









Supplemental Fig. S49. Details of assembled PKD1L2 and selected variants. (a) Assembled PKD1L2. Top panels: the inter-assembly collinear genes (colorized rectangles) are linked by gray lines, while genes not presented in all of ten assemblies are marked in black. PKD1L2 is denoted as a circle. Different scaffolds are distinguished by alternant white and gray backgrounds. Second panels: the higher coverage and depth for the longest gene model of the PKD1L2 by crossly mapping with reads from paired-end DNA libraries (insert sizes of 180 and 500 bp) of ten assemblies. This result suggests the slightly distinct structures of collinear missing genes among ten assemblies (See b), which are attributable to the limitations of short reads assembly. Therefore, the longest gene model was considered more reliable and used for subsequent analyses. (b) A selected missense mutation (C53-T) in *PKD1L2* between Chinese wild boars (n = 6) and domestic Erhualian pigs (n = 5). Top panels: $F_{\rm ST}$ and Heterozygosity / (1 - $F_{\rm ST}$), FDR (Arlequin)³⁹ and q values (BayeScan)⁴⁰ are plotted for 102 coding mutations (28 missenses and 74 synonymous mutations). Second panels: structures of PKD1L2 among ten assemblies and in various species. Boxes and lines indicate exons and introns, respectively. (c) An amino acid variant (Thr18-lle) in PKD1L2. Left panels: The phylogenetic tree of the orthologous protein sequences of pig and 10 vertebrates (Ensemble release 83) were derived from multiple alignment as implemented in the Clustal Omega tool³⁵, which were in accordance with the evolutionary distance with pig lineage. Right panels: Multispecies alignment of proteins for a selected missense mutation. The protein coordinate is based on longest gene model of inter-assembly collinear genes among ten assemblies (HAGENE20165). Dots indicate identities to the Chinese wild boar sequence and dashes indicate missing data. (d) Selected missense mutation (C53-T) is highly frequent in domestic pigs.