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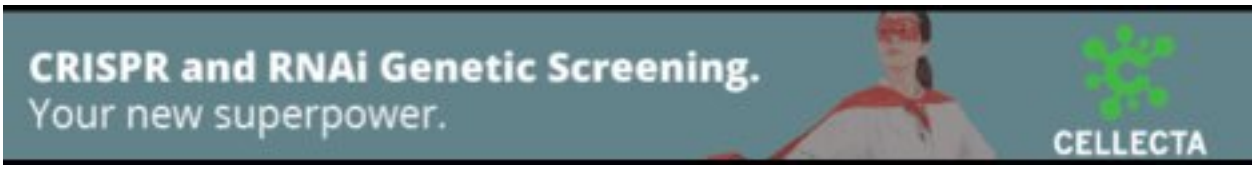

Comprehensive Variation Discovery and Recovery of Missing Sequence in the Pig Genome using Multiple *De Novo* Assemblies

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Abstract

Uncovering genetic variation through resequencing is limited by the fact that only sequences with similarity to the reference genome are examined. Reference genomes are often incomplete and cannot represent the full range of genetic diversity as a result of geographical divergence and independent demographic events. To more comprehensively characterize genetic variation of pigs (*Sus scrofa*), we generated *de novo* assemblies of nine geographically and phenotypically representative pigs from Eurasia. By comparing them to the reference pig assembly, we uncovered a substantial number of novel SNPs, structural variations, as well as 137.02 Mb sequences harboring 1,737 protein coding genes that were absent in the reference assembly, revealing variants left by selection. Our results illustrate the power of whole-genome *de novo* sequencing relative to resequencing, and provide valuable genetic resources that enable effective use of pigs in both agricultural production and biomedical research.

Introduction

Sus scrofa (i.e., pig or swine) is of enormous agricultural importance and also an attractive model for biomedical research and applications. There are over 730 distinct pig breeds worldwide, of which two thirds reside in Europe and China (Chen et al., 2007), whose diverse phenotypes are shaped by the combined effects of local adaptation and artificial selection (Ai et al., 2015). Efforts have been made to characterize the genetic variation that underlies this

phenotypic diversity using resequencing data and the genome of the European domestic Duroc pig as a reference (Choi et al., 2015; Groenen et al., 2012; Moon et al., 2015; Rubin et al., 2012). Nonetheless, resequencing is limiting in terms of capturing genetic variation and assessing gaps and misassigned regions of the reference genome (Weisenfeld et al., 2014). In contrast, multiple *de novo* assemblies of pig genomes from different regions and breeds promise a more accurate and comprehensive understanding of genetic variation within this species (Besenbacher et al., 2015; Chaisson et al., 2015b). Among populations of plants (coccolithophores (Read et al., 2013), *Arabidopsis thaliana* (Gan et al., 2011), soybean (Li et al., 2014) and rice (Zhang et al., 2014)), animals (mosquitoes (Neafsey et al., 2015) and macaques (Yan et al., 2011)) and even modern humans (Li et al., 2010a), surprisingly large amount of variation has been uncovered by *de novo* assemblies.

To advance the characterization of the genetic diversity of pigs, we generated *de novo* assemblies of nine geographically and phenotypically representative individuals from Eurasia. Combining this resource with genome assembly of the Tibetan wild boar (Li et al., 2013), we carried out in-depth comparisons between ten *de novo* assemblies and the reference genome. We uncovered a substantial number of single nucleotide polymorphisms (SNPs) and structural variations, as well as hundreds of millions of base pairs that are not present in the reference genome, including thousands of protein-coding genes that are either missing or fragmented in the reference genome, which contain potentially important genetic information pertaining to porcine evolution.

Results

De novo genome assemblies of nine pig breeds

We sequenced the genomes of nine female individuals from nine diverse breeds (five originated in Europe and four originated in China) to an average of ~100-fold coverage (~229.5 gigabase (Gb)) using Illumina sequencing technology and a whole-genome shotgun strategy (**Supplemental Fig. S1** and **Table S1**). The genomes were independently assembled using SOAPdenovo (Li et al., 2010b) (**Supplemental Methods**), which yielded contig N50 sizes of 28.99 to 42.66 kilobases (kb), scaffold N50 sizes of 1.26 to 2.45 megabases (Mb) and a total of 2.45 to 2.49 Gb of ungapped sequences that exhibited genomic features similar to those of the reference assembly (Groenen et al., 2012) (**Supplemental Figs. S2-S7** and **Tables S2-S6**). We also improved the available genome assembly of the Tibetan wild boar (Li et al., 2013) by increasing the contig N50 size from 20.69 kb to 22.54 kb and the ungapped genome assembly size from 2.43 Gb to 2.44 Gb (**Supplemental Table S4**).

Discovery and characterization of SNPs

We identified 8.86-15.95 million (M) SNPs in individual pig genomes using an assembly-versus-assembly approach (**Supplemental Methods**). These SNPs were consistent with more than 98% SNPs identified from the Illumina's porcine 60K Genotyping BeadChip (v.2) (**Supplemental Table S7**), and covered most SNPs identified by resequencing as implemented in SAMtools (Li et al., 2009) (98.78%) and GATK tool (McKenna et al., 2010) (97.65%) and

3.12-5.40 M SNPs (33.46-35.25%) in divergent regions that failed to be cataloged by these algorithms, where unassembled short reads are difficult to be mapped (**Fig. 1** and **Supplemental Figs. S8,9**).

Extensive inter-continental genomic divergence was reflected by the significantly larger amount of variation of Chinese pigs when compared to the reference Duroc genome of European origin (15.14 to 15.95 M SNPs; the Ts / Tv ratio: 2.13 to 2.15) than that between European pigs and Duroc (8.86 to 10.14 M SNPs; the Ts / Tv ratio: 1.95 to 1.99) ($P < 10^{-16}$, Mann-Whitney *U* test) (**Figs. 2A,B** and **Supplemental Fig. S10**), attributable to considerable divergence time between European and Asian lineages (at least 1 million years) and their independent domestication in multiple locations across Eurasia in the past ~10,000 years (Frantz et al., 2013; Groenen et al., 2012; Larson et al., 2005)**.**

We also observed higher genomic diversity of Chinese pigs than European pigs, reflected by the former's higher heterozygous SNP ratio (2.17×10^{-3} to 2.69×10^{-3} vs. 0.94×10^{-3} to 1.63×10^{-3}) and lower homozygosity (382 regions of homozygosity (ROHs) with a total size of 107.5 Mb vs. 907 ROHs with totaling 289.9 Mb per assembly) ($P < 10^{-16}$, Mann-Whitney *U* test) (**Fig. 2B** and **Supplemental Figs. S11,12**). Principal component analysis (PCA) and identity score (IS) analysis of pairwise breed genomes also recapitulated these findings (**Fig. 2C** and **Supplemental Fig. S13A**). This may be a reflection of the fact that European origin breeds have undergone intense selection in inbred commercial lines for economical traits, while Chinese breeds experienced weaker selection in scattered, individual farms, and exhibited

relatively weak linkage disequilibrium (LD) (White, 2011) (**Supplemental Fig. S13B**). Another possible explanation is that European wild boars (ancestors of European domestic pigs) may have suffered more pronounced population bottlenecks during the last glacial maximum (~20,000 years ago) compared to their Asian counterparts (Bosse et al., 2012; Groenen et al., 2012).

We pooled the SNPs of ten breeds into a non-redundant set of 33.60 M sites that account for ~81.25 % of the estimated repertoire of SNPs in the pig (**Supplemental Fig. S14**, **Table S8** and **Supplemental Methods**), of which 6.34 M (18.87% of 33.60 M) SNPs were considered to be novel based on their absence in the pig dbSNP (Build 143) entries (**Supplemental Fig. S15**). Compared with synonymous SNPs (122.44 k), missense SNPs (83.39 k) exhibited greater diversity among breeds (77.44% of the estimated repertoire compared to 80.61%), accounted for a larger proportion of breed-specific (and thus rare) SNPs (32.42% compared to 30.18%) and had a higher ratio of homozygous to heterozygous SNPs (0.37 compared to 0.32) (**Supplemental Figs. S14,16**), which may be associated with breed-specific adaptation.

Maps of structural variations

We detected 161.45-279.98 k insertions (15.99-23.07 Mb in length) and 137.89-269.55 k deletions (3.61-5.63 Mb in length) in individual genomes against the reference genome (**Fig. 2B**, **Supplemental Table S9** and **Supplemental Methods**). More than 80% of the insertions and deletions (indels) were 1-10 bp in length, and there was also a relatively high number of indels ~300 bp in length, due to the enrichment of indels of $tRNA^{Glu}$-derived short interspersed element ($SINE/tRNA^{Glu}$) (**Supplemental Fig. S17**) (Ai et al.,

2015). Repetitive elements (38.05% of the genome) comprised ~52.73% of indels, which are an important source of structural variation in pig genome. Moreover, the SINE/tRNA^{Glu} (290.47 Mb and containing 18.09% of indels) showed higher incidence of indels than the predominant long interspersed elements (LINE/L1) (636.50 Mb and containing 15.48% of indels) (**Supplemental Fig. S18**).

The indels appeared to be regulated by selection: most indels were located in intergenic regions (72.20% to 74.14%), the indel ratio was lower in the coding sequences than in introns (**Supplemental Fig. S19**) and more conserved genes showed fewer structural variations (**Supplemental Fig. S20**). We observed an enrichment of short indels (1-15 bp in length) in coding sequences (414 of 1,582, or 26.17%) that were multiples of 3 bp, which is expected to preserve the reading frame, and identified 1,152 frameshift mutations in 947 genes (**Supplemental Fig. S21** and **Table S10**), which mainly represented the cellular functions of 'binding of nucleoside, ATP, and cation' and 'neuron development' (**Supplemental Table S11**). As the SNPs, distribution of indels across the genome also reflected a deep phylogenetic split between European and Chinese pigs and higher genetic variability of Chinese pigs than European pigs (Bosse et al., 2012; Groenen et al., 2012) (**Supplemental Fig. S22**).

Signatures of diversifying selection in pig breeds

To uncover genetic variation underlying phenotypic diversity of pigs, we identified breed-specific signatures left by diversifying selection using a relative homozygous SNP density (RSD) algorithm (Atanur et al., 2013)

(**Supplemental Methods**). We identified 493 separate genomic regions of 20 to 150 kb (a total of 20.10 Mb and containing 308 genes) to be under selection (FDR < 0.05) (**Fig. 3A** and **Supplemental Table S12**). These putative selected regions also exhibited significantly strong LD and lower negative Tajima's *D* values ($P < 10^{-16}$, Mann-Whitney *U* test) (**Supplemental Fig. S23**), and distinct phylogenetic relationships compared to genomic background (**Supplemental Fig. S24**).

Most homozygous SNPs (88.60%) in the selected regions were unique to a particular breed (**Fig. 3A**), exhibiting lower degree of haplotype sharing with other breeds than pairwise between other breeds (**Fig. 3B** and **Supplemental Fig. S25**). These private SNPs were highly concentrated in a small number of discrete genomic regions (0.79% of the genome), and may be associated with phenotypes described by standard breed criteria (Wang et al., 2011): typically, 9 (out of 49, or 18.37%; $P = 0.004$, χ^2 test) and 6 (out of 59, or 10.17%; $P = 0.491$, χ^2 test) genes within or in the vicinity of the selected regions in the fatty Rongchang and Jinhua pigs were orthologous to well-established mammalian fat deposition genes (Kunej et al., 2013) (**Fig. 3B** and **Supplemental Fig. S25A**), including factors involved in the regulation of feed intake and energy homeostasis (*CEP120*, *GABRA2*, *NPPA*, *NPY1R* and *NYP5R*), lipid metabolism (*ABCC4*, *ANGPT2*, *LRPAP1* and *PRKAG2*) and indicators of obesity-induced hypertension, inflammatory signaling and insulin resistance (*ADD1*, *HSPD1*, *MMP2*, *PIK3R4, RAE1* and *TBCA*) (**Supplemental Table S13**). In contrast to highly inbred European pigs that have undergone selection for lean growth (high protein and low fat content; lean meat percentage of carcass ranging between 63-73%) as a response to demands for reduced

calorie intake in modern society, Chinese pigs have been selected for extreme fatness all along (typical lean meat percentage is under 45%) (**Supplemental Fig. S1**), driven by demand for energy-rich food in developing countries until ~10 years ago (Wang et al., 2011).

We also identified 16 (31.37%; $P = 8.21 \times 10^{-11}$, χ^2 test) out of 51 genes with strong selective sweep signals in the Tibetan wild boar (**Supplemental Fig. S25B** and **Table S13**) that were likely driven by the harsh and hypoxic environment of the Tibetan plateau and might have a role in the formation of characteristic phenotypes, such as an insulating layer formed by hard skin and long, dense hair, and larger lungs and hearts (Li et al., 2013).

Identifying missing sequences of the reference pig genome

There are considerable unidentified regions (289.24 Mb of 2.81 Gb, or 10.29%) in the reference pig assembly (Sscrofa10.2) (Groenen et al., 2012), of which 266.15 Mb (91.92%) is composed of 5,317 gaps of at least 50 kb long (**Supplemental Fig. S26**). To recover such missing genetic information, we retrieved ~9.17 G 'orphan reads' for which neither end mapped to the reference genome (**Supplemental Fig. S27**), and re-localized them to their respective assemblies of origin. Consequently, we identified 83.8 k sequences of ≥ 500 bp (137.02 Mb in length) that were missing in the reference genome (**Table 1** and **Supplemental Table S14**). Only a small portion of missing sequences was considered to be insertions (~0.91Mb) or copy number gains (~4.16%) (**Supplemental Tables S14,15** and **Supplemental Methods**). Compared with whole assemblies, these missing sequences exhibited similar heterozygous SNP ratio (2.67×10^{-3} vs. 2.56×10^{-3}; $P = 0.623$, Mann-Whitney

U test), but significantly higher GC content (43.07% vs. 41.41%; $P < 10^{-16}$, Mann-Whitney *U* test) and repeat ratio (47.57% vs. 38.38%; $P < 10^{-16}$, Mann-Whitney *U* test) (**Supplemental Fig. S28**).

Most sequences missing in the reference genome were common between different assemblies, as most orphan reads (95.04%) could crossly align to missing sequences of other assemblies with coverage (97.10% with depth ≥ 4 per base) comparable to that against their respective assemblies (mapping ratio = 95.83% and coverage = 99.51%) (**Supplemental Fig. S29**). Pairwise similarity of the orphan reads and the missing sequences between ten breeds revealed a clear distinction between European and Asian pigs, as well as relatively high genetic variability in Chinese pigs than in European pigs (Bosse et al., 2012; Groenen et al., 2012) (**Supplemental Fig. S30**), suggesting that these common sequences, which were absent in the reference genome, may be important sources of pig diversity and contain biologically meaningful information.

We were also able to fill in 71.37% (3,795 of 5,317) of the gaps in the reference genome by missing sequences of at least one breed **(Supplemental Fig. S27, Tables S14,16** and **Supplemental Methods**). These filled missing sequences were highly collinear across ten breeds and exhibited similar distribution over the reference assembly gaps (average of pairwise Person's $r = 0.89$, $P < 10^{-16}$) (**Supplemental Fig. S31**). Typical examples are shown in **Supplemental Fig. S32**.

Recovery of missing genes

Of the average 20,782 protein-coding genes (87.13% were supported by

evidence of transcription) predicted in each of the ten assemblies (**Supplemental Figs. S33-S35**, **Table S17** and **Supplemental Methods**), we found an average of 1,096 (5.27%) genes to be embedded or almost completely contained (> 50% to > 90% overlap of gene length, respectively) in the missing sequences of the reference assembly (**Table 1** and **Supplemental Table S18**), which we referred to as 'missing genes' (Genovese et al., 2013; Kidd et al., 2010).

To check whether these predicted missing genes are likely to be functional, we compared their conservation level across 19 mammalian genomes and found that they generally exhibited similar identity (81.55% vs. 83.60%) and coverage (96.32% vs. 97.37%) as annotated genes (**Supplemental Fig. S36**). Coding sequences of missing genes were enriched at high cross-species (human, cow and sheep) identity level (> 90%), also consistent with the sequence identity distribution of well-annotated coding sequences of the reference genome (**Supplemental Fig. S37**). We then retrieved ~0.59 G orphan reads against the reference genome from each of 96 pair-end RNA-seq libraries (7 to 10 libraries for each breed) and mapped them onto the missing genes in their respective assemblies (**Supplemental Figs. S38A,B**). Consequently, an average of 91.51% (1,003 of 1,096) missing genes in each assembly showed $\log_2$-transformed FPKM expression values (denoted as fragments per kb of transcript per Mb orphan reads) greater than 0.3 in at least one library (**Supplemental Fig. S38C**), suggesting that a considerable number of missing genes are functional and biologically important.

To determine the collinear relationships of missing genes among ten breeds, we separately aligned the protein sequences of nine assemblies to the

assembly of the Large White breed, which had the longest scaffold N50 size (2.45 Mb). Using MCScanX toolkit (Wang et al., 2012), we found that 10,313 of 10,959 (94.10%) missing genes in all of ten assemblies belonged to 1,091 inter-assembly collinear gene models, of which 871 (79.84%) models were present in all ten assemblies (**Table 1** and **Supplemental Tables S18,19**). There were a total of 646 missing genes (14 to 95 per assembly) assembled in only a single breed, which could be found in other assemblies when orphan reads from short-insert (180 and 500 bp) libraries were used for mapping (coverage 94.05% at least 1 × depth), suggesting that the absence of these singleton genes from other assemblies is likely an artifact of fragmentation or misassignment in short reads assembly (Alkan et al., 2011; Chaisson et al., 2015b) (**Supplemental Fig. S39**).

Together with the longest gene model of 1,091 inter-assembly collinear genes and 646 singleton genes, we obtained 1,737 missing gene models (**Table 1**). Aligning these missing genes to RefSeq proteins of pig, human, cow and mouse yielded 1,731 (99.65%) hits at least in one species (**Supplemental Table S19**), of which 359 (20.66%) missing genes could not be aligned to any known RefSeq proteins of pig, indicating that these genes have not been characterized in pig. Among hits that matched functionally classified proteins, the most abundant were members of olfactory receptors (65 hits, $P = 1.60 \times 10^{-12}$, χ^2 test), G-protein coupled receptors (104 hits, $P = 9.81 \times 10^{-6}$, χ^2 test) and those involved in neurological system processes (112 hits, $P = 4.26 \times 10^{-6}$, χ^2 test) (**Supplemental Table S20**), which are known to be rapidly evolving between species (Mainland et al., 2014). We also recovered genes corresponding to economically important traits that are valuable for future

functional analyses and improvements of pig as an important livestock species, such as genes related to pork production (74 of 1,515 fat deposition genes (Kunej et al., 2013), or 4.88%) and disease resistance (76 of 1,517 genes annotated with the GO: 0002376; immune system process, or 5.01%) (**Supplemental Table S19**). Typical examples are shown in **Supplemental Fig. S40**.

Selection in missing genes

To reveal variants left by selection, we measured pairwise the extent of population differentiation of the coding SNPs in the missing genes between Chinese wild boars (32.57 k coding SNPs) and seven Chinese domestic populations (23.02 k coding SNPs per population) using the *FDIST* approach as implemented in Arlequin (Excoffier and Lischer, 2010) (**Supplemental Figs. S41,42** and **Table S21**). A total of 605 non-redundant coding SNPs embedded in 328 missing genes were found to be under directional selection in seven Chinese domestic populations (FDR < 0.05, *FDIST* test) (**Supplemental Fig. S43** and **Table S22**), which also exhibited significantly lower *q* values in a Bayesian test (Foll and Gaggiotti, 2008) and F_{ST} values in a 'model-free' global F_{ST} test when compared to other unselected loci ($P < 10^{-16}$, Mann-Whitney *U* test) (**Supplemental Fig. S44**). The missing genes under selection in seven Chinese domestic populations were commonly enriched for biological processes related to 'binding of actin, calcium ion, and cytoskeletal protein' (**Supplemental Fig. S45A**). Intriguingly, 71 genes harboring 110 selected coding SNPs of domestic Erhualian pigs (one of the most prolific pig breeds known) (Wang et al., 2011) belonged predominantly to fertility-related

categories, such as 'sexual reproduction' (7 genes: *ADAM20*, *AKT1*, *GMCL1*, *MICALCL*, *NOTCH1*, *SPIN4* and *SPTBN4*; P = 0.001) and 'placenta development' (3 genes: *AKT1*, *RXRA* and *VWF*; P = 0.012), which may underlie the breed's markedly larger litters (~3 to 5 more piglets per litter) (**Supplemental Fig. 45B**).

The expression of missing genes under selection also showed remarkably higher tissue specificity, reflected by the lower Shannon entropy (H) values (a measure of the specificity of gene expression across tissues) (Schug et al., 2005) than unselected missing genes (1.98 vs. 2.37 per gene; $P < 10^{-16}$, Mann-Whitney U test) (**Supplemental Fig. S46**). As opposed to constitutive genes that are ubiquitously expressed and essential for basic cellular functions, tissue-specific genes are usually associated with the development of generally desirable traits, such as disease resistance, muscle growth, fat deposition and reproduction, and thus are more likely prone to be shaped by selection.

None of the selected coding SNPs was a nonsense mutation (resulting in premature stop codons in transcribed mRNAs) (**Supplemental Table S22**), supporting the idea that gene inactivation did not play a prominent role during pig domestication and consistent with the results from screens in chickens (Rubin et al., 2010), rabbits (Carneiro et al., 2014) and pigs (based on reference genome) (Rubin et al., 2012). Compared to synonymous substitutions, missense substitutions showed significantly lower genetic differentiation (global F_{st}, 0.05 compared to 0.10 per locus; $P < 10^{-16}$, Mann-Whitney U test) between Chinese wild boars and domestic pigs (**Supplemental Fig. S47**). Nonetheless, there were still 127 genes harboring

selected missense mutations, which were over-represented in the highly variable olfactory receptor family (12 genes; P = 0.02, χ^2 test)[34] (**Supplemental Table S22**). Of these, three missense mutations embedded in two genes related to the development of obesity were of interest: the closely linked Asn566-His (T1,696-G) and Ser578-Cys (G1,733-C) substitutions (D' = 1, r^2 = 0.975) found in *ALPK3* (alpha kinase 3) (**Fig. 4** and **Supplemental Fig. S48**)**,** and a Thr18-Ile (C53-T) substitution in *PKD1L2* (polycystin 1 like 2 (gene/pseudogene)) (**Supplemental Fig. S49**). These three missense mutations exhibited significant selection signals (FDR < 0.05, *FDIST* test) between Chinese wild boars and one of seven domestic populations (Min and Erhualian, respectively), but were nearly fixed in the more genetically homogenous European/North American domestic pigs, possibly as a result of stronger selective pressure in western societies, although larger sample sizes, inter-continental genetic discrepancy of pig genomes and functional analyses are required to validate the non-neutrality of these genes.

ALPK3 plays a role in cardiomyocyte differentiation; knockout of this gene in mice was associated with marked hypertrophic and dilated forms of cardiomyopathy (Van Sligtenhorst et al., 2012). *ALPK3* shows the strongest evidence of positive selection in the polar bear, which has a lipid-rich diet throughout life (Liu et al., 2014). Selection of *ALPK3* in domestic pigs suggests that potential protection against the chronically deleterious effects of a 'diabetogenic' environment (high calorie, atherogenic diet and little physical exercise) on the cardiovascular system may be favorable (Gerstein and Waltman, 2006; Koopmans and Schuurman, 2015). *PKD1L2* is primarily associated with fatty acid synthase in the skeletal muscle fiber; its

368 overexpression in mice provokes myofiber atrophy and suppressed
369 lipogenesis (Mackenzie et al., 2009). The triglycerides accumulated between
370 or within myofibers represent a large energy source (contributes up to 20% of
371 total energy turnover during physical exercise in human) (Roepstorff et al.,
372 2005). Selection of *PKD1L2* is likely related to the relatively weak athletic
373 performance of domestic pigs compared to wild boars due to limited active
374 space in pig farms.

375

376 ## Discussion

377 We describe an assembly-versus-assembly approach that relies on multiple
378 independently assembled genomes for improving the power of variant
379 detection, as opposed to the currently dominant resequencing approach. This
380 catalogue of variants, including SNPs, indels, and common and rare variants is
381 a valuable resource for further investigation of the genetic makeup of porcine
382 phenotypic diversity and adaptive evolution. We show that high-quality *de*
383 *novo* assembly of individual genomes followed by comparison with the
384 reference sequence is necessary for identifying novel genetic variation across
385 geographical ranges and different evolutionary histories. Such experimental
386 design is increasingly affordable with the advances in sequencing technology
387 (Zook and Salit, 2015), especially long-read sequencing (Chaisson et al.,
388 2015a) and single-molecule mapping (Koren et al., 2012) technologies.

389 Interpretation of the consequences of genetic variation has typically relied
390 on reference sequences, relative to which genes and variants are annotated
391 and examined. However, we recovered hundreds of millions of base pairs that

were not present in the pig reference genome, including thousands of protein-coding genes that are either missing or fragmented in the reference genome, which harbor abundant variants associated with economic traits that are likely subjected to artificial selection. These newly recovered genes can now be incorporated into genotyping platforms and expression microarrays to facilitate their functional characterization. Recovered sequences missing from the reference genome could also be the source of genetic signals that have been ascertained by linkage, association and copy number variation studies but not yet been mapped to causal mutations.

Methods

De Novo sequencing and assembly of pig genomes

We sequenced the genomes of nine geographically and phenotypically representative pig breeds using Illumina sequencing technology and a whole-genome shotgun strategy (**Fig. 2A** and **Supplemental Fig. S1**). Short-insert (180 and 500 bp) and long-insert (2, 5, 6 and 10 kb) DNA libraries were paired-end sequenced on the Illumina HiSeq 2500 platform (**Supplemental Fig. S2** and **Table S1**). We independently assembled nine genomes using SOAPdenovo (Li et al., 2010b), which is a *de Bruijn* graph algorithm–based *de novo* genome assembler (**Supplemental Methods**). We performed repeat annotation for ten breed assemblies and the reference genome using the same pipeline (**Supplemental Figs. S5,6** and **Supplemental Methods**).

SNP and indel calling using an assembly-versus-assembly method

We took advantage of an assembly-versus-assembly approach to identify candidate variations and further filtered out spurious variations by aligning short sequencing reads (**Supplemental Methods**). In brief, we first extracted candidate SNPs and small- and intermediate-scale indels (1-50 kb) in ten assemblies by pairwise gapped alignment of the ten assemblies and the reference genome assembly (Sscrofa10.2) using the LASTZ program. Then, the paired-end short-insert reads (180 and 500 bp) were separately aligned to the ten assembled genomes and the reference genome using the BWA software (v.0.7.12) (Li and Durbin, 2009). We filtered spurious SNPs and determined the heterozygous or homozygous mutations (depth ≥ 10) using SAMtools (v.1.3) (Li et al., 2009). With regard to indels, we eliminated spurious indel calls based on the calculation of read coverage for each indel locus with different criteria for indels ≤ 50 bp or > 50 bp (Li et al., 2011).

Identification of selected regions using RSD algorithm

To identify signatures of diversifying selection of pig breeds, relative homozygous SNP density (RSD) in nonoverlapping 10kb windows across the reference genome were calculated for each individual using a previously reported methodology (Atanur et al., 2013) (**Supplemental Methods**).

RNA-seq and data processing

The 92 strand-specific RNA libraries (7 to 10 tissue libraries for each of ten individuals, which were used for *de novo* genome assemblies) were sequenced on the Illumina HiSeq 2500 platform (**Supplemental Methods**). High-quality reads were mapped to their respective *de novo* assemblies (**Supplemental Figs. S35,46**) or the reference genome (**Supplemental Fig.**

S38) using TopHat (v.2.1.0) (Trapnell et al., 2009). Cufflinks (v.2.2.1) (Trapnell et al., 2012) was used to quantify gene expression.

Discovery of missing sequences and missing genes

We retrieved 'orphan reads' from paired-end DNA libraries with insert sizes of 180 and 500 bp for each of ten breeds, where both ends of the read cannot be uniquely mapped to the reference genome (**Supplemental Figs. S27**). We re-localized these orphan reads to their respective assemblies. Sequences (≥ 500 bp in length) absent from the public reference genome assembly that were mapped by at least four orphan reads per base were considered 'missing sequences' (Kidd et al., 2010).

To identify genes embedded in the missing sequences, we conducted annotation of protein coding genes in the ten assemblies separately, using a combination of evidences from reference assembly-guided approach, the *ab initio*- and the homology-based methods, as well as RNA-seq data (**Supplemental Figs. S33,34**, **Table S17** and **Supplemental Methods**). We considered genes that showed > 50% overlap of the gene length with missing sequences to be either missing or fragmented in the reference genome, and referred to them as 'missing genes'.

Determination of inter-assembly collinear genes

The protein sequences of genes in the nine assemblies were separately queried against the protein sequences of the Large White assembly, which had the longest scaffold N50 size (~2.45 Mb) using BLASTp with an E-value cutoff of 10^{-5} and restricting the output to a maximum of five hits per gene to serve as input for the MCScanX algorithm (Wang et al., 2012), which was used to detect

and classify high-confidence collinear blocks of coding genes (**Supplemental Tables S18,19**).

Detecting coding SNPs in missing genes under selection

To test whether the recovered genes missing in the reference genome were under selection, we retrieved ~365.55 Gb orphan reads against the reference genome from 117 publicly available pig genomes (in total 6.03 trillion base resequencing data) (Ai et al., 2015; Choi et al., 2015; Moon et al., 2015) and aligned them to the intact scaffolds harboring missing genes across the ten breed assemblies (~636.38 Mb per assembly) (**Supplemental Fig. S41**). Of these, 6 wild boars and 41 domestic pigs belonging to 7 populations in China that have high-coverage depth (27.29 × of the reference genome, 14.43 × of the missing genes embedded scaffolds by 3.91 Gb orphan reads per individual) (Ai et al., 2015) were used to test for differentiation and possibly selection (**Supplemental Fig. S42**). The remaining 70 individuals (including 10 Korean wild boars and 60 European/ North American domestic pigs) with intermediate-coverage (15.87 × of the reference genome, 6.99 × of missing gene embedded scaffolds by 2.60 Gb orphan reads per individual) (Choi et al., 2015; Moon et al., 2015) were used to investigate the patterns of selected loci (**Supplemental Fig. S41**).

We measured pairwise the extent of population differentiation of the coding SNPs in the missing genes between Chinese wild boars and seven Chinese domestic populations using the *FDIST* approach as implemented in Arlequin (v.3.5.2.2) (Excoffier and Lischer, 2010) (**Supplemental Fig. S43, Supplemental Table S21** and **Supplemental Methods**). We also measured

pairwise global F_{ST} values (**Supplemental Fig. S44A**) and performed a Bayesian test using the program BayeScan (v.2.1) (Foll and Gaggiotti, 2008) (**Supplemental Fig. S44B**) for every gene to detect highly differentiated SNPs between populations.

Data access

The nine pigs and Tibetan wild boar BioProjects are accessible at NCBI BioProject (http://www.ncbi.nlm.nih.gov/bioproject) under accession numbers PRJNA309108 and PRJNA186497, respectively. The assembled whole genome sequences have been deposited at NCBI GenBank (http://www.ncbi.nlm.nih.gov/genbank) under accession numbers LUXQ00000000.1 (Meishan), LUXR00000000.1 (Rongchang), LUXS00000000.1 (Hampshire), LUXT00000000.1 (Landrace), LUXU00000000.1 (Piétrain), LUXV00000000.1 (Bamei), LUXW00000000.1 (Berkshire), LUXX00000000.1 (Large White), LUXY00000000.1 (Jinhua) and AORO00000000.2 (Tibetan wild boar, v.2). The unassembled sequencing reads of nine pigs and Tibetan wild boar have been deposited in NCBI Sequence Read Archive (SRA: http://www.ncbi.nlm.nih.gov/sra) under accession numbers SRP068560 and SRA065461, respectively. All RNA-seq reads and the genotyping data of the Illumina's porcine 60K Genotyping BeadChip (v.2) have been deposited in NCBI Gene Expression Omnibus (GEO: https://www.ncbi.nlm.nih.gov/geo/) under accession numbers GSE77776 and GSE83910, respectively.

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Figure legends

Figure 1. Comparison of SNP calling between assembly-versus-assembly method and resequencing approaches based on read mapping.

Venn diagram with colors corresponding to bar chart shows the sharing of identified SNPs among assembly-versus-assembly method and two resequencing algorithms as implemented in SAMtools and GATK. An average of 4.25 M SNPs per breed were specifically identified by assembly-versus-assembly method (marked as yellow), while only 0.24 k SNPs per breed were categorized by resequencing approaches (marked as red). A significant fraction of the detected SNPs by the SAMtools (8.11 M per individual) and GATK (7.77 M per individual) was coincident (7.41 M; or 91.24% of SAMtools and 95.34% of GATK) (**Supplemental Fig. S8**).

Figure 2. Genomic variation between Chinese and European pigs.

(*A*) Geographic locations of the original pig breeds. The Duroc (donor of the reference genome; it is denoted by star) and Hampshire pigs were developed mainly in North America but originated in Europe.

(*B*) Neighbor-joining phylogenetic tree, number of SNPs, transition / transversion ratio (Ts / Tv), heterozygous SNP ratio, patterns of regions of homozygosity (ROHs), and length and number of indels in the ten breeds (left to right). Violin plots of the heterozygous SNP ratio and Ts / Tv ratio were generated using non-overlapping 1 Mb windows (the medians are shown). For ROH, the circled area indicates the total length of ROHs in each breed.

(*C*) Pairwise genomic similarity of Chinese and European pigs by identical score (IS) values within each 10 kb window across the genome (n = 259,511).

Figure 3. Identification of breed-specific selective sweeps.

(*A*) Number of homozygous SNPs in breed-specific selected regions. Of 74.21 k homozygous SNPs in 20.10 Mb selected regions, 65.75 k (88.60%) were unique to a particular breed, which highly concentrated in a small fraction (0.79%) of the genome and likely contributed to diversifying selection.

(*B*) Selective sweep regions identified in the Rongchang pig. Top panels, top half: genes residing within or in the vicinity (± 5 kb) of the selected regions are presented for each

chromosome and ordered according to their locations. Top panels, lower half: degree of haplotype sharing of selected regions in pairwise comparisons among the ten breeds. Homozygous SNP frequencies in individual breeds were used to calculate identity scores in 10 kb windows. Boxes (left) indicate pairwise comparison presented on that row (E, European pigs; C, Chinese pigs) according to the color assigned to each pig breed (right). Heat-map colors indicate identity scores. Second panels: Percentage stacked column showing RSD values in the Rongchang-specific selected regions across ten breeds sequenced. Rongchang showed predominantly higher RSD values than other breeds, indicating that only this breed shows SNPs against the reference genome in this region. Third panels: RSD in 10 kb windows for Rongchang plotted along chromosomes. Black lines indicate selected regions (FDR < 0.05). Nine selected genes orthologous to the mammalian fat deposition genes were marked in red.

Figure 4. Details of assembled *ALPK3* gene and selected variants.

(*A*) Structure of assembled *ALPK3*. Top panels: the inter-assembly collinear genes (colored rectangles) among ten assemblies are linked by gray lines, and the genes not present in all ten assemblies are marked in black. *ALPK3* is denoted by a circle. Different scaffolds are shown as alternating white and gray backgrounds. Second panels: comparison of structure of *ALPK3* among the ten assemblies. Boxes and lines indicate exons and introns, respectively.

(*B*) Coverage and depth for the longest gene model of *ALPK3* (Gene ID: RCGENE17759) by crossly mapping reads from paired-end DNA libraries (insert sizes of 180 and 500 bp) of the ten assemblies. The higher coverage depth (≥ 30 ×) suggests slightly different structures of *ALPK3* which is attributable to limitations of short read assembly; as such the longest gene model is considered more reliable and used for subsequent analyses.

(*C*) Two selected missense mutations (T1,696-G and G1,733-C) in *ALPK3* between Chinese wild boars (n = 6) and domestic Min pigs (n = 6). Top panels: F_{ST} and Heterozygosity / (1- F_{ST}), FDR (Arlequin) and q values (BayeScan) are plotted for 45 coding SNPs (18 missenses and 27 synonymous mutations). Second panels: LD pattern of 45 SNPs in 101 domestic pigs from China (n = 41), North America (n = 12) and Europe (n = 48). Squares shaded in pink or red indicate significant LD between SNP pairs (bright red indicates pairwise D' = 1), white squares indicate no evidence of significant LD, and blue squares indicate pairwise D' = 1 without statistical significance. The adjacent T1,696-G and G1,733-C are closely linked (D' = 1, r^2 = 0.975, LOD = 41.6).

Table 1. Summary of missing sequences and genes of the reference genome (Sscrofa10.2)

Assemblies	Missing sequence		Missing genes			
	Number	Length (Mb)	Number	Assigned by inter-assembly collinearity		
				Singleton	Assembled in 2-9 breeds	Assembled in all 10 breeds
Hampshire	82,824	136.33	1,105	67	167	
Berkshire	82,958	136.40	1,092	63	158	
Landrace	82,741	135.86	1,093	65	157	
Piétrain	82,472	135.75	1,096	60	165	
Large White	82,987	136.04	1,105	14	220	
Bamei	84,336	137.49	1,104	83	150	871
Jinhua	84,031	137.34	1,090	65	154	
Meishan	85,197	138.65	1,116	95	150	
Rongchang	84,062	137.88	1,064	49	144	
Tibetan wild boar	86,592	138.42	1,094	85	138	

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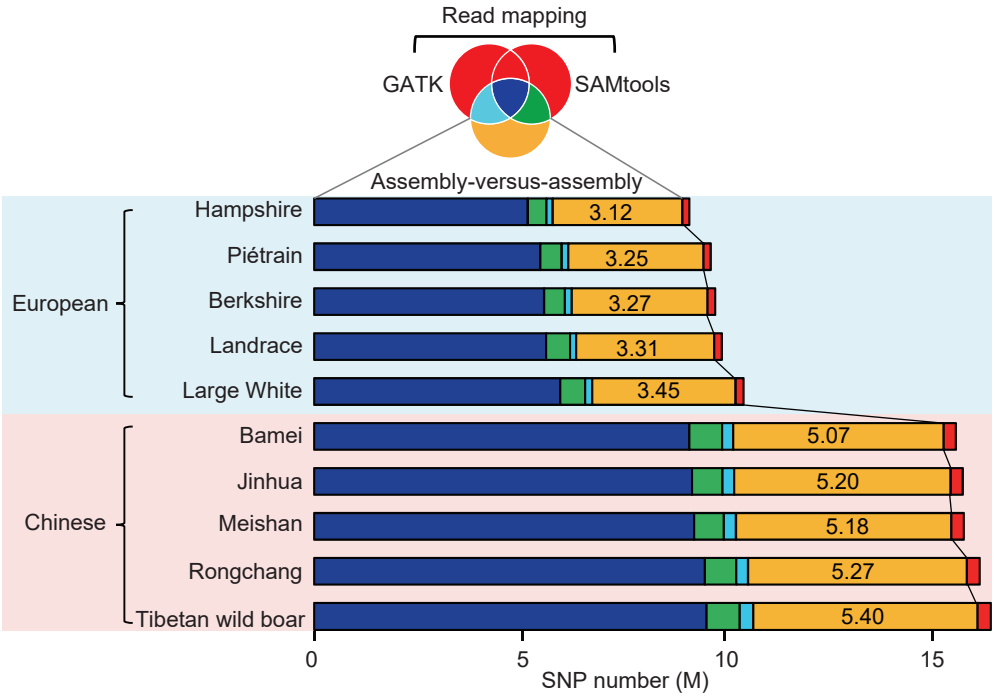
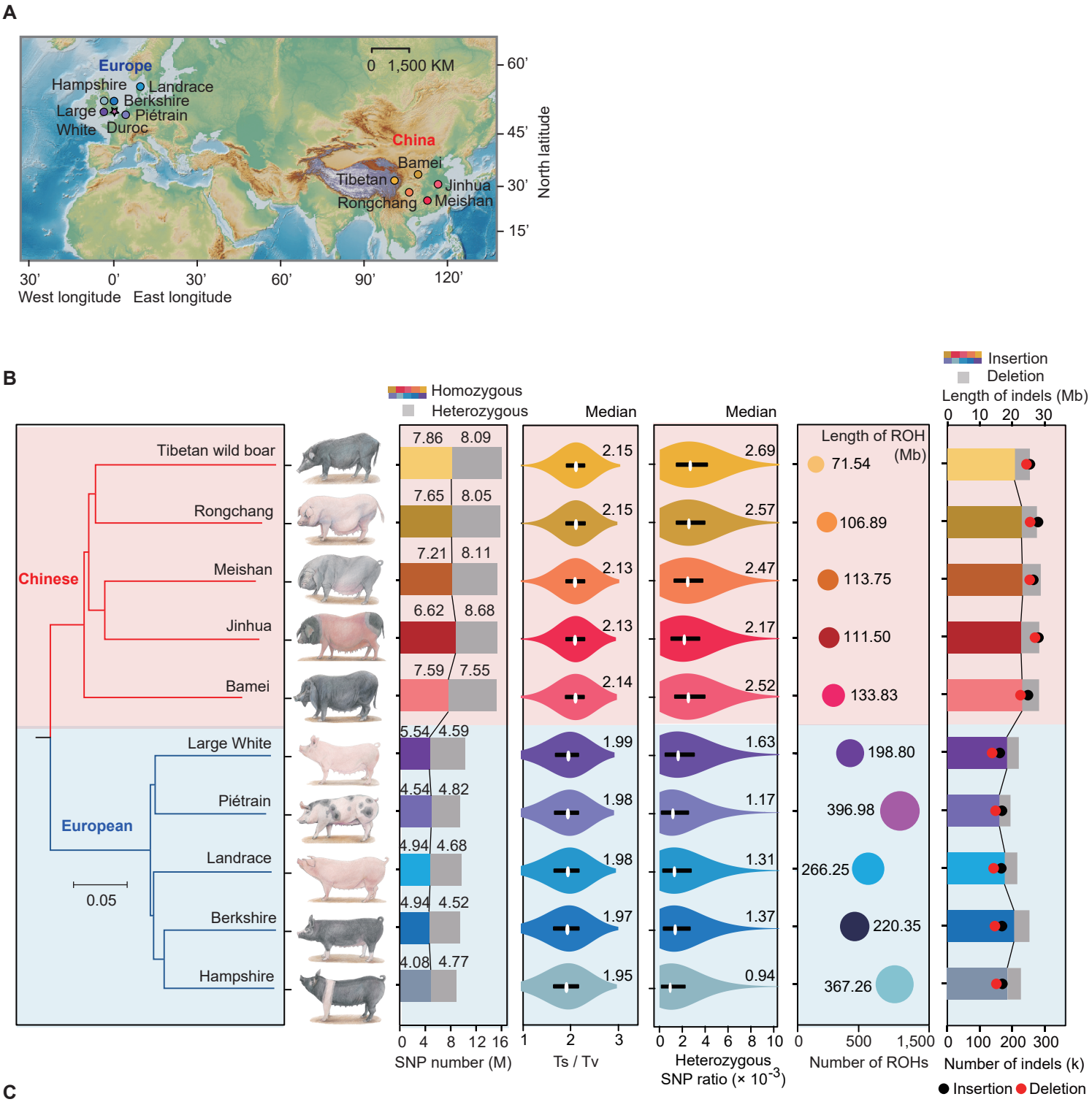
Fig. 1





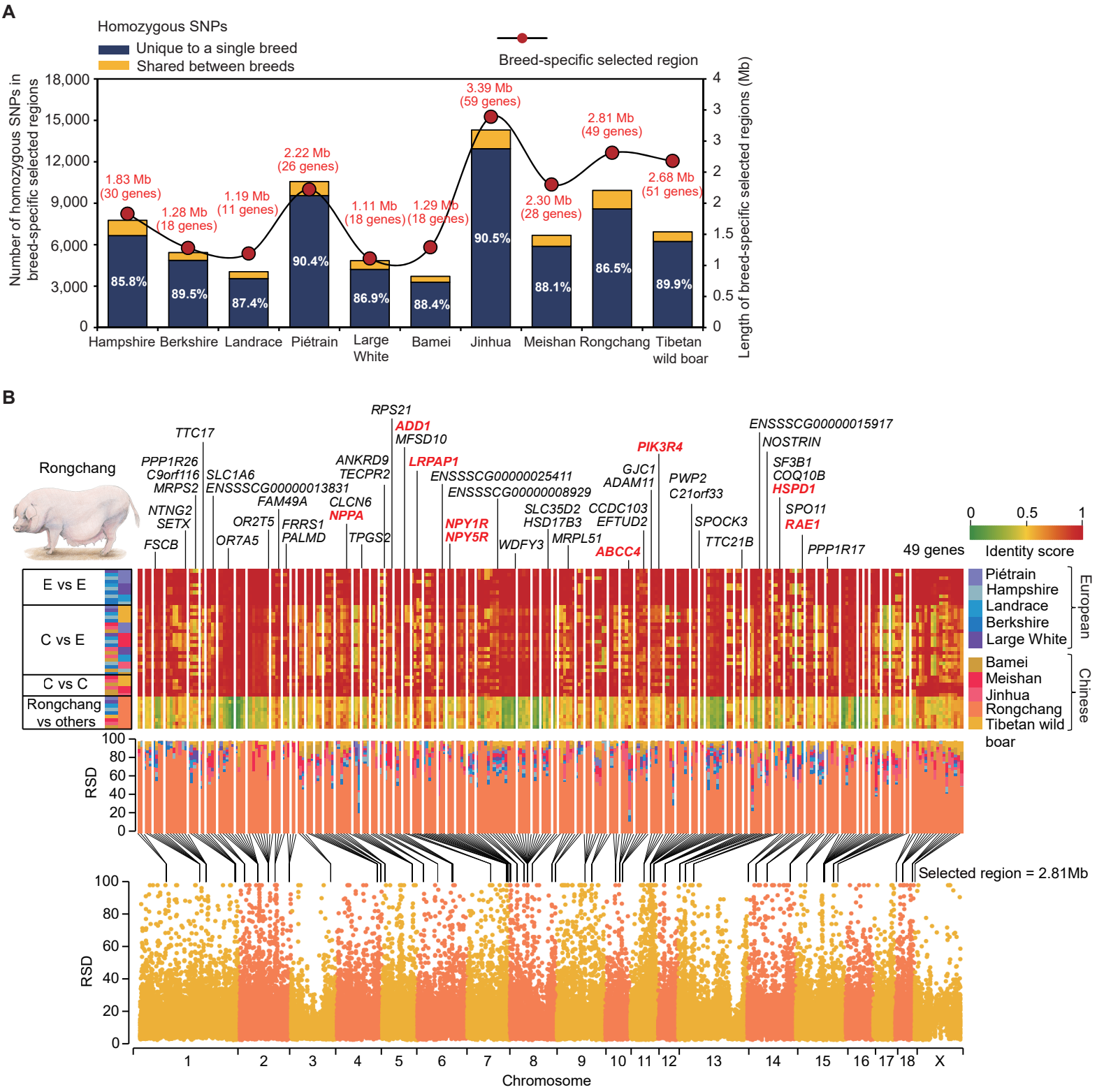


Fig. 4