High-density genetic map using whole-genome resequencing for fine mapping and candidate gene discovery for disease resistance in peanut

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Summary
Whole-genome resequencing (WGRS) of mapping populations has facilitated development of high-density genetic maps essential for fine mapping and candidate gene discovery for traits of interest in crop species. Leaf spots, including early leaf spot (ELS) and late leaf spot (LLS), and Tomato spotted wilt virus (TSWV) are devastating diseases in peanut causing significant yield loss. We generated WGRS data on a recombinant inbred line population, developed a SNP-based high-density genetic map, and conducted fine mapping, candidate gene discovery and marker validation for ELS, LLS and TSWV. The first sequence-based high-density map was constructed with 8869 SNPs assigned to 20 linkage groups, representing 20 chromosomes, for the ‘T’ population (Tifrunner × GT-C20) with a map length of 3120 cM and an average distance of 1.45 cM. The quantitative trait locus (QTL) analysis using high-density genetic map and multiple season phenotyping data identified 35 main-effect QTLs with phenotypic variation explained (PVE) from 6.32% to 47.63%. Among major-effect QTLs mapped, there were two QTLs for ELS on B05 with 47.42% PVE and B03 with 47.38% PVE, two QTLs for ELS on A05 with 47.63% and B03 with 34.03% PVE and one QTL for TSWV on B09 with 40.71% PVE. The epistasis and environment interaction analyses identified significant environmental effects on these traits. The identified QTL regions had disease resistance genes including R-genes and transcription factors. KASP markers were developed for major QTLs and validated in the population and are ready for further deployment in genomics-assisted breeding in peanut.

Introduction
Peanut, Arachis hypogaea (2n = 4x = 40), is an allotetraploid with an AABB genomic constitution which originated from a single recent hybridization and spontaneous tetraploidization of two ancestral diploid species, A. duranensis (A-genome) and A. ipaensis (B-genome) approximately 4000–6000 years ago (Bertioli et al., 2016; Halward et al., 1992; Moretzsohn et al., 2013). The relatively short evolutionary history of domesticated peanut and the presence of hybridization barriers between diploid and tetraploid species has been one of the major reasons for the narrow genetic base in cultivated peanuts. This low level of diversity has been a major hurdle for mining large-scale polymorphic markers, such as simple-sequence repeats (SSRs) for construction of high-density genetic maps and QTL studies (Pandey et al., 2016; Qin et al., 2012; Varshney et al., 2013).

The genetic yield potential of peanut cultivars has been continuously challenged by several diseases including early leaf spot (ELS) caused by Cercospora arachidicola, late leaf spot (LLS) caused by Cercosporidium personatum and Tomato spotted wilt virus (TSWV). These foliar diseases cause yield losses of up to 70%, resulting in approximately $600 million in losses (Food and Agriculture Organization (FAO), 2004; Ogwulumba et al., 2008). While insecticides and fungicides have been used as part of an integrated pest management approach, breeding disease-resistant cultivars with high yield and good agronomic performance is the most economical and sustainable solution (Guo et al., 2013; Pandey et al., 2012; Varshney et al., 2013).

Genomics-assisted breeding (GAB) has demonstrated great potential for accelerated development of improved varieties (Varshney et al., 2009) including peanut. However, it is worth mentioning that success of the diagnostic markers in breeding...
depends on the precise phenotyping and high-density genotyping. The earlier efforts towards identifying linked markers for leaf spot and TSWV resistance were based on SSR-based genetic maps using T-population (Tifrunner × GT-C20) (Pandey et al., 2014, 2017a; Qin et al., 2012; Wang et al., 2012). After making use of available SSR resources in the public domain, a total of 418 SSR loci could be mapped leading to identification of several QTLs with <20% phenotypic variation explained (PVE) for these diseases (Pandey et al., 2017a). The identified genomic regions were also too large to be exploited in diagnostic marker development.

The last decade has witnessed rapid progress in genome sequencing that greatly helped in high-resolution trait mapping, candidate gene discovery and breeding applications in many legumes (Pandey et al., 2016). Single nucleotide polymorphisms (SNPs) have high frequency of occurrence throughout the genome and are genetic markers of choice for several genetic and breeding applications. Recent availability of reference genomes for both the progenitor species, namely, A. duranensis (Bertioli et al., 2016; Chen et al., 2016) and A. ipaensis (Bertioli et al., 2016), has made the application of next-generation sequencing (NGS) approach a possibility in peanut. Bertioli et al. (2016) not only provided insights on genome architecture and evolution but also opened opportunities for developing other required genomic tools and technologies for deployment in different genetic and breeding applications. For example, these reference genomes allowed for the construction of a high-density genotyping array with >58K highly informative SNPs (Pandey et al., 2017b) and SNP identification using RNA-seq data (Nayak et al., 2017), which further opens an array of options for deeper exploration of genome and gene discovery. Further, the accelerated pace of developments in sequencing technologies has made sequencing more affordable due to increased throughput data generation and competition in the market.

We resequenced the parental genotypes and the RIL population for conducting high-resolution genetic mapping and development of diagnostic markers for resistance to leaf spots and TSWV for peanut breeding application. The detailed sequence analysis of a RIL population has facilitated QTL discovery for both the diseases followed by candidate gene discovery and marker identification. This study also developed and validated PCR-based KASP (Kovetitive allele-specific polymerase chain reaction) markers which can now be deployed in marker-assisted selection. This study also provided the first SNP-based high-density genetic map currently available for cultivated peanut which can be used for a variety of applications including the improvement of tetraploid peanut reference genome assemblies.

Results

Variation in disease severity in parents and RIL population

The evaluation of disease resistance was all conducted in the field based on natural infection. We did not experience any interactions among these three diseases. As the causal organism is different for each disease, there were no any studies to investigate the possible interaction among these diseases. The distributions of the ratings of disease severity for ELS, LLS and TSWV were relatively normal except a few instances, where some individuals of the population showed extreme phenotypes and were out of the normal curve (Figure S1). Overall, the quantitative nature of the investigated traits was observed. The phenotypes for disease severity ratings of two parental genotypes and RILs showed significant differences among environments across the environments (associated with major QTLs) (Table S1). Disease severity of ELS for the parental lines ranged from 2.3 to 3.3 for ‘Tifrunner’ and from 3.8 to 7.0 for GT-C20 during 2009–2013. ELS severity in RILs ranged from 1.0 to 8.0 over the same period. In case of LLS, disease ratings for Tifrunner ranged from 5.0 to 6.5, 7.2 to 10.0 for GT-C20 and 4.7 to 10.0 among RILs over the same period. TSWV severity in the parents ranged from 1.7 to 2.3 for Tifrunner and 4.7 to 5.7 for GT-C20, while the disease severity among RILs ranged from 1.0 to 6.7 (Figure S1).

Sequencing of the RIL population and SNP discovery

Over one Tb of filtered data (~23 billion reads) was generated for the parental lines and 91 selected RILs. The first parent, Tifrunner, was sequenced at 100X and the second parent, GT-C20, at 10X coverage, while the RIL population individuals were each sequenced at 2–5X coverage (Table S2). After quality filtering, an average read length of 93, 96 and 97 bp was obtained for Tifrunner, GT-C20 and individual RILs, respectively. In total, ~40% of sequence data were mapped on A-subgenome and ~60% on B-subgenome except for line T92 where more than 60% of reads were mapped on A-subgenome. Although the amount of data produced from each plant sample was different, the proportion of mapped reads on the respective genomes was similar for each of the individual plants with the exception of T92 (Figure S2). It could be due to the exchanges between A- and B-subgenomes, as reported by Leal-Bertioli et al. (2015) that autotetraploid-like tetrasomic recombination is quite frequent in RIL population derived from a cultivated peanut. As Tifrunner has considerable tetrasomic regions and the RIL population with Tifrunner as a parental line could observe these changes in the progeny. All the reads mapped to the reference genome were used for haplotype-based SNP calling. A total of 97 571 SNPs were detected between the two parental lines, and 18 252 of those remained after removing the low-quality SNPs (Table S3). The SNPs were distributed throughout the 20 linkage groups (LGs) with the highest number of SNPs occurring on chromosome A06 (2771) and the fewest on chromosome B07 (261). None even distribution of SNPs on the genomes could also be attributed to the presence of autotetraploid-like tetrasomic regions (either AAAAA or BBBB) in the genomes, where the chances of finding the SNPs are almost negligible (Clevenger et al., 2017; Leal-Bertioli et al., 2015). RILs thus obtained from Tifrunner as one of the parent are expected to have lesser or no markers identified in the regions with tetrasomy. Of the 18 252 high-quality SNPs, 16 674 SNPs could genotype the population and were polymorphic. The other 1578 SNPs could not be genotyped in the population, which could be due to the errors in sequencing of those parental genotypes or difference in sequence depth of the parents and the RILs. One of the parents, Tifrunner, was sequenced at 100X depth, and GT-C20 was at about 10X. However, the RILs were sequenced at much lower depth (2X to 5X). Of these, 10 274 SNPs showed less than 20% missing data and no segregation distortion, and were used for genetic mapping (Figure S3).

Most dense genetic map with homologous and translocated markers

Of the 10 274 SNP markers obtained using haplotype SNP mining, 8869 SNPs were mapped onto 20 LGs spanning a genetic map length of 3120.71 cM with map density of 1.45 cM/focus. Many SNPs occupied the same genetic loci on the LGs; therefore, a total of 2156 marker loci covering 8869 SNPs were mapped on
the 20 LGs (Figure S4). The mapped marker loci per LG ranged from 38 (B07) to 179 (A03) with an average of 107.8 loci/LG. A- and B-genome LGs were identified with 1219 marker loci covering 1637.8 cm (1.34 avg. marker distance) and 937 marker loci with 1484.91 cm genetic map distance (1.58 avg. marker distance), respectively (Table 1). Overall, the SNP markers were usually densely and homogeneously distributed along the 20 LGs except gaps of more than 10 cm each on LGs A01, A07, A08, B01 and B08, and five such gaps on B07 between two contiguous markers.

Homeologous and translocated SNP markers were also identified in the current genetic map. A total of 422 homeologous markers identified from A-subgenome were mapped on B-subgenome. Similarly, 317 homeologous SNPs from B-subgenome were mapped on the A-subgenome (Table 1, Figure 1a, b). A total of 309 and 104 translocated markers were identified on LGs A01-A10 and B01-B10, respectively. For example, LGs A03 and A06 have 99 (68 from A-chromosomes; 31 from B-chromosomes) and 69 (45 from A-chromosomes; 25 from B-chromosomes) translocated markers, respectively (Tables 1, S4). Largely, the markers were seen to have been moved from the end of each chromosome to corresponding and other LGs (Figure 1a, b).

We also considered the major QTL (PVE ≥ 10%) defining SSR markers reported in our earlier study (Pandey et al., 2017a) in the current map. Of the 19 major QTL-linked markers reported earlier, 15 were mapped to the current LGs. However, integration of these markers increased the genetic distance of the respective LGs (Table S5). Thirteen of these 15 markers were mapped to the same LGs as previously reported, but remaining two markers were mapped to LG B03 instead of A03 as reported earlier. The two SSR loci (TC38F01, GM1986-2) on A07 were the first and the last on that linkage group, and all the SNPs identified in the current study were harboured between these two SSR loci.

### Table 1 Summary of total number of SNP markers across 20 linkage groups

| Linkage groups | Total markers on map | A-markers | B-markers | Translocated markers | Av. Interval (cM) | Length (cM) | Loci/Homologous | Homeologous |
|----------------|----------------------|-----------|-----------|----------------------|------------------|-------------|----------------|-------------|
| A01            | 278                  | 260       | 13        | 5                    | 1.48             | 176.02      | 119            |             |
| A02            | 346                  | 296       | 27        | 23                   | 1.3              | 185.7       | 143            |             |
| A03            | 1222                 | 1047      | 76        | 99                   | 1.08             | 192.46      | 179            |             |
| A04            | 676                  | 623       | 33        | 20                   | 1.43             | 153.52      | 107            |             |
| A05            | 701                  | 647       | 36        | 18                   | 1.22             | 190.24      | 156            |             |
| A06            | 1644                 | 1529      | 46        | 69                   | 1.24             | 157.13      | 127            |             |
| A07            | 406                  | 368       | 15        | 23                   | 1.76             | 89.59       | 51             |             |
| A08            | 244                  | 210       | 13        | 21                   | 1.54             | 181.7       | 118            |             |
| A09            | 475                  | 426       | 33        | 16                   | 1.47             | 154.56      | 105            |             |
| A10            | 307                  | 267       | 25        | 15                   | 1.38             | 156.88      | 114            |             |
| B01            | 304                  | 29        | 272       | 3                    | 1.78             | 153.18      | 86             |             |
| B02            | 311                  | 37        | 260       | 14                   | 1.37             | 143.5       | 105            |             |
| B03            | 373                  | 153       | 215       | 5                    | 1.29             | 175.91      | 136            |             |
| B04            | 164                  | 26        | 134       | 4                    | 1.5              | 136.25      | 91             |             |
| B05            | 334                  | 40        | 282       | 12                   | 1.36             | 142.86      | 105            |             |
| B06            | 283                  | 41        | 237       | 5                    | 1.58             | 163.98      | 104            |             |
| B07            | 50                   | 6         | 35        | 9                    | 4.28             | 162.58      | 38             |             |
| B08            | 159                  | 8         | 135       | 16                   | 1.5              | 85.54       | 57             |             |
| B09            | 252                  | 28        | 204       | 20                   | 1.51             | 173.3       | 115            |             |
| B10            | 340                  | 54        | 270       | 16                   | 1.46             | 145.81      | 100            |             |

Highlighted in grey are the homeologous markers on the LGs of A- and B-subgenomes.

### QTLs associated with disease resistance

Quantitative trait locus mapping resulted in the identification of 35 QTLs with 6.3–47.6% PVE and LOD values between 2.5 and 11.3 associated with ELS, LLS and TSWV (Figures 2 and S5; Table 2). Of these 35 QTLs mapped onto 12 LGs, 31 were identified as major QTLs. Twenty QTLs were mapped onto six LGs of the A-subgenome, and the remaining 15 QTLs were mapped onto six LGs of B-subgenome. On the A-subgenome, LG A06 harboured one minor QTL while the remaining 16 major QTLs were mapped on LGs A01, A03, A04, A05, A06 and A08. The B-subgenome contained 15 major QTLs of the total 20 QTLs present on LGs B02, B03, B05, B06, B09 and B10.

For ELS, seven major QTLs were found on LGs A01, A03, A05, A06, A08, B03 and B05 with PVE ranging from 10.4% (qELS_T13_A01) to 47.4% (qELS_T09_B05). Five QTLs were present on the five LGs of A subgenome and the rest two on B03 and B05. Of all the seven major QTLs for ELS, qELS_T09_B05 exhibited 47.4% PVE covering 2.3 Mb on the physical map. Two other QTLs, qELS_T11_B03_2 and qELS_T11_B03_1, mapped on B03 showed 47.3% and 20.35% PVE covered 80.13 Kb and 99.2 Kb, respectively, on the physical map. Both the QTLs with more than 47% PVE were contributed by the resistant parent Tifrunner.

Interestingly, SSRs flanking the LLS QTL were found to be within the SSRs marking the ELS QTL in the current genetic map of A05 (Figure S6). We identified 308 SNPs between the SSRs (TC40D04–GM1878) from an earlier report (Pandey et al., 2017a) on the current genetic map. Six of these 308 SNPs were found associated with three major QTLs for leaf spot on LG A05. First QTL (qELS_T12_A05_4) was flanked by A05_20504728 and B09_138649848, which showed 10.27% PVE. The second major
QTL identified was qLLS_T12_A05_3 flanked by A05_20406182 and B05_20992208, which showed 47.8% PVE. The third QTL (qLLS_T13_A05_6) was flanked by A05_10286363 and A05_11746142 with 10.36% PVE. Seven of 15 QTLs including the one with maximum 47% PVE were found on LG A05, which is consistent with the earlier findings, where A05 contained the maximum number of QTLs (Pandey et al., 2017a).

In the case of TSWV, of the total, 13 QTLs were identified, and nine of these were major QTLs. These QTLs were present on LGs A03, A04, A08, B03, and B09 ranging from PVE of 10.8% (qTSWV_T10_B09_2) on B03 to 40.7% on B09 (Table 2). QTL on B03 with 10.8% PVE was flanked by SNPs identified from chromosome A03 (A03_131914876 and A03_131407286). Two common QTLs were identified between earlier study by Pandey et al. (2017a) and present study. These QTLs were qTSW_T10_A04 and qTSW_T10_B02 for July 2010 and August 2010. The other QTLs identified in the current study were novel.

Environment interaction QTLs associated with disease resistance

A total of 13 QTL x E (environment) interactions were detected for three traits, of which nine QTLs were falling in the same genomic region, where major QTLs were also detected (Table 56). A total of three QTL x E interactions were detected for TSWV, and two of these were mapped on B09 with 3.42 and 8.35% PVE and another on A08 with 2.59%. In case of LLS, three QTL x E interactions were also identified out of which two were on A05 with 2.17%–4.82% and another on B03 with 5.95%. A total of seven QTL x E interactions were identified for ELS on B05, A08, B03, B09, A04, B04 and A08 with 11.89, 4.95, 3.75, 3.00, 2.82, 2.67 and 2.66% PVE, respectively.

Epistatic QTLs associated with disease resistance

A total of 1048 epistatic QTL interactions were observed which included 268 for ELS, 505 for LLS and 275 for TSWV across the multiple environments. A maximum of 81 interactions for ELS_72_Tift_2013 and minimum of 28 interactions for LLS_93_Tift_2011 were identified. These interactions possessed both the positive and the negative additive effects on the traits. These epistatic interactions showed varied range of contribution towards the phenotypes as the PVE contributed by these detected environmental QTL (e-QTL) were up to 59.43% PVE for ELS (ELS_8_Tift_2011), 44.91% PVE for LLS (LLS_91_Tift_2011) and 55.30% for TSWV (TSW_8_Tift_2010) (Table S7).

Genomic region(s) and putative candidate genes associated with leaf spot and Tomato spotted wilt virus resistance

For ELS, two QTLs were identified with over 47% PVE on LG B03 (qELS_T11_B03) and B05 (qELS_T11_B03). The QTL mapped on LG B03 was flanked by markers A03_133651613 and A03_133731756, covering physical map distance of only 80.13 Kb. Of this 80 Kb region, only 10.5 Kb encompassed two genes and the remaining 69.5 Kb region was intergenic. The two genes in this region code for serine hydroxymethyl trans-ferase (SHMT) and a rhodanese cell cycle control phosphatase superfamily protein. To scan the nearby regions, we looked for the genes 200 Kb upstream and downstream of the QTL. Flanking regions contained genes coding for xylolucan endo-transglucosylase/hydrolase, S-adenosyl-L-homocysteine hydrolase and methyltransferase-like protein, ATP/DNA-binding protein, and small ubiquitin-like modifier. A total of 15 SNPs were identified within the genes identified in these genes with a maximum of
three SNPs each in two genes coding for UDP-glycosyltransferase superfamily protein (*Aradu.VB4ZI*) and rhodanese/cell cycle control phosphatase superfamily protein (*Aradu.C56U2*) (Table S8). Other QTL on B05 was flanked by B05_22527171 and B05_20207815, encompassing >2.0 Mb distance on physical map. The 2.0 Mb QTL harboured 83 genes, including pectin esterase inhibitor, protein kinase, pentatricopeptide, NB-ARC disease resistance, WRKY TF, F-box/LRR-repeat, MATE efflux family protein, cell wall protein-like structure involved in cellulose microfibril organization, phosphotransferases and pathogenesis-related genes as major defence-related genes. The identified genes harboured 18 SNPs with three of these SNPs in gibberellin 2-beta-dioxygenase 8-like protein-coding gene (*Araip.A06C0*).

For LLS, a QTL on LG B03 flanked by A03_134198144 and A03_134634474 was reported with a PVE of 34%. It covered 0.43 Mb on the physical map encompassing a total of 29 genes including six unknown/uncharacterized protein-coding genes. The other 23 genes code for proteins including a protein kinase family protein (leucine-rich repeat-containing N-terminal), receptor-like kinase (leucine-rich repeat), WRKY TF, and heat-shock transcription factor, glutathione S-transferase, NADH:ubiquinone oxidoreductase intermediate-associated protein, major intrinsic protein (MIP) family transporter (aquaporin like), zinc finger family protein, cinnamyl alcohol dehydrogenase, tetratricopeptide and transporter proteins. These genes are known to play an important role in plant defence against pathogen response including...
necrotrophic fungi. Genes within this QTL contained 10 SNPs. Another major QTL on LG A05 with 47% PVE covering 2 Mb harboured 78 genes coding for MATE efflux family protein, alanine and tryptophan aminotransferases, pathogenesis-related protein, histone deacetylase, proteasome inhibitor, serine/threonine phosphatase, receptor kinase, protein kinase, pyruvate kinase, serine carboxypeptidase, ABC transporter family protein, xylolucan xylantransferase and peptidase M50 family protein with 18 SNPs falling in the genes.

The major QTL for TSWV with more than 40% PVE on LG B09 was flanked by B09_5189475 and B09_6739506 and covered 1.55 Mb of physical distance, encompassing 114 genes including histone acetyltransferase, heat-shock protein, ATP-binding ABC transporter, glutathione S-transferase, cytochrome P450 superfamily protein, protein kinase superfamily protein, receptor-like serine/threonine kinase, root hair defective 3 homolog 1-like, MATE efflux, ethylene-insensitive 3 family protein, disease resistance (TIR-NBS-LRR), glutamate dehydrogenase and others.
involved in direct or indirect defence responses against pathogens (Table S8). A total of 25 SNPs were identified within the genes in this QTL with a maximum of three effective SNPs in a putative candidate gene coding for protein kinase family protein (Araip.RN7PY).

SNP efficiency and validation in RIL population

All SNPs present within the genes and in the flanking region of major QTLs were annotated for their effect and impact on gene function. SNPs within the coding region of a gene showed either a synonymous (does not change protein sequence) or a nonsynonymous (changes protein sequence) effect with moderate, high or low impact on gene function. Other SNPs were present either in the introns or untranslated regions (5' or 3') of a gene (Table S8). Five SNPs associated with ELS and LLS QTLs were also confirmed for their in silico genotype calls using KASP assay in the population. Overall, these SNPs showed efficiency of over 90%, implying that most of the in silico genotype calls were validated using the PCR-based KASP assay (Table S9). KASP assay could clearly distinguish between the allelic variations in the population (Figure 3). To study correlation between the phenotype and KASP genotyping, we considered the extreme resistant and susceptible lines based on their phenotype ratings. One of the five markers associated with ELS (B05_22527171) showed not only good correlation with the in silico identified SNP calls, but also showed strong correlation with the phenotyping data of the RILs genotyped using this marker. Phenotype for the resistant lines carrying allele from the resistant parent (Tifrunner) genotyped using B05_22527171 averaged 3.5 and the susceptible lines carrying the alleles from susceptible parent (GT-C20) averaged 6.2 (Figure 4). Other four markers (A03_134198144, A05_82270000, A05_20406182 and A06_14301316), however, did not show very strong correlation with the phenotyping ratings.

Discussion

Evolution in sequencing and genotyping technologies has allowed for rapid advances in conducting genetic and breeding studies. Currently, these technologies have become not only cost-effective and high throughput, both also provided opportunities to get deeper insights into the target genomic locations (Pandey et al., 2016). More and more deployment of these technologies has made available diagnostic markers for target traits for use in GAB towards accelerated crop improvement (Van Erp et al., 2013). The recent origins of cultivated, allotetraploid peanut, a lack of optimal genetic resources and narrow genetic base has so far hampered the development of dense genetic maps for candidate gene identification. However, with the recent release of the reference genomes for both the diploid peanut progenitors, namely, *A. duranensis* (A) and *A. ipaensis* (B) (Bertioli et al., 2016), the application of NGS technologies in marker discovery and high-density map generation is now possible for cultivated peanut. Therefore, here we used a WGRS strategy for genomewide SNP mining, genetic mapping and QTL analysis.

Most dense genetic map for genetics and breeding applications

This WGRS method and subsequent analyses have already been used successfully in other legumes including chickpea (Kale et al., 2015) and soya bean (Qi et al., 2014; Xu et al., 2013). Using this approach, we developed the densest genetic map currently available for cultivated peanut with 8,869 SNPs and 2156 mapped loci, which can be used for numerous applications including improvement of cultivated tetraploid peanut genome assembly (Peanutbase.org). While this genetic map has a high number of SNP markers (8869), only 24.3% (2156) of them represented the recombination as independent genetic loci. In comparison, other genetic maps such as that generated by Zhou et al. (2014) who used double-digest restriction site-associated DNA sequencing (ddRADseq) and successfully mapped 1621 SNPs and 64 SSRs, each representing an independent locus (1685 marker loci). The reduced loci:marker ratio may be due to the fact that these are haplotype-based SNPs located in close physical proximity to each other on the respective chromosomes, which are tightly linked with little recombination between them. In addition, the reduced number of RIL individuals (91) used in the present study, compared to the 166 RILs used by Zhou et al. (2014), may have resulted in reduced mapping resolution to break apart the obtained haplotypes across multiple marker loci. Despite this, the density of the current genetic map demonstrates the power of this WGRS approach in peanut as the first report of its application in a biparental mapping population for trait mapping in this crop.

B-subgenome is lengthier but less diverse than A-subgenome

More SNPs were identified and mapped on the A-subgenome LGs compared to the B-subgenome LGs despite the fact that B-genome chromosomes are physically larger than A-genome chromosomes. On this genetic map, a significant proportion of marker loci with assigned physical locations to a chromosome of one subgenome were mapped to respective homeologous positions on chromosomes of the other subgenome, indicating previous duplication events or mis-assignments in these regions of genomes of diploid peanut that are the progenitors of cultivated tetraploid peanut. Most of this homoeologous mapping of markers was seen between LGs A03 and B03, and the least were observed between A07, B07, and A08, B08 (Table S4, Figure 1a, b). Homeologous mapping has also been earlier reported in peanut between LGs A07 and B07, and A08 and B08 (Qi et al., 2012). This observation may also provide putative evidence for genetic exchanges occurring between the peanut subgenomes either during or following tetraploidization and the formation of cultivated peanut, a phenomenon also observed in earlier studies (Bertioli et al., 2016; Leal-Bertioli et al., 2015). Also, a significant number of markers were placed on the genetic map on different
chromosomes to which they were assigned. Most of these ‘translocated’ markers were seen between A06 and A10 and between A03 and A09, A10, fewer were seen on B04 (Table 1, Figure 1a, b). These ‘translocated’ markers observed might be due to reciprocal translocation occurring because of the exchange of two terminal segments between two nonhomologous chromosomes (Farré et al., 2011). As a result, recombination between loci around translocation breakpoints is suppressed leading to linkage between markers in these regions, which explains the linkage detected between markers lying on different chromosomes seen in the current study. Such translocations are well documented in various crops including barley (Farré et al., 2011), soya bean (Mahama and Palmer, 2003), Prunus spp. (Jauréguï et al., 2001), lentils (Tadmor et al., 1987), pea (Kosterin et al., 1999) and peanut (Qin et al., 2012). Some of these observed ‘translocated’ markers might be also due to artefacts and misassignments because of the highly repetitive structure of the genome (Bertioli et al., 2016).

QTLs and potential candidate genes related to defense against leaf spots and TSWV

Disease resistance is a highly heritable trait of great value to crop production systems. Efforts have been made to identify the QTLs/candidate genes for important diseases such as ELS, LLS and TSWV in peanut (Khera et al., 2016; Pandey et al., 2017a,c). In the present study, 20 QTLs were identified on A-subgenome LGs and 15 on B-subgenome LGs. Of the 35 identified QTLs, 24 (65%) were flanked by markers from A-subgenome. In contrast, Bertioli et al. (2016) identified 345 and 397 NB-LRR (nucleotide-binding-leucine-rich repeat) genes in the A and the B genomes sequenced, respectively.

In earlier studies with sparse genetic maps and few polymorphic markers, the maximum PVE of detected QTL was 27.35% (Pandey et al., 2017a; Wang et al., 2013) and up to 29.14% in another RIL population, SunOleic97R × NC 94022 (S-population) (Khera et al., 2016). All these studies resulted in broad QTL regions with no candidate gene identification being possible. The current study identified seven major QTLs for ELS with four of them having negative additive effects suggesting contribution by parent 1, Tifrunner (R), of which two were located on LGs B03 and B05 contributing over 47% PVE each (Table 2, Figure 2). In an earlier study, LG A03 was reported to harbour major ELS resistance-related QTL (Pandey et al., 2017a). The two major QTLs for ELS on LG B03 were flanked by homeologous markers from chromosome A03, suggesting that these regions of chromosome A03 may constitute a hot spot for genes responsible for ELS resistance. In a recent study using QTL-seq approach in a TAG24 × GPBD4 RIL population, a 2.98 Mb (131.67–134.65 Mb) genomic region on chromosome A03, which overlaps with the region indicated in the present study to contain these two major ELS-resistant QTLs (132.10–132.20 Mb and 133.65–133.73 Mb, respectively), was also found to contribute to LLS resistance (Pandey et al., 2017c). This supports the notion that this region on chromosome A03 has genes playing role in defence against leaf spots. However, it is also possible that although ELS and LLS classification was based on observations of predominant signs and symptoms in the fields used for disease phenotyping, both ELS and LLS likely co-occurred in the field, and the detected QTL may not be specific. Greenhouse-based phenotyping with controlled inoculations will be required to confirm this conclusion.

Major QTL regions on LG B05 for ELS, and A05 and B03 for ELS and LLS also contained leucine-rich repeat (LRR), NB-ARC, or receptor kinase genes which may function as putative R-genes. Interestingly, while the major QTL region found for ELS was also observed for LLS, but at a lesser PVE (qELS_T11_B03_2 and

Figure 4  Graph showing correlation between the average disease severity scores (phenotype) and the five KASP validated SNPs. Vertical axis represents the average of disease severity, and horizontal axis shows the markers. Disease scores for the parents, Tifrunner (P1) and GT-C20 (P2) for the season in that which QTL was detected are also shown for each marker. Unpaired t-test was performed, and P-values were estimated using the phenotyping data for number of alleles in population coming from Tifrunner and GT-C20. Extremely statistically significant correlations are marked using *** with P-value < 0.0001. Significant correlations are marked with *.

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Gaurav Agarwal et al.
qLLS_T13_B03_1), there was another major QTL for LLS on B03 associated with different markers which contained a putative R-
gene. This may indicate a degree of R-gene specificity for each
pathogen based on the observed PVEs associated with each QTL.
The association of markers from A05 and A03 based on diploid
reference sequences with major LLS QTL on LGs A05 and B03,
respectively, while the ELS R-gene containing QTL on LG B05 is
associated with markers from chromosome B05 in the diploid
reference genome on the current map may also indicate subgenome-associated specificity. Given the possibility, however,
of co-occurrence of both pathogens during the disease ratings,
future studies with controlled inoculation will be required to
explain this hypothesis more completely.

In addition to R-genes, genes coding for serine hydroxymethyl-
transferase (SHMT), rhodanese cell cycle control phosphatase
superfamily protein and ubiquitin-protein ligase were identified in
the region harboured by above-mentioned two major QTLs on LG
B03. It has been reported that mutation of shmt1 compromises
host resistance to biotrophic and necrotrophic foliar pathogens
(Moreno et al., 2005). Rhodanese is a detoxifying enzyme known
to detoxify the harmful effects of HCN produced in plants as an
detoxification of the harmful effects of HCN produced in plants as an
toxin. This may indicate a degree of R-gene specificity for each
pathogen based on the observed PVEs associated with each QTL.

Further, the epistatic QTL study detected hundreds of QTL
interactions across the genome for ELS, LLS and TSWV. Two SNP
loci (A05_20406717, A05_82270000) associated with LLS and
B05_22527171 associated with ELS were successfully validated
and KASP assay. These QTL–QTL interactions indirectly show the
cross-talk of the homeologous genes across the A and B
subgenome. This detailed study of QTL × E and QTL × QTL
along with the major QTL showed the importance of the genomic
interaction and suggest to introgress favourable genomic loci
including main effect, epistasis and QTL × Environment interac-
tions for achieving desired phenotype through the molecular
breeding programme.

Validated diagnostic markers available for use in
molecular breeding

Robust KASP genotyping assays were also developed for ELS-
and LLS-related disease traits using the SNP markers. It remarkably
demonstrated the robustness and accuracy of KASP markers as
these markers were validated in RIL population in correlation with
in silico genotype calls. Studies in other legumes like pea, pigeon
pea, chickpea, and soybean using KASP markers have shown
promising results (Boutet et al., 2016; Hiremath et al., 2012; Patil
et al., 2017; Saxena et al., 2014). Studies in chickpea, pigeon pea
and peanut identified 66.8%, 75.86% and 80% of the KASP
markers to be polymorphic (Saxena et al., 2014). Also, in the
current study, a very high level of consistency (~90%) between the
in silico called SNPs and KASP validated data was reported.
While significant, though numerically limited separation between
resistant and susceptible genotyped plants when considering
average phenotypic disease ratings could be observed for single
markers, utilizing multiple major QTL-associated KASP markers
will provide a means of improved predictive selection. These
markers can be implemented in peanut breeding programmes,
which can augment GAB of disease-related traits with greater
efficiency and accuracy. The validated KASP markers can also be
used for allele mining, marker-assisted selection and forward
breeding programmes. Further studies will be conducted to
implement these markers as diagnostic markers in other resistant
and susceptible genetic backgrounds.

Conclusions

The development of high-density genetic maps is a requirement for
the fine mapping of loci contributing to quantitative traits
such as disease resistance. Here, we have presented the densest
genetic map currently available for cultivated peanut and the first
to be generated using a WGRS approach in peanut. With this
high-density map, 35 major QTLS were identified for important
diseases of peanut including ELS (47.4% PVE), LLS (47.6% PVE)
and TSWV (40.7% PVE). This also allowed for the development of
KASP markers for SNPs associated with ELS and LLS QTLS. These
PCR-based markers provide peanut breeders a useful tool to
improve marker-assisted selection in breeding programmes. In
addition, the generation of this high-density genetic map also
allows for the correction of future genome assemblies for
tetraploid peanut.

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Materials and methods

Plant materials and generation of phenotyping data

Two peanut inbred lines, Tifrunner and GT-C20, were selected as parents to develop a RIL mapping population. Tifrunner, the female parent (runner market type) has a high level of resistance to TSWV, moderate resistance to ELS and LLS, and late maturity (Holbrook and Culbreath, 2007). GT-C20, the male parent (Spanish market type), is susceptible to all three diseases (Liang et al., 2012; Wang et al., 2012), primarily using SSR markers. For this study, a subset of 91 RILs were selected for whole-genome resequencing.

The whole RIL population was phenotyped in field for ELS, LLS and TSWV disease severity from 2009 to 2013. There were two plantings for 2011, 2012 and 2013, an early planting in April and a late planting in May in each year. Overall, there were a total of eight plantings each with three replications using a randomized complete block design. Disease severity ratings for the three diseases (ELS, LLS and TSWV) were performed at three different dates in growing seasons in July, August and September. Ratings for ELS and LLS were measured using the Florida 1–10 scale (Chiteka et al., 1988). TSWV disease severity was measured on a 1–10 disease severity in plots exhibiting typical symptoms such as stunting, ringspot, leaf necrosis and chlorosis (Baldessari, 2008; Culbreath et al., 2003).

Library construction and sequencing

Young leaf tissues from each of the 91 selected RILs along with the parental lines were used for genomic DNA isolation using the CTAB method and quantified as described by Wang et al. (2016). A whole-genome shotgun sequencing strategy was then used to construct the paired-end libraries. Paired-end sequencing libraries were sequenced with read length of 100 bp using an Illumina HiSeq 2000 platform (Illumina, San Diego, CA). Parental genotypes were sequenced separately at a high sequencing depth, ~100× for Tifrunner with insert size ranging from 170 bp to 40 kb, and ~10× for GT-C20 with an insert size of 500 bp. Individual RILs were sequenced at ~2–5× coverage. Filtered reads were used for alignment to the reference genome assemblies of A. duranensis (v1, peanutbase.org) and A. ipaensis (v1, peanutbase.org) separately and used for SNP identification and genotyping. The binary alignment mapping (BAM) files using the read data have been submitted to the SRA database at NCBI under the SRA accession: SRP134206.

Sequence analysis and SNP discovery for genetic analysis

Filtered reads from Tifrunner, GT-C20 and the individual RILs (fastq formatted) were mapped to the genome assemblies of A. duranensis (v1, peanutbase.org) and A. ipaensis (v1, peanutbase.org) separately using the Burrows-Wheeler alignment (BWA) tool with default parameters (Li and Durbin, 2009). Alignment files were used to identify polymorphic haplotypes using a novel pipeline (Clevenger et al., unpublished), which is an improved version of SWEEP described by Clevenger and Ozias-Akins (2015). The haplotypes were generated within a read, that is less than 100 bp. Briefly, all potential polymorphisms were called using Samtools mpileup. A sliding window strategy was used to visit every two base haplotype of potential polymorphisms that were physically within 100 bp. At each haplotype locus, every distinct haplotype along with observed counts for each haplotype within each genotype was collected. A polymorphic haplotype was identified when meeting the following criteria: (i) All haplotypes for both genotypes were observed more than once, (ii) at least one haplotype differed between the genotypes at one base while the other base remained constant and (iii) the haplotype with the least number of observations had within 25% of the observations of the haplotype with the most observations. Identified putative polymorphic haplotype SNPs were then used to genotype the population. Each individual was mapped to the A. ipaensis and A. duranensis genome separately. For each individual, the A-genome-derived haplotypes were scored in the alignment mapped to A. duranensis and the B-genome-derived haplotypes were scored in the alignment mapped to A. ipaensis. At each potential polymorphic locus, if the polymorphic haplotype from one parent is observed more than once and the haplotype from the other parent is not observed, the individual was scored to have the allele from the parent that the observed haplotype is derived. If both haplotypes were observed or one haplotype was only observed one time, the individual was scored as missing data.

Linkage map construction, identification of homeologous and translocated markers

Filtered SNPs with less than 20% missing data were used to construct the genetic maps using QTL IciMapping v4.1 (Meng et al., 2015). The chi-square (χ²) values calculated for each SNP marker were used to determine the goodness of fit to the expected 1:1 segregation ratio. Highly distorted and unlinked markers were not considered for the linkage map construction. The markers were grouped at LOD ≥ 4 and ordered using the nntTwoOpt algorithm. Kosambi’s mapping function was used for converting the recombination frequency into map distance in centiMorgan (cm). Homeologous SNPs are defined as the ones that were identified from A-genome chromosome of the reference diploid genome, but mapped on the corresponding B-genome LG. For example, when a SNP called on chromosome A01 was mapped to LG B01 or a SNP called on chromosome B01 was mapped to LG A01, they were regarded as homeologous. Translocated SNPs are defined as the ones that are identified on A-genome chromosome of the reference diploid genome; however, on the linkage map, the same SNPs were mapped on any A- or B-genome linkage groups except the corresponding B-genome LG. For example, SNPs called on chromosome A01 when mapped to any of the A- or B-LG except A01 and B01 were regarded as translocated. Circa (http://omgenomics.com/circa/) was used to plot circos to demonstrate the homeologous and translocated markers.

Epistatic and environmental QTL analysis

The genotyping and phenotyping data of population were used for QTL analysis using the inclusive composite interval mapping (ICIM) function of QTL IciMapping v4.1. A QTL was considered to be major only if had a LOD ≥ 3 and PVE explained >10%. The genetic map information together with phenotyping data was used for the identification of epistatic and environmental QTL interaction studies for ELS, LLS and TSWV using ICIM mapping. BIP (biparental) and MET (QTL by environment) functionalities of inclusive composite interval mapping (ICIM) was used for identification of epistasis and environmental QTLs, respectively. The additive (two-dimensional scanning, ICIM-EPI) method with 5 cm
step and 0.001 probability mapping parameters in stepwise regression were employed in QTL analysis. For environmental QTL analysis, one input file for each in multiple environments was used. Also, the largest P-value for removing variables was assumed to be two times the value of P-value for entering variables in stepwise regression. The missing phenotypic data were replaced by the phenotypic mean of the trait built-in function.

KASP assay development
The KASP genotyping assay is a fluorescence-based assay for identification of biallelic SNPs. Two allele-specific forward primers along with tail sequences and one common reverse primer were synthesized (Table S10). The reaction mixture was prepared following the manufacturer’s instructions (KBioscience; http://www.lgcgroup.com/products/kasp-genotyping-chemistry/#.Vs2K7PkrKMB). KASP reaction mixture with 10 µL final reaction volume contained 5 µL KASP master mix, 0.14 µL primer mix, 2 µL of 10–20 ng/µL genomic DNA and 2.86 µL of water. PCR conditions used were as follows: 15 min at 95°C followed by 10 touchdown cycles of 20 s at 94°C and 1 min at 61–55°C (dropping 0.6°C per cycle), and then 26 cycles of 20 s at 94°C and 1 min at 55°C. KASP marker data were then analysed using SNPviewer software (LGCGenomics; http://www.lgcgroup.com) to generate genotype calls for each RIL and parental line, and were correlated with observed disease ratings.

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Author contributions
GA performed most of the experiments, conducted sequence alignment, SNP calling and drafted the manuscript. JC developed the programme, conducted sequence analysis and called the SNPs. MKP contributed in planning and designing the experiment, data analysis and manuscript revision. HW and YC prepared the DNA. HW, CCH and AKC performed field phenotyping. HW, JCF, YS and DC participated the mapping and designed primers. MKP and YS did the environment and QTL interaction studies. XL and GH generated the sequence data. RD helped in statistical analyses. XW, DIB, SAJ and PO provided inputs and discussion. BG conceived the plan; BG and RKV designed and finalized the manuscript.

Conflict of interest statement
The authors declare that there is no conflict of interests.

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Supporting information

Additional Supporting Information may be found online in the supporting information tab for this article:

**Figure S1** Phenotypic distribution of ELS, LLS and TSWV in T-pop RILs during different seasons.

**Figure S2** Percentage reads mapped to the diploid reference A- and B-genome in each RIL and the two parents.

**Figure S3** Frequency histogram of the percentage of missing data points in WGRS of 91 RILs with 16 674 polymorphic SNPs in the population.

**Figure S4** Distribution of markers on linkage groups.

**Figure S5** QTL maps showing the major QTL peaks at different LODs on vertical axis.

**Figure S6** Genetic and QTL map of major QTLs (>10% PVE) comprising SNP and SSR markers in Tifrunner x GT-C20 population in peanut (Pandey et al., 2017a).

**Table S1** Phenotypic variation of diseases (ELS, LLS and TSWV) in T-population parents and RILs.

| Table S2 | Overview of the WGRS data and alignment to the reference genome. |
|---------|------------------------------------------------------------------|
| Table S3 | Summary of SNPs detected between Tifrunner and GT-C20 and SNPs used in RIL population. |
| Table S4 | Number of homeologus and translocated markers. |
| Table S5 | Effect of major QTL associated SSR markers reported in earlier study on the QTL and length of current genetic map. |
| Table S6 | Summary of environment QTLs detected in T-population. |
| Table S7 | Summary of the epistatic QTLs detected across different environments and locations for the ELS, LLS and TSWV in the RILs. |
| Table S8 | SNPs in putative candidate genes underlying the major QTLs for ELS, LLS and TSWV resistance with their effect and impact. |
| Table S9 | SNP efficiency of markers validated using KASP assay. |
| Table S10 | List of primer sequences for KASP assay of SNPs developed and validated for Early Leaf Spot (ELS) and Late Leaf Spot (LLS). |