

Supporting Information for:

**Development of a High-Performance Sliding
Microneedle-Lateral flow immunoassay strip Device
for Ultra-Rapid Point-of-Care Diagnosis**

Soo-bin Yu and Jae Hwan Jung*

Department of Pharmaceutical Engineering, Dankook University, Cheonan, Korea

*** Corresponding Author**

E-mail: jjahwan@dankook.ac.kr

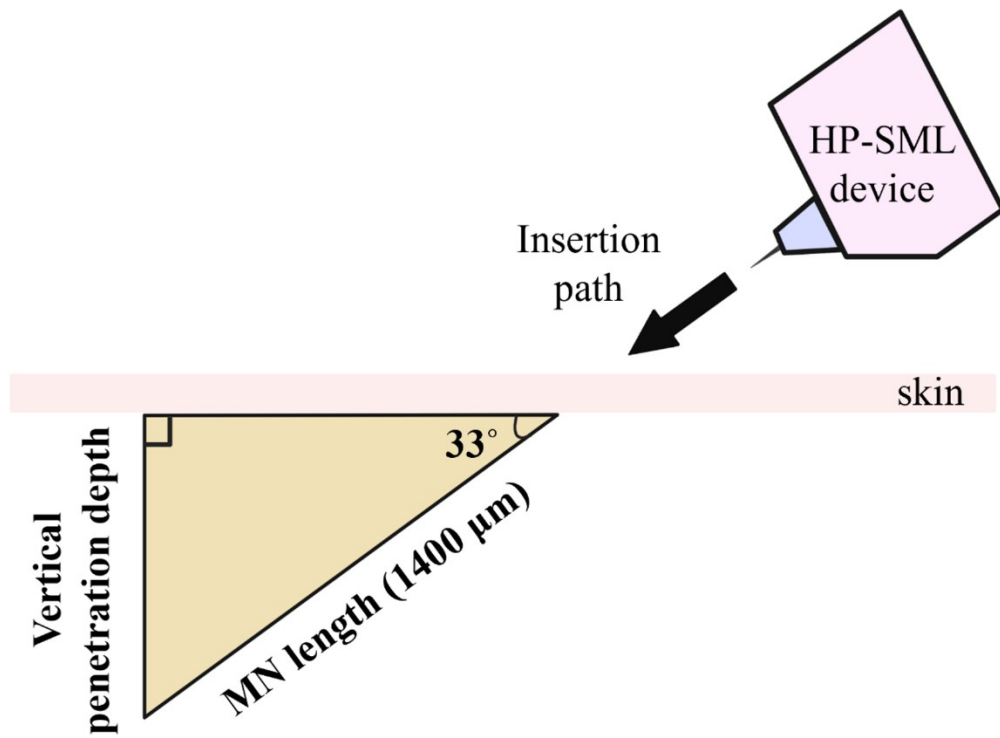


Figure S1. Figure S1. A schematic diagram illustrating microneedle insertion parameters using the HP-SML device. The vertical penetration depth is 750 μm , and the insertion angle is 33° . The schematic was created using BioRender.

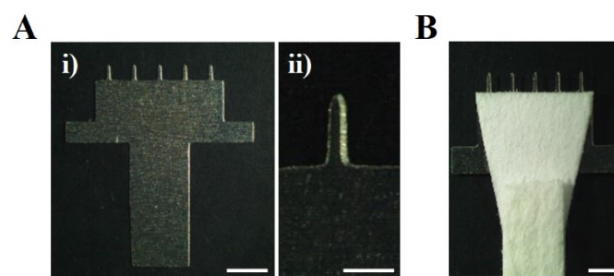


Figure S2. (A) Optical images of a 750 μm stainless Microneedle (MN) array: (i) Overview showing the array (scale bar = 2 mm) and (ii) Magnified view of a single MN (scale bar = 1 mm). (B) Optical images of filter paper-affixed MNs with lengths of 750 μm (scale bar = 1 mm).

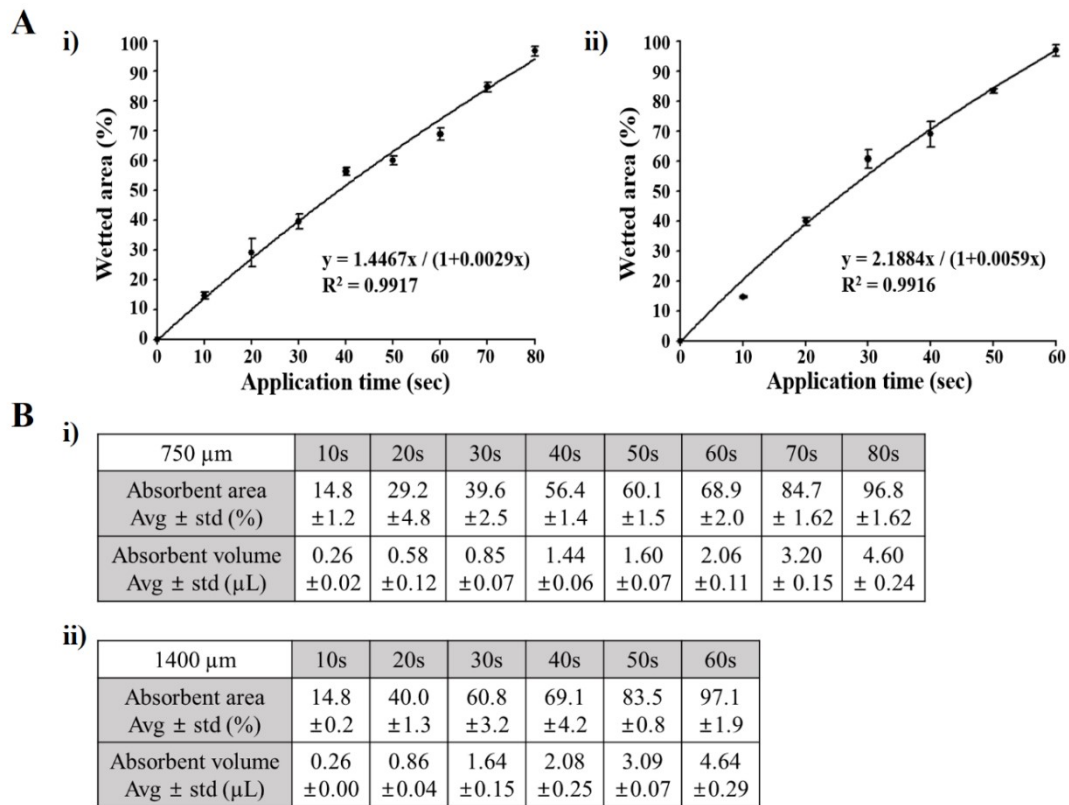


Figure S3. (A) Graphs and trend formulas showing the relationship between the wetted area of the filter paper on the MN and the application time of the HP-SML device incorporating (i) 750 μm MNs and (ii) 1400 μm MNs to SRB-containing agarose gel. (B) Tables presenting the absorption area and absorption volume over time using the HP-SML device incorporating (i) 750 μm MNs and (ii) 1400 μm MNs. The tables display averages ± standard deviations derived from three replicates.

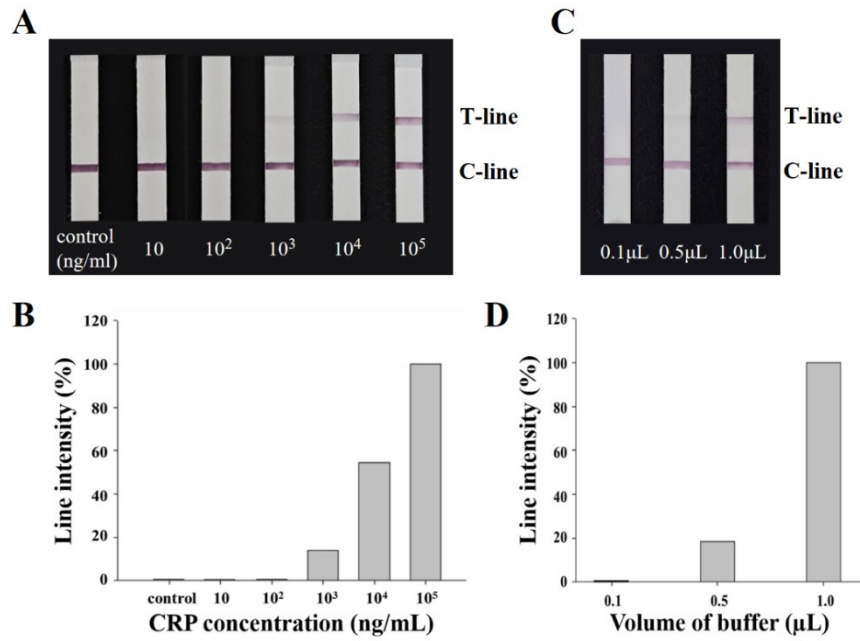


Figure S4. (A) Optical images of LFAs showing CRP detection with 1 μ L CRP solution and 99 μ L running buffer using HP-SML device. (B) The detection limit of LFAs illustrated by the intensity of the test line (A), analyzed using ImageJ software. (C) Optical images of LFAs showing CRP detection at a concentration of 10³ ng/mL with different CRP sample volumes of 0.1, 0.5, and 1 μ L (the remaining volume, excluding the CRP solution, up to 100 μ L was used as a running buffer). (D) The detection limit of the LFA illustrated by the intensity of the test line (C), analyzed using ImageJ software.

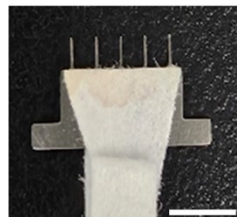


Figure S5. Image of ISF extraction by MNs after applying the HP-SML device to rat skin for 35 seconds. (scale bar = 2 mm).