

## Supporting information

# Fabrication of optomicrofluidics for real-time bioassays based on hollow sphere colloidal photonic crystals with wettability patterns

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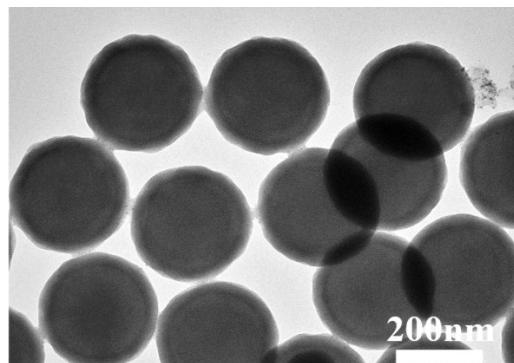
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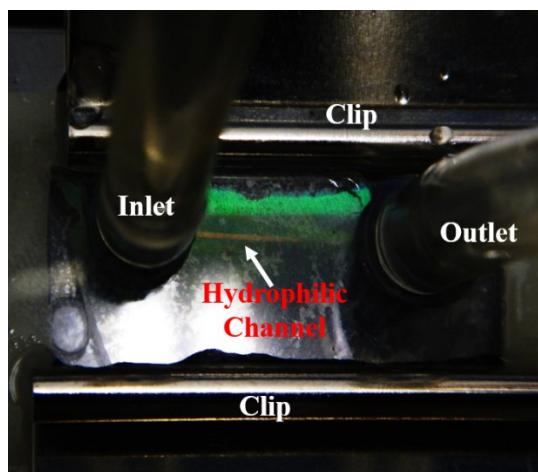
The sequence of two complementary single-stranded DNA used in this work as shown below:

Probe ss-DNA1: 5'-NH<sub>2</sub> -C<sub>6</sub>H<sub>12</sub> -AAA AAA ACC CCT GCA GCC CAT GTA TAC CCC CGA ACC-3'

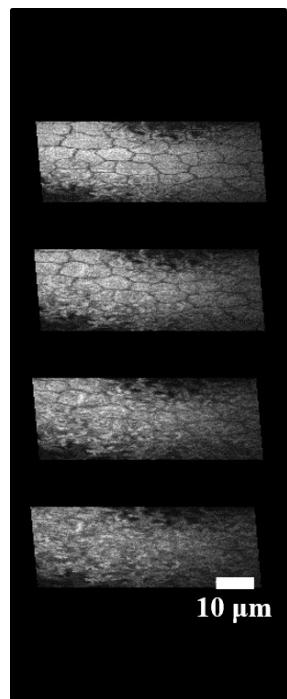
Target ss-DNA2: 5'- Cy3 -C<sub>9</sub>H<sub>18</sub> -GGT TCG GGG GTA TAC ATG GGC TGC AGG GG -3'



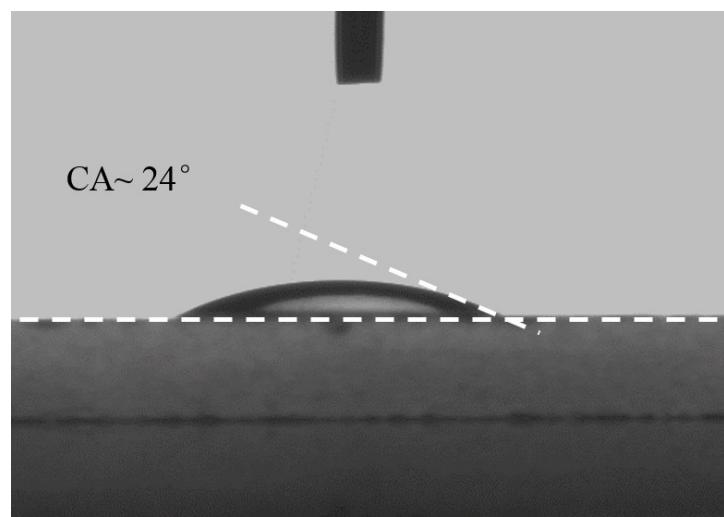
**Fig. S1** TEM image of the PS@SiO<sub>2</sub> core-shell spheres used to produce the hollow spheres, which corresponds to the TEM image of the hollow spheres as shown in Fig.2A.



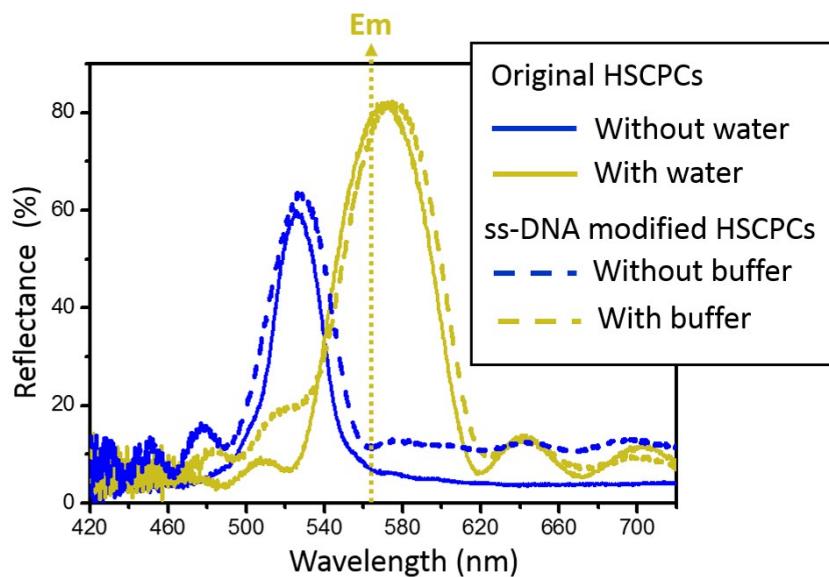
**Fig. S2** Photography of the device contructed from the superhydrophobic HSCPCs with a superhydrophilic channel filled with water for visualization



**Fig. S3** Sliced two-photon fluorescence 3D images of the hydrophilic channel in the HSCPCs. The interval is  $1\mu\text{m}$



**Fig. S4** The CA measurement of water droplets on sample surface of the ss-DNA1 modified channel region.



**Fig. S5** The reflectance spectra of the HSCPCs with (dashed lines) and without (solid lines) ss-DNA modification before (blue color) and after (yellow color) solution infiltration. The yellow arrow indicates the emission spectrum of the fluorophore (Cy3)

1. Z.-F. Liu, T. Ding, G. Zhang, K. Song, K. Clays and C.-H. Tung, *Langmuir*, 2008, **24**, 10519.
2. W. Stober, A. Fink and E. Bohn, *J. Colloid Interface Sci.*, 1968, **26**, 62.