Microbiome- metabolomics deciphers the effects of Cistanche deserticola

polysaccharides on the aged constipation rats

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Supplementary files

Figure S1. Body weight of rats (**A**), and the time c/time (a + c) ratio of rats in the novel object recognition (NOR) test at 0 week (**B**). NC, the negative control group; MS, the aged constipated group; PC, the Mosapride group; CDPS, the *Cistanche deserticola* Polysaccharides group. *p<0.05, **p<0.01, significant differences as compared to the NC group. #p<0.05, ##p<0.01, significant differences as compared to the MS group (n=8).

Figure S2. Rarefaction curves **(A)**, the number of species at different levels **(B)** of the rats in the negative control group **(NC)**, the aged constipated rats **(MS)**, the Mosapride group **(PC)**, and the *Cistanche deserticola* Polysaccharides group **(CDPS)**.

Figure S3. Score plot of principal component analysis (PCA) (**A**), partial least-squarediscrimination analysis (PLS-DA) (**B**), orthogonal partial least squares discriminant analysis (OPLS- DA) (**C**) and the model permutation (**D**), and the corresponding S-Plot (**E**) of the aged constipated rats (**MS**) and the *Cistanche deserticola* Polysaccharides group (**CDPS**) (n=8). The PLS-DA model was validated by using 200 random permutations ($R^2 = 0.80$, $Q^2 = 0.66$).

Figure S4. Enrichment analysis of pathways involving in the pathology of the aged constipation (the negative control group (NC) versus the aged constipated rats (MS), and the effects of *Cistanche deserticola* Polysaccharides group (CDPS) (the MS group versus the CDPS group). Color and size represent p and impact, respectively. Pathways with p < 0.05 and impact > 0.1 represent a significant impact of this pathway.

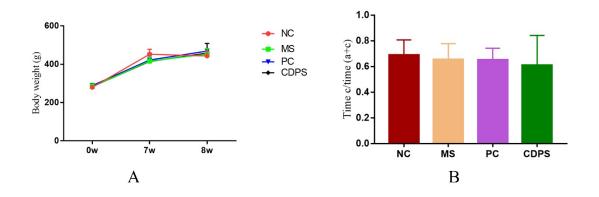


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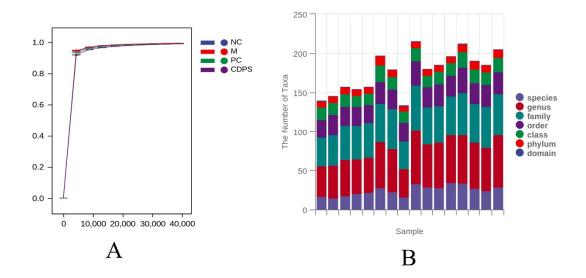


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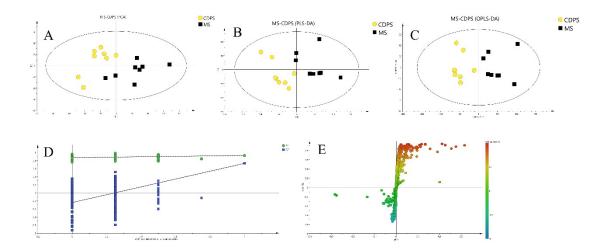


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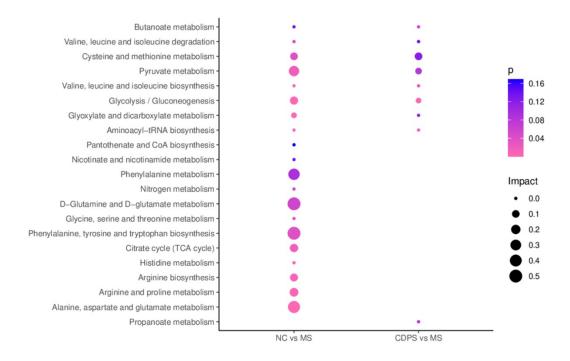


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