Figure S1. Flow gating strategies for each cell population.
Figure S2. Effects of BBR against DSS-induced colitis after pre-treatment with ABX.
(A) Schematic diagram illustrates the experimental design. (B) DAI changes of each group. (C) Measurement of the length of colons harvested from mice in each group. (D) H\&E staining (Bar=200um above, Bar=50um below) sections and histological scores of colon tissue from mice in each group. Data are presented as mean $\pm S D$. ${ }^{*} P<$ 0.05 , significantly different as indicated.

Figure S3. BBR repaired the integrity of intestinal epithelial barrier in colitis mice.
(A) Paneth staining(Bar=200um above, Bar=50um below) sections of colon tissue from mice in each group. (B) The MLNs and liver FITC-dextran concentrations of mice in each group. (C) The colonic mRNA level of Muc2 in each group were measured by qPCR. (D) The mRNA levels of Occludin and Zo-1 in Caco2 cells from each group were measured by qPCR. Data are presented as mean $\pm S D . * P<0.05$, significantly different as indicated.

Figure $\mathbf{S 4}$. BBR regulates the expression of cytokines in the colon.
(A) The concentration of inflammatory cytokines in each group were measured by ELISA. Data are presented as mean $\pm \mathrm{SD} .{ }^{*} P<0.05$, significantly different as indicated.

Figure S5. BBR treatment significantly alter the proteomics of colon tissue.
(A) Volcano graph of the distribution of the different protein in DSS-BBR- and DSS + BBR- groups, the red dots represents up-regulated proteins and green dots represent down-regulated proteins. (B) Hierarchical clustering of proteins in DSS-BBR- and DSS+BBR- groups. (C) GO biological process analysisinDSS-BBRand DSS+BBR- groups. (D) KEGG pathways analysis in DSS-BBR- and DSS+BBRgroups.

Figure S1


## Figure S2

A

B

C

D



## Figure S3



## Figure S4



Figure S5
A

B

C

D


