

Supporting Information

Liquid-like Nanofluids Mediated Modification of Solar-assisted Sponge for Highly Efficient Cleanup and Recycling of a Viscous Crude Oil Spill

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Section S1: The preparation of SiO₂ nanofluids (SiO₂ nfs)

1.0 g SiO₂ powder were dispersed in 50 mL of deionized water and ultrasonicated for 30 min. Then 10 mL of DC5700 in methanol (40 mass %) were dropwise added and aged for 24 h at room temperature by gently shaking it periodically. Next, the solution was separated by extraction filtration and washed three times in turn with water and methanol. The separated light yellow precipitate was dried at 65 °C for 24 h in a vacuum oven to obtain the SiO₂ chlorine salt. Subsequently, the SiO₂ chlorine salt was added in 15 mass % solution of NPEP in water/chloroform (volume ratio = 2:3) while stirring at 50 °C for 24 h. Finally, the resultant solution was dried and extracted three times in turn. The solid was also dispersed in chloroform, centrifuged, and then dried in a vacuum oven at 65 °C for 48 h to obtain yellow nanofluids at room temperature..

Section S2: Supporting figures

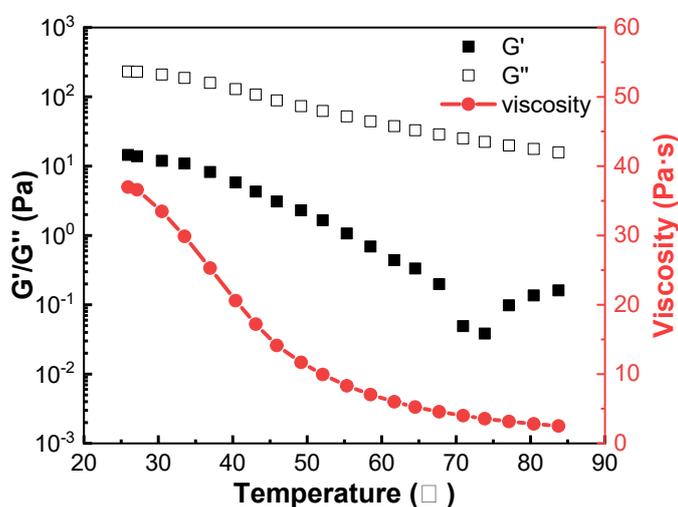


Figure S1. Rheological behavior of SiO₂ nfs. All the above tests were conducted at an environmental temperature of 25 °C.

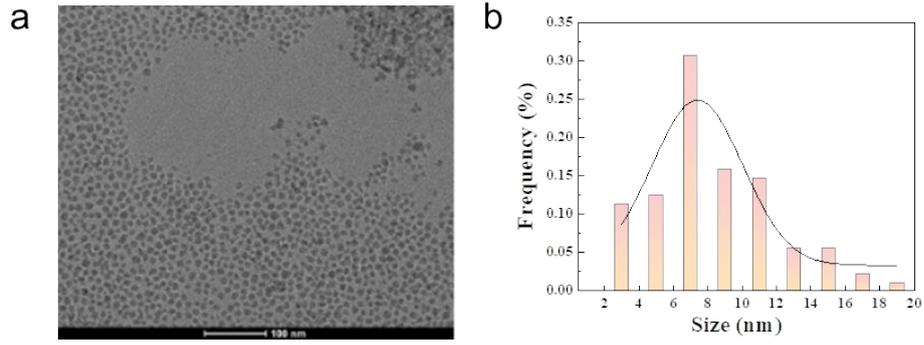


Figure S2. (a) TEM images of SiO₂ nfs. (b) Size distribution of SiO₂ nfs.

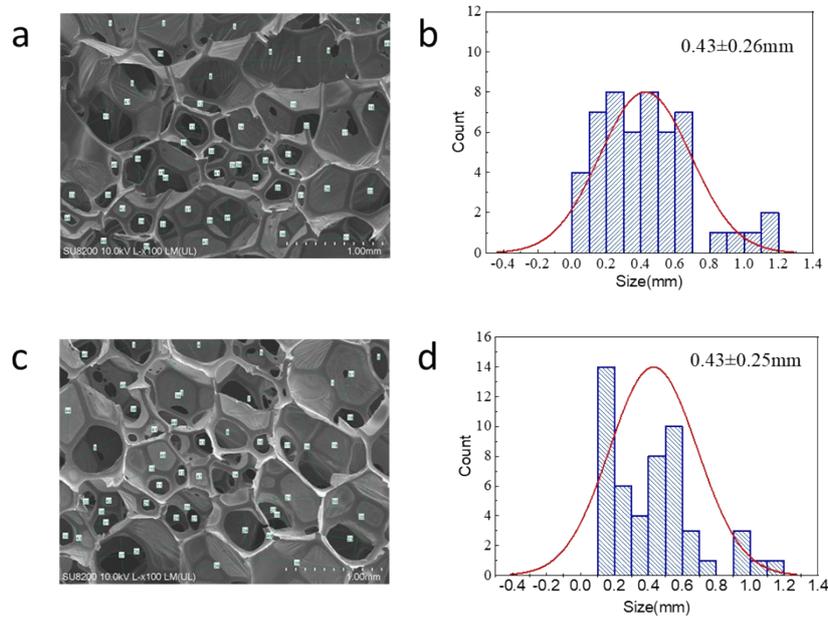


Figure S3. SEM images of the (a) PUS sponges and (c) PSiNFs@PU, Statistical data of pore size distribution of the (b) PUS sponges and (d) PSiNFs@PU obtained from SEM.

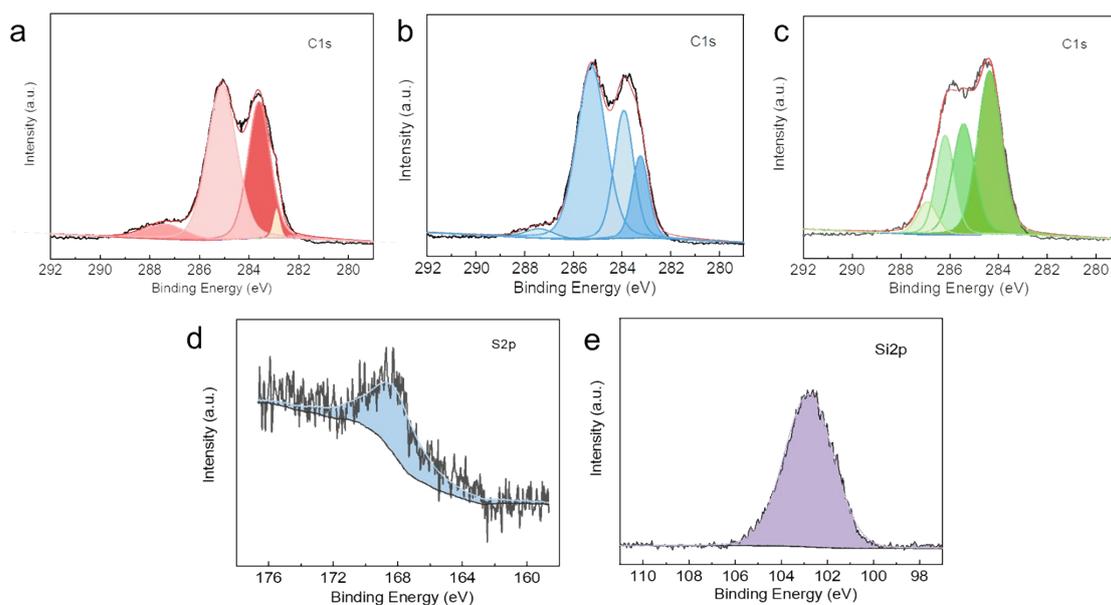


Figure S4. High resolution XPS spectra of C1s of (a) PUS, (b) PDA@PU, and (c) PSiNFs@PU. High resolution XPS spectra of (d) S2p and (e) Si2p of PSiNFs@PU. All the above tests were conducted at an environmental temperature of 25 °C.

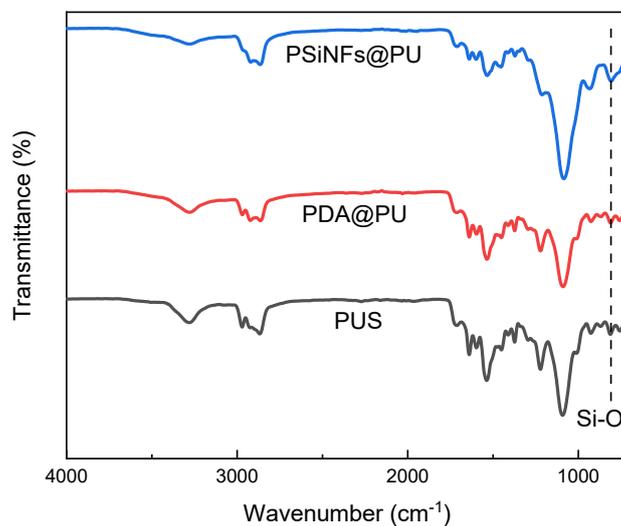


Figure S5. FTIR curves of PUS, PDA@PU, and PSiNFs@PU, respectively. All the above tests were conducted at an environmental temperature of 25 °C.

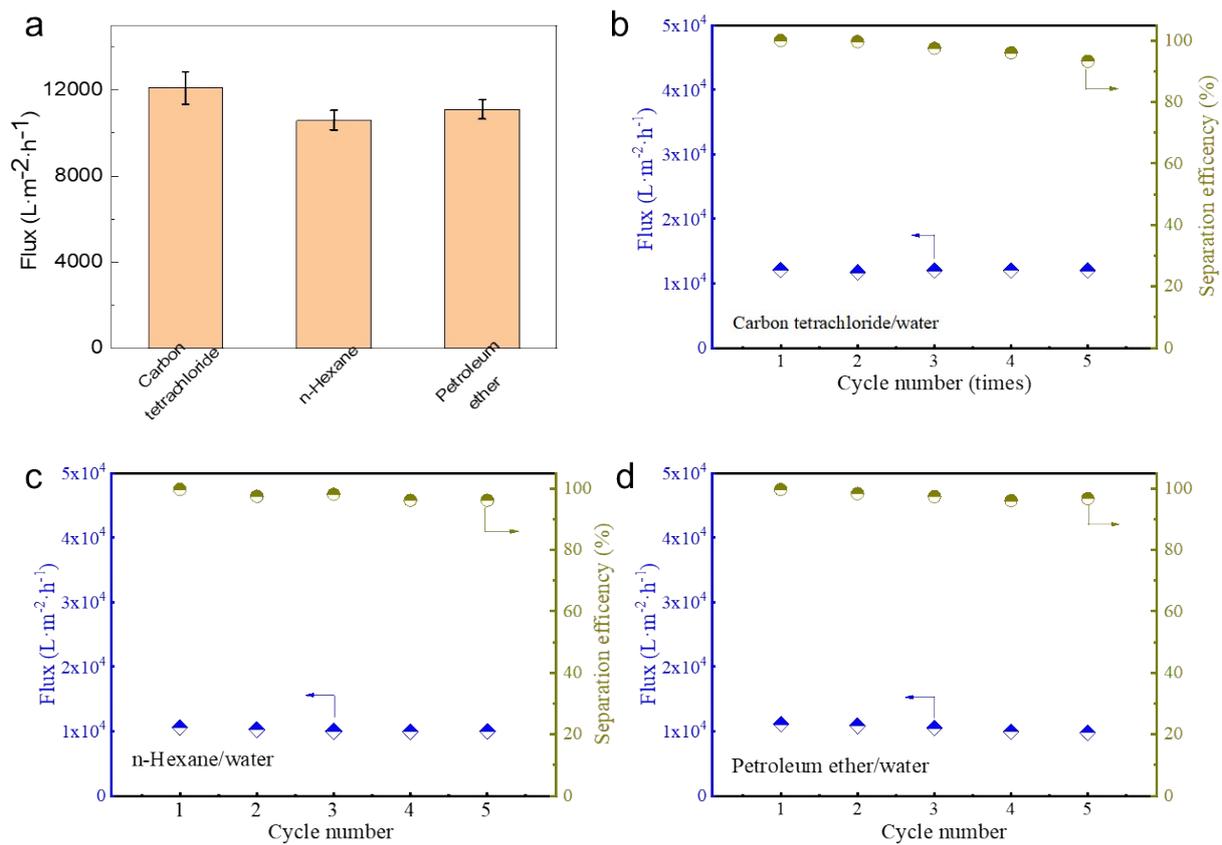


Figure S6. Separation performance of the PSiNFs@PU sponges. (a) Surfactant-stabilized water in oil emulsions separation. Cycling performance of the PSiNFs@PU sponges: (b) carbon tetrachloride/water emulsion, (c) n-hexane/water emulsion, and (d) petroleum ether/water emulsion.

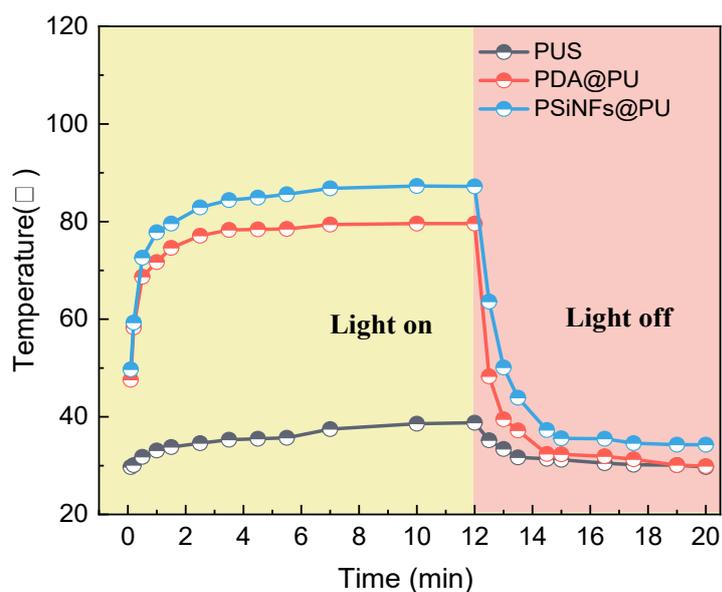


Figure S7. Time-dependent temperature evolution curves of different sponges under one sun illumination (power density: 1.00 kW /m^2 , the yellow and pink region represent temperature-rise and temperature-fall period). All the above tests were conducted at an environmental temperature of $25 \text{ }^\circ\text{C}$.

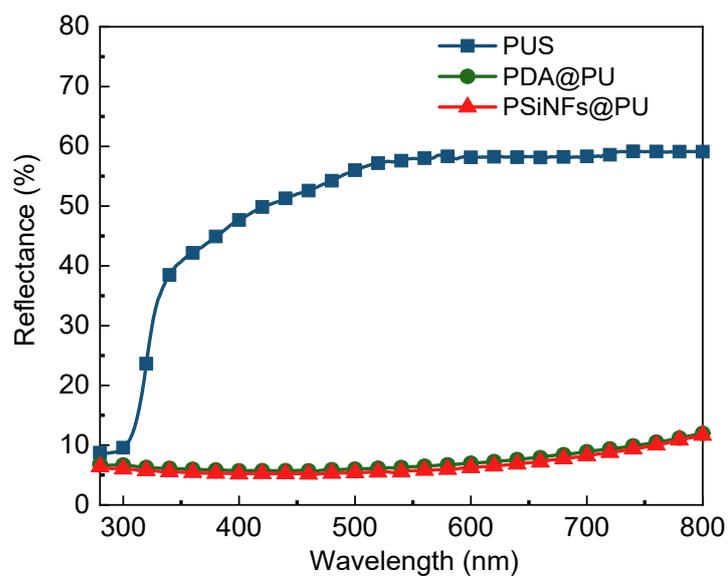


Figure S8. Reflectance spectra of the PUS, PDA@PU, and PSiNFs@PU. All the above tests were conducted at an environmental temperature of $25 \text{ }^\circ\text{C}$.

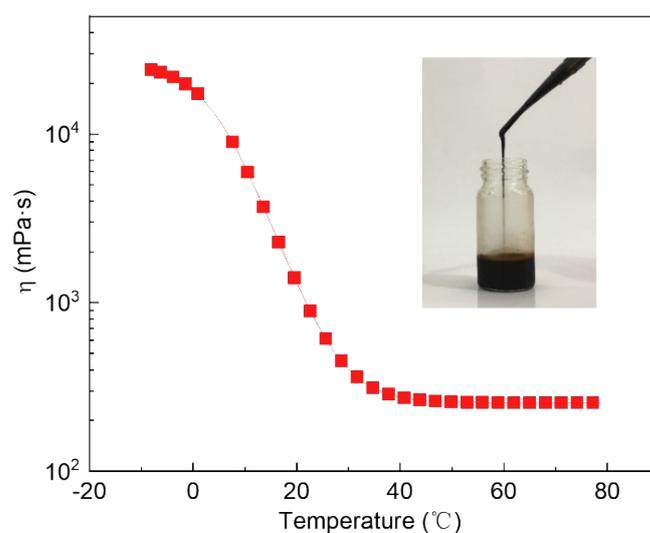


Figure S9. Viscosity changes of crude oil as a function of oil temperature (the shear rate is 1 s^{-1}). The inset image is the crude oils at $0 \text{ }^\circ\text{C}$. The test was conducted at an environmental temperature of $25 \text{ }^\circ\text{C}$.

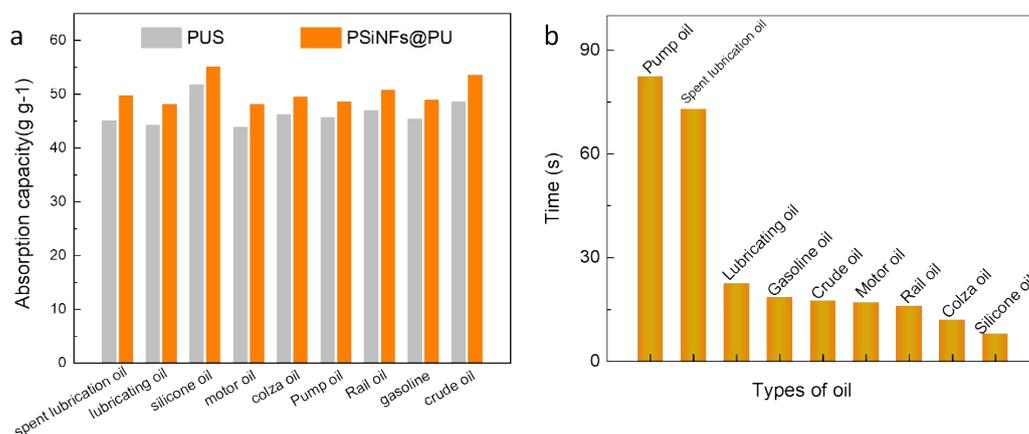


Figure S10. (a) Mass-based absorption capacity of the PUS and PSiNFs@PU sponges for various oils (power density: 1.00 kW / m^2). (b) The time required for the PSiNFs@PU sponges to absorb all types of oil to saturated state (power density: 1.00 kW / m^2). All the above tests were conducted at an environmental temperature of $25 \text{ }^\circ\text{C}$.

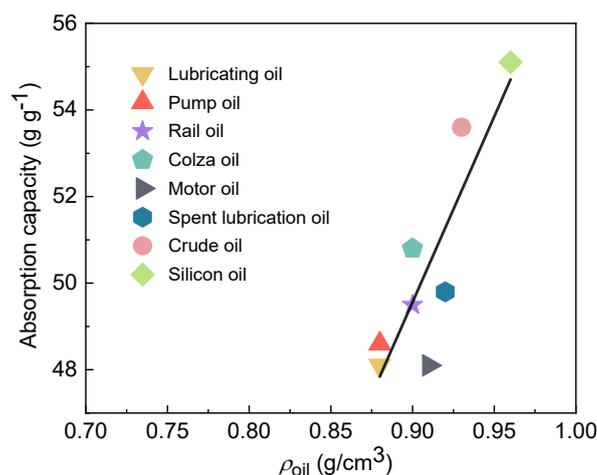


Figure S11. The relationship between the saturation capacity of oil absorption and the density of oil of the PSiNFs@PU sponges (the solid line is the fitted curve). All the above tests were conducted at an environmental temperature of 25 °C.

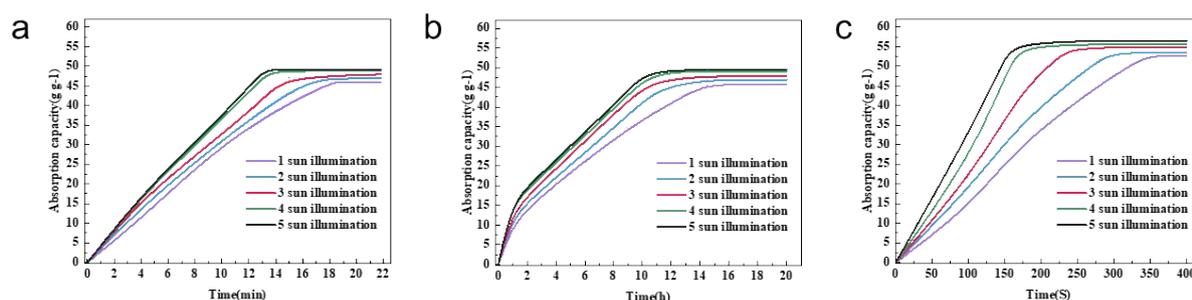


Figure S12. The oil absorption capacity versus time for the PUS (a), PDA@PU (b), and PSiNFs@PU (c), respectively. All the above tests were conducted at an environmental temperature of 25 °C.

Table S1 Adsorption capacity of different adsorbents reported in the literature

Material	Absorbed organics	Capacity ^k	
Fe ₃ O ₄ /HDPE PU sponge	Toluene, hexane, dichloromethane, chloroform, cook oil, carbon tetrachloride	15-52	1
Fe ₃ O ₄ -TMHFDS, TEOS/PU sponge	Peanut oil, pump oil, and silicone oil	39-46	2
Fe ₃ O ₄ @GO@OTS PU sponge	Lubricating oil, peanut oil, hexadecane, octane, hexane, heptane	9-27	3
F-MoS ₂ -PU sponge	Hexane, diethyl ether, ethanol, toluene, rapeseed oil, acetonitrile, dichloromethane and chloroform	25-90	4
NDS-PDA-PFDT-PU sponge	Chloroform, diesel, pump oil, gasoline, ethanol, toluene, hexadecane, hexane	4-60	5
CNT/PDMS-PU sponge	Soybean oil, used motor oil, diesel oil, n-hexadecane, gasoline, n-hexane	15-25	6
Polymer molecular brush/PU sponge	Crude oil, Peanut oil, Toluene, Acetone Chloroform, n-hexane Petroleum ether	17-40	7
Fe ₃ O ₄ -graphene PU sponge	lubricating oil, paraffin oil, hydraulic oil, peanut oil, n-hexadecane, heptane	9-27	3
PDA-ZIF-8 melamine sponge	Petroleum ether, Chloroform, Tetrahydrofuran, Gasoline, Toluene, Dodecane	10-38	8
PSiNFs@PU	Crude oil, gasoline, lubricating oil, Pumping oil, Colza oil, Rail oil, motor oil	45-55	Present work

^k Refers to the times of the weight of adsorbed substances vs. the weight of adsorbent.

HDPE: high-density polyethylene, TMHFDS: trimethoxy (1H, 1H, 2H, 2H-heptafluorodecyl) silane, TEOS: tetraethoxysilane, GO: graphene oxide, OTS: octadecyltrichlorosilane, PU: polyurethane, NDS: nanodiamonds, PDA: polydopamine, PFDT: 1H,1H,2H,2H-perfluorodecanethiol, CNT: carbon nanotubes, PDMS: polydimethylsiloxane.

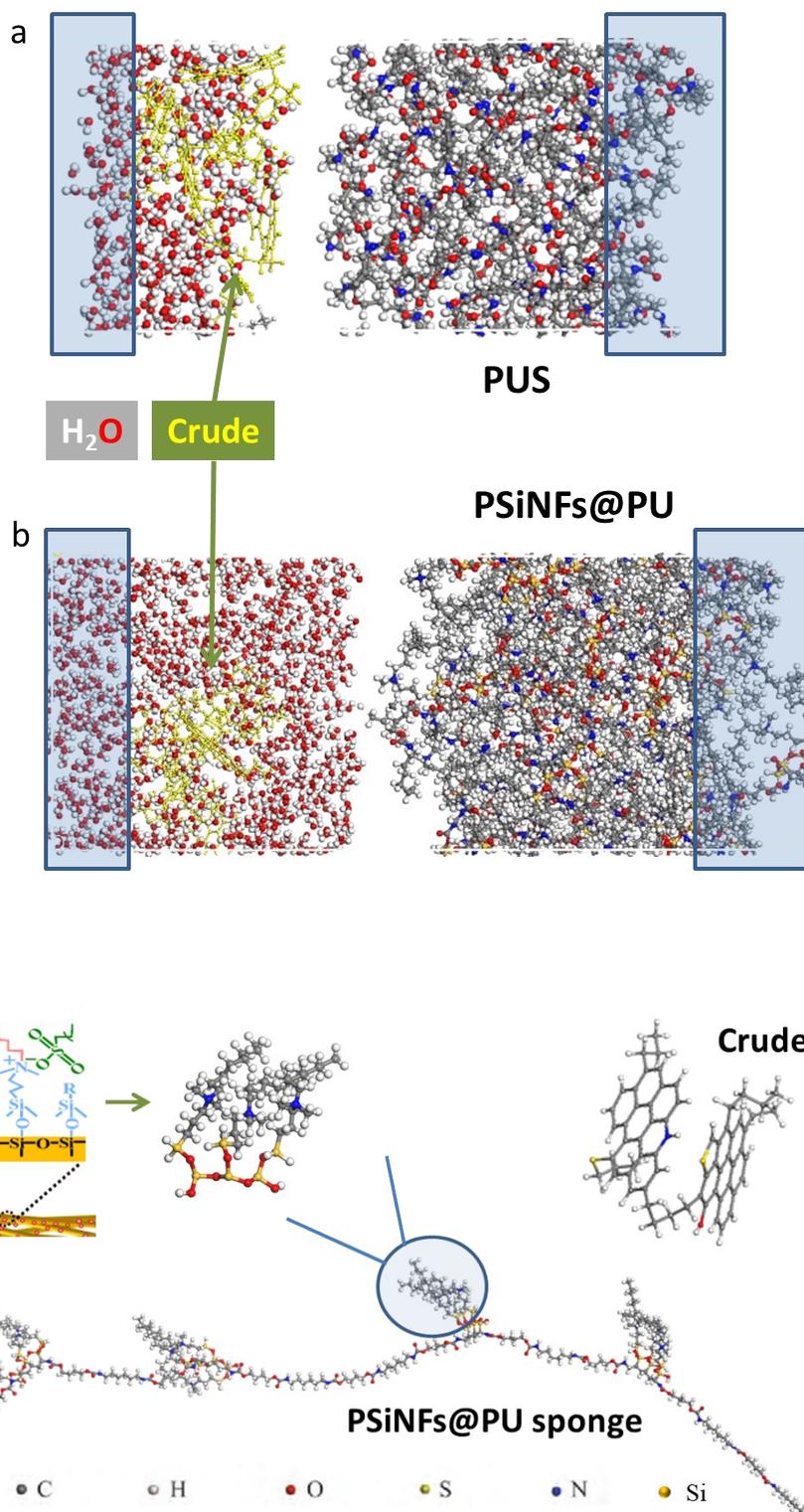


Figure S13. Interface model of crude oil diffusion from water to (a) PUS and (b)

PSiNFs@PU.

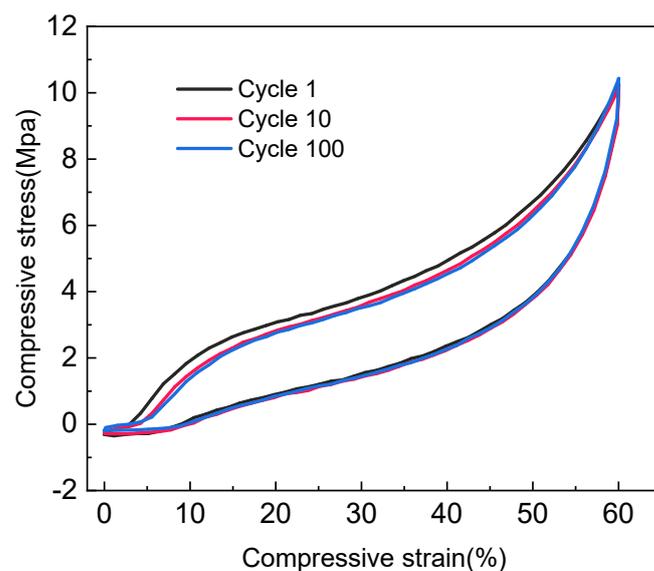


Figure S14. Strain–stress curves of the PSiNFs@PU sponges with cyclic compression under 60% compressive strains. The tests were conducted at an environmental temperature of 25 °C.



Figure S15. Compression recovery of the PSiNFs@PU sponge at 60 % compression strain.

The tests were conducted at an environmental temperature of 25 °C.

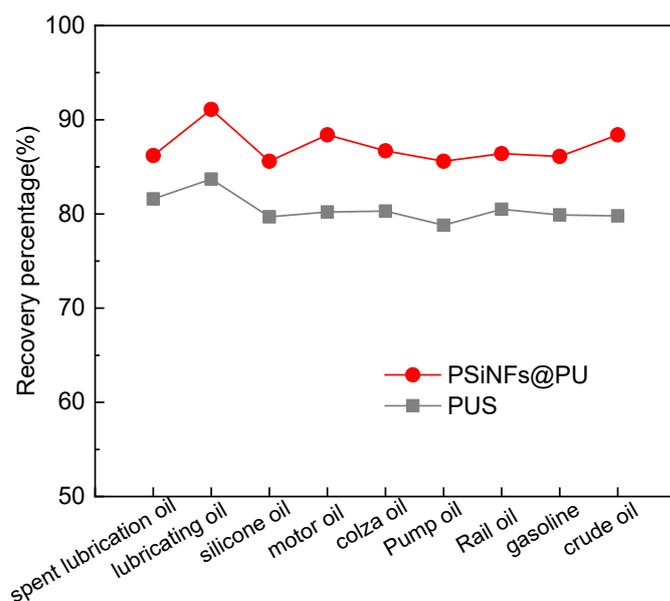


Figure S16. Mass-based squeeze capacity of the PSiNFs@PU sponges for various oils without sun condition. The tests were conducted at an environmental temperature of 25 °C.

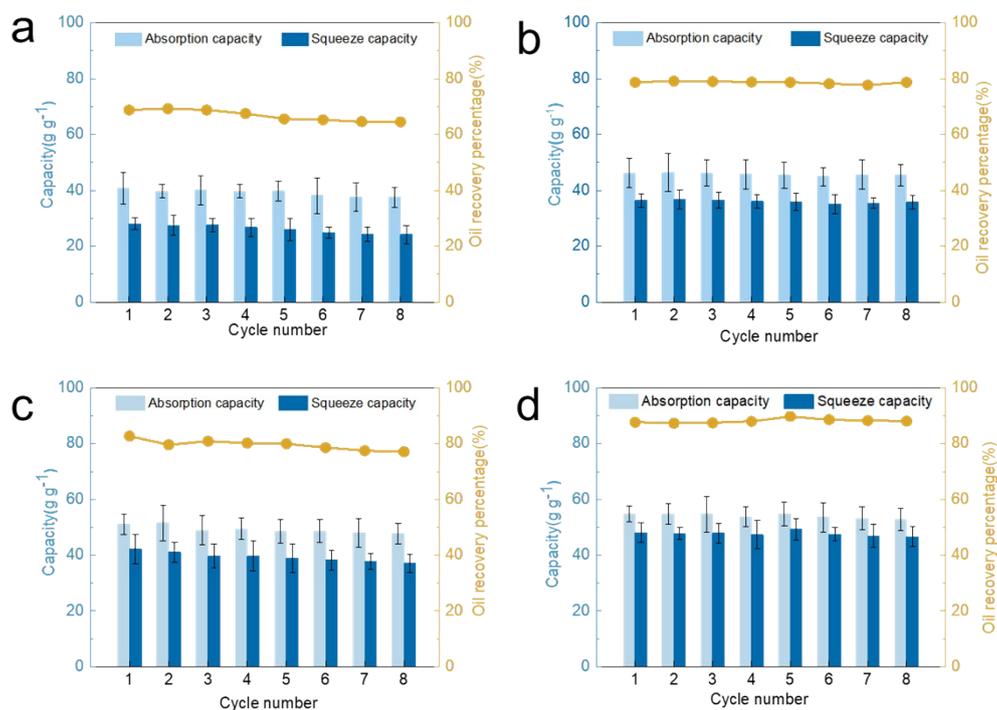


Figure S17. The amount of oil absorbed, squeezed and the recovery rate of the PUS (a) and PSiNFs@PU (b) at no sun illumination. The amount of oil absorbed, squeezed and the recovery

rate of the PUS (c) and PSiNFs@PU (d) at one sun illumination. The tests were conducted at an environmental temperature of 25 °C.

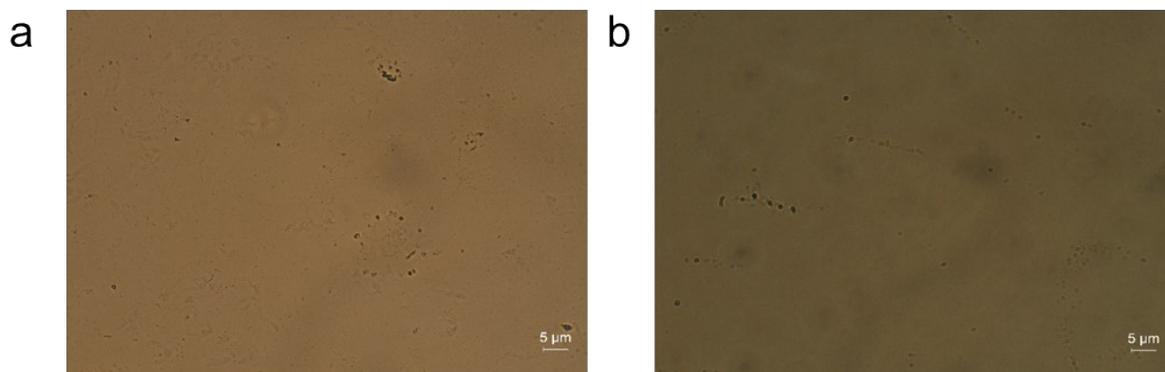


Figure S18. POM images of (a) original crude oil and (b) recovery crude oil extruded from the PSiNFs@PU sponges. The tests were conducted at an environmental temperature of 25 °C.

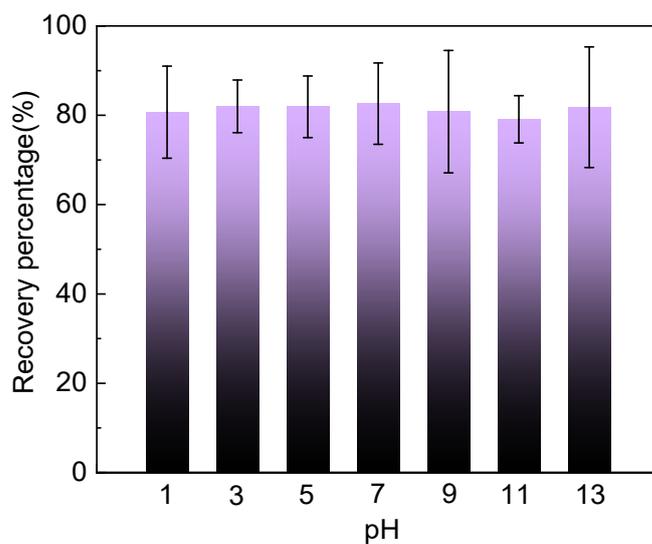


Figure S19. Crude oil recovery performance of the PSiNFs@PU at different pH values without sun illumination. The tests were conducted at an environmental temperature of 25 °C.



Figure S20. Photograph of continuous oil absorption of the PSiNFs@PU sponge under light condition (power density: 1.00 kW /m²). The tests were conducted at an environmental temperature of 25 °C.

Reference

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