Electronic Supplementary Information (ESI)

Hierarchically porous electrospun nanofibrous mats produced from intrinsically microporous fluorinated polyimide for the removal of oils and non-polar solvents

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Experimental Section.

Membrane Preparation. A 6FDA-TrMPD dense film was prepared through solvent evaporation from (3 wt/vol%) its polymer solution in chloroform. The solution was filtered through 0.45 μ m PTFE filter and poured onto a flat glass Petri dish. The solvent was evaporated slowly at RT over 24 h. Thereafter, the obtained film was further dried at 120 °C for 24 h under vacuum. The resulting robust film with a thickness of ~ 40 μ m was used for tensile measurement and for BET surface area analysis.

Characterization. The infrared spectrum was recorded for polyimide powder using a Varian 670-IR FT-IR spectrometer.

Table S1. The characteristics of the 6FDA-TrMPD polyimide. Weight (M_w) and number (M_n) average molecular weights, polydispersity index (PDI), degradation temperature $T_{d,5\%}$, and surface area (S_{BET}).

	$M_{ m w}$	$M_{ m n}$	PDI	T _{d, 5%}	S _{BET}
Polymer	(g mol ⁻¹)	(g mol ⁻¹)	(-)	(°C)	$(m^2 g^{-1})$
6FDA-TrMPD	176,606	110,987	1.59	509	450 ±15

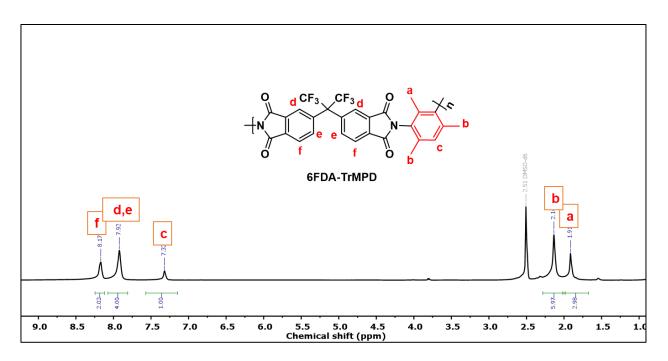


Fig. S1. ¹H NMR spectrum of the 6FDA-TrMPD in DMSO-d₆.

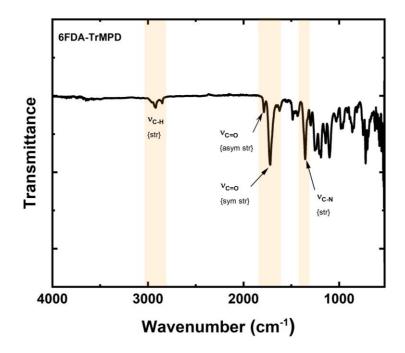


Fig. S2. Attenuated total reflectance Fourier-transform infrared spectrum (ATR-FTIR) spectrum of the 6FDA-TrMPD.

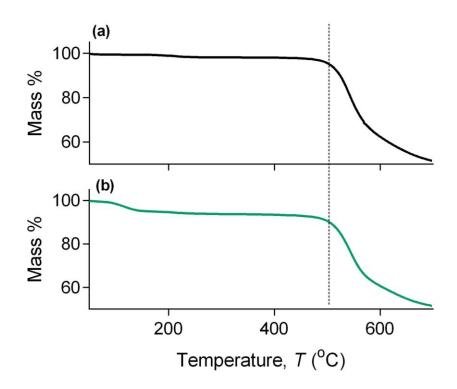


Fig. S3. TGA thermograms of the 6FDA-TrMPD (a) powder and (b) electrospun mat.

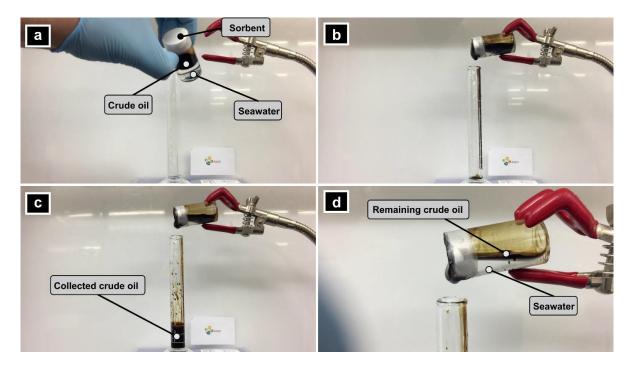


Fig. S4. The use of the electrospun mat for water-oil separation. (a) Before, and (b-d) during the separation process.

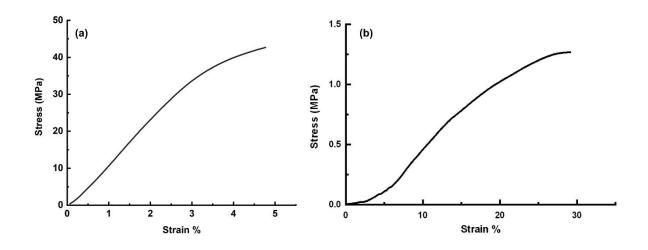


Fig. S5. Stress-strain curves of the free standing film made truly from the 6FDA-TrMPD powder (a) and its electrospun mat (b).

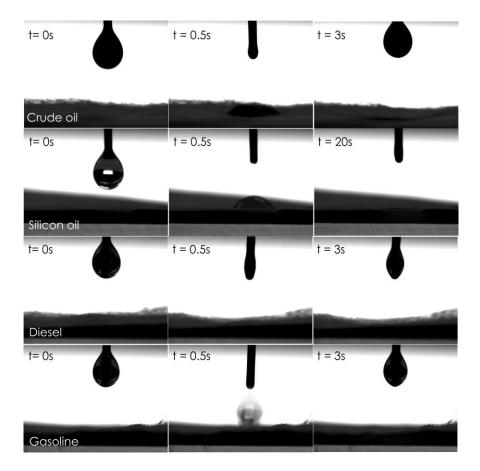


Fig. S6. Contact angle measurements of the fiber mat using various oils. The mat imbibed oil droplets immediately upon contact with the droplet (*i.e.* oil contact angle (OCA) $< 1^{\circ}$), demonstrating superoleophilicity.

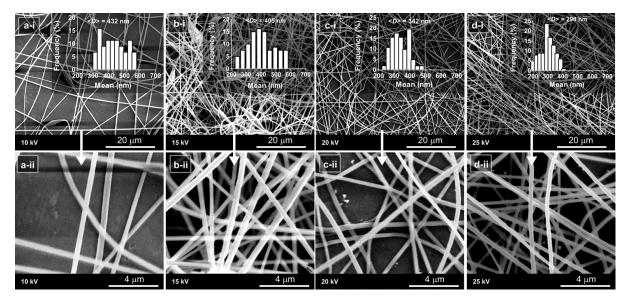


Fig. S7. The influence of the applied voltage on the morphology of the 6FDA-TrMPD nanofibers. $c_{6FDA-TrMPD}=10\%$ (w/v). The tip-to-collector distance kept at 15 cm and the flow rate set to 0.5 mL/h. Insets show the statistical distribution for the diameter of the respective nanofibers.

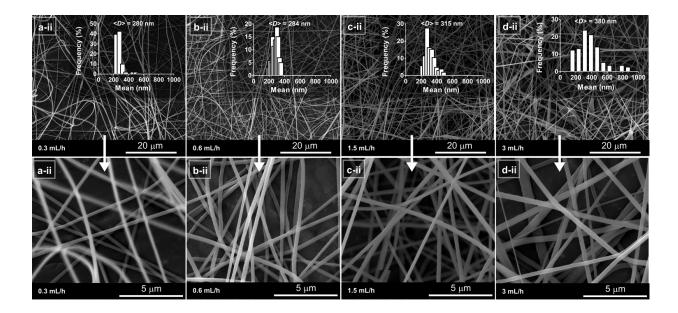


Fig. S8. The influence of the flow rate on the morphology of the 6FDA-TrMPD nanofibers. $c_{6FDA-TrMPD}=10\%$ (w/v). The applied voltage was 20 kV and the tip-to-collector distance kept at 15 cm. Insets show the statistical distribution for the diameter of the respective nanofibers.

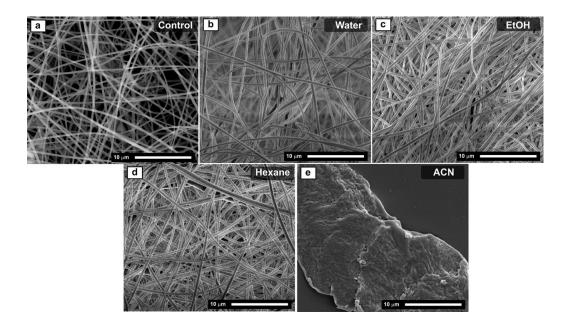


Fig. S9. Scanning electron micrographs of the 6FDA-TrMPD nanofibers exposed to various solvents for 24 h. c_{6FDA} . TrMPD= 10% (w/v). (a) Control sample without any solvent treatment, (b) water, (c) ethanol (EtOH), (d) hexane, and (e) acetonitrile (ACN).

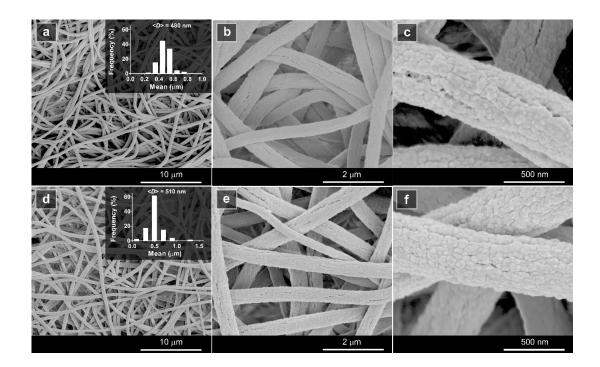


Fig. S10. Scanning electron micrographs of the electrospun mats exposed to toluene (a-c) and m-xylene (d-e) for 24 h. Insets show the statistical distribution for the diameter of the respective nanofibers.

Sorbent	Material form	BET surface area (m ² g ⁻¹)	Sorbate	Sorption capacity (g g ⁻¹)	Ref.
			Crude oil	34.6	
Intrinsically porous fluorinated polyimides	Fibrous mat	560	Diesel	55.76	This study
			Gasoline	31.25	
A porous superhydrophobic SiO2@polystyrene	Foam	n.d.	Crude oil	32.1	1
Bio-based oil gelling agent	Powder	n.d.	Crude oil	4.7	2
SiO ₂ decorated cotton fibers	Fibers	n.d.	Crude oil	57.0	_ 3
			Diesel	25.61	
Porous PS fibers	Fibers	n.d.	Diesel	7.13	4
Carbon shoot sponge	Sponge	440	Crude oil	~30	5
Cellulose-based aerogels	Aerogel	n.d.	Crude oils	18.4-20.5	6
Graphene sponge	Sponge	n.d.	Crude oil	85-90	7
CNF/carbon foam	Foam	n.d	Diesel	21	8
			Gasoline	16	
Lignin-based polyurethane/graphe ne oxide foam	Foam	n.d.	Crude oil	25.4	9
Poly(dimethylsiloxa ne)-TiO2 coated polyurethane sponge	Sponge	n.d.	Diesel	14.2	10

Table S2. Oil sorption performance of various polymeric adsorbents.

n.d.: Not determined

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