Supplementary Information for

Optical Tweezer-Assisted Cell Pairing and Fusion for Somatic Cell

Nuclear Transfer within an Open Microchannel

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1. Fabrication of microchannel (Fig. S1)

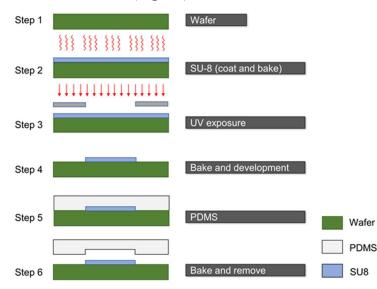


Fig. S1 Fabrication process of PDMS structure. Step 1: prepare a silicon wafer. Step 2: SU-8 photoresist is spin coated on a silicon wafer and prebaked on a hotplate. Step 3: expose the SU8 to UV light with the mask of the upper microchannel on top to make it harden. Step 4: rinse off excess photoresist. Step 5: a mixture of PDMS and curing agent with a ratio of 10:1 is dropped onto the master mold and baked in an oven for 1.5 h. Step 6: peel off the master mold.

2. Oocyte reaches the wrong position (Fig. S2)

The height of PDMS pillars determines whether cells can accurately halt at the target position under the influence of DEP forces. When the height of the microchannel is lower than the radius of the oocyte, the contact point between the oocyte and the PDMS pillars is below the height of the oocyte center. The oocytes easily roll over the PDMS pillars due to the torque generated by the DEP force exerted on the oocytes, as shown in Fig. S2.

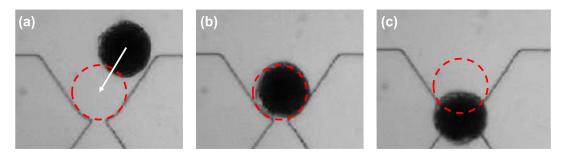


Fig. S2 The oocyte rolls over the PDMS pillars to the wrong position during the transfer process. (a) The initial location of the oocyte before capture. (b) Under the action of DEP force, the oocyte moves towards the narrowest part of the hourglass groove, with the red dashed line indicating the target position of the oocyte. However, due to the height of the PDMS pillars (60 μ m), it is difficult to stop the larger oocytes with a diameter of 130-150 μ m stopping at the target position. (c) The oocyte passes the target position, rolls over the PDMS pillars, and finally stops in the wrong position.