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1

Supplementary information

- 2 Title
- 3 Formation, structural characteristics and physiochemical properties of beeswax oleogel prepared with
- 4 tea polyphenols loaded gelator

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15 Supplementary data



Fig. S1 Polarized micrographs of TP gelator before (A) and after (B) freeze-drying. Microscopic
images were captured at 200× magnification.

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In order to analyze the structure of solid W/O emulsions during the preparation of TP gelator, 20 the microstructure of samples before and after freeze-drying was observed with PLM, as shown in 21 Fig. S1. Beeswax and TP aqueous solution were shear emulsified to form a W/O emulsion. The X-22 ray diffractogram signified the crystalline nature of beeswax, which was determined by its chemical 23 composition. When removed from high temperature, beeswax could be rapidly cooled because of the 24 long saturated fatty acid chains in the structure to ensure the size of the TP solution dispersed in it. 25 From Fig. S1(A), it could be clearly observed that the dispersed TP droplets were wrapped in the 26 network structure formed by beeswax crystallization. The beeswax crystals prevented the aggregation 27 of TP droplets and ensured the stability of the emulsion. And the TP gelator after the removal of water 28 by lyophilization from solid W/O emulsion was shown in Fig. S1(B). 29

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Fig. S2 Fourier-transform infrared spectra of TP (A), Soybean lecithin (B), Physical mixture of TP
and soybean lecithin (C) and TP-SL complex (D), respectively.

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TP-SL complex was prepared according to the method in Section 2.2, and beeswax was replaced 35 by deionized water. The characteristic absorption peaks of soybean lecithin appeared at 2925 and 36 2853 cm⁻¹ for C-H stretching of long fatty acid chain, 1738 cm⁻¹ for C=O stretching vibration of fatty 37 acid ester, 1232 and 1086 cm⁻¹ for P=O and P-O-C stretching ^{1, 2}. The infrared spectrum of physical 38 mixture was only the superposition of TP and soybean lecithin. It was similar to the spectra of soybean 39 lecithin without peak shift, while the characteristic absorption peak of TP was not obvious due to the 40 small amount of addition. The TP-SL complex spectra showed that the peak corresponding to the 41 functional group N-H (3374 cm⁻¹) was shifted to lower wavenumber and the intensity decreased. In 42

addition, the absorption peaks corresponding to the P=O (1227 cm⁻¹) and P-O-C (1066 cm⁻¹)
stretching was shifted to lower wavenumber by 5 cm⁻¹ and 20 cm⁻¹, respectively. The shifting of peaks
in TP-SL complex compared with soybean lecithin indicated that the presence of intermolecular
interactions between the phenolic hydroxyl group in TP and the quaternary ammonium nitrogen in
the choline portion of soybean lecithin ³.

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Peak	Compound	Content (%)
1	catechin (C)	32.01
2	epigallocatechin-3- gallate (EGCG)	28.80
3	gallocatechin-3-gallate (GCG)	9.82
4	epicatechin-3-gallate (ECG)	19.36

49 Table S1 Composition of the Tea Polyphenols (HPLC).

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62