Supporting Information for

Sorting Gold and Sand (Silica) Using Atomic Force Microscope-Based Dielectrophoresis

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Supplementary Figures

Fig. S1 COMSOL simulation $\log (|\mathbf{E}|^2)$ of various phases of electrodes with the gap (a), (b), and (c) and without the gap (d). Only (a) case separate the maximum and the minimum at the edges and the center, respectively. Therefore, we apply the same phase of AC voltage of (a).



Fig. S2 Same simulation with Fig. 2(c),2(d) with the NDEP condition ($f_{CM} = -0.5$). (a) Force of the direction toward the center that is NDEP force. (b) shows particles under NDEP near electrodes go upward by NDEP force, but DEP energy is not bigger than Brownian motion energy near the center.



Fig. S3 Experimental process of DEP. AC voltage is applied while approaching before the formation of a water channel when the distance between the orifice and the substrate is almost 750 nm.



Fig. S4 Amplitude signals of QTF when applying AC voltage onto the quadrupole electrodes after liquid nanochannel was formed. (a) Au nanoparticles, PDEP. (b) Silica nanoparticles, PDEP. (c) Silica nanoparticles, NDEP. (d) Silica and au nanoparticles (selective deposit). In all cases, $3\sim4$ oscillations of the same period (~5 s) occurred. After those oscillations, a liquid channel gradually vanished in (a) Au case, because aggregated Au nanoparticles increase current and generated thermal energy vaporize the meniscus. However, amplitude signals in (b), (c) silica case show solidification behavior caused by connecting electrodes by the aggregated silica nanoparticles such as the inset image in (c). (d) mixed case also shows solidification behavior, but it is not as fast as (b), (c) because of the gold nanoparticles.



Fig. S5 COMSOL simulation of trap energy depending on the width of the gap. In this simulation, the diameter of the pipette is 2.1 μ m, and $V_{peak} = 1.0$ V, 15 MHz was used. When the width of the gap is 0.01 μ m or 3.91 μ m, the gradient of trap energy does not exist inside the orifice of the pipette, so DEP force does not affect. Also, when the width of the gap is 0.61 μ m, PDEP spot (potential minimums) is much wider than the NDEP spot (a potential maximum at the center), so the DEP effect does not work properly as shown in Fig. S6(b). Therefore, the ratio of the diameter of the pipette and the gap distance should be near 1.



Fig. S6 SEM images of inappropriate ratio results. Both images do not show the expected result. (a) A smaller pipette case. 100 nm diameter silica nanoparticles and 20 nm diameter gold nanoparticles, 1 μ m diameter pipette and 2 μ m gap, $V_{peak} = 2.5$ V, and 30 MHz frequency are used. (b) A smaller gap case. 50 nm diameter gold nanoparticles, ~ 1.5 μ m diameter pipette, and ~ 0.35 μ m gap, $V_{peak} = 1.0$ V and 15 MHz frequency are used.



Fig. S7 SEM images of no gap cases. Three images do not show a particular regularity. (a) An au nanoparticle solution case. 50 nm diameter gold nanoparticles, 1.5 μ m diameter pipette, $V_{peak} = 1.0$ V and 15 MHz frequency are used. (b) A silica nanoparticle solution case. 100 nm diameter silica nanoparticles, 3 μ m diameter pipette, $V_{peak} = 2.5$ V and 30 MHz frequency are used. (c) A mixed solution case. 100 nm diameter silica nanoparticles, 1.8 μ m diameter pipette, $V_{peak} = 2.5$ V and 30 MHz frequency are used. (c) A mixed solution case. 100 nm diameter silica nanoparticles, 1.8 μ m diameter pipette, $V_{peak} = 2.5$ V and 30 MHz frequency are used.

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Fig. S8 AC Amplitude dependence of PDEP effect of Au nanoparticles. (a, b) SEM images of PDEP experiment with the different amplitude of AC voltage. (a) $V_{peak} = 3.0$ V, 15 MHz frequency (b) $V_{peak} = 5.0$ V 15 MHz frequency (c) Schematic diagram of simulation region of electrothermal effect. We assumed that electrical thermal loss goes to the water in the orange square region. (d) Graph of the particle rate (black squares) and the ETE effect rate (a dashed line) versus amplitude. The particle rate means the number rate of not appropriately deposited particles and the ETE effect rate means the theoretical calculation of the ETE effect versus the DEP effect rate. Typically, it is expected that Higher amplitude makes higher spatial deposit accuracy because of stronger DEP force. However, the opposite results are produced because of the ETE effect.