RSC advance

## **Electronic Supplementary Information (ESI)**

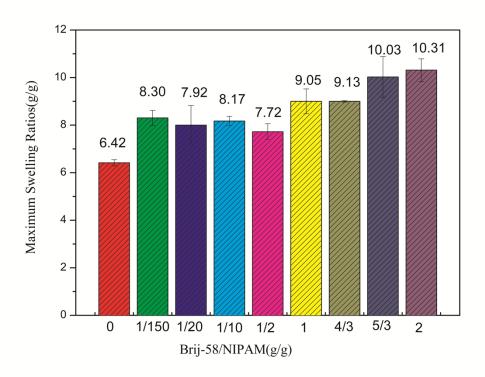
# Trail of Pore shape and Temperature-sensitivity of Poly(N-isopropylacrylamide) Hydrogels before and after Removing Brij-58 Template and Pore Formation Mechanism

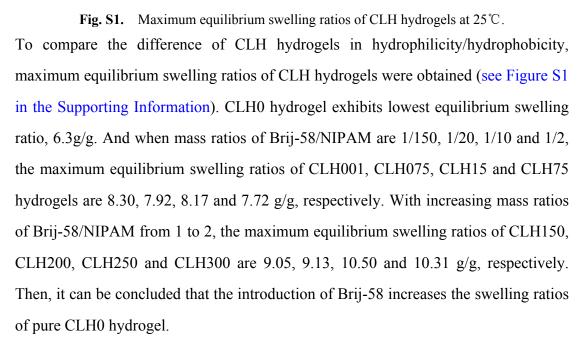
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#### Characterization

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In general, the swelling equilibrium of hydrogel network depends on the osmotic pressure between polymer chains and water molecules, and restricted by cross-linking degree of balance among network flexibility. In fact, no reaction or cross-linking occurs between Brij-58, NIPAM and MBA. Then it is assumed that Brij-58 plays a key role in increasing maximum equilibrium swelling ratios of CLH hydrogels. On the one hand, the chemical structure of Brij-58 is with hydrophilic group like

polyoxyethylene (20). Zhang et al has pointed out that Brij-58 presents highest hydrophilicity in contrast to Brij-52 and Brij-56. On the other hand, it is supposed that Brij-58 serves as a porogenic agent.

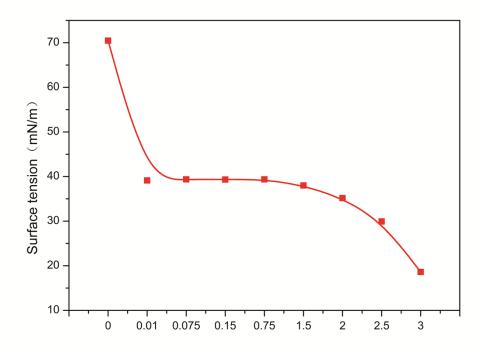
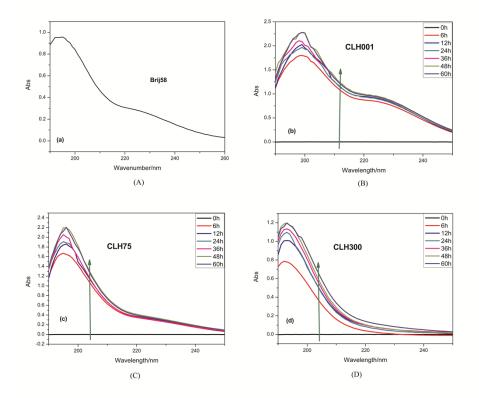


Fig. S2. Surface tension of Brij-58 aqueous solution at different concentration.

The size and shape of Brij-58 template have close relationship with Brij-58 concentration, and then surface tension of Brij-58 aqueous solution with various concentrations was investigated from Fig. S2. It can be observed that the surface tension of pure distilled water is 70.5mN/m, while with introducing Brij-58, surface tension dramatically decreases to 39.1mN/m and then keeps approximately at 39mN/m at concentration from 1 to 75g/L. With further increasing Brij-58 concentration from 150 to 200, 250 and 300g/L, surface tension occurs second decrease and becomes 39.2, 35.2, 29.9 and 18.6mN/m, respectively. The rapid decrease of surface tension at beginning can be attributed to decrease of surface free energy of water resulting from the introduction of hydrophilic/hydrophobic Brij-58. The subsequent equilibrium indicates that Brij-58 molecules still manifest as the formation of micelle. With increasing Brij-58 concentration, Brij-58 molecules form

hexagonal and cubic phases, leading to continuous decrease of surface tension.



**Fig. S3.** The absorbance change of CLH hydrogels during the process of removing Brij-58. To illustrate template action of CLH hydrogels, the UV spectra of Brij-58, CLH0, CLH075 and CLH300 under different removing time are displayed in Fig. S3. Here, the removing time means required time that Brij-58 was completely removed from CLH hydrogel by immersing into the aqueous solution. For Brij-58, an apparent absorption peak occurs at 195 nm. However, for CLH001, CLH75 and CLH300 hydrogels, the absorption peak at 195 nm increases with increasing removing time from 0 to 60 h. It reveals that Brij-58 molecules were gradually removed from CLH hydrogel matrix by immersing into aqueous solution. It is also particularly noteworthy that the two UV/Vis lines of CLH001 and CLH75 at 48 h and 60 h almost coincide with each other, which suggesting that Brij-58 has been basically removed from CLH hydrogel matrix after 48 h.



#### Fig. S4. The chemical structure of Brij-58.

As shown in Fig. S4, "Brij-58" is an ICI trademark name for polyoxyethylene 20 cetyl ether, a nonionic surfactant commonly used in biochemical applications. On a hydrophilic-lipophilic scale (HLB) of 0-20, on which 20 is very hydrophilic (polar), this surfactant has a calculated HLB value of 15.7.<sup>1</sup> The critical micelle concentration (CMC) is reported as 0.007 mM to 0.077 mM; CMC values vary with the salt concentration and temperature.<sup>2</sup>

### References

- 1. Neugebauer, J.M. Methods in Enzymology 1990, 182, 239-253
- 2. Helenius, A.; Simons, K. Biochim. Biophys. Acta 1975, 415, 29-79