

## Supplementary materials

**Table S1** Alpha diversity indexes inter-group difference table.

Item	Groups		SEM	<i>P</i>
	T1	T3		
observed_species	716.00	748.20	32.54	0.504
shannon	6.95	6.82	0.17	0.626
simpson	0.98	0.97	0.01	0.227
chao1	821.38	901.17	43.31	0.229
ACE	830.98	897.99	38.63	0.255
goods_coverage	99.52	99.42	0.04	0.143

T1: the broilers were fed a basal diet; T3: the broilers were fed a basal diet supplemented with AGE at 0.4%.

**Table S2** Differential metabolites identified among T1 and T3 groups from the data set of the feces samples

No.	Differential metabolites	Score <sup>1</sup>	RT (min)	Formula	<i>P</i> -Value <sup>2</sup>	Fold change <sup>2</sup>
1	p-cresol sulfate	0.48	0.402	C <sub>7</sub> H <sub>8</sub> O <sub>4</sub> S	0.018	0.22
2	cholesterol sulfate	0.66	0.405	C <sub>27</sub> H <sub>46</sub> O <sub>4</sub> S	0.049	0.53

1) The values were the degree of secondary spectrum (MS<sup>2</sup>) matching between the analytical compounds of base peak intensity chromatogram and an in-house MS<sup>2</sup> database (Biotree DB, Shanghai, China) in positive ion mode. The property of the value is between [0,1], and the larger the value, the higher the matching.

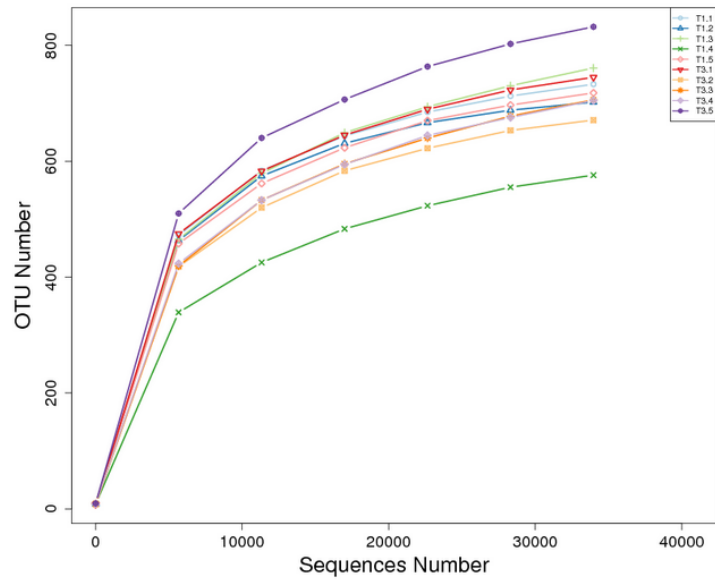
2) Upregulation ( $p < 0.05$ , FC > 1.10); Downregulation ( $p < 0.05$ , FC < 0.90).

**Table S3** MS data and identification of dihydromyricetin metabolites in broiler feces.

Name	RT (min)	Ion mode <sup>1</sup>	Metabolic pathway	Formula	Fragment ions
M1	1.004	[M-H]	Dehydration, Reduction	C <sub>15</sub> H <sub>12</sub> O <sub>7</sub>	125.02383,151.03963,177.01920
M2	3.174	[M-H]	Dehydration, Nitro Reduction, Glycine Conjugation	C <sub>17</sub> H <sub>15</sub> N O <sub>6</sub>	190.01439,166.04994,117.03430
M3	3.409	[M-H]	Nitro Reduction, Acetylation	C <sub>17</sub> H <sub>18</sub> O <sub>5</sub>	257.11786, 185.09554,93.03414

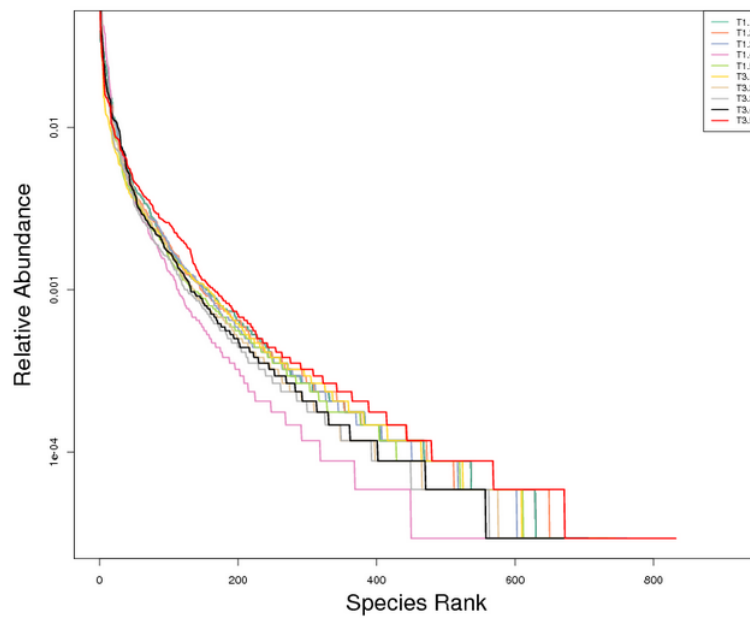
**Table S4** MS data and identification of myricetin metabolites in broiler feces.

Name	RT(min)	Ion mode <sup>1</sup>	Metabolic pathway	Formula	Fragment ions
M1	1.004	[M-H]	Nitro reduction oxidation	C <sub>15</sub> H <sub>12</sub> O <sub>7</sub>	125.02383,151.03963,178.99844
M4	3.636	[M-H]	Nitro reduction, methylation	C <sub>16</sub> H <sub>12</sub> O <sub>6</sub>	255.06613,135.00806,119.04892
M5	0.871	[M+H]	Nitro reduction, methylation	C <sub>16</sub> H <sub>16</sub> O <sub>4</sub>	229.08534,147.04385



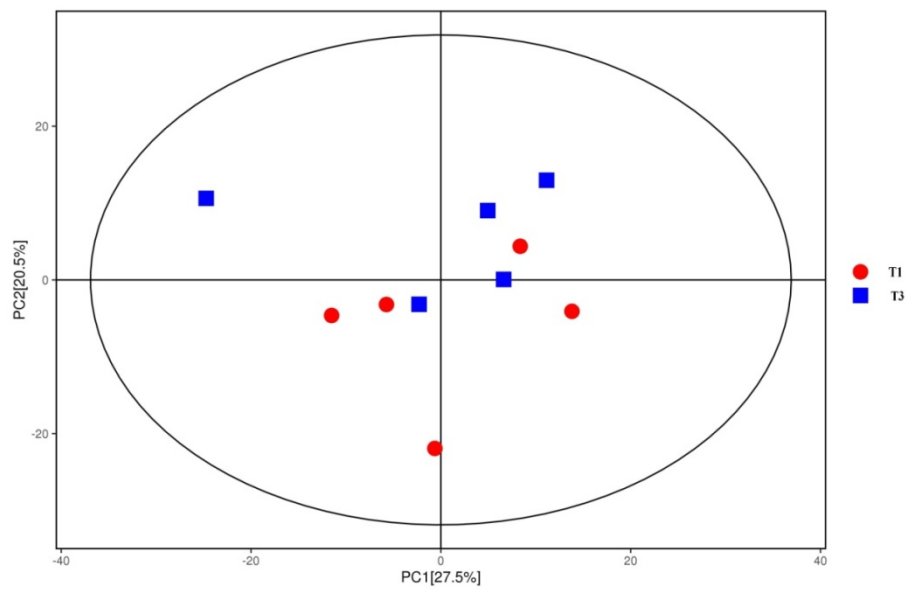
**Figure S1** Dilution curves

T1: the broilers were fed a basal diet; T3: the broilers were fed a basal diet supplemented with AGE at 0.4%.



**Figure S2** Rank abundance curves

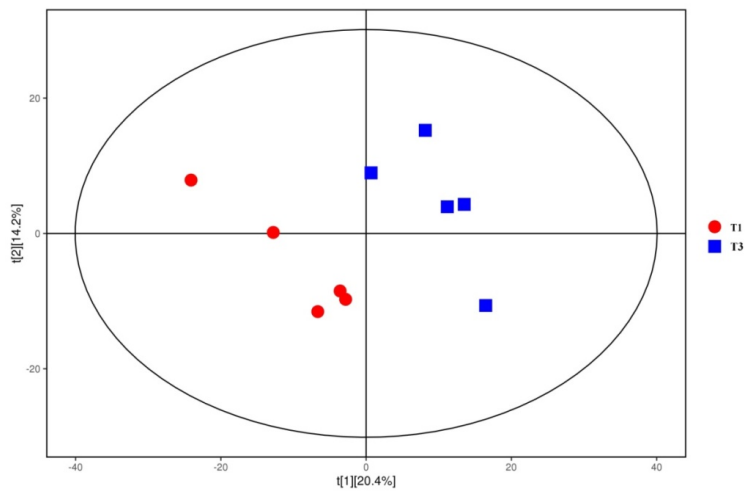
T1: the broilers were fed a basal diet; T3: the broilers were fed a basal diet supplemented with AGE at 0.4%.



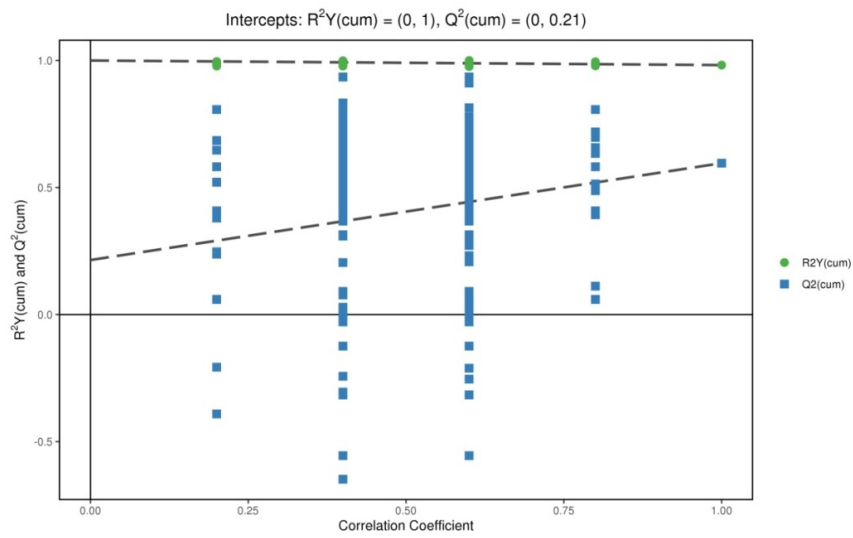
**Figure S3** PCA score plot.

T1: the broilers were fed a basal diet; T3: the broilers were fed a basal diet supplemented with AGE at 0.4%.

# A PLS-DA scores



# B PLS-DA permutation



**Figure S4** PLS-DA analysis.

A) PLS-DA scores: The abscissa represents the first principal component PC1, and the ordinate represents the second principal component PC2. T1: the broilers were fed a basal diet; T3: the broilers were fed a basal diet supplemented with AGE at 0.4%.

B) Permutation diagram:  $R^2$  stands for model verification, and the Y matrix of original classification and  $N$  times of different arrangement are linearly regressed with  $R^2Y$  and  $Q^2Y$ , and the intercept values of regression line and y-axis are  $R^2$  and  $Q^2$ , respectively; used to measure whether the model is over-fitted.