Supporting Information

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Fig. S1. Effect of charge on cellular internalization. (*A*) The chemical transformation, leading to changes in surface charge. (*B*) Particle internalization after 4 h of incubation with HeLa cells with *positively* charged particles. (*C*) Particle internalization after 4 h of incubation with HeLa cells with *negatively* charged particles.



Fig. S2. Confocal laser scanning microscopy images of HeLa cells after a 1-h incubation period at 37°C with 200-nm cylindrical particles (A), 1- μ m cylindrical particles (AR = 1) (B), 2- μ m cubic particles (C), and 3- μ m cubic particles (D) (scale bar: 10 μ m.)

DNA NG



Fig. S3. Transmission electron microscopy image of HeLa cells at 37°C incubated with 200 nm (AR = 1) cylindrical particles (15-min incubation time).

ANd

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DNA



Fig. S5. Transmission electron microscopy image of HeLa cells at 37°C incubated with 200-nm (AR = 1) cylindrical particles (4-h incubation time).

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Fig. S6. Transmission electron microscopy image of HeLa cells at 37°C incubated with 200-nm (AR = 1) cylindrical particles (4-h incubation time). Red arrows depict particle location.

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Fig. S7. Transmission electron microscopy image of HeLa cells at 37°C incubated with 150-nm (AR = 3) cylindrical particles (1-h incubation time). Red arrow depicts particle location.

DNA S



Fig. S8. Transmission electron microscopy image of HeLa cells at 37°C incubated with 150-nm (AR = 3) cylindrical particles (1-h incubation time).

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Fig. S9. Transmission electron microscopy image of HeLa cells at 37°C incubated with 150-nm (AR = 3) cylindrical particles (4-h incubation time). Red arrow depicts particle location.

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Fig. S10. Transmission electron microscopy image of HeLa cells at 37°C incubated with 150-nm (AR = 3) cylindrical particles (4-h incubation time). Red arrow depicts particle location.

SANG SAT



Fig. S11. Transmission electron microscopy image of HeLa cells at 37° C incubated with $1-\mu$ m (AR = 1) cylindrical particles (1-h incubation time).

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Fig. S12. Transmission electron microscopy image of HeLa cells at 37° C incubated with $1-\mu$ m (AR = 1) cylindrical particles (1-h incubation time).

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Fig. S13. Transmission electron microscopy image of HeLa cells at 37° C incubated with $1-\mu$ m (AR = 1) cylindrical particles (4-h incubation time).

DNA C

Table S1. Particle size characterization as determined by scanning electron microscopy measurements

| Particle size | Height, μ m | Width, μ m |
|-----------------------------------|-----------------|-----------------------------------|
| 5-μm cubes | 4.60 ± 0.07 | 4.67 ± 0.20 |
| $3-\mu m$ cubes | 2.50 ± 0.10 | $\textbf{2.63} \pm \textbf{0.09}$ |
| $2-\mu m$ cubes | 1.56 ± 0.09 | 1.86 ± 0.04 |
| $1-\mu m$ cylinders (AR = 1) | 0.58 ± 0.05 | 0.90 ± 0.01 |
| $0.5-\mu m$ cylinders (AR = 2) | 0.38 ± 0.02 | 0.77 ± 0.09 |
| $0.2-\mu m$ cylinders (AR = 1) | 0.217 ± 0.006 | 0.159 ± 0.007 |
| $0.15 - \mu m$ cylinders (AR = 3) | 0.479 ± 0.026 | 0.134 ± 0.026 (top) |
| | | 0.159 ± 0.012 (bottom) |
| 0.1- μ m cylinders (AR = 3) | 0.277 ± 0.014 | 0.075 ± 0.003 (top) |
| | | 0.118 ± 0.005 (bottom) |

Particle labels describe master cavity size.

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Table S2. Characterization of the surface charge of PRINT particles

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| Particle size | Zeta potential, mV |
|----------------------------------|--------------------|
| 5-μm cubes | +26 ± 3 |
| 3-µm cubes | +21 ± 3 |
| 2-μm cubes | +21 ± 3 |
| 1- μ m cylinders (AR = 1) | $+22 \pm 3$ |
| 0.5- μ m cylinders (AR = 2) | $+32 \pm 3$ |
| $0.2 - \mu m$ cylinders (AR = 1) | $+42 \pm 3$ |
| 0.15- μ m cylinders (AR = 3) | $+35 \pm 3$ |
| 0.1- μ m cylinders (AR = 3) | +41 ± 3 |