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Supplementary Information

- 2 Microalgae Aqueous Extracts Exert Intestinal Protective Effects in
- 3 Caco-2 Cells and Dextran Sodium Sulphate-Induced Mouse Colitis
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11 Method

12 MTT Assay

- 13 To assay the cytotoxicity of MAEs in Caco-2 cells, wells were set up with each containing 1×10^4 cells in a 96-well
- 14 plate. Cells were cultured for 24 h and treated with 8 μ g/mL MAEs in complete medium for another 24 h. The
- 15 MTT solution (0.5 mg/mL in DMEM) was then added and the plate was incubated for further 4 h. Following
- 16 medium removing, 150 μ L DMSO was added to each well and the plate was agitated for 10 min on a plate shaker.
- 17 The absorbance was measured at 570 nm.

Table S1 The criteria for DAI

Score	Weight loss	Stool consistency	Stool consistency Rectal bleeding	
0	None	Normal stools	Negative	
1	1-5%	Soft stools	Negative	
2	6-10%	Soft stools	oft stools Positive	
3	11-15%	Very soft	Visible in stool	
4	> 15%	Watery stool	Gross bleeding	

$21 \quad \hbox{\bf Table S2} \ \hbox{Chemical Compositions of the Microalgae Powders}$

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	Protein	Lipid	Total carbohydrate	Ash
Chlorella pyrenoidosa	59.36±0.44 ^b	14.82±0.33 ^a	14.20±0.44 ^a	7.09±0.17 ^b
Spirulina platensis	67.25±0.52 ^a	9.10±0.15 ^b	11.56±0.44 ^b	5.97±0.20 ^c
Synechococcus sp. PCC 7002	65.79±0.20 ^a	8.21±0.35 ^b	9.30±0.28 ^c	13.23±0.21 ^a

²² Data were expressed in % of dry weight (means \pm standard deviations, n = 3). Different superscript

²³ letters (a-c) in the same column denote statistically significant differences (p < 0.05).

- Figure S1 Viabilities of Caco-2 cells following an incubation with 8 μ g/mL of MAEs for 24 h. Data were expressed as
- 26 means \pm standard deviations (n = 6).

