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## **Supplementary Material Legends**

Fig. S1 Analyses of gut microbial NR gene numbers and diversity of gut microbial NR genes in pigs. A The number of gut microbial NR genes quantified in 112 pigs from seven breeds. B Shannon indexes of gut microbial NR genes quantified in 112 pigs from seven breeds. C PCoA of gut microbial NR genes quantified in 112 pigs from seven breeds. Data are presented as mean  $\pm$  SEM (n = 8) and were evaluated by the Kruskal-Wallis test. \*\*p < 0.01.

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Fig. S2 Comparative analyses of gut microbial KEGG pathways in pigs. 9 Comparative analyses of the relative abundances of microbial genes involved in KEGG 10 pathways, including biosynthesis of unsaturated fatty acids (A), secondary bile acid 11 12 biosynthesis (B), tryptophan metabolism (C), phenylalanine metabolism (D), propanoate metabolism (E), butanoate metabolism (F), purine metabolism (G), 13 pyrimidine metabolism (H), methane metabolism (I), and nitrogen metabolism (J), 14 benzoate degradation (K), and dioxin degradation (L), respectively. Data are presented 15 16 as the mean  $\pm$  SEM (n = 8). Significantly differentially abundant KEGG pathways were identified using the Kruskal-Wallis test. \*\*p < 0.01, \*p < 0.05; ns, not significant. 17

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Fig. S3 Correlation analysis of gut microbial species and antimicrobial resistance genes. Heatmap for the Spearman's correlation analysis of microbial species (relative abundance >0.5 %) and antimicrobial resistance genes. \*\*p < 0.01, \*p < 0.05.

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Fig. S4 Correlation analysis of gut microbial species and antimicrobial resistance types. Heatmap for the Spearman's correlation analysis of microbial species (relative abundance >0.5 %) and antimicrobial resistance types. \*p < 0.01, \*p < 0.05.

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Fig. S5 Rarefaction curves and heatmap analysis of gut bacterial community as

28 revealed using 16S rRNA gene sequencing. A Rarefaction curves for Chao index. B

29 Heatmap analysis of bacterial genera.

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Fig. S6 Gut microbial taxonomic compositions in seven pig breeds. A-D Gut microbial taxonomic compositions, including kingdom (A), class (B), order (C), and family (D), as revealed using metagenomics. E-G Gut bacterial taxonomic compositions, including class level (E), order level (F), and family level (G), as revealed using 16S rRNA gene sequencing.

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Fig. S7 Heatmap analysis of relative abundances of three core-predominant microbial species. A Heatmap analysis of relative abundances of *P. succinatutens*, *P. copri*, and *O. valericigenes* in the feces of the mice (FMT group) revealed by metagenomics. B Heatmap analysis of relative abundances of *O. valericigenes* in the feces of the mice (SPF group) revealed using 16S rRNA gene sequencing.

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Fig. S8 Effects of core-predominant microbes on composition of fecal metabolites 43 in germ-free mice. A-C OPLS-DA for fecal metabolites in mice from groups (Ctrl and 44 45 PS) (A), groups (Ctrl and PC) (B), groups (Ctrl and OV) (C) (Ctrl, control; PS, P. succinatutens; PC, P. copri; OV, O. valericigenes). D-F Volcano plot analysis of fecal 46 metabolites in mice from groups (Ctrl and PS) (D), groups (Ctrl and PC) (E), and groups 47 (Ctrl and OV) (F). G Venn diagram analysis of the fecal metabolites that are upregulated 48 49 by the treatments with the three core gut microbes, respectively. H Venn diagram analysis of the fecal metabolites that are downregulated by the treatments with the three 50 core gut microbes, respectively. I-K The KEGG enrichment analyses of the fecal 51 metabolites altered by the treatments with P. succinatutens (I), P. copri (J), and O. 52 53 valericigenes (K), respectively. L UpSet plot comparing the KEGG pathways enriched 54 with the fecal differentially metabolites induced by the treatments with *P. succinatutens*, P. copri, and O. valericigenes, respectively. The differentially abundant metabolites 55 were assessed based on VIP  $\geq 1$  from the OPLS-DA, absolute Log2 (fold change)  $\geq 1$ , 56 and p value < 0.05. 57

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59 Fig. S9 Analyses of complete genome sequences of core-predominant microbes. A

Circular representation of genome of P. succinatutens (PS, P. succinatutens). B KEGG 60 pathway classification of P. succinatutens at level 2. C Venn diagram analysis of the 61 KEGG pathways (level 3) annotated in *P. succinatutens* and the KEGG pathways (level 62 3) enriched by the fecal metabolites in GF mice that were orally administrated with P. 63 succinatutens. D Circular representation of genome of P. copri (PC, P. copri). E KEGG 64 pathway classification of P. copri at level 2. F Venn diagram analysis of the KEGG 65 pathways (level 3) annotated in *P. copri* and the KEGG pathways (level 3) enriched by 66 67 the fecal metabolites in GF mice that were orally administrated with P. copri. G Circular representation of genome of O. valericigenes (OV, O. valericigenes). H KEGG 68 pathway classification of O. valericigenes at level 2. I Venn diagram analysis of the 69 KEGG pathways (level 3) annotated in O. valericigenes and the KEGG pathways (level 70 71 3) enriched by the fecal metabolites in GF mice that were orally administrated with O. valericigenes. 72

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