Supporting Information

A novel cell membrane-cloaked magnetic nanogripper with enhanced stability for drug discovery

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Samples preparation for TEM：

1. Use 300 mesh carbon coated grids.

2. Prepare α_{1A}/MNGs solution and ultrasonic dispersed evenly.

3. Place a drop (approx. 20 μL) of α_{1A}/MNGs solution on the grid.

4. Dry overnight in a Petri dish and view the next day in TEM.

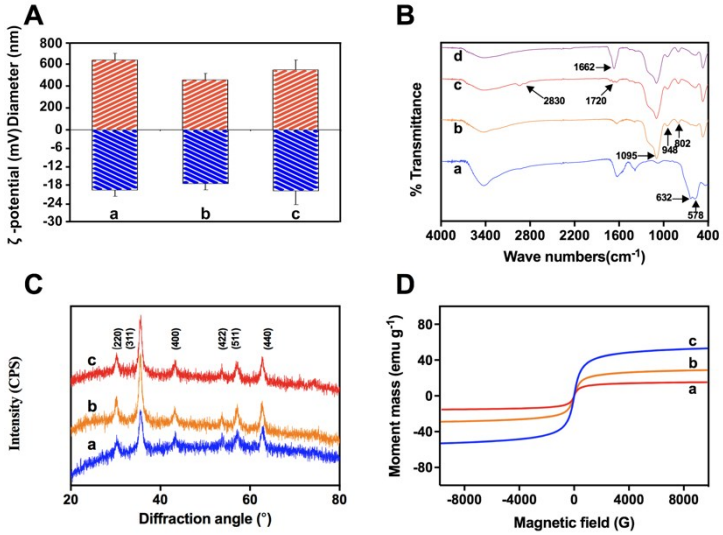

Figure S1. Size and zeta potential results of high α_{1A}-AR expression HEK293 cell membrane-derived vesicles (a), Fe_3O_4-CHO nanoparticles (b) and α_{1A}/MNGs (c) (A); FT-IR spectra of Fe_3O_4 (a), Fe_3O_4-SiO_2 (b), Fe_3O_4-CHO (c) and α_{1A}/MNGs (d) (B); XRD patterns of Fe_3O_4 (a), Fe_3O_4-CHO (b) and α_{1A}/MNGs (c) (C) and VSM curves of α_{1A}/MNGs (a), Fe_3O_4-CHO (b) and Fe_3O_4 (c) (D).

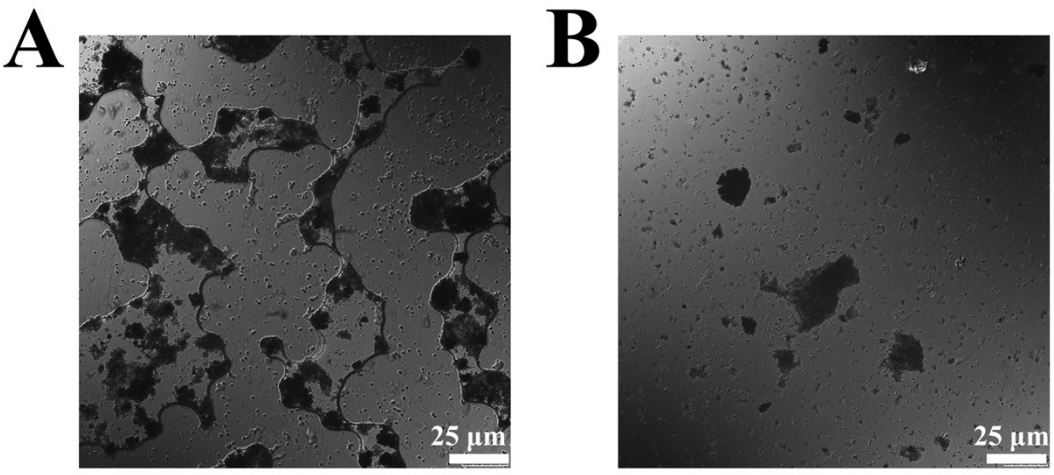

Figure S2. Bright-field images of confocal microscopy images of MNGs cores (A) and α1A/MNGs (B).

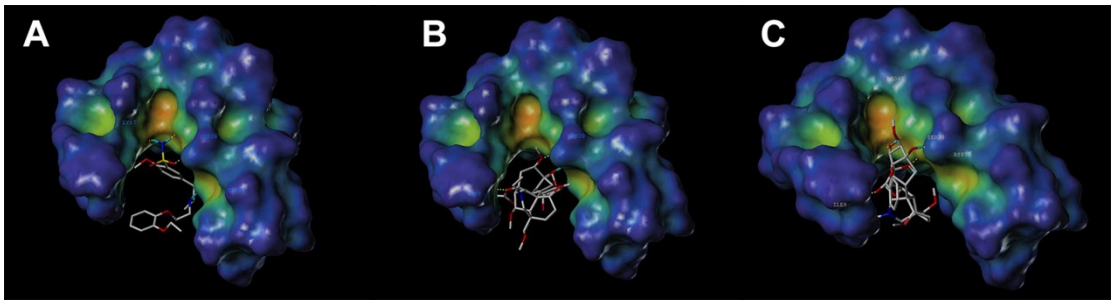

Figure S3. The binding model of compounds tamsulosin (A), bulleyaconitine A (B) and benzoylhypacoitine (C) with α_{1A} AR (PDB ID: 4iye).