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Supplementary Information

Polymer Microneedles with Interconnected Porous Structure via Phase Inversion Route for Transdermal Medical Applications

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Supplementary Figures:



Fig. S1 Characterization of porous microneedles (MNs) made of PES ($a_1 \sim a_5$), PLGA ($b_1 \sim b_5$), PLA ($c_1 \sim c_5$), and PVDF ($d_1 \sim d_5$), respectively. Optical microscopy images of the MN arrays ($a_1 \sim d_1$). Surface ($a_2 \sim d_2$ and $a_3 \sim d_3$) and cross-sectional ($a_4 \sim d_4$ and $a_5 \sim d_5$) SEM images of PES, PLGA, PLA and PVDF MNs with different magnifications. The optical microscopy images show that MNs with good tapered structure and sharp needles could be prepared. Surface and cross-sectional scanning electron microscope (SEM) images of single needle demonstrated that porous structure with interconnected pores is obtained.



Fig. S2 Cross-sectional SEM images of CA MNs made of 35 wt % CA solution at different position of the MNs. Inset in (a) shows the schematic illustration of the cross-sectional position of the MNs in (b-d). It can be observed that the microstructures throughout the needles are consistent.



Fig. S3 Characterization of porous MNs made of CA $(a_1 \sim a_4)$, PES $(b_1 \sim b_4)$, PLGA $(c_1 \sim c_4)$, PLA $(d_1 \sim d_4)$, PVDF $(e_1 \sim e_4)$, and PSF $(f_1 \sim f_4)$, respectively. Surface $(a_1 \sim f_1 \text{ and } a_2 \sim f_2)$ and cross-sectional $(a_3 \sim f_3 \text{ and } a_4 \sim f_4)$ SEM images of the base of CA, PES, PLGA, PLA, PVDF and PSF MNs with different magnifications.



Fig. S4 Optical microscopy image of CA MN after 1 year. The morphology of CA MN remained unchanged, indicating the good stability of the CA MN.



Fig. S5 Cross-sectional SEM images with varying magnifications of porous CA MNs made of different initial concentrations: 10 wt % ($a_1 \sim a_3$), 15 wt % ($b_1 \sim b_3$), 20 wt % ($c_1 \sim c_3$), 25 wt % ($d_1 \sim d_3$), 30 wt % ($e_1 \sim e_3$), 35 wt % ($f_1 \sim f_3$), respectively.



Fig. S6 The water contact angle of a smooth CA film obtained by spin-coating.



Fig. S7 Snapshots shows the water spreading process on the porous CA MN over time. Water droplet with the size of 2 μ L spread over the surface within 65 s, indicating that water can be absorbed by porous CA rapidly.



Fig. S8 Confocal laser scanning microscopy (CLSM) images of porcine cadaver skin after administration of sodium fluoresceinloaded CA MN made of 35 wt % CA solution for 10 min. The scale bars can be applied to the other images. The CLSM images show that CA MN could penetrate the stratum corneum and reach the dermis.



Fig. S9 The cytotoxicity study of the MNs made of different polymer materials. The MNs were incubated with NIH-3T3 fibroblasts for 24 h.



Fig. S10 The ISF extraction amount of CA MNs over time in mice. The increase of mass indicates the diffusion of ISF from the skin to MNs.