Genotyping-by-sequencing and SNP-arrays are complementary for detecting quantitative trait loci by tagging different haplotypes in association studies

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Abstract

Background: Single Nucleotide Polymorphism (SNP) array and re-sequencing technologies have different properties (*e.g.* calling rate, minor allele frequency profile) and drawbacks (*e.g.* ascertainment bias). This lead us to study their complementarity and the consequences of using them separately or combined in diversity analyses and Genome-Wide Association Studies (GWAS). We performed GWAS on three traits (grain yield, plant height and male flowering time) measured in 22 environments on a panel of 247 F1 hybrids obtained by crossing 247 diverse dent maize inbred lines with a same flint line. The 247 lines were genotyped using three genotyping technologies (Genotyping-By-Sequencing, Illumina Infinium 50K and Affymetrix Axiom 600K arrays).

Results: The effects of ascertainment bias of the 50K and 600K arrays were negligible for deciphering global genetic trends of diversity and for estimating relatedness in this panel. We developed an original approach based on linkage disequilibrium (LD) extent in order to determine whether SNPs significantly associated with a trait and that are physically linked should be considered as a single Quantitative Trait Locus (QTL) or several independent QTLs. Using this approach, we showed that the combination of the three technologies, which have different SNP distributions and densities, allowed us to detect more QTLs (gain in power) and potentially refine the localization of the causal polymorphisms (gain in resolution).

Conclusions: Conceptually different technologies are complementary for detecting QTLs by tagging different haplotypes in association studies. Considering LD, marker density and the combination of different technologies (SNP-arrays and re-sequencing), the genotypic data

45 available were most likely enough to well represent polymorphisms in the centromeric
46 regions, whereas using more markers would be beneficial for telomeric regions.

47 **Keywords**: GWAS, linkage disequilibrium, genome coverage, maize, high-throughput
48 genotyping technologies.

49 Background

50 Understanding the genetic bases of complex traits involved in the adaptation to biotic and
51 abiotic stress in plants is a pressing concern, with world-wide drought due to climate change
52 as a major source of human food and agriculture threats. Recent progress in next generation
53 sequencing and genotyping array technologies contribute to a better understanding of the
54 genetic basis of quantitative trait variation by allowing Genome-Wide Association Studies
55 (GWAS) on large diversity panels [1]. Single Nucleotide Polymorphism (SNP)-based
56 techniques became the most commonly used genotyping methods for GWAS because SNPs
57 are cheap, numerous, codominant and can be automatically analysed with SNP-arrays or
58 produced by genotyping-by-sequencing (GBS), or sequencing [2-4]. The decreasing cost of
59 genotyping technologies has led to an exponential increase in the number of markers used for
60 the GWAS in association panels, thereby raising the question of computation time to perform
61 the association tests. Computational issues were addressed by using either approximate
62 methods by avoiding re-estimating variance components for each SNP [5] or exact methods
63 using mathematical tools for sparing time in matrix inversion [6, 7]. It is noteworthy that
64 using approximate computation in GWAS can produce inaccurate p-values when the SNP
65 effect size is large or/and when the sample structure is strong [8].

66 Several causes may impact the power of Quantitative Trait Locus (QTL, locus involved in
67 quantitative trait variation) detection in GWAS. Highly diverse panels have in general

undergone multiple historical recombinations, leading to a low extent of linkage disequilibrium (LD). However, these panels can present different average and local patterns of LD [9-11]. A high marker density and a proper distribution of SNPs are therefore essential to capture causal polymorphisms. Furthermore, minor allele frequencies (MAF), population stratification and cryptic relatedness are three other important parameters affecting power and false positive detection [12, 14]. These last two factors are substantial in several cultivated species such as maize [15] and grapevine [16], and their impact on LD can be statistically evaluated [17]. Population structure and kinship can be estimated using molecular markers [18-21] and can be modelled to efficiently detect marker-trait associations due to linkage only [12, 22, 23]. These advances have largely increased the power and effectiveness of linear mixed models that can now efficiently account for population structure and relatedness in GWAS [8, 12].

In maize, an Illumina Infinium HD 50,000 SNP-array, named MaizeSNP50 (hereafter 50K) was developed by Ganal *et al.* [3] and has been used extensively for diversity and association studies [24, 25]. For example, GWAS were conducted to unravel the genetic architecture of phenology, yield component traits and to identify several flowering time QTLs linked to adaptation of tropical maize to temperate climate [26, 27]. In the same way, Rincent *et al.* [11] showed that LD occurs over a longer distance in a dent than in a flint panel, with appreciable effects on the power of QTL detection. Low LD extent and relationship between allelic frequencies with population and pedigree structure at some SNPs reduce the power of GWAS [14, 27]. Therefore, higher marker densities are desirable because the maize genome size is large (2.4 Gb), the level of diversity is high (more than one substitution per hundred nucleotides), and LD extent is low [28]. As a consequence, an Affymetrix Axiom 600,000

SNP-array (hereafter 600K) was developed and used in association genetics [29, 30] and detection of selective sweeps [4]. Another possibility is whole genome sequencing, but this is currently impractical for large genomes such as maize because of the associated cost. Hence, a Genotyping-By-Sequencing (GBS) procedure has been developed [2] that targets low-copy genomic regions by using restriction enzymes. Genotyping-by-sequencing technology is cost-effective and has been successfully used in maize for genomic prediction [31]. Romay *et al.* [32] and Gouesnard *et al.* [33] highlighted the interest of the GBS for (i) deciphering and comparing the genetic diversity of the inbred lines in seedbanks and (ii) identifying QTLs by GWAS for kernel colour, sweet corn and flowering time.

Few studies in plants have compared datasets from different high-throughput genotyping technologies [34-36]. Elbasyoni et al. [34] used GBS and a 90K SNP-array in winter wheat. They highlighted strong positive correlations between the population structure matrices and kinships identified by both technologies. They showed that GBS-SNPs led to higher genomic prediction accuracy compared to Array-SNPs. Torkamaneh and Belzile [36] used GBS and a 50K SNP-array in soybean. They estimated *ca.* 98% accuracy of genotype called by their GBS pipeline and showed that the accuracy of imputation for missing genotypes was hardly affected by the chosen the MAF and only moderately affected by the rate of missing values. Li et al. [35] created a reliable integrated variation map using a 600K and 50K SNP-array, GBS and RNA sequencing to dissect regulatory causality and its link to maize kernel variation. These authors used a fixed physical distance (<10 kb) for grouping associated SNPs into QTLs despite the variable LD pattern along the genome. None of these studies compared QTL detection between the different technologies.

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The main drawback of the DNA arrays is that they do not allow the discovery of new SNPs. This possibly leads to some ascertainment bias in diversity analysis when the SNPs selected for building arrays come from (i) the sequencing of a set of individuals that did not represent well the diversity explored in the studied panel, (ii) a subset of SNPs that skews the allelic frequency profile towards the intermediate frequencies [27, 37]. Ascertainment bias can compromise the ability of the SNP-arrays to reveal an exact view of the genetic diversity [37]. Genotyping-by-sequencing can overcome ascertainment bias since it is based on sequencing and therefore allows the discovery of alleles in the diversity panel analysed. It can be generalized to any species at a low cost providing that numerous individuals have been sequenced in order to build a representative library of short haplotypes to call SNPs [38]. Non-repetitive regions of genomes can be targeted with two- to three-fold higher efficiency, thereby considerably reducing the computationally challenging problems associated with alignment in species with high repeat content. However, GBS may have low coverage leading to a high missing data rate (65% in both studies; [32, 33]) and heterozygote under-calling, depending on genome size and structure, and on the multiplexing level per sequencing flow-cell. Furthermore, GBS requires the establishment of demanding bioinformatic pipelines and imputation algorithms [39]. Pipelines have been developed to call SNP genotypes from raw GBS sequence data and to impute the missing data from a haplotype library [38, 39].

Here, we investigated the impact of using GBS and SNP-arrays on the quality of the genotyping data, together with the biological properties of data generated by each technology, and the potential complementarity of these approaches. In particular, we analyzed the impact of marker density and genotyping technologies (sequencing *vs* array) on (i) the estimates of relatedness and population structure, and (ii) the detection of QTLs (power). To address these issues, we performed a GWAS based on genotypic datasets obtained using either GBS or

SNP-arrays with low (50k) or high (600k) densities on a diversity panel of maize hybrids obtained by crossing a panel of dent lines with a common flint tester lines. Three traits were considered, namely grain yield, plant height and male flowering time (day to anthesis), measured in 22 different environments (sites × years × treatments) over Europe. We developed an original approach based on LD extent in order to determine whether SNPs significantly associated with a trait should be considered as a single QTL or several independent QTLs.

Results

Combining Tassel and Beagle imputations improved the genotyping quality for GBS

We estimated the genotyping and imputation concordance of the GBS based on common markers with the 50K or 600K arrays (Additional file 1: Figure S1 and Table 1). The genotyping concordance of the 600K with the 50K was extremely high (99.50%), although slightly lower for residual heterozygotes (92.88%). After SNP calling from sequencing reads using AllZeaGBSv2.7 database (direct reads, GBS_1, Additional file 1: Figure S1), the call rate was 33.81% for the common SNPs with the 50K, *vs* 37% for the whole GBS dataset. The genotyping concordance rate between the direct reads of GBS and the 50K was 98.88% (Table 1). After imputation using *TASSEL* by Cornell Institute (GBS_2), the concordance rate was 96.04% on the common markers with the 50K and 11.91% of missing data remained for the whole GBS dataset. In GBS_3, all missing data were imputed by *Beagle* and the remaining missing genotypes in GBS_2 were excluded here to be comparable with *TASSEL*. This method yielded a lower concordance rate (93.04% and 92.84% with 50K and 600K, respectively). In

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an attempt to increase the concordance rate of the genotyping while removing missing data, we tested two additional methods, namely GBS_4 where the missing data and heterozygotes of Cornell imputed data (GBS_2) were replaced by Beagle imputation, and GBS_5 where Cornell homozygous genotypes (GBS_2) were completed by imputations from GBS_3 (Additional file 1: Figure S1 and Table 1). GBS_5 displayed a slightly better concordance rate than GBS_2 (96.25% vs 96.04%) and predicted heterozygotes with a higher quality than GBS_4. GBS_5 was therefore used for all genetic analyses and named GBS hereafter.

Table 1: Percentage of GBS concordance and call rates (in parentheses).

	Reference	Total	Homozygotes	Heterozygotes
GBS_1 Direct Read	50K	98.88 (33.81)	99.03 (33.72)	45.09 (0.09)
	600K	98.99 (35.58)	99.21 (35.47)	28.67 (0.11)
GBS_2 Cornell Imputation	50K	96.04 (91.56)	98.66 (88.79)	12.51 (2.78)
	600K	95.50 (93.41)	98.69 (90.14)	7.75 (3.28)
GBS_3 Beagle Imputation	50K	93.04 (*91.56)	93.23 (91.30)	30.54 (0.26)
	600K	92.84 (*93.41)	93.07 (93.12)	22.50 (0.29)
GBS_4 Beagle Imputation on the missing data and heterozygotes after Tassel Imputation (GBS2)	50K	96.46 (*97.64)	96.46 (97.63)	<0.01 (<0.01)
	600K	96.21 (*99,97)	96.21 (>99.99)	<0.01 (<0.01)
GBS_5 Compilation of Homoz. genotypes from Tassel Imputation (GBS2) and Imputation by Beagle for Other Data (GBS3)	50K	96.25 (*97.65)	96.36 (97.47)	39.07 (0.18)
	600K	95.98 (*99.97)	96.11 (99.78)	32.02 (0.22)

The 50K and 600K SNP-arrays were considered as reference genotypes. * After Beagle inference of missing data, the call rate was 100%. Here the call rate is <100% because the comparison was made against the 50K and the 600K that include few missing data. For GBS_3, the remaining missing genotypes in GBS_2 were also excluded to obtain comparable results.

GBS displayed more rare alleles and lower call rate than SNP-arrays

The SNP call rate was higher for the SNP-arrays (average values of 96% and >99% for the 50K and 600K, respectively), than for the GBS (37% for the direct reads). The MAF

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distribution differed between the technologies (Additional file 2: Figure S2): while the use of SNP-arrays resulted in a near-uniform distribution, GBS resulted in an excess of rare alleles with a L-shaped distribution (22% of SNPs with MAF < 0.05 for the GBS *versus* 6% and 9% for the 50K and 600K, respectively). This can be explained by the fact that the 50K was based on 27 sequenced lines for SNPs discovery [3], the 600K was based on 30 lines [4], whereas GBS was based on 31,978 lines, thereby leading to higher discovery of rare alleles. Consistent with MAF distribution, the average gene diversity (*He*) was lower for GBS (0.27) than for arrays (0.35 and 0.34 for the 50K and 600K arrays, respectively). The distribution of SNP residual heterozygosity of inbred lines was similar for the three technologies, with a mean of 0.80%, 0.89% and 0.22 % for the 50K, 600K and GBS, respectively. The residual heterozygosity of inbred lines was highly correlated between technologies with large coefficients of Spearman correlation: $r_{50K\text{-}600K} = 0.90$, $r_{50K\text{-}GBS} = 0.76$, $r_{600K\text{-}GBS} = 0.83$. The distribution of the SNPs along the genome was denser in the telomeres for the GBS and in the peri-centromeric regions for the 600K, whereas the 50K exhibited a more uniform distribution (Figure 1 and top graph in Additional file 3: Figure S3).

Population structure and relatedness were consistent between the three technologies

We used the ADMIXTURE software to analyse the genetic structure within the studied panel based on SNPs from the three technologies, by considering two to ten groups. Based on a K-fold cross-validation, the clustering in four genetic groups ($N_Q = 4$) was identified as the best for the three datasets. Considering a threshold of 0.5 for ancestral fraction, the assignation to the four groups was identical except for a few admixed inbred lines (Additional file 4: Figure S4). Based on the 50K, the four groups were constituted by (i) 39 lines in the Non Stiff Stalk

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(Iodent) family traced by PH207, (ii) 46 lines in the Lancaster family traced by Mo17 and Oh43, and (iii) 55 lines in theStiff Stalk family traced by B73 and (iv) 107 lines that did not fit into these three primary heterotic groups, such as W117 and F7057. This organization appeared consistent with the organization of breeding programs into heterotic groups, generally related to few key founder lines.

We compared two estimators of relatedness between inbred lines, IBS (Identity-By-State) and *K_Freq* (Identity-By-Descent), calculated per technology. For IBS, pairs of individuals were on average more related using GBS than SNP-arrays (mean IBS: 0.66, 0.67 and 0.73 for 50K, 600K and GBS, respectively). As expected, mean IBD was close for the three technologies (*K_Freq*: -0.004). Relatedness estimates with the two SNP-arrays were highly correlated: r = 0.95 and 0.98 for IBS and *K_Freq*, respectively (Additional file 5: Figure S5b and d). Likewise, relatedness estimates between arrays and GBS were strongly correlated (between 0.94 and 0.98, Additional file 5: Figure S5b and d)..

We further carried out diversity analyses by performing Principal Coordinate Analyses (PCoA) on IBD (*K_Freq*, weights by allelic frequency) estimated from the three technologies (Figure 2). The three first PCoA axes explained 12.9%, 15.6% and 16.3% of the variability for the GBS, 50K and 600K, respectively (Figure 2). The same pattern was observed regardless of the technology with the first axis separating the Stiff Stalk from all other groups (Iodent and Lancaster lines, see illustration with the 50K kinship, Figure 2). Key founder lines of the three heterotic groups (Iodent: PH207, Stiff Stalk: B73, Lancaster: Mo17) were found at extreme positions along the axes, which was consistent with the admixture groups previously described.

Long distance linkage disequilibrium was removed by taking into account population structure or relatedness

In order to evaluate the effect of kinship and the genetic structure on linkage disequilibrium (LD), we studied genome-wide LD between 29,257 PANZEA markers from the 50K within and between chromosomes before and after taking into account the kinship (*K_Freq* estimated from the 50K), structure (Number of groups = 4) or both (Additional file 6: Figure S6). Whereas inter-chromosomal LD was only partially removed when the genetic structure was taken into account, it was mostly removed when either the kinship or both kinship and structure were considered (Additional file 6: Figure S6b and c). Accordingly, long distance intra-chromosomal LD was almost totally removed for all chromosomes by accounting for the kinship, structure or both. Interestingly, some pairs of loci located on different chromosomes or very distant on a same chromosome remained in high LD despite correction for genetic structure and kinship (Additional file 6: Figure S6). This can be explained either by genome assembly errors, by chromosomal rearrangements such as translocations or by strong epistatic interactions. Linkage disequilibrium decreased with genetic or physical distance Figure 3. The majority of pairs of loci with high LD (r^2K>0.4) in spite of long physical distance (>30Mbp), were close genetically (<3cM), notably on chromosome 3, 5, 7 and to a lesser extent 9 and 10 (data not shown). These loci were located in centromeric and peri-centromeric regions that displayed low recombination rate, suggesting that this pattern was due to variation of recombination rate along the chromosome. Only very few pairs of loci in high LD were genetically distant (>5cM) but physically close (<2Mbp). Linkage disequilibrium (r^2K and r^2KS) was negligible beyond 1 cM since 99% of LD values were less than 0.12 in this case. Note that some unplaced SNPs remained in LD after taking into account the kinship and structure with some SNPs with known positions on chromosome 1, 3 and 4 (Additional File 6:

Figure S6). Therefore, LD measurement corrected by the kinship can help to map unplaced SNPs.

Linkage disequilibrium strongly differed between and within chromosomes

We combined the three technologies together to calculate the r^2K for all pairs of SNPs which were genetically distant by less than 1 cM. For any chromosome region, LD extent in terms of genetic and physical distance showed a limited variation over the 100 sets of 500,000 loci pairs (cf. Material). This suggests that the estimation of LD extent did not strongly depend on our set of loci. LD extent varied significantly between chromosomes for both high recombinogenic (>0.5 cM/Mbp) and low recombinogenic regions (<0.5 cM/Mbp, Table 2). Chromosome 1 had the highest LD extent in high recombination regions (0.062 ±0.007 cM) and chromosome 9 the highest LD extent in low recombinogenic regions (898.6±21.7 kbp) (Table 2). Linkage disequilibrium extent relative to genetic and physical distances was highly and positively correlated in high recombinogenic regions (r = 0.86), whereas it was not in low recombinogenic regions (r = -0.64).

Table 2: Variation of LD extent, and percentage of genome covered.

		Chromosome										Whole Genome
		1	2	3	4	5	6	7	8	9	10	
Physical Size (Mbp)		301	237	232	241	217	169	176	175	156	150	2,058
Genetic Size (cM)		268	211	188	150	205	129	148	182	145	139	1766
Physical LD extent (kbp) in low recombination regions		306	491	846	808	658	418	547	497	899	815	629
Genetic LD Extent (cM) in high recombination regions		0.062	0.027	0.033	0.022	0.031	0.019	0.012	0.038	0.023	0.019	0.029
Percent of physical genome covered	50K	81%	72%	76%	77%	74%	67%	71%	73%	71%	71%	74%
	600K	98%	88%	91%	89%	90%	84%	81%	90%	87%	84%	89%
	GBS	92%	81%	84%	83%	83%	77%	77%	81%	79%	76%	82%
	ALL	98%	90%	92%	90%	91%	87%	83%	92%	88%	85%	90%

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Percent of genetic map covered	50K	72%	41%	44%	38%	41%	32%	24%	46%	32%	27%	42%
	600K	96%	71%	76%	68%	72%	62%	47%	78%	63%	53%	71%
	GBS	86%	58%	61%	53%	61%	48%	37%	63%	48%	40%	58%
	ALL	97%	74%	78%	72%	74%	65%	51%	81%	66%	57%	74%

Genetic and Physical LD extent were obtained by adjusting Hill and Weir model's on 100 different sets of 500,000 loci randomly sampled in high (>0.5 cM / Mbp) and low (<0.5 cM / Mbp) recombination regions, respectively. The value represented the average across these 100 sets. The percentage of genome coverage was estimated using markers with MAF > 5% and E(r^2k) = 0.1, for each technology and for the three technologies combined (ALL: GBS+600K+50K).

Large differences in genome coverage between technologies

We estimated the percentage of the genome that was covered by LD windows around SNPs, calculated by using either physical or genetic distances (Figure 3, Table 2). We observed a strong difference in coverage between the three technologies at both genome-wide and chromosome scale, as illustrated in Figure 1 on chromosome 3 (Table 2, and Additional file 3: Figure S3). For a LD extent of r^2K = 0.1, 74%, 82% and 89% of the physical map, and 42%, 58% and 71% of the genetic map were covered by the 50K, the GBS and the 600K, respectively (Table 2). For the combined data (ALL: 50K + 600K + GBS), the coverage strongly varied between chromosomes, ranging from 83% (chromosome 7) to 98% (chromosome 1) of the physical map, and from 51% (chromosome 7) to 97% (chromosome 1) of the genetic map (Table 2). For the physical map, increasing the LD extent threshold to r^2K=0.4 reduced the genome coverage from 89% to 49% for 600K, 82% to 28% for GBS, 74% to 20% for 50K and 90% to 52% for the combined data. Increasing the MAF threshold reduced slightly the genome coverage, with smaller reduction for the physical map than genetic map. Surprisingly, increasing the SNP number by combining the markers from the arrays and GBS did not strongly increase the genome coverage as compared to the 600K,

282 regardless of the threshold for LD extent (Figure 1and Additional file 3: Figure S3).
283 We observed a strong variation of genome coverage along each chromosome with contrasted
284 patterns in low and high recombinogenic regions (Figure 1Additional file 3: Figure S3).
285 While low recombinogenic regions were totally covered with all the technologies (except for
286 few intervals using the 50K), the genome coverage in high recombinogenic regions varied
287 depending on both technology and SNP distribution. 47% of the 2Mbp intervals in high
288 recombination regions were better covered by the 600K than the GBS against only 1%, which
289 were better covered by GBS than 600K. When exploring smaller window sizes (20, 100, 500
290 kb), the number of intervals better covered by 600K than GBS decreased strongly when the
291 intervals were shortened (17.1% of 20kbp-intervals *vs* 47.1% of 2Mbp-intervals). In the
292 contrary, the intervals better covered by GBS than 600K increased slightly (4.1% *vs* 1.1% of
293 2Mbp-intervals). The number of interval with no or weak coverage differences between GBS
294 and 600K increased strongly: 84.5% of 20kbp-intervals *vs* 68% of 2Mbp-intervals with
295 coverage differences inferior to 10%. Interestingly, the proportion of interval with strong
296 coverage differences (>50%) increased when the intervals were shortened (7.8% of 20kbp-
297 intervals *vs* 0% of 2Mbp-intervals).
298

Number of QTLs detected using genome-wide association studies increases with markers density

299 We observed a strong variation in the number of SNP significantly associated with the three
300 traits across the 22 environments (Table 3). The mean number of significant SNPs per
301 environment and trait was 3.7, 44.7, 17.9 and 62.4 for the 50K, 600K, GBS and the three
302 technologies combined, respectively (Table 4). Considering the p-value threshold used, 28,

303 303 and 204 false positives were expected among the 243, 2,953 and 1,182 associations
304 detected for 50K, 600K and GBS, respectively. False discovery rate appeared therefore higher
305 for GBS (17.2%) than for DNA arrays (11.5% and 10.2% for 50K and 600K, respectively). It
306 could be explained by the higher genotyping error rate of GBS due to imputation and/or by its
307 higher number of markers with a low MAF. Both reduce the power of GBS compared to DNA
308 arrays and therefore lead to a higher false discovery rate. Proportionally to the SNP number,
309 the 50K and 600K resulted in 1.5- and 1.7-fold more associated SNPs per situation
310 (environment × trait) than GBS (p-value<$2x10^{-6}$, Table 4). This difference between SNP-
311 arrays and GBS was higher for grain yield (GY) and plant height (plantHT) than for male
312 flowering time (DTA, Table 4).
313
314 **Table 3:** Number of significant SNPs per environment, per technology and for the combined
315 technologies.

	Flowering Time					Plant Height					Grain Yield				
Env.	Herit.	50K	600K	GBS	ALL	Herit.	50K	600K	GBS	ALL	Herit.	50K	600K	GBS	ALL
Cam12R	0.40	0	3	1	4	0.57	23	209	88	289	0.37	21	286	102	381
Cam12W	0.60	1	18	13	31	0.39	22	270	72	339	0.54	41	525	167	684
Cam13R	0.46	0	9	7	16	0.21	0	2	3	5	0.19	0	1	3	4
Cra12R	0.69	1	40	18	59	0.32	0	6	3	9	0.25	3	57	45	103
Cra12W	0.72	3	25	19	43	0.19	10	69	16	83	0.53	12	98	53	150
Deb12R	0.63	1	14	16	30	0.33	0	1	0	1	0.57	2	14	0	16
Deb12W	0.71	0	25	38	61	0.37	0	6	7	7	0.47	0	6	2	8
Deb13R	0.59	1	17	5	23	0.08	0	33	9	42	0.37	1	22	15	37
Gai12R	0.64	8	80	24	104	0.18	1	47	41	89	0.31	0	23	8	31
Gai12W	0.66	5	42	15	59	0.46	0	1	3	4	0.58	3	71	14	85
Gai13R	0.62	0	24	8	31	0.66	0	6	6	11	0.63	0	4	5	9
Gai13W	0.73	1	45	9	54	0.33	0	1	3	4	0.76	2	7	1	9
Kar12R	0.72	4	30	21	52	0.30	0	4	3	7	0.71	0	5	6	11
Kar12W	0.77	8	60	10	73	0.23	1	10	4	14	0.53	2	19	11	29
Kar13R	0.68	3	65	11	77	0.31	0	4	2	6	0.75	4	37	24	62
Kar13W	0.73	0	17	12	29	0.26	0	2	7	9	0.67	4	12	6	19
Mur13R	0.83	3	48	19	68	0.32	7	61	7	68	0.84	14	90	28	116
Mur13W	0.76	0	11	8	19	0.32	3	4	2	9	0.74	10	80	25	104
Ner12R	0.72	7	23	18	45	0.34	0	7	3	10	0.54	1	10	6	16

Ner12W	0.81	1	80	30	107	0.24	0	2	2	4	0.59	1	8	6	15
Ner13R	0.76	3	60	26	88	0.26	1	25	13	38	0.32	0	13	7	20
Ner13W	0.81	2	23	17	42	0.20	0	8	5	13	0.73	2	28	4	32
Average	0.68	2.4	34.5	15.7	50.7	0.31	3.1	35.4	13.6	48.2	0.55	5.6	64.4	24.5	88.2
Median	0.71	1	25	15.5	47.5	0.32	0	6	4.5	9.5	0.56	2	20.5	7.5	30
SD	0.11	2.6	22.7	8.7	27.7	0.13	6.8	69.6	23.2	90.2	0.18	9.6	120.1	39.5	156.8

The average, median and standard deviation (SD) per environment are calculated for each trait (male Flowering Time, Plant Height, Grain Yield). "Herit.": narrow sense heritability. "Env." : Environment.

Table 4: Comparison of associated SNPs and QTLs detected between traits and three technologies.

		Significant SNPs				QTLs			
Technology		50K	600K	GBS	ALL	50K	600K	GBS	ALL
Marker Nb		42046	459191	308929	810580	42046	459191	308929	810580
Total Nb	DTA	52	759	345	1115	20	130	133	226
	plantHT	68	778	299	1061	16	96	90	160
	GY	123	1416	538	1941	33	166	120	238
	Per trait	81	984	394	1372	23	131	114	208
Average per envir.	DTA	2.4	34.5	15.7	50.7	0.9	5.9	6.0	10.3
	plantHT	3.1	35.4	13.6	48.2	0.7	4.4	4.1	7.3
	GY	5.6	64.4	24.5	88.2	1.5	7.5	5.5	10.8
	Per trait	3.7	44.7	17.9	62.4	1.0	5.9	5.2	9.5

QTLs were obtained by grouping associated SNPs with overlapping LD windows (*LD_win*) for the three traits (DTA: male flowering time; PlantHT: plant height; GY: grain yield). "*Marker Nb"* indicates the number of markers tested in GWAS. "*Total number"*: is the sum of associated SNPs or QTLs across environments. "*Average per envir"* indicates the average number of QTLs obtained in 22 environments for three traits (66 trait-environments combinations). *Average per SNP tested* indicates the number of associated SNPs or QTLs detected divided by the number of SNP tested.

We used two approaches based on LD for grouping significant SNPs (Figure 3): (i) considering that all SNPs with overlapping LD windows for $r^2K=0.1$ belong to the same QTL (*LD_win*) and (ii) grouping significant SNPs that are adjacent on the physical map and are in LD ($r^2K > 0.5$, *LD_adj*). The QTLs defined by using the two approaches were globally

334 consistent since significant SNPs within QTLs were in high LD whereas SNPs from different
335 adjacent QTLs were not (Additional file 7: Figure S7-LD-Adjacent and Additional file 8:
336 Figure S8-LD-Windows). *LD_adj* detected more QTLs than *LD_win* for flowering time (242
337 vs 226), plant height (240 vs 160) and grain yield (433 vs 237). The number of QTLs detected
338 with the *LD_adj* approach increased strongly when the LD threshold was set above 0.5.
339 Differences in QTL groupings between the two methods were observed for specific LD and
340 recombination patterns. This occurred for instance on chromosome 6 for grain yield
341 (Additional file 7: Figure S7-LD-Adjacent and Additional file 8: Figure S8-LD-Windows).
342 Within this region, the recombination rate was low and the LD pattern between associated
343 SNPs was complex (Additional file 1: Figure S1). While *LD_adj* splitted several SNPs in high
344 LD into different QTLs (for instance QTL 232, 235, 237, 249), *LD_win* grouped together
345 associated SNPs that are genetically close but displayed a low LD (Additional file 7: Figure
346 S7-LD-Adjacent and Additional file 8: Figure S8-LD-Windows). Reciprocally, for flowering
347 time, we observed different cases where *LD_win* separated distant SNPs in high LD into
348 different QTLs whereas *LD_adj* grouped them (QTL 25 and 26, 51 to 53, 95 to 97, 208 and
349 209, 218 and 219). As these differences were specific to complex LD and recombination
350 patterns, we used the *LD_win* approach for the rest of the analyses.
351
352 Although a large difference in number of associated SNPs was observed between 600K and
353 GBS, little difference was observed between QTL number after grouping SNPs (Table 3,
354 Table 4). The mean number of QTLs was indeed 1.0, 5.9, 5.2 and 9.5 for the 50K, 600K,
355 GBS, and the three technologies combined, respectively (Table 4). Note that the number of
356 QTLs continued to increase with marker density when SNPs from GBS, 50K and 600K were
357 combined (Additional file 9: Figure S9). The number of SNPs associated with each QTL

varied according to the technology (on average 3.7, 7.6, 3.4 and 6.6 significant SNPs for the 50K, 600K, GBS, and the combined technologies, respectively). The total number of QTLs detected over all environments by using the 600K and GBS was close for flowering time (130 *vs* 133) and plant height (96 *vs* 90). It was 1.4-fold higher for the 600K than GBS for grain yield (166 *vs* 120).

The 600K and GBS were highly complementary for association mapping

Seventy eight percent, 76% and 71% of the QTLs of flowering time, plant height and grain yield were specifically detected by 600K or GBS, respectively (Figure 4). On the contrary, 50K displayed very few specific QTLs. When we combined the GBS and 600K markers, 7% of their common QTLs had $-log_{10}(Pval)$ increased by 2 and 21% by 1, potentially indicating a gain in accuracy of the position of the causal polymorphism (Additional file 10: Table S1).

This complementarity between GBS and 600K is well exemplified with two strong association peaks for flowering time on chromosome 1 (QTL32) and 3 (QTL95) detected in several environments (Additional file 10: Table S1 and Figure 5a). In order to better understand the origin of the complementarity between GBS and 600K technologies for GWAS, we scrutinized the LD between SNPs and the haplotypes within these two QTLs (Figure 5b and c, and Additional file 11: Figure S10 for other examples). QTL95 showed a gain in power. It was only identified by the 600K although the region included numerous SNPs from GBS close to the associated peak. None of these SNPs was in high LD with the most associated marker of the QTL95 (Figure 5b). QTL32 was detected by 1 to 10 GBS markers in 9 environments with *–log(p-value)* ranging from 5 to 7.6, whereas it was detected by only two 600K markers in one environment (Ner12W) with *–log(p-value)* slightly above

18

the significance threshold (Additional file 10: Table S1 and Figure 5b).

Haplotype analyses showed that the SNPs from the GBS within QTL95 were not able to discriminate all haplotypes (Figure 5c). In QTL95, the 600K markers discriminated the three main haplotypes (H1, H2, H3), whereas using the GBS markers did not discriminate H3 against H1 + H2. As H1 contributed to an earlier flowering time than H2 or H3, associations appeared more significant for the 600K than for GBS (Figure 5c). In QTL32, the use of GBS markers identified late individuals that mostly displayed H1, H2 and H3 haplotypes, against early individuals that mostly displayed H4 and H5 haplotypes (Figure 5c). The gain of power for GBS markers as compared to 600K markers for QTL32 originated from the ability to discriminate late individuals (black alleles) from early individuals (red alleles) within H4 haplotypes (Figure 5c).

To further decipher which differences between 600K and GBS impacted GWAS, we used a resampling approach to explore the interplay between (i) MAF distribution and (ii) SNP distribution along the genome, at different SNP densities. We detected more SNP associations but less QTL with MAF distribution skewed towards low than high MAF. This difference increased as marker density increased (Additional file 12: Figure S11). As GBS has a MAF distribution skewed towards low MAF compared to 600K, GBS detected more QTLs but less associated SNPs than 600K. This discrepancy between association and QTL detection came from the fact that QTLs with low MAF were identified by less associated SNP than those with high MAF (Additional file 13: Figure S12).

Regarding distribution along the genome, SNPs distributed similarly to GBS detected more QTLs but less significant SNPs than those following the distribution of 600K and 50K, notably for the highest SNP density (Additional file 13: Figure S12). We observed

404 that SNP evenly distributed according to the physical distance detected more associations but
405 less QTLs than all other SNP distributions along the genome. It was the contrary for SNP
406 evenly distributed according to genetic distance (Additional file 12: Figure S11 C and D).
407 This is consistent with QTL distribution along the genome being more correlated to the
408 genetic than physical distance (see below), and the fact that recombination is higher in gene
409 rich regions, leading to less associated SNPs per QTL. Superiority of QTL detection by GBS
410 distribution as compared to 600K and 50K SNP distributions came from the higher proportion
411 of SNPs in high recombinogenic regions for GBS than for 600K and 50K (Figure 1). This
412 suggests that the complementarity of 600K and GBS in terms of QTL detected and SNP
413 associations came also from their specificities for both SNP distributions along the genome
414 and MAF distribution. In the end, we studied the impact of genomic coverage differences
415 between 600K and GBS on QTL detection along the genome. QTLs detected by both 600K
416 and GBS were located in intervals with large differences in coverage less frequently than their
417 proportion in the entire genome (0.8% vs 7.8%, respectively). Intervals with specific QTLs
418 showed an enrichment in such intervals with high differences in coverage (3.5%) but still
419 below the proportion in the entire genome. It confirmed that most specific QTLs showed no
420 strong genomic coverage differences between GBS and 600K and therefore that
421 complementarity of QTL detection between these two technologies came from ability to tag
422 different haplotypes.
423

Colocalization of QTLs between environments and traits and distribution of QTL along the genome

424 After combining the three technologies, we identified 226, 160, 238 QTLs for flowering time,

plant height and grain yield, respectively (Table 4 and Additional file 10: Table S1). We highlighted 23 QTLs with the strongest effects on flowering time, plant height and grain yield ($-log_{10}(Pval) \geq 8$, Table 5). The strongest association corresponded to the QTL95 for flowering time ($-log_{10}(p\text{-}value) = 10.03$) on chromosome 3 (158,943,646 – 159,005,990 bp), the QTL135 for GY ($-log_{10}(p\text{-}value) = 18.7$) on chromosome 6 (12,258,527 – 29,438,316 bp) and QTL78 on chromosome 6 (12,258,527 – 20,758,095 bp) for plant height ($-log_{10}(p\text{-}value) = 17.31$). The QTL95 for flowering time trait was the most stable QTLs across environments since it was detected in 19 environments (Additional file 10: Table S1). Moreover, this QTL showed a colocalization with QTL74 for grain yield in 5 environments and QTL30 for plant height in 1 environment suggesting a pleiotropic effect. More globally, 472 QTLs appeared trait-specific whereas 70 QTLs overlapped between at least two traits (6,3%, 5.2% and 3.0% for GY and plantHT, GY and DTA, and DTA and plantHT, respectively) suggesting that some QTLs may be pleiotropic (Additional file 14: Figure S13). This was not surprising since average corresponding correlations within environments for these traits were moderate (0.47, 0.54 and 0.45, respectively). Only 0.7% overlapped between the three traits (Additional file 14: Figure S13). Twenty percent of QTLs were detected in at least two environments and 9% in at least three environments (Additional file 15: Table S2). We observed no significant differences of stability between the three traits ($p\text{-}value = 0.2$). However, 6 out 7 most stable QTLs (Number of environments >5) were found for flowering time. This was consistent with higher average correlations between environments observed flowering time than for plant height and grain yield (0.76, 0.43, 0.48, respectively). We observed that QTLs that displayed a significant effect in more than one environment had larger effects and *-log(p-value)* values than those significant in a single environment. This difference in *-log(p-value)* values was stronger for grain yield and plant height than flowering time.

21

Table 5: Summary of the main QTLs ($-log_{10}(Pval) \geq 8$) identified for the three traits.

Trait	QTL	Chr	Pos	Lower Limit	Upper Limit	R2	Effect	Log	Minor All	Major All	MAF	EnvMax	Nb DiffEnv
DTA	95	3	158,974,594	158,943,646	159,005,990	0.15	1.27	10.03	G	C	0.41	Ner13R	19
plantHT	21	2	130,441,738	129,971,437	130,912,039	0.15	-6.74	9.21	A	G	0.14	Cam12W	3
	71	6	6,614,012	6,593,785	6,636,807	0.13	-4.76	8.39	G	A	0.18	Cam12R	2
	72	6	6,807,230	6,793,841	6,837,747	0.14	-4.87	8.77	T	G	0.18	Cam12R	2
	78	6	20,330,595	12,258,527	20,758,095	0.27	-8.99	17.31	C	T	0.26	Cam12W	4
	79	6	22,905,376	21,037,721	23,951,687	0.19	-5.45	11.42	T	G	0.31	Cam12R	3
	80	6	25,3178,25	24,184,017	26,606,537	0.13	-4.41	8.17	T	C	0.2	Cam12R	2
	81	6	28,130,108	26,695,327	28,659,766	0.16	-5.14	9.84	C	G	0.44	Cam12R	2
	94	6	101,482,646	101,463,249	101,501,936	0.14	-5.98	8.22	T	A	0.17	Cam12W	2
	110	8	12,782,777	12,767,198	12,798,330	0.19	-7.67	12.44	C	T	0.22	Cam12W	3
GY	65	3	141,621,777	140,505,559	144,210,207	0.12	0.42	8.13	A	C	0.27	Gai12W	5
	85	3	187,028,970	186,994,852	187057,772	0.12	-0.48	8.49	A	C	0.28	Kar12W	1
	120	6	5,155,708	5,131,927	5,177,694	0.12	-0.58	8.24	C	T	0.42	Cam12W	2
	122	6	5,638,516	5,623,945	5,659,803	0.12	-0.57	8.11	T	G	0.23	Cam12W	2
	124	6	5,871,000	5,855,407	5,887,383	0.12	-0.56	8.4	T	G	0.42	Cam12W	2
	127	6	6,612,654	6,593,785	6,636,807	0.16	-0.65	10.44	C	A	0.32	Cam12W	2
	128	6	6,807,462	6,793,841	6,837,747	0.12	-0.54	8.41	A	C	0.26	Cam12W	2
	129	6	6,890,199	6,878,877	6,930,838	0.12	0.63	8.06	T	A	0.48	Cam12W	3
	130	6	7,046,773	7,027,497	7,088,575	0.13	0.62	8.71	A	G	0.4	Cam12W	3
	131	6	7,159,714	7,113,662	7,200,479	0.12	0.59	8.24	T	C	0.39	Cam12W	3
	135	6	18,528,943	12,258,527	29,438,316	0.28	-0.78	18.7	G	C	0.31	Cam12W	6
	147	6	101,482,646	101,463,249	101,501,936	0.22	-0.65	15.04	T	A	0.17	Cam12W	5
	173	8	12,782,777	12,767,198	12,798,330	0.17	-0.61	11.8	C	T	0.22	Cam12W	4

"Pos" indicates the physical position in base pair of the SNP with the strongest association on the V2 of reference genome. *LowerLimit* and *UpperLimit* indicate the lower and upper physical limits estimated by LD windows (*LD_win*) for each QTL. The proportion of the variance explained (R^2), the effect of the major allele (*Effect*) as outputted by FastLMM, $-log_{10}(\mathbf{Pval})$ (*Log*), the minor and major alleles (Minor All and Major All) and the minor allele frequency (*MAF*) of the most significant SNP within the QTL are shown. The following columns

22

represent environment for which the most associated QTL was observed (EnvMax) and the number of different environments in which QTL are detected (NbDiffEnv) are shown. Note that QTLs 71-72 for the plant height and QTLs 129-130 for the grain yield are genetically close (<1cM) and display high mean LD (r^2K>0.5). Hence, QTLs 71-72 and 129-130 can potentially be merged.

23

457 The distribution of QTLs was not homogeneous along the genome since 82%, 77% and 79%
458 of flowering time, plant height and grain yield QTLs, respectively, were located in the high
459 recombinogenic regions, whereas they represented 46% of the physical genome (Additional
460 file 16: Table S3). The QTLs were more stable (≥ 2 environments) in low than in high
461 recombinogenic regions (12.8% vs 5.8%, *p-value* = 0.03).

462 **Discussion**

GBS required massive imputation but displayed similar global trends than DNA arrays for genetic diversity organization

463 In order to reduce genotyping cost, GBS is most often performed at low depth leading to a
464 high proportion of missing data, thereby requiring imputation in order to perform GWAS.
465 Imputation can produce genotyping errors that can cause false associations and introduce bias
466 in diversity analysis [33]. We evaluated the quality of genotyping and imputation obtained by
467 different approaches, taking the 50K or 600K as references. The best imputation method that
468 yielded a fully genotyped matrix with the lowest error rate for the prediction of both
469 heterozygotes and homozygotes was the approach merging the homozygous genotypes from
470 TASSEL and the imputation of Beagle for the other data (GBS_5 Table 1). The quality of
471 imputation was high with 96% of allelic values consistent with those of the 50K and 600K.
472 This level of concordance is identical to a study of USA national maize inbred seed bank by
473 Romay *et al.* [32]. It is higher than in a diversity study of European flint maize collection
474 (93%) by Gouesnard *et al.* [33], which was more distant from the reference AllZeaGBSv2.7
475 database than for the panel presented here. For further studies, integrating genotyping data
476 from the three technologies may reduce imputation errors for missing data of GBS [35].
477

24

478 The ascertainment bias of SNP-arrays due to the limited number of lines used for SNP
479 discovery was reinforced by counter-selection of rare alleles during the design process of
480 DNA arrays [3, 4]. For GBS, the polymorphism database to call polymorphisms included
481 thousands of diverse lines [38]. In our study, we used AllZeaGBSv2.7 database. After a first
482 step of GBS imputation (GBS_2), missing data dropped to 11.9% *i.e.* only slightly more than
483 in Romay *et al.* (10%) [32]. This confirms that the polymorphism database (AllZeaGBSv2.7)
484 covered adequately the genetic diversity of our genetic material.
485
486 Although, we observed differences of allelic frequency spectrum between GBS and DNA
487 arrays, these technologies revealed similar trends in the organization of population structure
488 and relatedness (Figure 2, Additional file 4: Figure S4) suggesting no strong ascertainment
489 bias for deciphering global genetic structure trends in the panel. However, although highly
490 correlated, level of relatedness differed between GBS and DNA arrays, especially when the
491 lines were less related as showed by the deviation (to the left) of the linear regression from the
492 bisector (Additional file 5: Figure S5).

The extent of linkage disequilibrium strongly varied along and between chromosomes

493 Linkage disequilibrium extent in high recombinogenic regions varied to a large extent among
494 chromosomes, ranging from 0.012 to 0.062 cM. Similar variation of genetic LD extent
495 between maize chromosomes has been previously observed by Rincent et al. [14], but their
496 classification of chromosomes was different from ours. This difference could be explained by
497 the fact that we analyzed specifically high and low recombination regions. According to Hill
498 and Weir Model [40], the physical LD extent in a genomic region increased when the local
499 recombination rate decreased. As a consequence, chromosome 1 and 9 had the lowest and

highest physical LD extent and displayed the highest and one of the lowest recombination rate in pericentromeric regions, respectively (0.26 vs 0.11 cM / Mbp, Table 2 and Additional file 16: Table S3). Unexpectedly, the genetic LD extent also correlated negatively with the recombination rate. It suggested that chromosomes with a low recombination rate also display a low effective population size. Background selection against deleterious alleles could explain this pattern since it reduces the genetic diversity in low recombinogenic regions [41, 42]. Finally, we observed a strong variation of the LD extent along each chromosome . As we used a consensus genetic map [43] that represents well the recombination within our population, it suggested, according to Hill and Weir's model, that the number of ancestors contributing to genetic diversity varied strongly along the chromosomes. This likely reflects the selection of genomic regions for adaptation to environment or agronomic traits [41], that leads to a differential contribution of ancestors according to their allelic effects. Ancestors with strong favorable allele(s) in a genomic region may lead ultimately to large identical by descent genomic segments [44].

SNPs were clustered into QTL highlighting interesting genomic regions

In previous GWAS studies, the closest associated SNPs were grouped into QTLs according to either a fixed physical distance [1] or a fixed genetic distance [30, 43]. These approaches suffer of two drawbacks. First, the physical LD extent can vary strongly along chromosomes according to the variation of recombination rate (Figure 1 and Additional file 3: Figure S3). Second, the genetic LD extent depends both on panel composition and the position along the genome (Table 2). These approaches may therefore strongly overestimate or underestimate the number of QTLs. To address both issues Cormier *et al.* [46] proposed to group associated SNPs by using a genetic window based on the genetic LD extent estimated

by Hill and Weir model in the genomic regions around the associated peaks [40]. In our study, we improved this last approach (*LD_win):*

- First, we used r^2K that corrected r^2 for kinship rather than the classical r^2 since r^2K reflected the LD addressed in our GWAS mixed models to map QTL [17].

- Second, we took advantage of the availability of both physical and genetic maps of maize to project the genetic LD extent on the physical map. This physical window was useful to retrieve the annotation from B73 reference genome, decipher local haplotype diversity (Figure 5) and estimate physical genome coverage (Table 2, 1, Additional file 3: Figure S3).

- Third, we considered an average LD extent estimated separately in the high and low recombinogenic genomic regions. This average was estimated by using several large random sets of pairs of loci in these regions rather than the local LD extent in the genomic regions around each associated peaks.

We preferred this approach rather than using local LD extent in order to limit the effect of (i) the strong variation of marker density along the chromosome (Additional file 3: Figure S3), (ii) the local ascertainment bias due to the markers sampling (iii) the poor estimation of the local recombination rate using a genetic map, notably for low recombination regions [3, 44] (iv) errors in locus order due to assembly errors or chromosomal rearrangements.

We compared *LD_win* with *LD_adj,* another approach based on LD to group the SNPs associated to trait variation into QTL. The discrepancies between the two approaches can be explained by the local recombination rate and LD pattern. Since *LD_adj* approach was based on the grouping of contiguous SNPs according to their LD, this approach was highly sensitive to (i) error in marker order or position due to genome assembly errors or structural variations,

546 which are important in maize [47] (ii) genotyping or imputation errors, which we estimated at
547 *ca.* 1% and *ca.* 4%, respectively, for GBS (Table 1), (iii) presence of allelic series with
548 contrasted effects in different experiments which are currently observed in maize [45], (iv)
549 LD threshold used. On the other hand, *LD_win* lead either to inflate the number of QTLs in
550 high recombinogenic regions in which SNPs were too distant genetically to be grouped, or
551 deflated their number by grouping associated SNPs in low recombinogenic regions. Since
552 *LD_win* considered the average LD extent, this method could conduct either to separate or
553 group abusively SNPs when local LD extent was different than the global LD extent.
554 Simulations will be carried out in further research to better understand the properties of
555 *LD_win* and *LD_adj* approach.
556
557 Note that LD windows should not be considered as confidence intervals since the
558 relationship between LD and recombination is complex due to demography, drift and
559 selection in association panels, contrary to linkage based QTL mapping [17]. The magnitude
560 of the effect of causal polymorphism in the estimation of these intervals, which is well
561 established for linkage mapping, should be explored further [48]. Other approaches have been
562 proposed to cluster SNPs according to LD [45, 46]. These approaches aim at segmenting the
563 genome in different haplotype blocks separating by high recombination regions. These
564 methods are difficult to use for estimating putative windows inside which the causal
565 polymorphisms are located because such approaches are not centered on the associated SNP.
566 Several QTLs identified by *LD_win* in our study correspond to regions previously
567 identified: in particular six regions associated with female flowering time [27] and 30 regions
568 associated with different traits in the Cornfed dent panel [11]. Conversely, we did not identify
569 in our study any QTL associated to the florigen *ZCN8*, which showed significant effect in

570 these two previous studies. One of the explanation is that we narrowed the flowering time
571 range in our study, in particular by eliminating early lines. This reduced the representation of
572 the early allele at the Zcn8 locus, leading to a MAF of 0.27 in our study vs. 0.35 in Rincent *et*
573 *al.* [11], which can slightly diminish the power of the tests [14]. Also, this effect may have
574 strengthened by frequency evolution at loci involved in epistatic interactions with Zcn8 (see
575 [47] for a recent demonstration of such effects).

Complementarity of 600K and GBS for QTL detection resulted mostly from the tagging of different haplotypes rather than the coverage of different genomic regions.

576 Number of significant SNPs and QTLs increased with the increase in marker number (Table
577 4, Additional file 9: Figure S9). This could be explained partly by a better coverage of some
578 genomic regions by SNPs, notably in high recombinogenic regions which showed a very short
579 LD extent and were enriched in QTLs (Additional file 16 Table S3). Numerous new QTLs
580 identified by the 600K and GBS as compared with those identified by the 50K were detected
581 in high recombinogenic regions that were considerably less covered by the 50K than the 600K
582 or GBS (Figure 1 and Additional file 3: Figure S3).
583
584 The high complementarity for QTL detection between GBS and 600K was only explained to a
585 limited extent by the difference of the SNP distribution and density along the genome, since
586 these two technologies targeted similar regions as showed by coverage analysis (Figure 1 and
587 Additional file 3: Figure S3). However, at a finer scale, SNPs from the 600K and GBS could
588 tag close but different genomic regions around genes. SNPs from the 600K were mostly
589 selected within coding regions of genes [4], whereas SNP from GBS targeted more largely
590 low copy regions, which included coding but also regulatory regions of genes [32, 38]. To

591 further analyse the complementarity of the technologies, we analysed local haplotypes and the
592 effect of genome coverage differences between technologies on QTL detection. We showed
593 that both technologies captured different haplotypes when similar genomic regions were
594 targeted (Figure 5). In this figure, two QTLs were specifically detected by markers from
595 either 600K or GBS although there are several markers from other technology very close from
596 the most associated marker, considering the size of LD windows around it . Additionally, we
597 did not observe an enrichment of QTL specifically detected by one technology in 20kbp-
598 intervals with high genomic coverage difference between 600K and GBS Hence, we
599 pinpointed that GBS and DNA arrays are highly complementary for QTL detection because
600 they tagged different haplotypes rather than different regions (Figure 5). Based on the L-
601 shaped MAF distribution, which suggests no ascertainment bias, and the high number of
602 sequenced lines used for the GBS, we expect a closer representation of the variation present in
603 our panel by this technology compared to the 600K, but this comes to the cost of an
604 enrichment in rare alleles. Both factors tend to counterbalance each other in terms of GWAS
605 power (Additional file 13: Figure S12).
606
607 Our results suggest that we did not reach saturation with our *c.* 800,000 SNPs because (i)
608 some haplotypes certainly remain not tagged (ii) the genome coverage was not complete, and
609 (iii) the number of significant SNPs and QTLs continued to increase with marker density
610 (Additional file 9: Figure S9). Considering LD and marker density, the genotypic data
611 presently available were most likely enough to well represent polymorphisms in the
612 centromeric regions, whereas using more markers would be beneficial for telomeric regions.
613 New approaches based on resequencing of representative lines and imputation are currently
614 developed to achieve this goal.

Methods

Plant Material and Phenotypic Data

The panel involves 247 maize inbred lines, further referred to as DROPS panel (Additional file 17: Table S4). They include 164 lines from a wider panel of lines from Europe and America [11] and 83 additional lines derived from public breeding programs in Hungary, Italy and Spain and recent lines free of patent from the USA. All lines belong to the dent genetic group, which can be subdivided in different sub-groups (see [11, 30]). Lines were selected within a restricted flowering time window (10 days) in order to limit the effect of drought escape due to flowering time variation in the identification of genomic regions involved in drought tolerance [30]. Candidate lines with poor sample quality, i.e. high level of heterozygosity, or high relatedness with other lines were discarded in this selection. The lines selection was also guided by pedigree to avoid as far as possible over-representation of some ancestral materials.

The 247 inbred lines were all crossed with a common line (UH007) from the Flint genetic groups to obtain 247 hybrids (hybrid panel). Dent and Flint genetic groups are known to be complementary to produce hybrids [48]. Further, as UH007 is unrelated to any line in the panel, no hybrid is affected by inbreeding depression. This guarantees that hybrids have a level of performance and an overall physiology comparable to that of varieties used in agriculture. Conversely, field evaluation of inbred lines per se would have diminished yield by more than 50%.

Experimental design and model used for obtaining adjusted means for male flowering time (Day To Anthesis, DTA), plant height (plantHT), and grain yield (GY) were previously

31

637 described [30]. While DTA and GY were previously analyzed in [30], PlantHT was not.
638 Briefly, the hybrid panel were evaluated for these three traits in 22 experiments (combination
639 year x site x water regime), *i.e.* at seven sites in Europe, during two years (2012 and 2013),
640 and for two water treatments (watered and rainfed) [30]. Experiments were designed as alpha-
641 lattice designs with two and three replicates for watered and rain-fed regimes, respectively.
642 Grain yield (t ha^{-1}) was adjusted to 15% moisture. The adjusted mean (Best Linear Unbiased
643 Estimation, BLUEs, https://doi.org/10.15454/IASSTN) of the three traits were estimated per
644 environment (site × year × water regime) using a mixed model based on fixed hybrid and
645 replicate effects, random spatial effects (rows and columns), and spatially correlated errors in
646 order to take into account spatial variation of micro-environment in each field trial (see [30]
647 for more details). The same model, but with random hybrids effects, was used to estimate
648 variance components. Models were fitted with ASReml-R [49]. Narrow-sense heritability of
649 each trait in each environment were also estimated as in [30] (Additional file 18: Table S5).
650 As all hybrids share a common parent (UH007), adjusted means (BLUEs) of hybrids were
651 combined with genotyping data of the corresponding dent inbred lines of the panel to perform
652 GWAS, following a usual practice in maize genetics [11].

Genotyping and Genotyping-By-Sequencing Data

653 The 247 inbred lines were genotyped using three technologies: a maize Illumina Infinium HD
654 50K array [3], a maize Affymetrix Axiom 600K array [4], and Genotyping-By-Sequencing [2,
655 38]. In the arrays, DNA fragments are hybridized with probes attached to the array
656 (Additional file 19: Notes S1 for the description of the data from the two SNP-arrays).
657 Genotyping-by-sequencing technology is based on multiplex resequencing of tagged DNA
658 using restriction enzyme (Keygene N.V. owns patents and patent applications protecting its

659 Sequence Based Genotyping technologies) [2]. Cornell Institute (NY, USA) processed raw
660 sequence data using a multi-step Discovery and a one-step Production pipeline (*TASSEL-*
661 *GBS*) in order to obtain genotypes (Additional file 19: Notes S1). An imputation step of
662 missing genotypes was carried out by Cornell Institute [39], which utilized an algorithm that
663 searches for the closest neighbour in small SNP windows across the haplotype library [38].
664
665 We applied different filters (heterozygosity rate, missing data rate, minor allele frequency) for
666 a quality control of the genetic data before performing the diversity and association genetic
667 analyses. For GBS data, the filters were applied after imputation using the method
668 "Compilation of Cornell homozygous genotypes and Beagle genotypes" (GBS_5 in Additional
669 file 1: Figure S1; See section "Evaluating Genotyping and Imputation Quality"). We
670 eliminated markers that had an average heterozygosity and missing data rate higher than 0.15
671 and 0.20, respectively, and a Minor Allele Frequency (MAF) lower than 0.01 for the diversity
672 analyses and 0.05 for the GWAS (Additional file 20: Table S6). Individuals which had
673 heterozygosity and/or missing data rate higher than 0.06 and 0.10, respectively, were
674 eliminated.
675

Evaluating Genotyping and Imputation Quality

676 Estimation of genotyping and imputation quality was performed using the entire panel except
677 two inbred lines that had different seedlots between technologies. The 50K and the 600K were
678 taken as reference to compare the concordance of genotyping (genotype matches) with the im-
679 putation of GBS based on their position. While SNP positions and orientation from GBS were
680 called on the reference maize genome B73 AGP_v2 (release 5a) [50], flanking sequences of

33

SNPs in the 50K were primary aligned on the first maize genome reference assembly B73 AGP_v1 (release 4a.53) [51]. Both position and orientation scaffold carrying SNPs from the 50K can be different in the AGP_v2, which could impair correct comparison of genotype between the 50K and GBS. Hence, we aligned flanking sequences of SNPs from the 50K on maize B73 AGP_v2 using the Basic Local Alignment Search Tool (BLAST) to retrieve both positions and genotype in the same and correct strand orientation (forward) to compare genotyping. The number of common markers between the 50K/600K, 50K/GBS, GBS/600K and 50K/600K/GBS was 36,395, 7,018, 25,572 and 5,947 SNPs, respectively. The comparison of the genotyping and imputation quality between the 50K/GBS, 50K/600K and 600K/GBS was done on 5,336 and 24,286 and 26,154 common markers, respectively. The comparison for the 50K involved PANZEA markers, prefixed as "PZE" [52]. In order to achieve these comparisons, we considered the direct reads from GBS (**GBS_1**) and four approaches for imputation (GBS_2 to GBS_5, Additional file 1: Figure S1). **GBS_2** approach consisted of one imputation step from the direct read by Cornell University, using *TASSEL* software, but missing data was still present. **GBS_3** approach consisted of imputation by *Beagle v3* [13] of the missing data of GBS_1. To compare data from GBS_3 and GBS_2 to those of the 50K and 600K, missing data in GBS_2 were excluded from GBS_3. In **GBS_4,** genotype imputation by Beagle was performed on Cornell imputed data after replacing the heterozygous genotypes with missing data. **GBS_5**, consisted of homozygous genotypes of GBS_2 completed by values imputed in GBS_3, no missing data remained (Additional file 1: Figure S1).

Diversity Analyses

After excluding the unplaced SNPs and applying the filtering criteria for the diversity analyses (MAF > 0.01), we obtained the final genotyping data of the 247 lines with 44,729

34

703 SNPs from the 50K, 506,662 SNPs from the 600K array, and 395,024 SNPs from the GBS
704 (Additional file 20: Table S6). All markers of the 600K and GBS_5 that passed the quality
705 control were used to perform the diversity analyses (estimation of Q genetic groups and K
706 kinships). For the 50K, we used only the PANZEA markers (29,257 SNPs) [52] in order to
707 reduce the ascertainment bias noted by Ganal *et al.* [3] when estimating Nei's index of
708 diversity [53] and relationship coefficients. Genotypic data generated by the three
709 technologies were organized as *G* matrices with *N* rows and *L* columns, *N* and *L* being the
710 panel size and number of markers, respectively. Genotype of individual *i* at marker *l* ($G_{i,l}$) was
711 coded as 0 (the homozygote for an arbitrarily chosen allele), 0.5 (heterozygote), or 1 (the
712 other homozygote). Identity-By-Descent (IBD) was estimated according to Astle and Balding
713 [19]:

714 $$K\_Freq_{i,j} = \frac{1}{L}\sum_{l=1}^{L}\frac{\left(G_{i,l}-p_l\right)\left(G_{j,l}-p_l\right)}{p_l\left(1-p_l\right)} \quad ,$$

715 where p_l is the frequency of the allele coded 1 of marker *l* in the panel of interest, *i*
716 and *j* indicate the inbred lines for which the kinship was estimated. We also estimated the
717 Identity-By-State (IBS) by estimating the proportion of shared alleles. For GWAS, we used
718 *K_Chr* [14] that are computed using similar formula as *K_Freq*, but with the genotype data of
719 all the chromosomes except the chromosome of the SNP tested. This formula provides an
720 unbiased estimate of the kinship coefficient and weights by allelic frequency assuming Hardy-
721 Weinberg equilibrium. Hence, relatedness is higher if two individuals share rare alleles than
722 common alleles.

723

724 Genetic structure was analysed using the sofware *ADMIXTURE v1.22* [18] with a
725 number of groups varying from 2 to 10 for the three technologies. We compared assignation
726 by *ADMIXTURE* of inbred lines between the three technologies by estimating the proportion

of inbred lines consistently assigned between technologies two by two (50K *vs* GBS_5, 50K *vs* 600K, 600K *vs* GBS_5) using a threshold of 0.5 for admixture.

Expected heterozygosity (He) [53] was estimated at each marker as $2p_l(1 - p_l)$ and was averaged on all the markers for a global characterization of the panel for the three technologies. Principal Coordinate Analyses (PCoA) were performed on the genetic distance matrices [54], estimated as $1_{N,N} - K\_Freq$, where $1_{N,N}$ is a matrix of ones of the same size as *K_Freq*.

Linkage Disequilibrium Analyses

We first analyzed the effect of the genetic structure and kinship on linkage disequilibrium (LD) extent within and between chromosomes by estimating genome-wide linkage disequilibrium using the 29,257 PANZEA SNPs from the 50K. Four estimates of LD were used: the squared correlation (r^2) between allelic dose at two markers [55], the squared correlation taking into account global kinship with *K_Freq* estimator (r^2K), the squared correlation taking into account population structure (r^2S), and the squared correlation taking into account both (r^2KS) [17].

To explore the variation of LD decay and the stability of LD extent along the chromosomes, we estimated LD between a non-redundant set of 810,580 loci from the GBS, the 50K and 600K. To save computation time, we calculated LD between loci within a sliding window of 1 cM. Genetic position was obtained by projecting the physical position of each locus using a *smooth.spline* function R calibrated on the genetic consensus map of the Cornfed Dent Nested Association Mapping (NAM) design [56]. We used the estimator r^2 and r^2K using 10 different

36

749 kinships *K_Chr*. This last estimator was calculated because it corresponds exactly to the LD
750 used to map QTL in our GWAS model. It determines the power of GWAS to detect QTL
751 considering that causal polymorphisms were in LD with some polymorphisms genotyped in
752 our panel [17]. To study LD extent variation, we estimated LD extent by adjusting Hill and
753 Weir's model [40] using non-linear regression (*nls* function in R-package *nlme*) against both
754 physical and genetic position within each chromosome. Since recombination rate (cM / Mbp)
755 varied strongly along the genome (Figure 1 andAdditional file 3: Figure S3), we defined high
756 (>0.5 cM / Mbp) and low (<0.5 cM / Mbp) recombinogenic genomic regions within each
757 chromosome. We adjusted Hill and Weir's model [40] separately in low and high
758 recombinogenic regions (Additional file 16: Table S3)by randomly sampling 100 sets of
759 500,000 pairs of loci distant from less than 1 cM. This random sampling avoided over-
760 representation of pairs of loci from low recombinogenic regions due to the sliding-window
761 approach (Figure 3). 500,000 pairs of loci represented 0.36% (Chromosome 3 / High rec) to
762 1.20% of all pairs of loci (Chromosome 8 / High rec).
763 For all analyses, we estimated LD extent by calculating the genetic and physical distance for
764 the fitted curve of Hill and Weir's Model that reached r^2K=0.1, r^2K=0.2 and r^2K=0.4.

Genome coverage estimation

765 In order to estimate the genomic regions in which the effect of an underlying causal
766 polymorphisms could be captured by GWAS using LD with SNP from three technologies, we
767 developed an approach to define LD windows around each SNP with MAF ≥ 5% based on LD
768 extent (Figure 3). To set the LD window around each SNP, we used LD extent with r^2K=0.1
769 (negligible LD), r^2K=0.2 (intermediate LD) and r^2K=0.4 (high LD) estimated in low and high
770 recombinogenic regions for each chromosome. We used the global LD decay estimated for

771 these large chromosomal regions rather than local LD extent (i) to avoid bias due to SNP
772 sampling within small genomic regions, (ii) to reduce computational time, and (iii) to limit the
773 impact of possible local error in genome assembly. In low recombinogenic regions, we used
774 the physical LD extent, hypothesizing that recombination rate is constant along physical
775 distance in these regions. In high recombinogenic regions, we used the genetic LD extent
776 since there is a strong variation of recombination rate by base pair along the physical position
777 (Figure 1 and Additional file 3: Figure S3). We then converted genetic LD windows into
778 physical windows by projecting the genetic positions on the physical map using the
779 *smooth.spline* function implemented in R, calibrated on the NAM dent consensus map [56].
780 Reciprocally, we obtained the genetic positions of LD windows in low recombinogenic
781 regions by projecting the physical boundaries of LD windows on the genetic map.
782
783 To estimate coverage of the three technologies to detect QTLs based on their SNP distribution
784 and density, we calculated cumulative genetic and physical lengths that are covered by LD
785 windows around the markers, considering different LD extents for each chromosome
786 (r^2K=0.1, r^2K=0.2, r^2K=0.4). In order to explore variation of genome coverage along the
787 chromosome, we estimated the proportion of genome covered using a sliding-windows
788 approach based on variable physical distances (20, 100, 500, 2000 kbp) considering LD extent
789 for a r2K =0.1.

Statistical Models for Association Mapping

790 We used four models to determine the statistical models that control best the confounding
791 factors (*i.e.* population structure and relatedness) in GWAS (Additional file 21: Notes S2). We
792 tested different software implementing either approximate (EMMAX) [8] or exact

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793 computation of standard test statistics (ASReml and FaST-LMM) [6, 49] for computational
794 time and GWAS results differences. Single-trait, single-environment GWAS was performed
795 for each marker for each environment and all traits using FaST-LMM. We selected the mixed
796 model using *K_Chr*, estimated from PANZEA markers of the 50K to perform GWAS on 66
797 situations (environment × trait) (Additional file 21: Notes S2). We developed a GWAS
798 pipeline in *R v3.2.1* [55] calling FaST-LMM software and implementing [14] approaches to
799 conduct single trait and single environment association tests.
800
801 To take into account multiple tests in GWAS and their dependence, we applied the methods of
802 Moskvina and Schmidt [56] and Gao *et al.* [57, 58] to infer the number of independent tests to
803 be considered in the Bonferroni formula. Using the Gao et al. [57, 58] approaches, we
804 estimated the number of independent tests for GWAS at 15,780 for the 50K, 92,752 for the
805 600K, 109,117 for the GBS_5 and 191,026 for the combined genetic data (i.e. merging of 50K,
806 600K, GBS), leading to different $-log_{10}(p\text{-}value)$ thresholds: 5.49, 6.27, 6.34 and 6.58,
807 respectively. Because of these differences, we used two thresholds of $-log_{10}(p\text{-}value) = 5$ (less
808 stringent) *and 8* (hightly conservative and slightly above Bonferroni) for comparing GWAS to
809 avoid the differences of identification of significant SNPs between the technologies due to the
810 choice of the threshold.
811

Methods for grouping associated SNPs into QTLs

812 We used two approaches based on LD for grouping significant SNPs. The first approach
813 (*LD_win*) used LD windows, previously described, to group significant SNPs into QTLs
814 considering that all significant SNPs with overlapping LD windows of $r^2K=0.1$ belong to the

39

same QTL (Figure 3). We hypothesized that significant SNPs with overlapping LD windows at $r^2K=0.1$ captured the same causal polymorphism and were therefore a single and unique QTL. For the second approach (*LD_adj*), significant SNPs were grouped into a same QTL if they were connected in terms of LD (r^2K between adjacent significant SNPs superior to *0.5*). We used LD heatmaps for comparing the SNP grouping produced by the two approaches on the three different traits across all environments (Additional file 7: Figure S7-LD-Adjacent and Additional file 8: Figure S8-LD-Windows). All scripts are implemented in R software [59].

Resampling approach to analyze effect of MAF distribution, SNP distribution along the genome, SNP density on QTL detection

To study the effect of SNP density, MAF distribution and SNP distribution along the genome on association and QTL detection, we used a resampling approach of several sets of SNPs displaying different MAF distribution and SNP distribution along the chromosome. We compared these modalities with different SNP densities (50,000, 100,000, 150,000, 200,000, 250,000 markers). In this resampling approach, we considered all markers together and that both associations and QTLs detected by the whole SNP sets are true. We selected only markers having MAF above 5%. To study the effect of MAF distribution on QTL detection SNPs were classified in 5 MAF classes (0-0.1, 0.1-0.2, 0.2-0.3, 0.3-0.4 and 0.4, 0.5) and SNP were randomly selected in each classes according to MAF distribution t1) similar to GBS (GBS_MAF) 2) similar to 600K (600K_MAF) 3) with equal frequency for 5 MAF classes (Flat_MAF) 4) skewed towards high MAF (High_MAF) with SNP frequency of 0, 0, 0.2, 0.4, 0.4 in (0-0.1], (0.1-0.2], (0.2-0.3], (0.3-0.4], (0.4-0.5] MAF classes, respectively 5)

40

835 skewed towards low MAF (Low_MAF) with SNP frequency of 0, 0, 0.2, 0.4, 0.4 in (0-0.1],
836 (0.1-0.2], (0.2-0.3], (0.3-0.4], (0.4-0.5] MAF classes, respectively.

837 To study the effect of SNP distribution along the genome on QTL detection, we compared
838 5 different SNP distributions along the chromosome: 1) evenly distributed according to the
839 physical distance (Dens_Phys), 2) evenly distributed according to the genetic distance
840 (Dens_Gen), 3) distributed like GBS (Dens_GBS), 4) distributed similarly to 600K
841 (Dens_600K) 5) distributed like 50K (Dens_50K). SNPs were sampled randomly according to
842 the different densities in contiguous windows of 10Mbp.

843 **List of abreviations**

844 DTA = Day to Anthesis
845 GY = Grain Yield adjusted at 15% moisture
846 plantHT = Plant Height
847 GBS = Genotyping By Sequencing
848 LD = Linkage disequilibrium
849 GWAS = Genome-Wide Association Studies
850 MAF = Minimum Allelic Frequency
851 SNP = Single Nucleotide Polymorphism
852 HRR = High Recombinogenic Regions
853 LRR = Low Recombinogenic Regions
854 QTL = Quantitative Trait Locus
855

Declarations

Ethics approval and consent to participate

Not applicable.

Consent for publication

Not applicable.

Availability of data and material

The following links toward the data will be available upon publication of this paper.

All the genotyping data used in this study can be found at https://doi.org/10.15454/AEC4BN.

Genotyping data will become publicly available upon the publication with link above.

Genotyping data will be accessible anonymously with following private link for reviewers:

https://data.inra.fr/privateurl.xhtml?token=dc792926-6996-4767-bbbe-bb347fd1edd4

The GWAS results can be found at https://doi.org/10.15454/6TL2N4.

GWAS results will become publicly available upon the publication with link below. GWAS results will be accessible anonymously with following private link for reviewers:

https://data.inra.fr/privateurl.xhtml?token=1b99f286-94c8-40d1-a289-c18db1d6e646

The phenotypic dataset can be found at https://doi.org/10.15454/IASSTN.

Competing interests

The authors declare that they have no competing interests.

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Authors' contributions

S.S.N., S.D.N. and A.C., designed the studied and wrote the article. S.S.N. performed genotyping data quality control, imputation and genetic analyses. S.D.N. developed and performed LD analyses. A.C. designed the association panel with the help of S.D.N. and C.W. C.B. participated in assembling the dent inbred lines panel, organizing the germplasms and field work for seeds production. E.J.M., C.W. and F.T. collected and analysed the phenotypic data. V.C. and D.M. performed DNA extraction and prepared the samples. All authors critically reviewed and approved the final manuscript.

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43

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Authors' information (optional)

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Figure legends

Figure 1: Variation of the markers density, the recombination rate and the genome coverage in non-overlapping 2 Mbp windows along chromosome 3. Markers have MAF above 5%. Top panel shows the variation of SNP number. In the bottom panel, dotted line represents the variation of recombination rate (cM / Mbp) and solid lines the proportion of genome covered by the SNPs using the cumulated length of physical LD windows around each SNP in each 2Mbp-windows. In these two panels, green, blue, red and black lines represent variation for GBS, 600K, 50K and combined technologies, respectively. Vertical dotted gray lines indicate limits of centromeric regions. Vertical lines between the two panels indicate the position of QTLs for flowering time (DTA), grain yield (GY) and Plant Height (PHT). Green, blue, red

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vertical lines indicate QTLs detected only by GBS, 600K and 50K technologies, respectively. Grey lines indicated QTLs detected by at least two technologies. Only QTL including a marker associated with -$\log_{10}$(pval) above 6 were shown.

Figure 2: Principal coordinate analysis (PCoA) of the DROPS panel. The PCoA was based on the covariance matrix *K_Freq* estimated from the 50K Illumina array. The genetic groups identified by ADMIXTURE (N_Q = 4) are colored. Three key founders are indicated (Iodent: PH207 in red, Stiff Stalk: B73 in blueviolet, Lancaster: Mo17 in turquoise).

Figure 3: Linkage disequilibrium based approach to delineate a physical window around each SNP, exemplified with chromosome 3. Linkage disequilibrium (LD) windows were defined for each SNP based on physical LD extent in low recombinogenic regions (left part) and based on genetic LD extent in high recombinogenic regions (right part). These LD windows were used (i) to group significant SNPs into QTLs when they overlapped, (ii) to estimate genome coverage region covered by LD windows around SNPs, and (iii) identify putative genes underlying QTLs involved in trait variations.

Figure 4: Complementarity of the three technologies to detect QTLs. The numbers of specific QTLs detected by each technology for the three traits (flowering time, plant height, grain yield) are shown.

Figure 5: Complementarity of QTLs detection between the 600K array and the GBS for two regions (QTL 32/QTL95). (**a**) Manhattan plot of the -log_{10}(*p*-value) along the genome. Dotted red lines correspond to QTL32 and QTL95 located on chromosome 1 and 3, respectively, for the flowering time in one environment (Ner13R). (**b**) Local manhattan plot of the -log_{10}(*p-value*) (top) and linkage disequilibrium corrected by the kinship (r^2K) (bottom) of all SNPs with the strongest associated marker within QTL 32 (left) and QTL 95 (right). Colored vertical lines between manhattan plot and linkage disequilibrium plot represents the

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distribution of markers for different technologies. Dotted lines between panels b and c linked the first marker, the most associated marker, and the last marker of each QTL (**c**) Local haplotypes displayed by all SNPs within the QTLs 32 (left) and 95 (right) with MAF>5%. Inbred lines are in rows and SNPs are in columns. Inbred lines were ordered by hierarchical clustering based on local dissimilarity estimated by all SNPs within each QTL. Genotyping matrix is colored according to their allelic dose at each SNP. Red and black represent homozygotes and gray represent heterozygotes. The associated peaks (red vertical lines) and other associated SNPs with $-log_{10}(p\text{-}value) > 5$ (orange vertical lines) are indicated above the genotyping matrix. H1, H2, H3, H4, H5 represent the 5 and 3 haplotypes obtained by cutting the dendograms with the most 5 and 3 dissimilar clusters within QTL32 and QTL95, respectively.

Additional file legends

Additional file 1 (.docx):

Figure S1: Different approaches used to impute missing data of the GBS. We considered the direct reads from GBS (**GBS_1**) and four approaches for imputation (GBS_2 to GBS_5). **GBS_2** approach consisted in one imputation step from the direct read by Cornell University, using *TASSEL* software, but missing data was still present. **GBS_3** approach consisted in a genotype imputation of the whole missing data of the direct read by *Beagle v3*. In **GBS_4,** genotype imputation by Beagle was performed on Cornell imputed data after replacing the heterozygous genotypes into missing data. **GBS_5**, consisted in homozygous genotypes of GBS_2 completed by values imputed in GBS_3.

Additional file 2 (.docx):

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Figure S2: Comparison of genotyping data between 50K and 600K arrays, and GBS. (a) Distribution of minor allele frequency per SNP before filtering (monomorphic SNPs removed). (b) Distribution of SNP missing data proportion for the 50K array, 600K array, GBS direct reads (GBS_1) and GBS after imputation by Cornell Institute (GBS_2, note that the scale of the x-axes is different). (c) Relatedness distribution (Identity-By-State, IBS) after QC filtering with MAF≥1% (IBS using GBS_1 was not estimated because of the low calling rate).

Additional file 3 (.pdf):

Figure S3: Variation of the markers density, the recombination rate and the genome coverage in non-overlapping 2 Mbp windows along each chromosome except chromosome 3 (presented in Figure 1). Markers have MAF above 5%. Top panel shows the variation of SNP number. In the bottom panel, dotted line represents the variation of recombination rate (cM / Mbp) and solid lines the proportion of genome covered by the SNPs using the cumulated length of physical LD windows around each SNP in each 2Mbp-windows. In these two panel, green, blue, red and black lines represent variation for GBS, 600K, 50K and combined technologies, respectively. Vertical dotted gray lines indicate limits of centromeric regions. Vertical lines between the two panels indicate the position of QTLs for flowering time (DTA), grain yield (GY) and Plant Height (PHT). Green, blue, red vertical lines indicate QTLs detected only by GBS, 600K and 50K technologies, respectively. Grey vertical lines indicate QTL detected by at least two technologies. Only QTL including a marker associated with $-\log_{10}$(pval) above 6 were shown.

Additional file 4 (.docx):

Figure S4: Contribution of four ancestral populations to 247 inbred lines after ADMIXTURE analysis. Markers from the 50K (top), 600K (middle) and GBS (bottom) were used. One vertical bar corresponds to one individual. Lines were ordered according to contributions

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observed for the 50K. From left to right, we have Stiff Stalk lines type B73 and B14a (blue), Iodent lines type PH207 (red), Lancaster lines type Mo17 and Oh43 (turquoise), a group of lines assembling W117, F7057 type lines (green).

Additional file 5 (.docx):

Figure S5: Correlation between kinship matrix estimated by different technologies. Correlation (*r*) between the IBS and IBD (*K_Freq*) for each technology (A) and correlation of IBD (B) and IBS (D) between the three technologies (after imputation). (C) Correlation of IBD between the three technologies after removing the excess of rare alleles in the GBS to have the same distribution of MAF as in the 50K and the 600K. The red line is the bisector.

Additional file 6 (.docx):

Figure S6: Heatmap of genome-wide linkage disequilibrium (LD) between all markers within and between chromosomes using PANZEA SNPs from the 50K. All SNPs were ordered according to their position on the genome. Dots represent LD between two loci and were colored according to their strength. Classical LD measurement r^2 between loci were represented within triangle below the diagonal. Linkage disequilibrium corrected for structure (r^2S, A), relatedness (r^2K, B) or both (r^2KS, C) were represented within triangle above the diagonal.

Additional file 7 (.pdf):

Figure S7: QTL limits obtained by the *LD_Adj* approach projected on heatmaps representing the level of LD between associated SNPs for each trait (DTA: male flowering time, plantHT: plant height and GY: grain yield) for each chromosome. Upper and lower triangles on the heatmaps represent the r^2 and r^2K values between associated SNPs, respectively. Linkage disequilibrium between loci was colored according to values from weak LD (yellow) to high LD (red). The significant markers were ordered according to their physical positions on the

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chromosome and were represented by ticks on the four sides of the heatmaps. Limits of QTLs were displayed by gray dotted lines. QTL numbers were indicated in gray on the top and the right of each heatmap.

Additional file 8 (.pdf):

Figure S8-LD_Windows: QTL limits obtained by the *LD_win* approach projected on heatmaps representing the level of LD between associated SNPs for each trait (DTA: male flowering time, plantHT: plant height and GY: grain yield) and each chromosome. Upper and lower triangles on the heatmaps represented the r^2 and r^2K values between associated SNPs, respectively. Linkage disequilibrium between loci was colored according to values from weak LD (yellow) to high LD (red). The significant markers were ordered according to their physical positions on the chromosome and were represented by ticks on the four sides of the heatmaps. Limits of QTLs were displayed by gray dotted lines. QTL numbers were indicated in gray on the top and the right of each heatmap.

Additional file 9 (.docx):

Figure S9: Number of significant SNPs (blue line) and QTLs (red line) identified as a function of SNP density (x-axis) for the three traits (DTA, male flowering time; plantHT, plant height; GY, grain yield).

Additional file 10 (.csv):

Table S1: Summary of all the QTLs identified for the male flowering time (DTA), plant height (plantHT) and grain yield (GY). “LowerLimit” and “UpperLimit” columns are the lower and upper physical limits for each QTL. The “Rec” column indicates if the QTL is located in a high or low region of recombination. “NbSNP50”, “LogPvaMax50”, “NbSNP600”, “LogPvaMax600”, “NbSNPGBS”, “LogPvaMaxGBS” are the number of significant SNPs and the most significant $-log_{10}(Pval)$ within the QTL for each technology

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1225 across all environments. The physical position ("PosMax"), the proportion of the variance
1226 explained ("R2_LDMax") and the effect ("EffectMax") of the most significant SNP within
1227 the QTL is shown. "NbDiffEnv" gives the number of different situations that detected the
1228 QTL.

1229 **Additional file 11 (.docx):**

1230 **Figure S10**: Examples of QTL detection on Chromosome 3, 6 and 8 for the different traits.
1231 The top panel represents the distribution of the QTLs along the chromosome of interest, for
1232 the different technologies. The vertical red line in this panel localizes the SNP chosen as
1233 reference for the QTL (marker with the strongest association). The middle panel is a zoom in
1234 the vicinity of the reference SNP, showing the Local distribution of the $-log_{10}(p\text{-}value)$. The
1235 bottom panel is the same zoom as the middle panel and shows the local linkage disequilibrium
1236 corrected by the kinship (r^2k) of all SNPs, within this region, within the reference SNP. Ticks
1237 on different x-axes show the marker density of the three technologies (red for the 50K, blue
1238 for the 600K and green for the GBS).

1239 **Additional File 12 (.pdf)**

1240 **Figure S11**: Effect of minor allelic frequency distribution, SNP distributions along the
1241 genome and SNP densities on the number of associated SNP and QTL detected. Boxplot were
1242 drawn on 100 sets of 50 000 to 250 000 markers sampled according to different MAF
1243 distributions (A, B) and different SNP distributions along the genome (C, D). A, C: number of
1244 SNP associated; B, D: Number of QTL detected. In A and B, 600K_MAF (yellow),
1245 GBS_MAF (green), Low_MAF (cyan), Flat_MAF (blue), High_MAF (pink) on x axis
1246 indicate boxplots corresponding to MAF distribution similar to 600K, similar to GBS, skewed
1247 towards low MAF, flat MAF and skewed toward high MAF, respectively. In C and D,
1248 Dens_50K (red), Dens_600K (yellow), Dens_GBS (cyan), Dens_Gen (blue), Dens_Phys

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(pink) on x axis indicate distribution of SNPs along the genome corresponding to 50K, GBS, 600K, even genetic and physical distances, respectively. For A, B, C and D, modalities indicated as "Random" in x axis correspond to random sample of SNP. Number of markers for each boxplot are indicated after the point.

Additional file 13 (.pdf)

Figure S12: Distribution of markers, associations and QTLs according to MAF for 50K, 600K GBS, and ALL technologies. A) Number of markers, B) Proportion of markers, C) Proportion of Association, D) Proportion of QTLs

Additional file 14 (.docx):

Figure S13: Colocalization of QTLs between the traits. Number of QTLs specific and shared by the three traits across all environments. Note that several QTLs from one trait were sometimes included in a single QTL of another trait.

Additional file 15:

Table S2: Stability of QTLs across environments for the three traits (DTA: male flowering time, plantHT: Plant Height, GY: Grain Yield) and all traits. "Env. Nb" indicates the number of environment in which a QTL was detected. Next four columns indicate the number of QTL corresponding to each cartegory

Additional file 16:

Table S3: Proportion of low and high recombination regions, recombination rate and percentage of QTLs located in these regions for the three traits.
"Chr" indicates the chromosome. Physical and genetic size columns indicated the size of each chromosome in bp and cM, respectively. Average recombination rate ("RecRate") and proportion of the physical ("Phys") and genetic ("Genetic) map in high recombination regions ("HighRec", >0.5 cM / Mbp) for each chromosome are shown. Percentage of QTL in high recombination regions were displayed for three traits (DTA PlantHT, GY)

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Additional file 17 (.xlsx):

Table S4: Description of inbred lines. Variety and accession along with the breeders, seeds providers and genetic groups obtained using ADMIXTURE for K=4 (Stiff Stalk, Iodent, Lancaster, Other).

Additional file 18 (.docx):

Table S5: Narrow sense heritability (h^2) and variance components (V_g, genetic variance; V_e, residual variance). The heritability and variance components were estimated for all traits (grain yield, male flowering time and plant height) using the R package Heritability [1].

Additional file 19(.docx):

Notes S1: Differences between arrays and GBS discrovery / pipelines and.

Additional file 20:

Table S6: Number of SNPs called, after QC filtering (MAF>1%) and useful for GWAS (MAF≥5%). Note that GBS1 have SNPs with 100% missing genotypes which were removed while GBS2 used external haplotype library which allow to impute loci with 100% missing data. It conducted to a smaller number of SNPs for GBS1 than GBS2.

Additional file 21:

Notes S2: GWAS statistical models and effects of confounding factors on GWAS.

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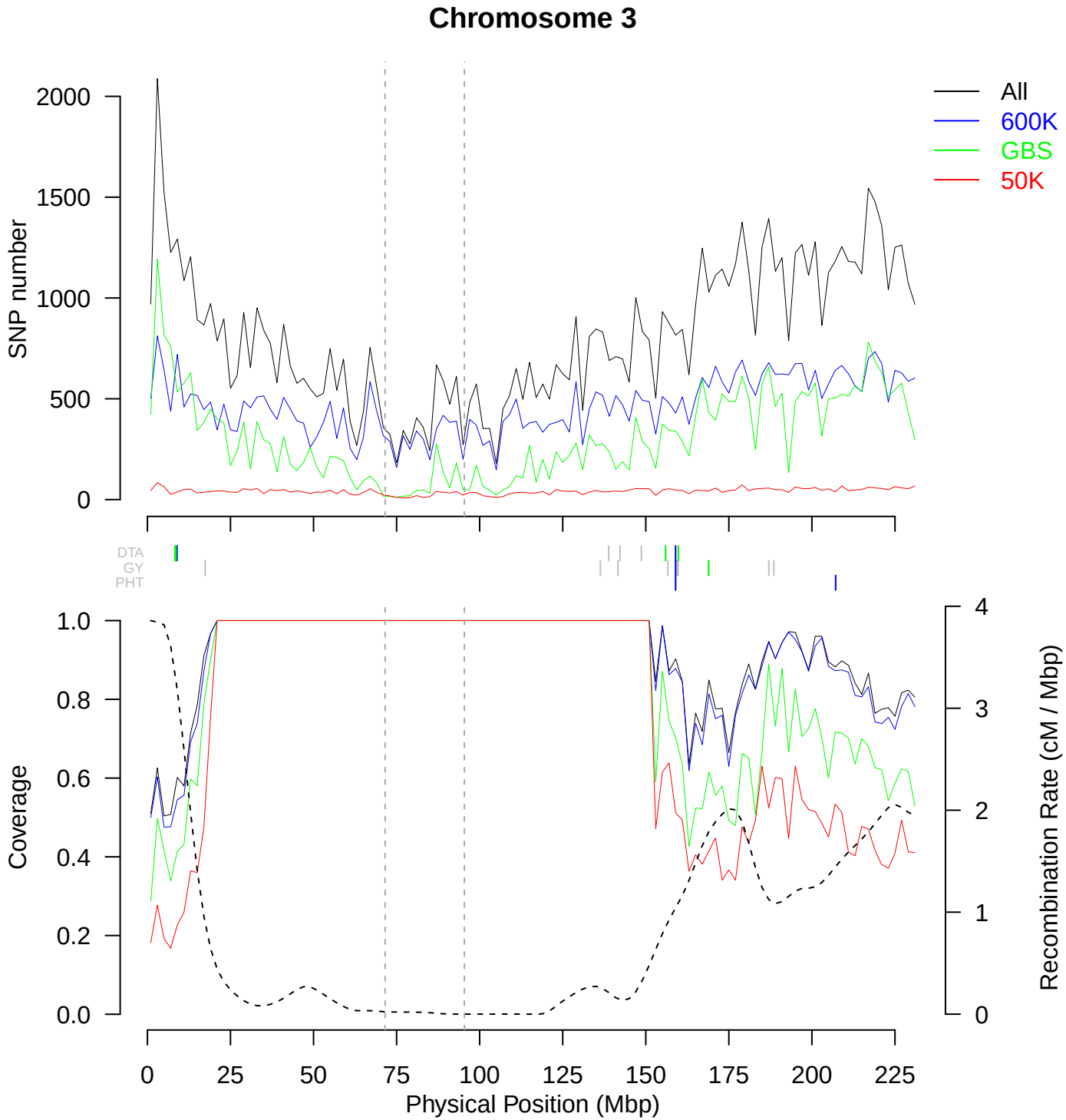

Chromosome 3
All
600K
GBS
50K
SNP number
2000
1500
1000
500
0
DTA
GY
PHT
Coverage
1.0
0.8
0.6
0.4
0.2
0.0
Recombination Rate (cM / Mbp)
4
3
2
1
0
0
25
50
75
100
125
150
175
200
225
Physical Position (Mbp)

Figure 1: Variation of the markers density, the recombination rate and the genome coverage in non-overlapping 2 Mbp windows along chromosome 3. Markers have MAF above 5%. Top panel shows the variation of SNP number. In the bottom panel, dotted line represents the variation of recombination rate (cM / Mbp) and solid lines the proportion of genome covered by the SNPs using the cumulated length of physical LD windows around each SNP in each 2Mbp-windows. In these two panels, green, blue, red and black lines represent variation for GBS, 600K, 50K and combined technologies, respectively. Vertical dotted gray lines indicate limits of centromeric regions. Vertical lines between the two panels indicate the position of QTLs for flowering time (DTA), grain yield (GY) and Plant Height (PHT). Green, blue, red vertical lines indicate QTLs detected only by GBS, 600K and 50K technologies, respectively. Grey lines indicated QTLs detected by at least two technologies. Only QTL including a marker associated with $-\log_{10}$(pval) above 6 were shown.

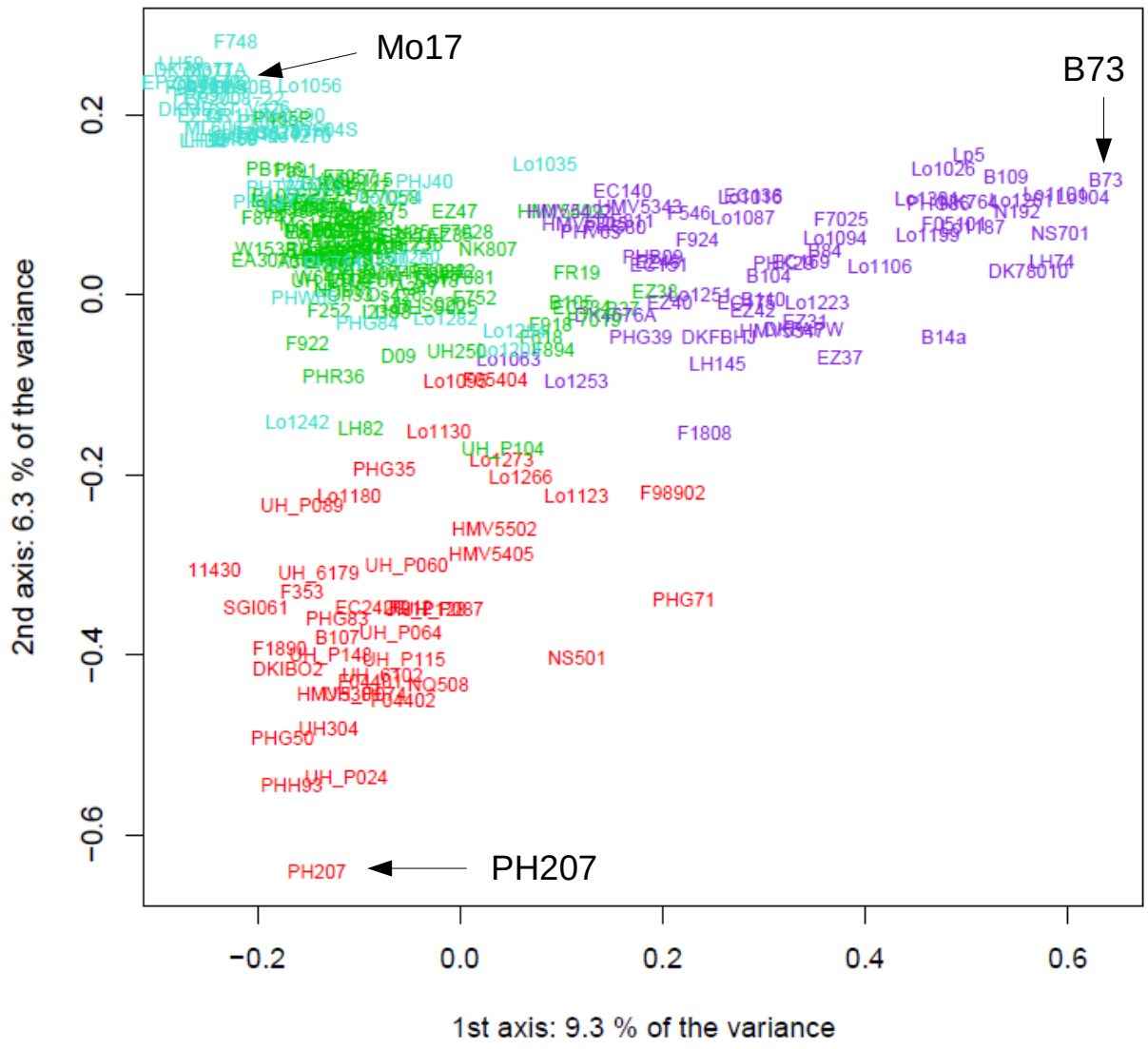
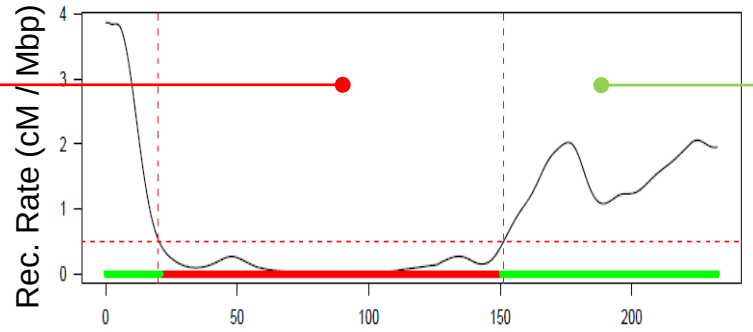


Figure 2: Principal coordinate analyses (PCoA) of the DROPS panel. The PCoA were based on the covariance matrix *K_Freq* estimated from the 50K Illumina array. The genetic groups identified by ADMIXTURE (N_Q = 4) are colored. Three key founders are indicated (Iodent: PH207 in red, Stiff Stalk: B73 in blueviolet, Lancaster: Mo17 in turquoise).

< 0.5cM / Mbp

> 0.5cM / Mbp

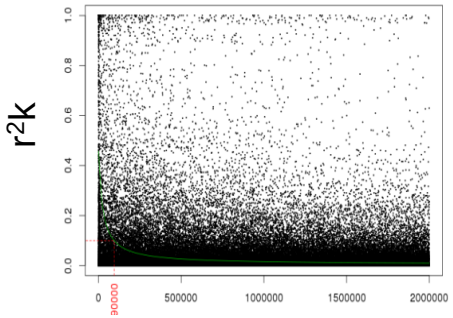


Position on chromosome 3

Low recombinogenic regions

High recombinogenic regions

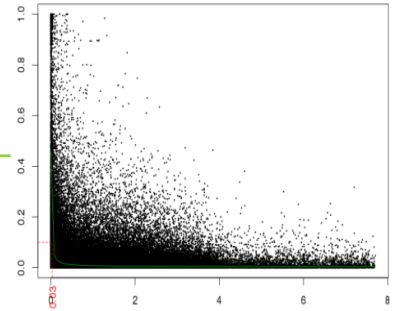
Projection of SNP on genetic map



Physical distance (bp)

$E(r^2k) = 0.1$ for d_p=846 kbp

$E(r^2k) = 0.1$ for d_g=0.03 cM



Genetic distance (cM)

SNP

d_p d_p

Physical interval

SNP

d_g d_g

Genetic interval

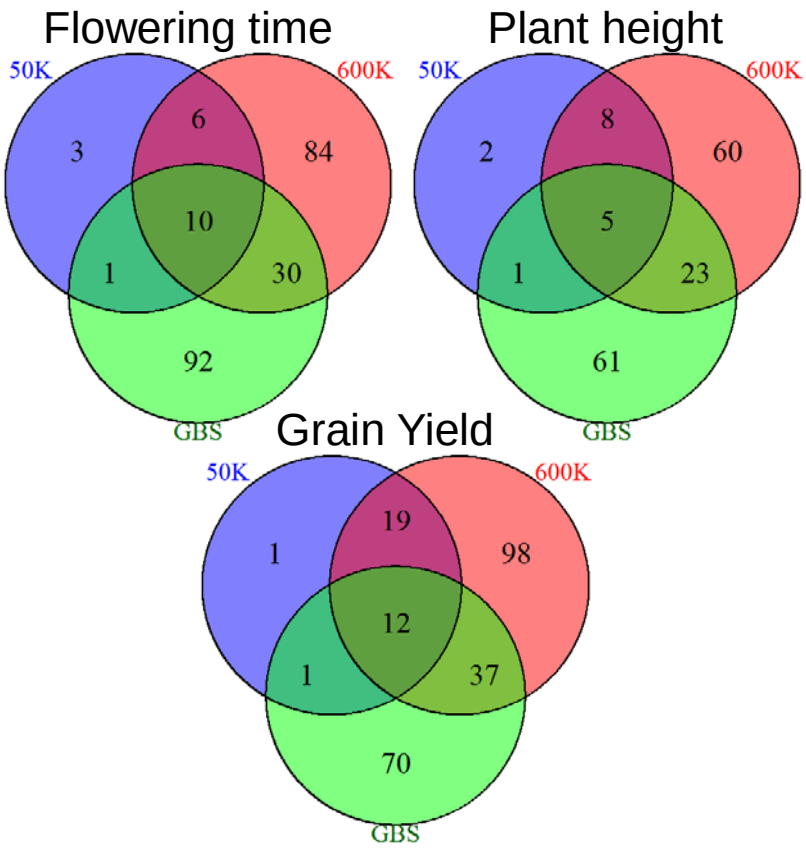
Projection of genetic interval on physical map

SNP

d_p d_p

Physical interval for each SNP based on LD extent

Figure 3: Linkage disequilibrium based approach to delineate a physical window around each SNP, exemplified with chromosome 3. Linkage disequilibrium (LD) windows were defined for each SNP based on physical LD extent in low recombinogenic regions (left part) and based on genetic LD extent in high recombinogenic regions (right part). These LD windows were used (i) to group significant SNPs into QTLs when they overlapped, (ii) to estimate genome coverage region covered by LD windows around SNPs, and (iii) identify putative genes underlying QTLs involved in trait variations.



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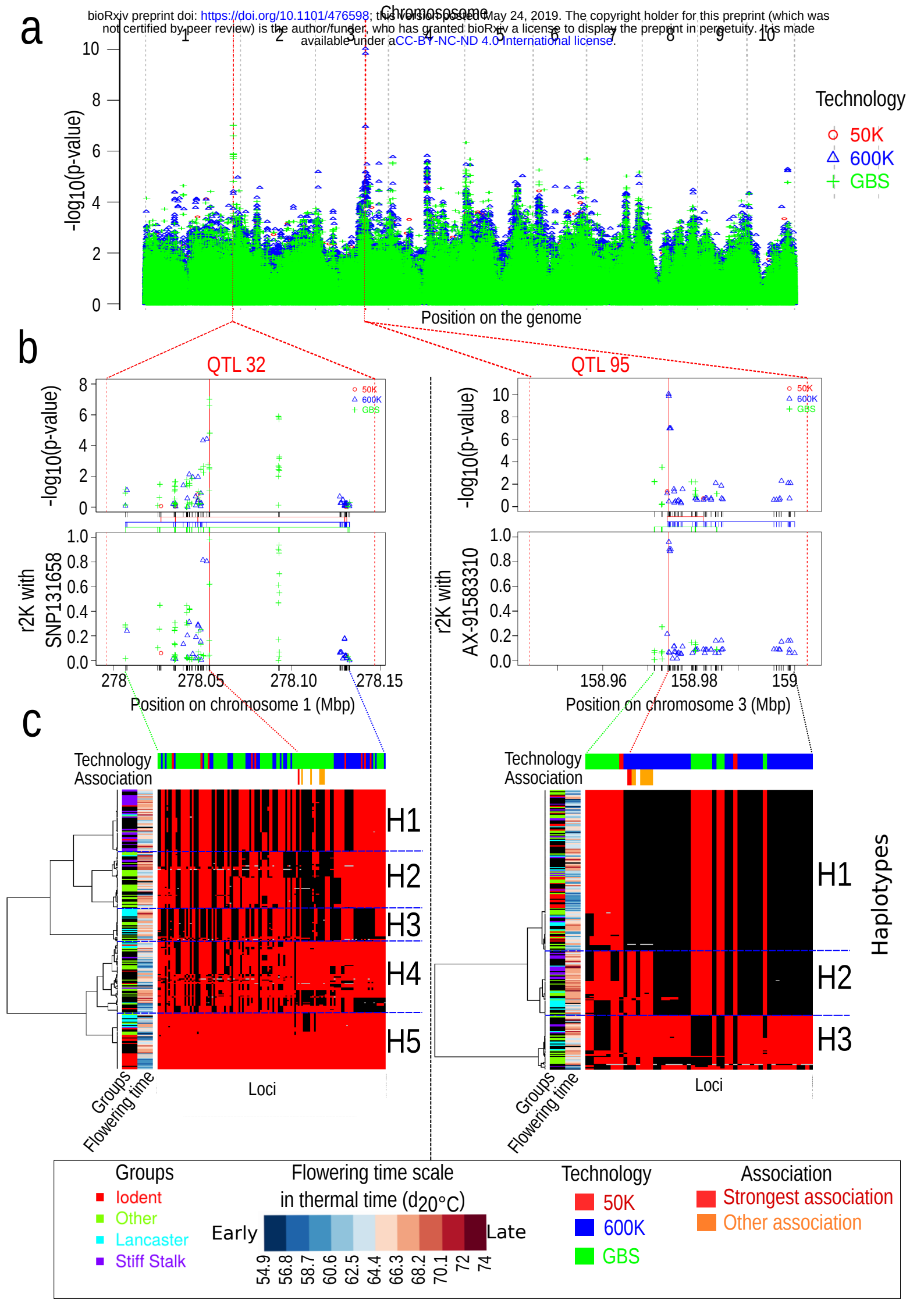

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