |
|             |            |                   | Seed Arg 1-g4                | NA            | (Zhang et al. 2018)  |
|             |            |                   | Seed coat luster 1-g1.3       | NA            | (Fang et al. 2017)   |
| RF          | 1          | 54647498          | First flower 4-g2            | NA            | (Mao et al. 2017)    |
|             | 7          | 329800            | Phytoph 2-g32                 | NA            | (Qin et al. 2017)    |
|             |            |                   | Phytoph 2-g7                  | NA            | (Qin et al. 2017)    |
|             | 18         | 12945778          | SCN 4-g14                    | NA            | (Vuong et al. 2015)  |
|             | 19         | 40218800          | Seed weight 9-g5.1           | NA            | (Copley et al. 2018) |
| SVR         | 7          | 1032587 \(^2\)    | WUE 2-g18                    | 2             | (Kaler et al. 2017)  |
|             | 19         | 40218800          | Seed weight 9-g5.1           | NA            | (Copley et al. 2018) |

\(^a\) Detected in separate environments in addition to the combined environment. 1) 2018Ridgetown, 2) 2019Ridgetwon, 3) 2018Palmyra, 4) 2019Palmyra, NA) Not found in any separate environment.

MLM: Mixed Linear Model, FarmCPU: Fixed and random model circulating probability unification, RF: Random Forest, SVR: Support Vector Regression
Table 5. The list of detected QTL for soybean total number of reproductive nodes per plant (RNP) using different GWAS methods in the tested soybean population.

| GWAS Method | Chromosome | Peak SNP position | Co-located QTL          | Environment a | Reference               |
|-------------|------------|-------------------|-------------------------|---------------|-------------------------|
| RF          | 9          | 40285014          | Shoot Fe 1-g8.1         | NA            | (Dhanapal et al. 2018) |
|             |            |                   | Shoot Fe 1-g8.2         | NA            | (Dhanapal et al. 2018) |
|             |            |                   | Shoot Fe 1-g8.3         | NA            | (Dhanapal et al. 2018) |
|             |            |                   | Shoot Fe 1-g9           | NA            | (Dhanapal et al. 2018) |
|             |            |                   | Shoot Fe 1-g10          | NA            | (Dhanapal et al. 2018) |
|             |            |                   | Shoot Fe 1-g11          | NA            | (Dhanapal et al. 2018) |
|             |            |                   | Soybean mosaic virus 2-g5 | NA           | (Che et al. 2017)      |
| 15          | 34958361   | SCN 5-g35         | NA                      | (Li et al. 2016) |
| SVR         | 7          | 1032587           | WUE 2-g18               | NA            | (Kaler et al. 2017)    |
| 15          | 34958361   | SCN 5-g35         | 1                       | (Li et al. 2016) |

a Detected in separate environments in addition to the combined environment. 1) 2018Ridgetown, 2) 2019Ridgetower, 3) 2018Palmyra, 4) 2019Palmyra, NA) Not found in any separate environment.

MLM: Mixed Linear Model, FarmCPU: Fixed and random model circulating probability unification, RF: Random Forest, SVR: Support Vector Regression
Table 6. The list of detected QTL for soybean total number of pods per plant (PP) using different GWAS methods in the tested soybean population.

| GWAS Method | Chromosome | Peak SNP position | Co-located QTL                  | Environment | Reference            |
|-------------|------------|-------------------|--------------------------------|-------------|----------------------|
| RF          | 7          | 15331676          | Seed weight, SoyNAM 14-g11     | NA          | (Xavier et al. 2016b) |
|             | 19         | 42300695          | First flower 4-g77             | NA          | (Mao et al. 2017)    |
|             |            |                   | Lodging, SoyNAM 4-g17          | NA          | (Cook et al. 2014)   |
| SVR         | 9          | 39366957          | Pod number 1-g4.1              | NA          | (Fang et al. 2017)   |
|             |            |                   | Pod number 1-g4.2              | NA          | (Fang et al. 2017)   |
|             |            |                   | Pod number 1-g4.3              | NA          | (Fang et al. 2017)   |
|             |            |                   | Seed thickness 2-g4             | NA          | (Fang et al. 2017)   |
|             | 9          | 39372117          | Seed Thr 2-g1                  | NA          | (Li et al. 2018b)    |
|             |            |                   | Seed Ser 2-g1                  | NA          | (Li et al. 2018b)    |
|             |            |                   | Seed Tyr 2-g2                  | NA          | (Li et al. 2018b)    |
|             |            |                   | Seed Lys 2-g2                  | NA          | (Li et al. 2018b)    |
|             |            |                   | Seed Ileu 2-g2                 | NA          | (Li et al. 2018b)    |
|             |            |                   | Seed Ile 2-g2                  | NA          | (Li et al. 2018b)    |
|             |            |                   | Seed Ala 2-g2                  | NA          | (Li et al. 2018b)    |
|             |            |                   | Seed Gly 2-g2                  | NA          | (Li et al. 2018b)    |
|             | 11         | 5245870           | Ureide content 1-g29           | NA          | (Ray et al. 2015)    |
|             |            |                   | Pod number 1-g6                | NA          | (Fang et al. 2017)   |
|             | 18         | 55645699          | Leaflet shape 1-g4.1           | NA          | (Fang et al.         |
| ID          | Descriptor                        | Value | Reference                      |
|-------------|-----------------------------------|-------|--------------------------------|
| 55469601    | Leaflet shape 1-g4.2              | NA    | (Fang et al. 2017)             |
|             | Leaflet shape 1-g4.3              | NA    | (Fang et al. 2017)             |
|             | Seed stearic 4-g5                 | NA    | (Li et al. 2015)               |
|             | Node number 1-g6.1                | NA    | (Fang et al. 2017)             |
|             | Node number 1-g6.2                | NA    | (Fang et al. 2017)             |
|             | Pode number 1-g1.1                | NA    | (Fang et al. 2017)             |
|             | Pode number 1-g1.2                | NA    | (Fang et al. 2017)             |
|             | Pode number 1-g1.3                | NA    | (Fang et al. 2017)             |
|             | WUE 3-g31                         | NA    | (Dhanapal et al. 2015a)        |
|             | Seed weight, SoyNAM 14-g28        | NA    | (Xavier et al. 2016b)          |
|             | Lodging, SoyNAM 4-g15             | NA    | (Cook et al. 2014)             |
|             | Branching 1-g1.1                  | NA    | (Fang et al. 2017)             |
|             | Plant height 5-g4.2               | NA    | (Fang et al. 2017)             |
|             | Plant height 5-g4.3               | NA    | (Fang et al. 2017)             |
|             | Shoot p 1-g30                     | NA    | (Dhanapal et al. 2018)         |
|             | Seed yield, SoyNAM 7-g19          | NA    | (Cook et al. 2014)             |
|             | R8 full maturity, SoyNAM 13-g19   | NA    | (Cook et al. 2014)             |
|             | Plant height 5-g4.3               | NA    | (Fang et al. 2017)             |
| 19          | 43077182                          | Seed weight 9-g5.2 | NA | (Copley et al. 2018) |
| Trait                          | Value | Reference                  |
|-------------------------------|-------|----------------------------|
| Seed weight 5-g21             | NA    | (Copley et al. 2018)       |
| First flower 5-g3             | NA    | (Fang et al. 2017)         |
| First flower 5-g17            | NA    | (Fang et al. 2017)         |
| 47235604 First flower 4-g77   | NA    | (Mao et al. 2017)          |
| Seed palmitic 1-g19           | NA    | (Priolli et al. 2015)      |
| 47350110 Leaf carotenoid content 1-g14 | NA | (Dhanapal et al. 2015b) |
| Ureide content 1-g50.3        | NA    | (Ray et al. 2015)          |
| Ureide content 1-g50.4        | NA    | (Ray et al. 2015)          |
| 47224293 Node number 1-g2.3   | NA    | (Fang et al. 2017)         |

\(^a\) detected in separate environments in addition to the combined environment. 1) 2018Ridgetown, 2) 2019Ridgetwon, 3) 2018Palmyra, 4) 2019Palmyra, NA) Not found in any separate environment.

MLM: Mixed Linear Model, FarmCPU: Fixed and random model circulating probability unification, RF: Random Forest, SVR: Support Vector Regression

**Figures**
Figure 1

LD decay distance in the tested 227 soybean genotypes.
Figure 2

The distribution of seed yield (A), maturity (B), NP (C), NRNP (D), RNP (E), and PP (F) in 227 soybean genotypes across four environments. The estimated heritability is provided for each of the six traits. RNP: Total number of reproductive nodes per plant, NRNP: The total number of non-reproductive nodes per plant, NP: The total nodes per plant, PP: The total number of pods per plant.
Figure 3

The distributions and Pearson correlations among the soybean seed yield, maturity, and yield component traits. RNP: Total number of reproductive nodes per plant, NRNP: The total number of non-reproductive nodes per plant, NP: The total nodes per plant, PP: The total number of pods per plant. The heat map scale for values is provided by colour for the panel.
Figure 4

Structure and kinship plots for the 227 soybean genotypes. The x-axis is the number of genotypes used in this GWAS panel, and the y axis is the membership of each subgroup. G1-G7 stands for the subpopulation.
Figure 5

Genome-wide Manhattan and quantile-quantile plots for GWAS studies of A) maturity and B) seed yield in soybean using MLM, FarmCPU, RF, and SVR methods, from top to bottom, respectively.

Figure 6
Genome-wide Manhattan and quantile-quantile plots for GWAS studies of A) the total number of nodes (NP) and B) the total number of non-reproductive nodes (NRNP) in soybean using MLM, FarmCPU, RF, and SVR methods, from top to bottom, respectively.

Figure 7

Genome-wide Manhattan and quantile-quantile plots for GWAS studies of A) The total number of reproductive nodes (RNP) and B) the total number of pods (PP) in soybean using MLM, FarmCPU, RF, and SVR methods, from top to bottom, respectively.
Figure 8

The average effects of reference allele and alternative allele from the detected SNP’s peak for maturity (A), seed yield (B), NP (C), NRNP (D), RNP (E), and PP (F) in 227 soybean genotypes across four environments. RNP: Total number of reproductive nodes per plant, NRNP: The total number of non-reproductive nodes per plant, NP: The total nodes per plant, PP: The total number of pods per plant.

Supplementary Files

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- TableS1.pdf