

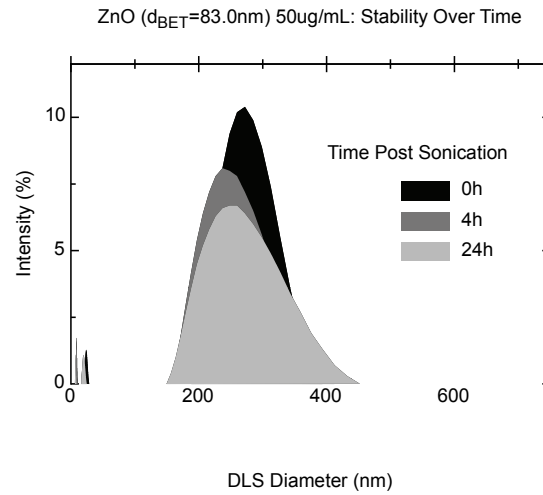
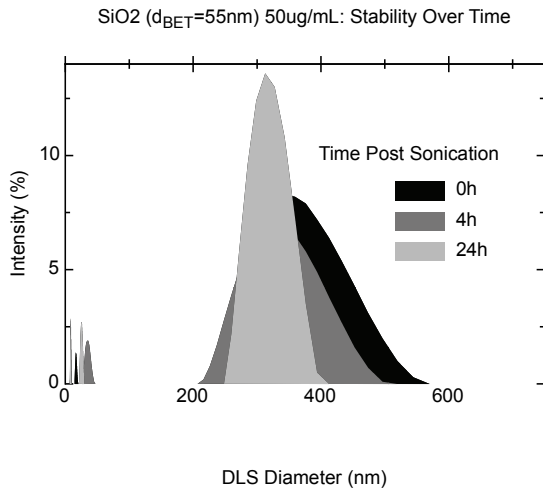
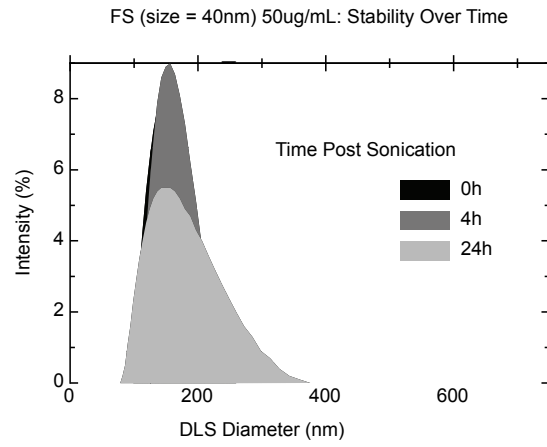
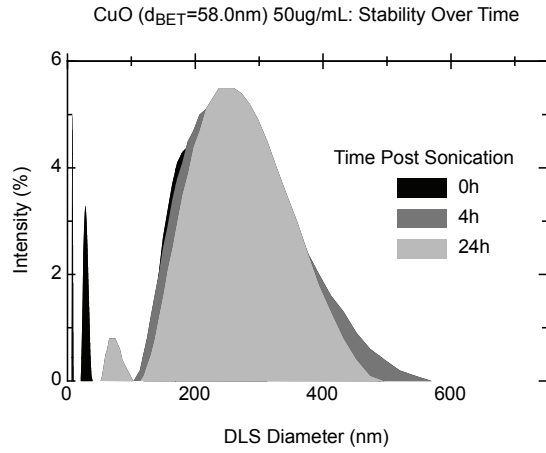
Supplement materials

Engineered nanoparticles impair wound healing of the human cornea

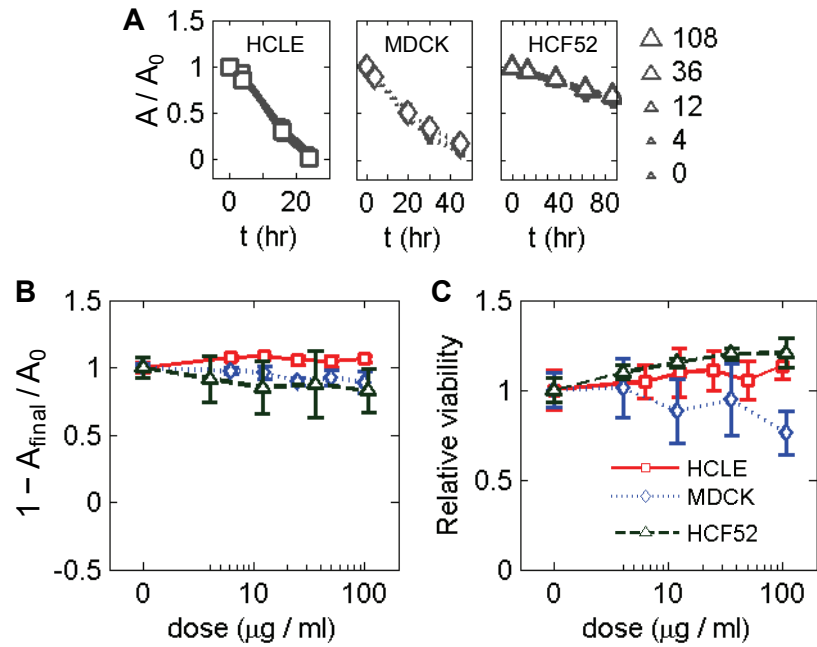
By Zhou et al.

Supplement Table 1. The dry powder properties for 4 ENPs. SSA: (specific surface area) by nitrogen adsorption/Brunauer-Emmett-Teller (BET) method. d_{BET} : particle diameter determined from SSA and particle density, ρ , as described in methods. d_{XRD} : particle diameter by X-ray diffraction.

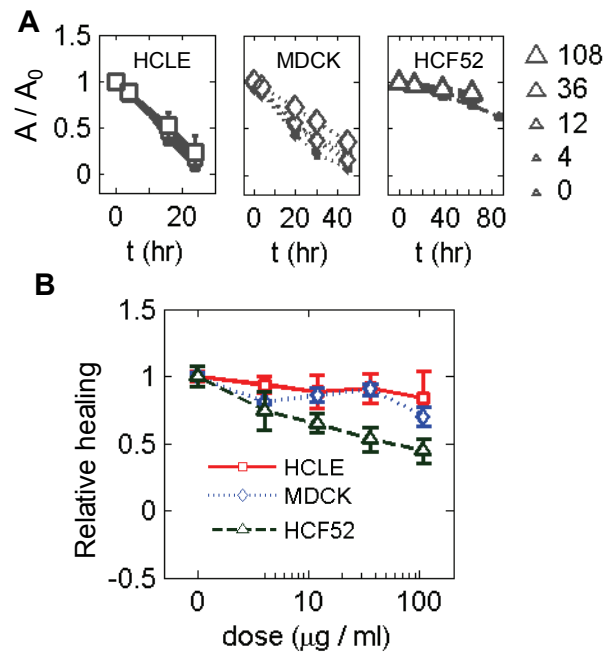
Material	SSA (m ² /g)	d_{BET} (nm)	d_{XRD} (nm)	ρ (g/cm ³)
EVONIK SiO ₂	50	55	NA	2.2
Sigma Aldrich CuO	17.3	58.0	22	6.31
Alfa Aesar ZnO	13	83	66.8	5.606
EVONIK TiO ₂	50	21	33	4.23



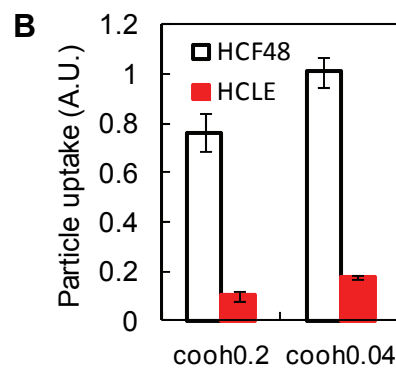
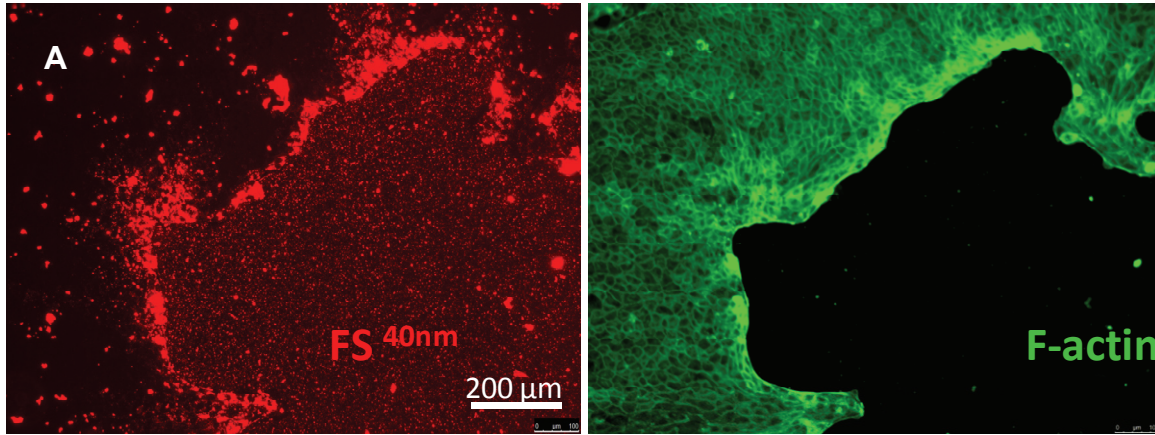
Supplement Fig. 1 Suspensions of CuO, SiO₂, ZnO, and FS^{40nm} ENPs exhibited stable particle size over time.



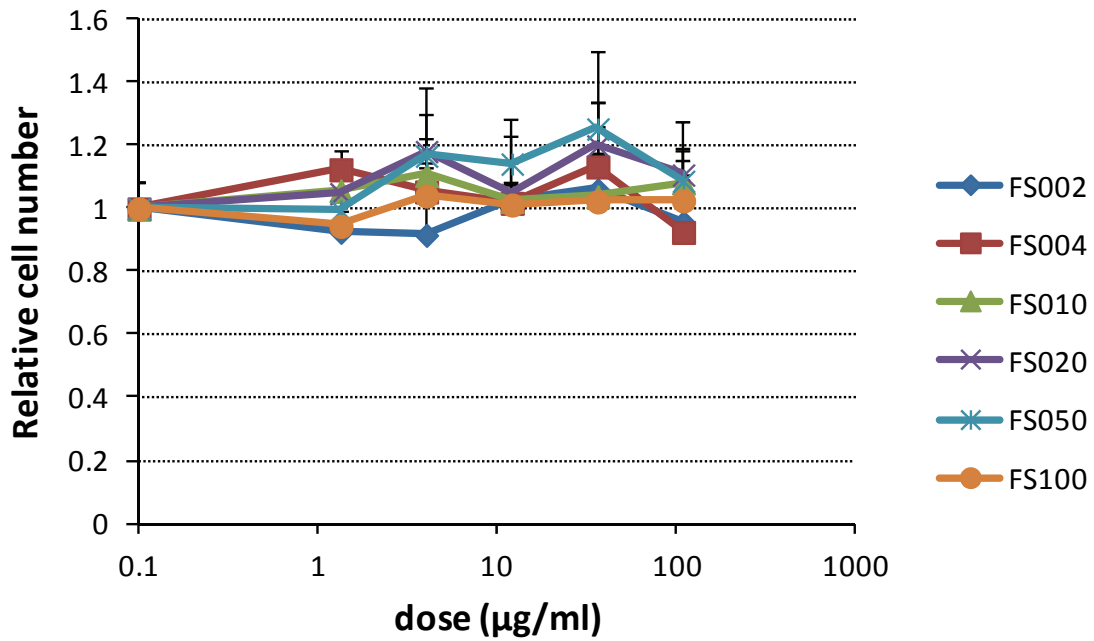
Supplement Fig. 2. SiO₂ ENPs exerted little impact on the wound healing rate (A, B) or the viability (C) of HCLE, MDCK and HCF52 cells. Mean \pm 1.96 s.e.m. with $n \geq 6$ wells from 2 experiments.



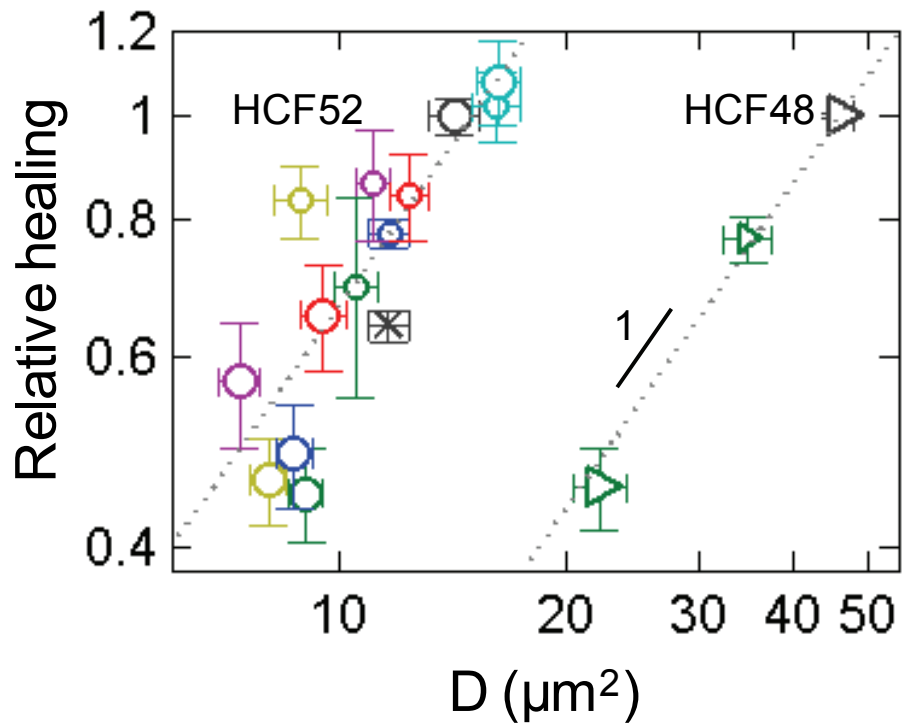
Supplement Fig. 3. Non-fluorescent, carboxylated polystyrene ENPs did not alter the wound healing rate of HCLE or MDCK cells but dose-dependently slowed down that of HCF52 cells. Mean \pm 1.96 s.e.m. with $n \geq 6$ wells from 2 experiments.



Supplement Fig. 4. (A) MDCK cells only took up FS^{40nm} at the wound edge. There was minimal particle deposition on the interior cells. Left: fluorescent particles; right: F-actin staining. (B) Particle uptake is higher for HCF48 cells than for HCLE cells. Confluent cells were treated for one day with 12 μg/ml of FS of two sizes, and then subjected to fixation before fluorescent particles are quantified using a plate reader. Compare these data with Fig. 3E, where cells were prepared identically except for receiving fixation/permeabilization. Mean ± 1.96 s.e.m.; non-overlapping error bars indicate statistically significant difference (P < 0.05); n = 6-12 wells from 2 experiments.



Supplement Fig. 5. Carboxylated Fluospheres do not impede cell proliferation. HCF cells were plated in a 96 well plate at 1600/w, treated with 4/3 to 108 µg/ml for two days, and counted using the cyquant assay following manufacturer’s instruction. Mean ± SD; n = 4 wells from 2 experiments.



Supplement Fig. 6. The wound healing rate relative to control is proportional to cell speed, D . The fitted lines have a power-law slope of about 1 in a log-log plot. For HCF52, $y = (0.066 \pm 0.005) * x$ and for HCF48, $y = (0.022 \pm 0.002) * x$; 95% confidence interval is reported here.