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Table S1. Composition of the animal diet.

Ingredients	Chow diet	High-fat diet
	(D12450H)	(D12451)
Casein	18.96%	23.31%
L-Cystine	0.28%	0.35%
Corn Starch	42.86%	8.48%
Maltodextrin 10	7.11%	11.65%
Sucrose	16.38%	20.14%
Cellulose, BW200	0.47%	5.83%
Soybean oil	2.37%	2.91%
Lard	1.90%	20.68%
Mineral Mix	0.95%	1.17%

Vitamin Mix	0.95%	1.17%
Choline Bitartrate	0.19%	0.23%
DiCalcium Phosphate	1.23%	1.57%
Calcium Crabonate	0.52%	0.64%
Potassium Citrate,1 H ₂ 0	1.56%	1.92%
Total calories (kcal/g)	3.85	4.73

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Name	HFD (%)	Anti (%)
p_Bacteroidetes, g_Alistipes	0.0224 ± 0.0277	0**
p_Firmicutes, g_Erysipelatoclostridium	0.1012 ± 0.2211	0*
pFirmicutes, gLachnospiraceae_NK4A136_group	1.2911±1.3494	$0.0013 \pm 0.0032^{**}$
p_Firmicutes, g_Enterococcus	1.7343 ± 0.9913	$0.0018 \pm 0.0022^{**}$
p_Actinobacteria, g_Enterorhabdus	1.7484 ± 0.7628	0**
p_Firmicutes, g_Lactobacillus	$2.7300{\pm}1.5076$	$0.0189 \pm 0.0425^{**}$
p_Firmicutes, g_Kurthia	$7.8084{\pm}13.0511$	0**
p_Actinobacteria, g_Bifidobacterium	8.2843±6.4787	$0.0009 \pm 0.0022^{**}$
p_Firmicutes, g_Faecalibaculum	18.1254±4.6443	0.0004±0.0011
p_Bacteroidetes, g_unclassified_o_Bacteroidales	0	0.0150±0.0129**
p_Proteobacteria, g_Parasutterella	$0.0004{\pm}0.0011$	14.8707±6.2449**
p_Proteobacteria, g_Escherichia- Shigella	0.0026 ± 0.0024	8.6522±5.4739**
p_Proteobacteria, g_Citrobacter	0.0040 ± 0.0040	27.1351±6.8712**
p_Proteobacteria, g_Klebsiella	$0.0040 {\pm} 0.0072$	3.6478±2.7194**
p_Bacteroidetes, g_Parabacteroides	$0.0374 {\pm} 0.0538$	41.9350±14.2825**

Table S2. The relative abundance of gut microbiota in genus level between HFDand Anti groups.

Data were expressed as mean \pm SD, and statistical analysis by Wilcoxon rank sum test, *p < 0.05, **p < 0.01.

Supplementary Figure Captions

Figure S1. Effect of FTES on obese mice. (A-C) The weight of epididymis fat, mesenteric fat and retroperitoneal fat. (D-E) Concentrations of LDL-C, HDL-C in serum. (F) The level of blood glycemia in oral glucose tolerance test. Data were expressed as mean \pm SD, and statistically analyzed by One-way ANOVA, *p < 0.05, **p < 0.01, ***p < 0.001; ns, not significant.

Figure S2. Effect of FTES on antibiotic treated mice. (A-C) The weight of epididymis fat, mesenteric fat and retroperitoneal fat. (D-E) Concentrations of LDL-C, HDL-C in serum. (F) The level of blood glycemia in oral glucose tolerance test. Data were expressed as mean \pm SD, and statistically analyzed by Student's t test, *p < 0.05, **p < 0.01, ***p < 0.001; ns, not significant.

Figure S3. (A) Food intake, (B) water intake, (C) OTU rank-abundance curves, (D) Partial least squares-discriminate analysis (PLS-DA) on OUT level. Data were expressed as mean \pm SD, and statistically analyzed by One-way ANOVA. Different letters a, b represent significant difference, p < 0.05.

Figure S4. (A) Hierarchical cluster analysis by Spearson-Approx method. (B) Nonmetric multidimensional scaling (NMDS) on OUT level. (C) The heatmap of differential genera among Chow, HFD and FTES groups, which were analysised by the Wilcoxon rank sum test (p < 0.05).

Figure S5. Fecal transplantation improved lipid metabolism in high-fat diet fed mice. (A) The weight of liver, mesenteric fat and retroperitoneal fat. (B) Serum concentrations of TC, LDL-C, HDL-C. (C) Food and water intake. (D) Alpha diversity was presented as the index of Chao, Shannon. Data were expressed as mean \pm SD, and statistically analyzed by One-way ANOVA, *p < 0.05, **p < 0.01, ***p < 0.001; ns, not significant.

Figure S6. Heatmap showing significantly different metabolites between the HFD group and Chow group or FTES group based on Student's t test (p < 0.05).

Figure S7. Differential metabolites and enriched metabolic pathways after treated by FTES in negative ion model. (A) Heatmap showing significantly different metabolites between the HFD group and FTES group based on Student's t test. (B) Metabolome view maps of the significantly different metabolites-related metabolic pathways between the HFD group FTES group. Pathway impact for topology analysis and *p*value for enrichment analysis were performed using MetaboAnalyst 4.0. The size and color of each circle represent the pathway impact value and *p*-value, respectively.









Figure S3.



Figure S4.







Figure S6.





