Gut microbiota and metabolites in estrus cycle and their changes in a menopausal transition rat model with typical neuroendocrine aging

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Methods:

SAMPLE COLLECTION:

Young (Y; 2-3-month-old) and middle-aged (MA; 9-10-month-old, retired breeders) female rats (Sprague Dawley, Charles River, Beijing) were fed with radiationsterilized lab rodent feed (number 1010086; Jiangsu Xietong Pharmaceutical Bioengineering Co., Ltd.) and water ad libitum. A 12-hour light/12-hour dark cycle (lights on at 8 a.m.) and the room temperature at 23°C was held. Estrous periodicity was detected by vaginal smear for at least consecutive two cycles (ten days). Only rats in possession of two regular estrous periods (four to five days) were included for the subsequent assays. On the basis of age and estrous stage, the rats were assigned to one of the following subgroups respectively: young, proestrus and diestrus (Y-P and Y-D); middle-aged, proestrus and diestrus (MA-P and MA-D). All fecal specimens were harvested and quickly chilled in liquid nitrogen and finally transferred to -80°C refrigerator for subsequent experiments.