

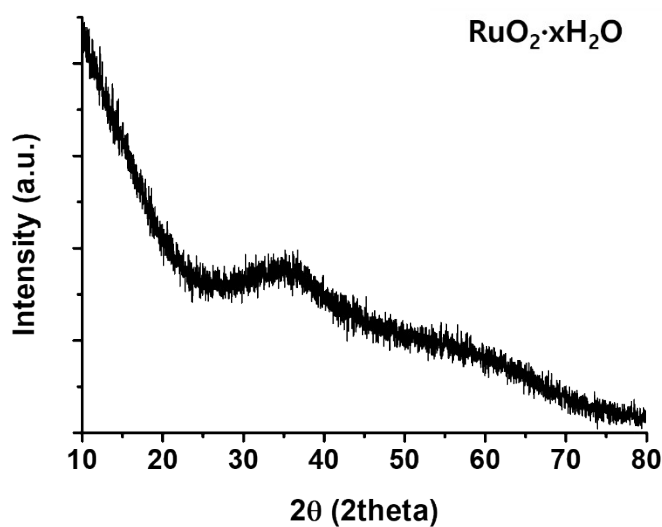
## Supplementary Information

### Monodisperse RuO<sub>2</sub> nanoparticles for highly transparent and rapidly responsive supercapacitor electrodes

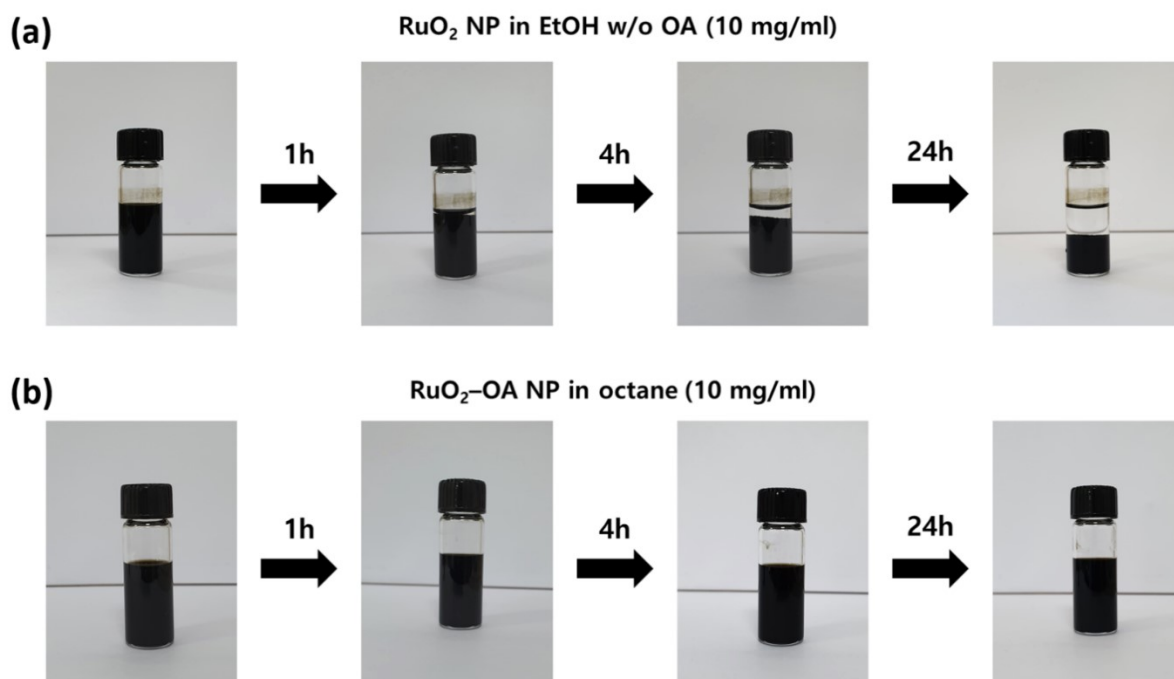
*Ilhwan Ryu<sup>†</sup>, Dongju Kim<sup>†</sup>, Guenpyo Choe, Sohyun Jin, Dajung Hong\* and Sanggyu Yim\**

Department of Chemistry, Kookmin University, Seoul 02707, South Korea

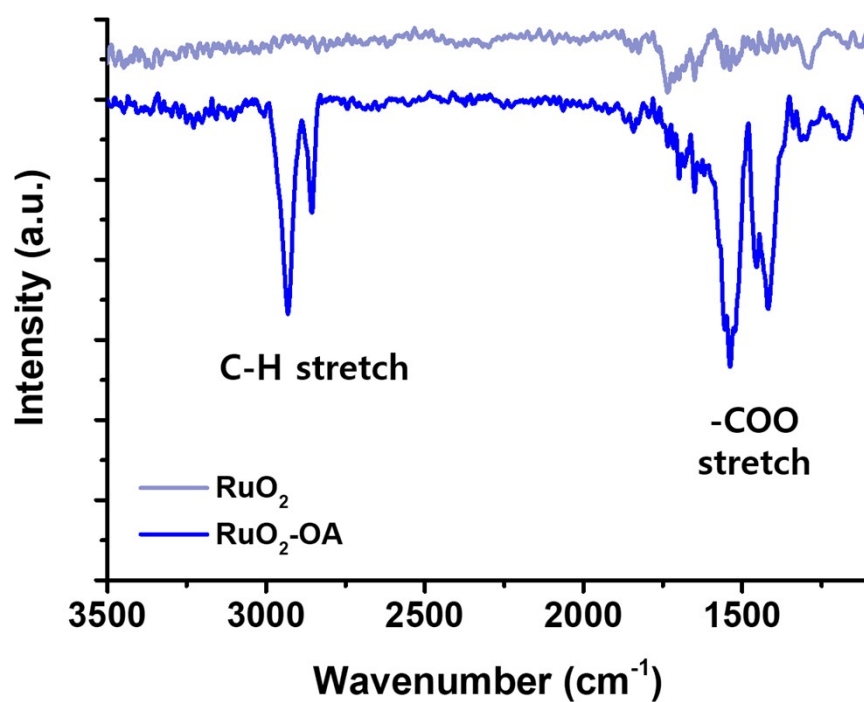
\* Corresponding authors: Dajung Hong (solar@kookmin.ac.kr), Sanggyu Yim (sgyim@kookmin.ac.kr)



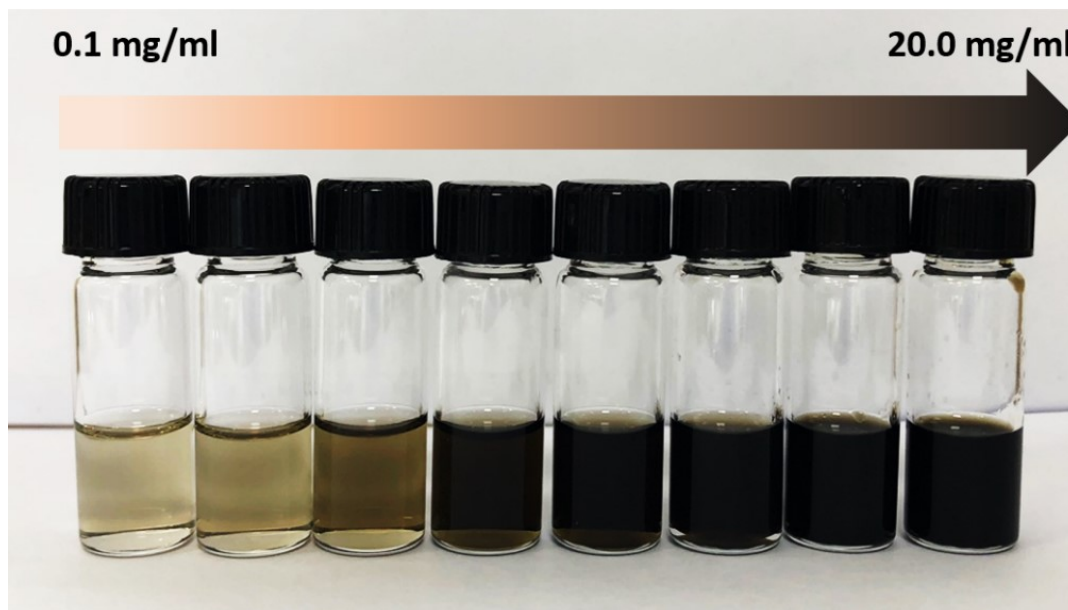
**Figure S1.** Powder X-ray diffraction pattern of the synthesized pristine RuO<sub>2</sub> NPs.



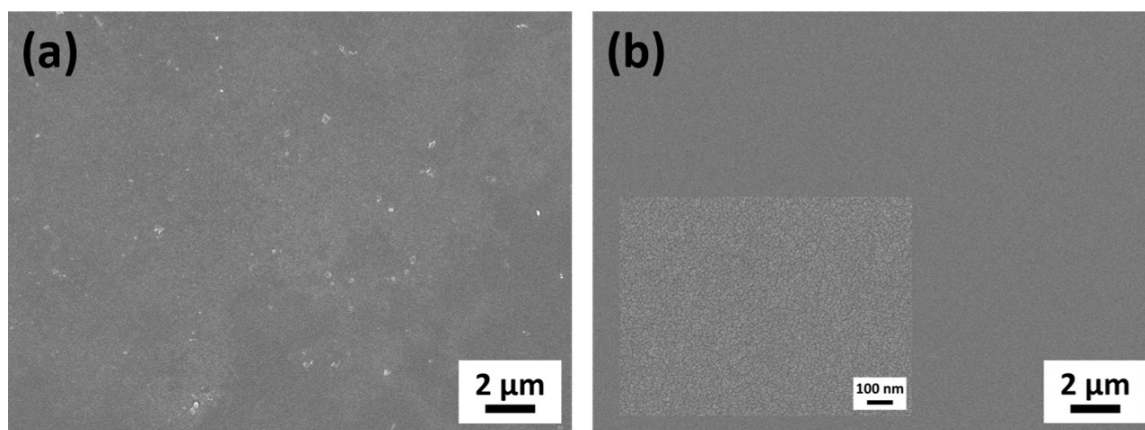
**Figure S2.** Photographs of (a) pristine RuO<sub>2</sub> NPs dispersed in ethanol and (b) OA-decorated RuO<sub>2</sub> NPs dispersed in octane and taken at various times after dissolution.



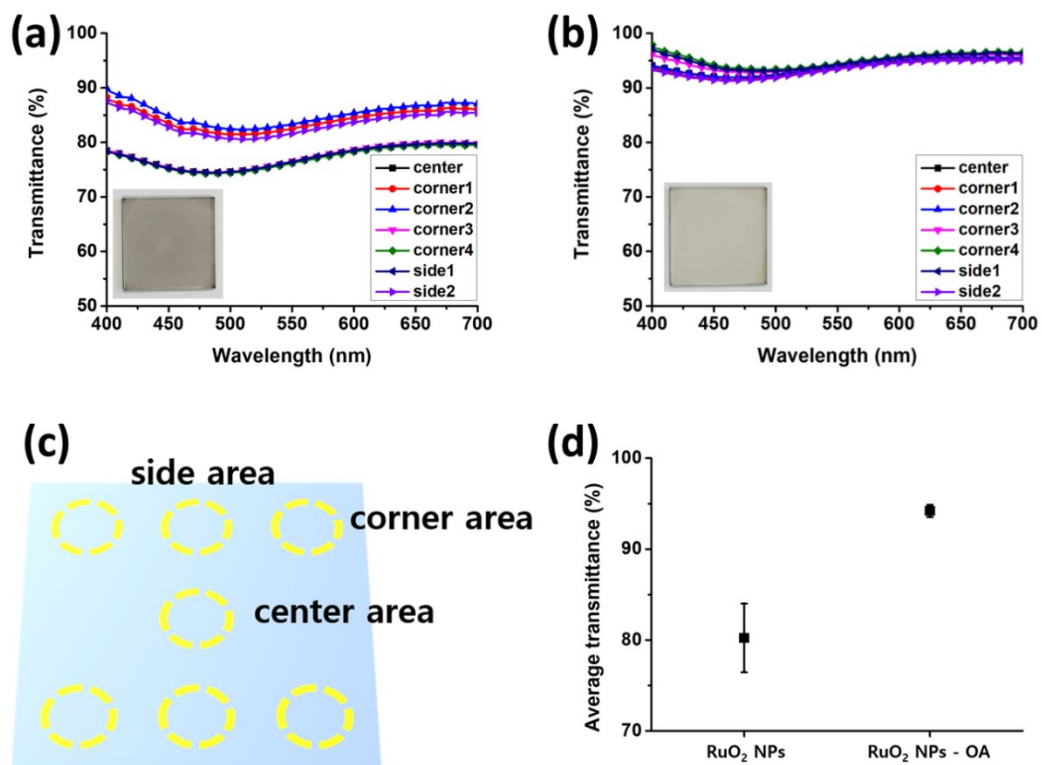
**Figure S3.** FTIR spectra of the pristine and OA-decorated RuO<sub>2</sub> NP-based electrodes.



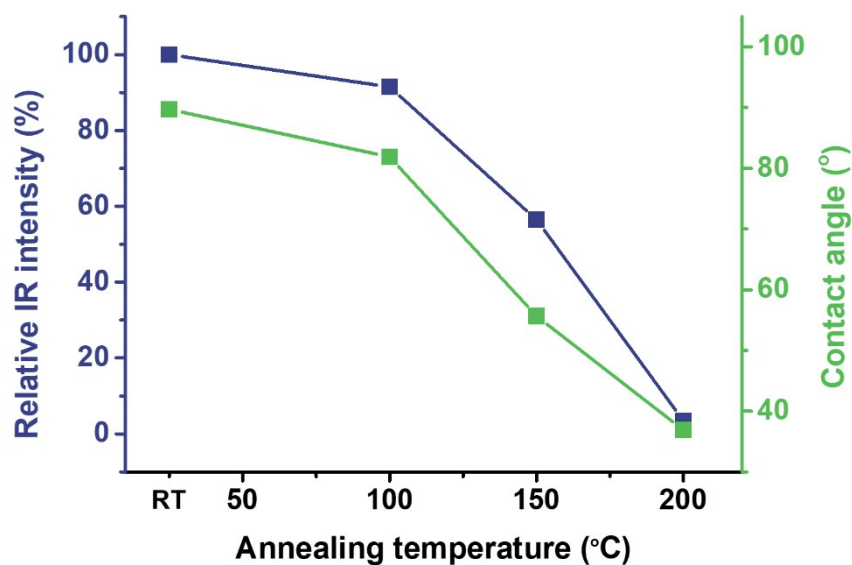
**Figure S4.** Photograph of OA-decorated RuO<sub>2</sub> NPs of various concentrations (from left to right: 0.1, 0.2, 0.5, 1.0, 2.0, 5.0, 10.0, and 20.0 mg/ml) dispersed in octane. Taken after 24 h of dissolution.



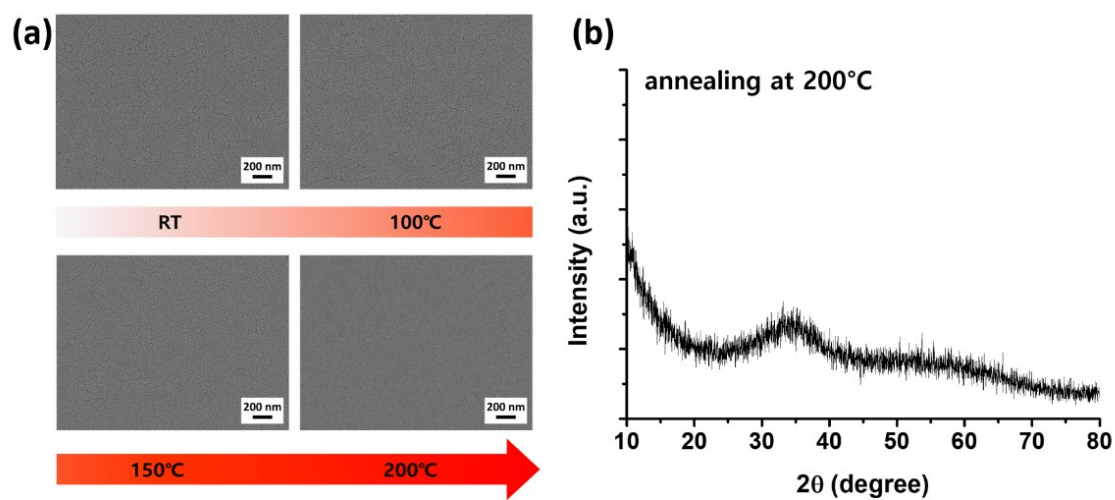
**Figure S5.** Surface FE-SEM images of thin films spin-coated with 10 mg/ml solution of (a) pristine RuO<sub>2</sub> NPs dispersed in ethanol and (b) OA-decorated RuO<sub>2</sub> NPs dispersed in octane.



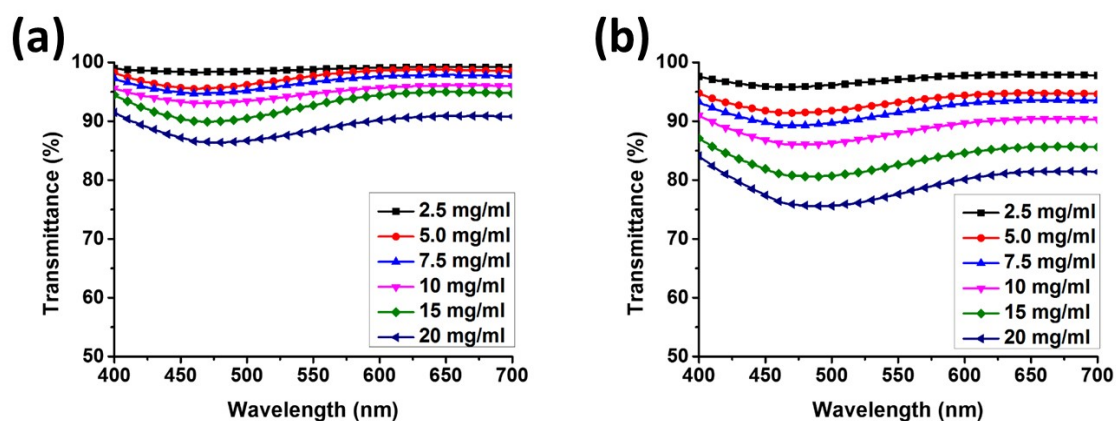
**Figure S6.** Transmittance spectra of thin films spin-coated with 10 mg/ml solution of (a) pristine RuO<sub>2</sub> NPs dispersed in ethanol and (b) OA-decorated RuO<sub>2</sub> NPs dispersed in octane. (c) Schematic illustrating measurement points on the films. (d) Mean and spread of the average transmittance over the visible wavelength range for pristine RuO<sub>2</sub> and RuO<sub>2</sub>-OA NP-coated films.



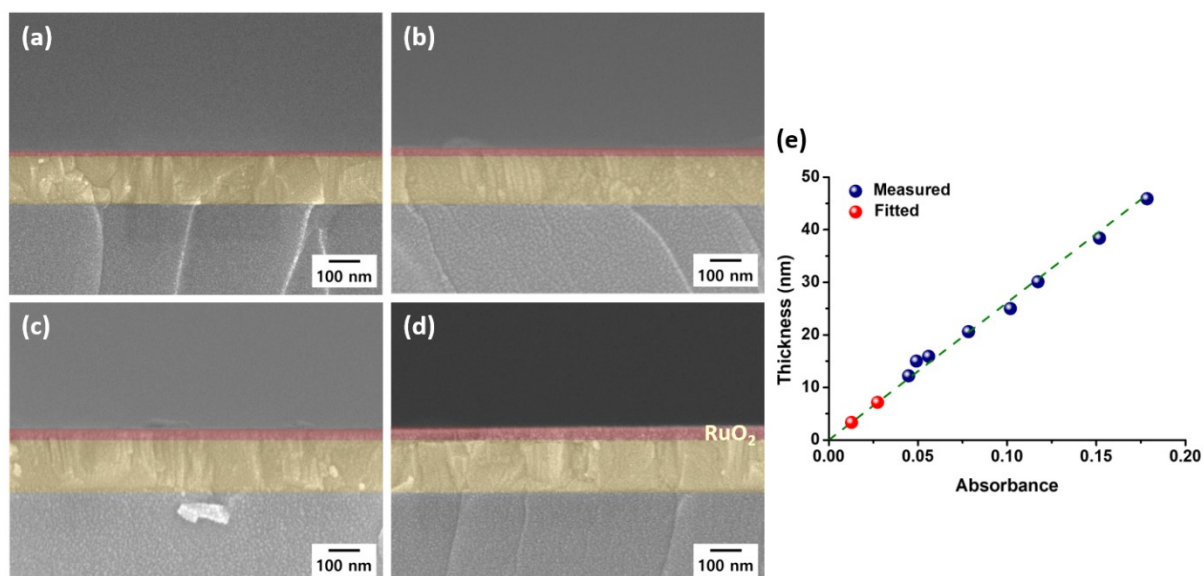
**Figure S7.** Plots of C–H stretching band intensity in FT-IR and water contact angle of the RuO<sub>2</sub>–OA NP-coated films as a function of the annealing temperature.



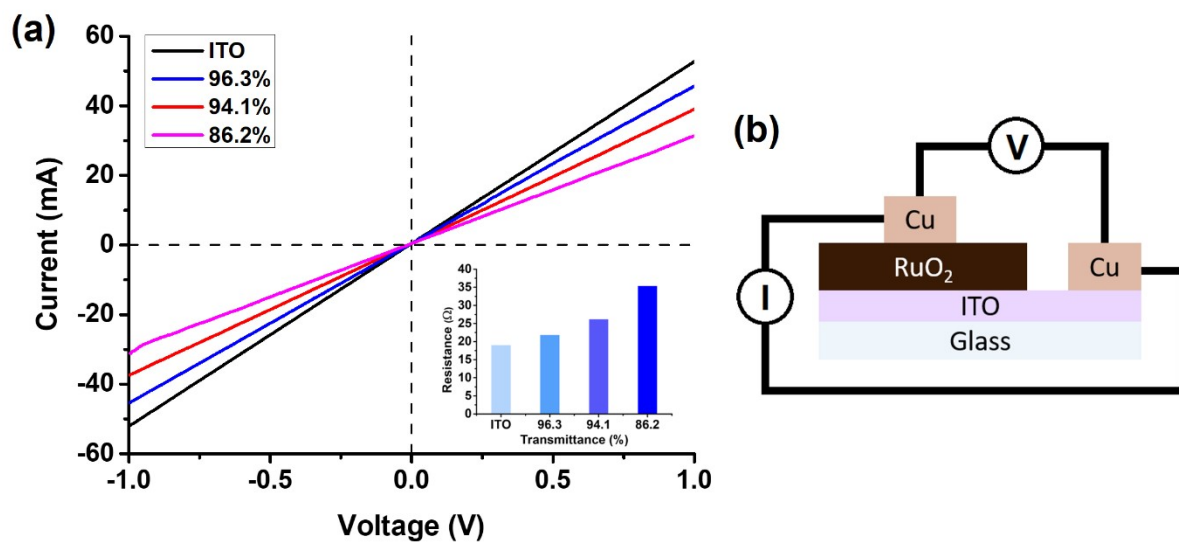
**Figure S8.** (a) Surface FE-SEM images of RuO<sub>2</sub>–OA NP-coated films after annealing at various temperatures for 1 h. (b) Powder X-ray diffraction pattern of the RuO<sub>2</sub>–OA NP-coated films after annealing at 200°C for 1 h.



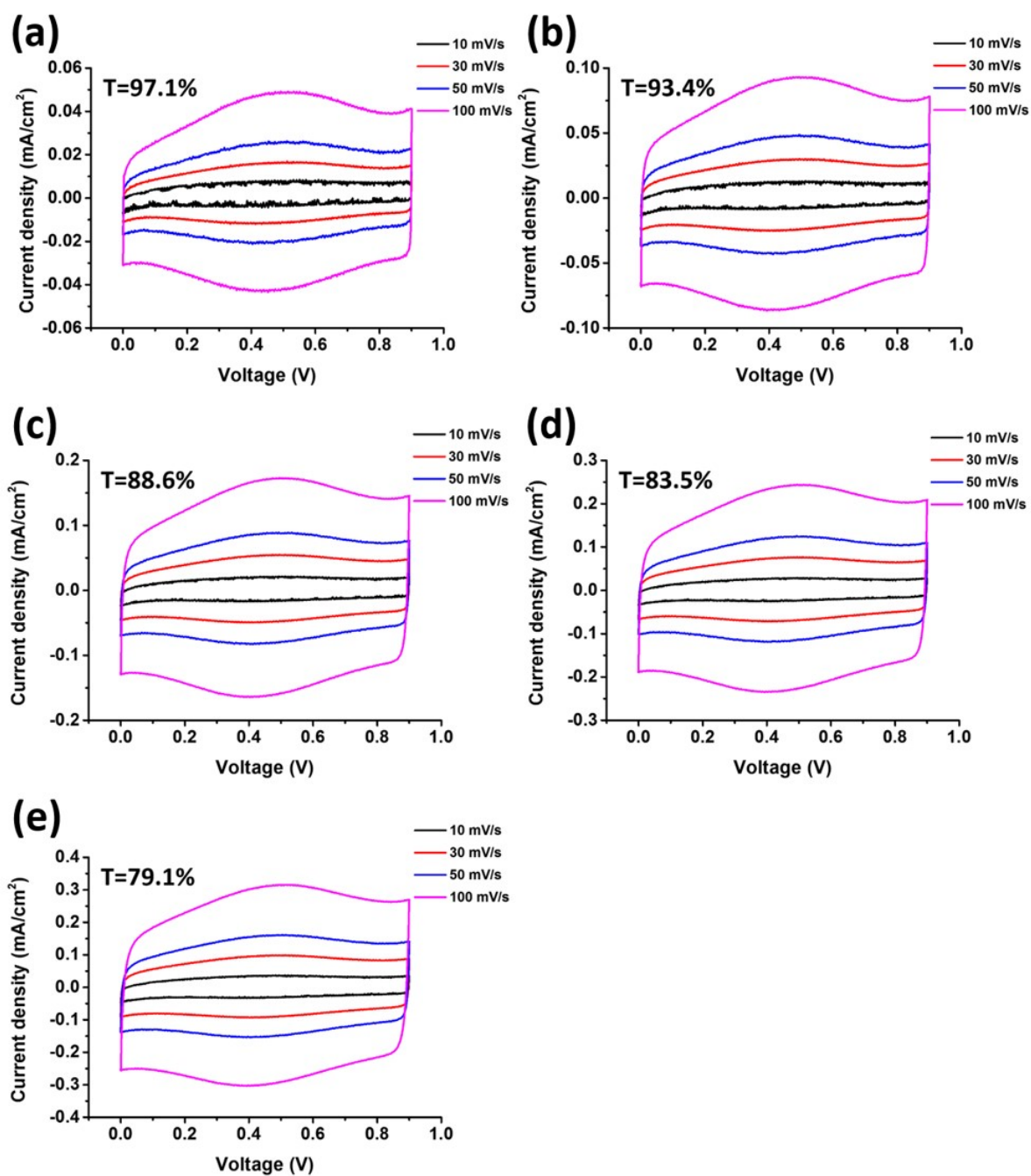
**Figure S9.** Transmittance spectra of RuO<sub>2</sub> NP-based electrodes prepared by (a) single and (b) double deposition of various NP concentration solutions.



**Figure S10.** Representative cross-sectional FE-SEM images of RuO<sub>2</sub> NP-based electrodes. The RuO<sub>2</sub> film thicknesses were (a) 15.0, (b) 22.3, (c) 38.4 and (d) 44.5 nm. (e) Plot of measured thicknesses as a function of optical absorbance. The thicknesses of the thinnest two films were determined using their absorbance and the fitting line from the measured data.

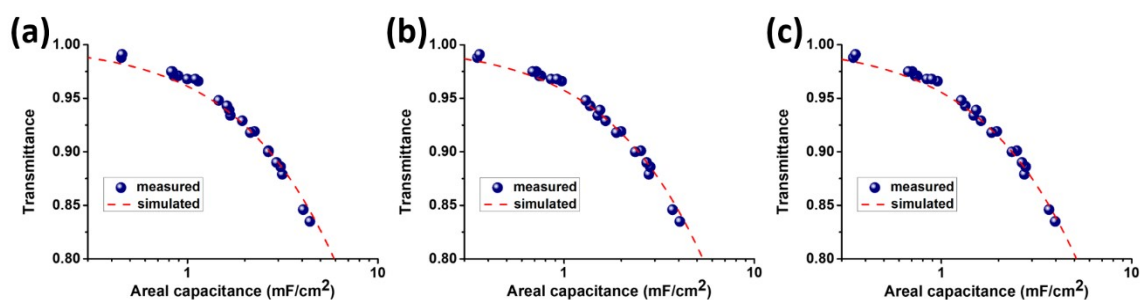


**Figure S11.** (a)  $I$ - $V$  characteristics of RuO<sub>2</sub> NP-based electrodes with various optical transmittances ( $T_{\text{vis}}$ ). (b) A schematic illustration of the  $I$ - $V$  measurement system.

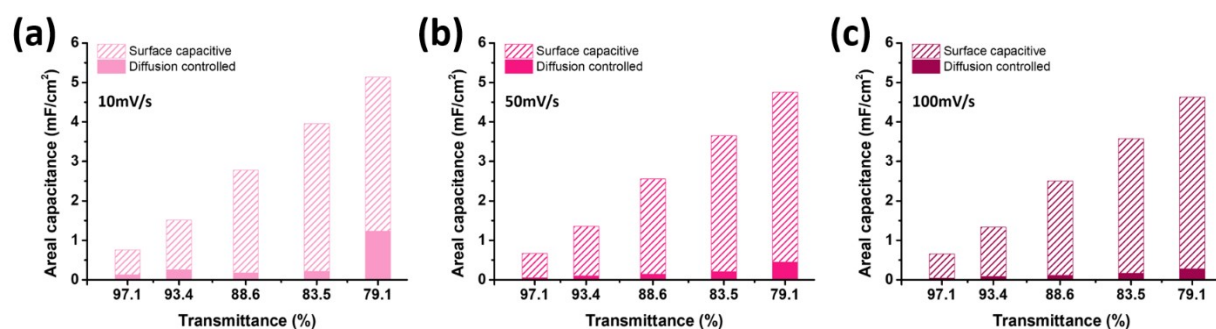


**Figure S12.** Cyclic voltammograms of RuO<sub>2</sub> NP-based transparent electrodes with  $T_{vis}$  of (a) 97.1%, (b) 93.4%, (c) 88.6%, (d) 83.5%, and (e) 79.1% measured at various scan rates.

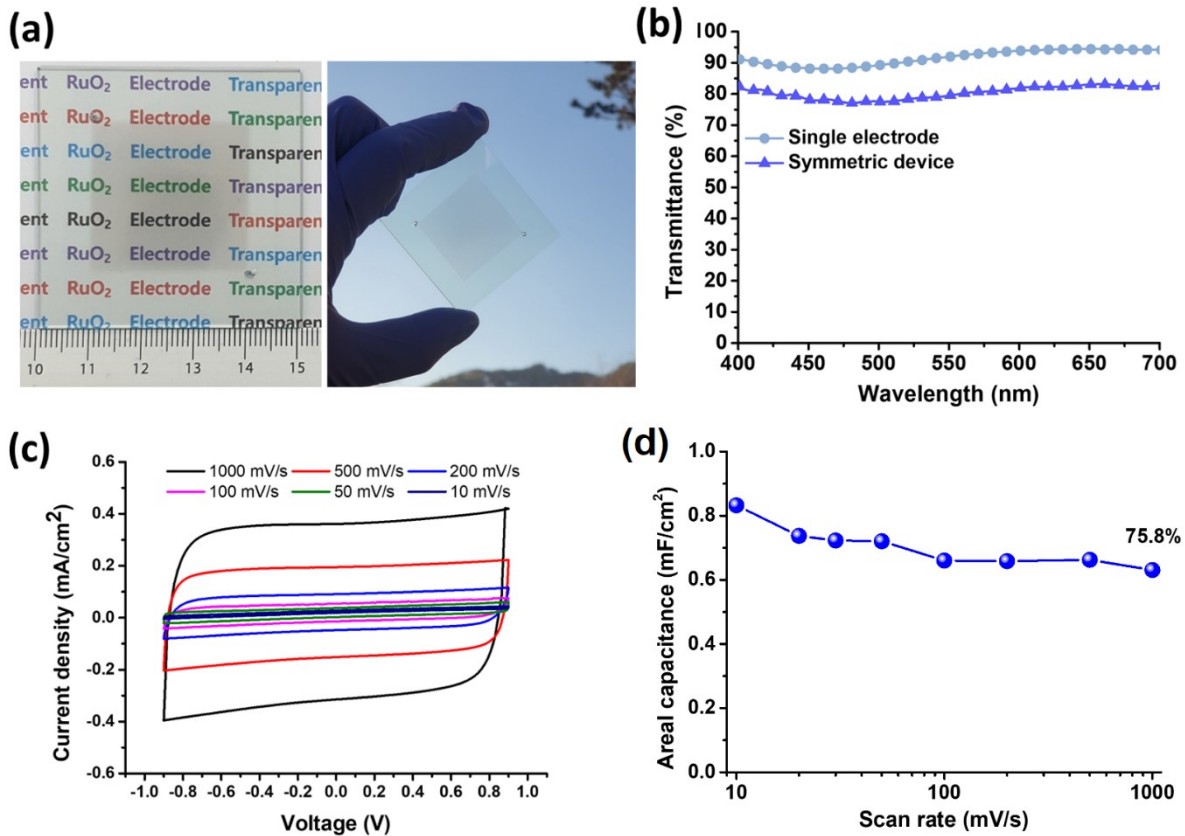




**Figure S13.** Plots of optical transmittance obtained from experiments (blue points) and simulations using Eq. 2 (red dashed lines) as a function of the areal capacitance at scan rates of (a) 10, (b) 50, and (c) 100 mV/s.



**Figure S14.** Deconvolution of areal capacitances measured at scan rates of (a) 10, (b) 50, and (c) 100 mV/s for RuO<sub>2</sub> NP-based electrodes with various transmittances. The shaded and solid regions represent surface capacitive and diffusion-controlled insertion elements, respectively.



**Figure S15.** (a) Photographs of the large-area full-cell device fabricated by spray-coating of RuO<sub>2</sub> NPs. (b) Transmittance spectra of a single electrode and symmetric device. (c) Cyclic voltammograms of the device measured at various scan rates. (d) Capacitance retention as a function of the scan rate.

**Table S1.** Summary of the relative C–H stretching band intensity in FT-IR and the water contact angle of RuO<sub>2</sub>–OA NP-coated films after annealing at various temperatures for 1 h.

Annealing temperature (°C)	Relative C-H stretching intensity in FT-IR (%)	Water contact angle (°)
RT	100	89.7
100	91.5	81.9
150	56.5	55.7
200	3.4	36.9

**Table S2.** Comparison of the optical transmittance and areal capacitance of transparent supercapacitor electrodes and devices in published literature.

Active material	Transmittance (%)	Areal capacitance (mF/cm <sup>2</sup> )	Reference
Graphene QD*	93.0	0.009	Nano Energy <u>26</u> (2016) 746
Graphene*	75.0	0.101	Chem. Mater. <u>27</u> (2017) 3621
SWCNT	92.0	0.552	Nanotech. <u>27</u> (2016) 235403
Carbon nanocup*	71.0	1.22	Sci. Rep. <u>2</u> (2012) 773
PEDOT:PSS*	65.0	1.18	J. Mater. Chem. A <u>4</u> (2016) 10493
Au/MnO <sub>2</sub> network	76.0	3.23	Small <u>13</u> (2017) 1701906
RuO <sub>2</sub> /N-doped graphene*	34.1	1.57	Sustain. Energy Fuels <u>2</u> (2018) 1799
RuO <sub>2</sub> /PEDOT	93.0	1.20	Nano Energy <u>28</u> (2016) 495
<b>RuO<sub>2</sub> NP</b>	<b>97.9</b>	<b>0.85</b>	<b>This work</b>
	<b>93.4</b>	<b>1.66</b>	

\*Data for devices. The others are data for electrodes