

Supporting information (SI)

TiO₂ sol-embedded in electroless Ni-P coating: A novel approach for ultra-sensitive sorbitol sensor

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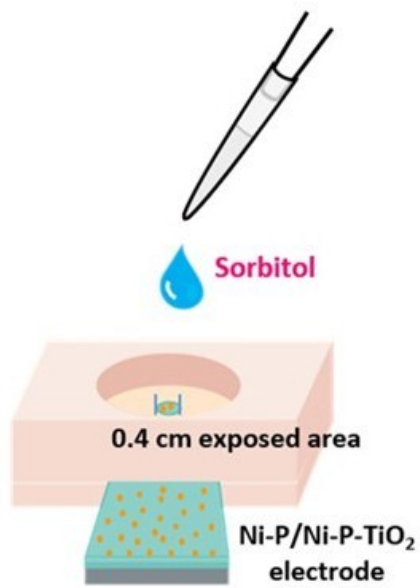


Fig. S1 The configuration of an electrochemical cell used in this work.

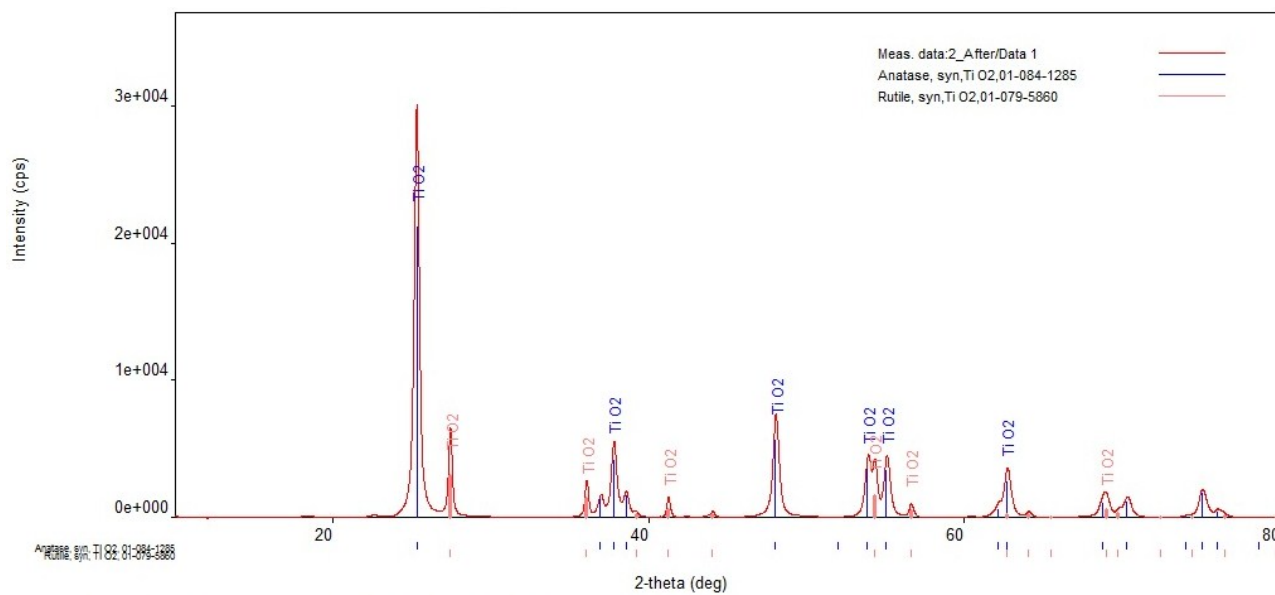


Fig. S2 An XRD spectrum of white TiO₂ powder after calcination at 600 °C for 1 hour (Rigaku, SmartLab, scan rate 10-80 degree, speed 1 degree/min, step 0.01 degree).

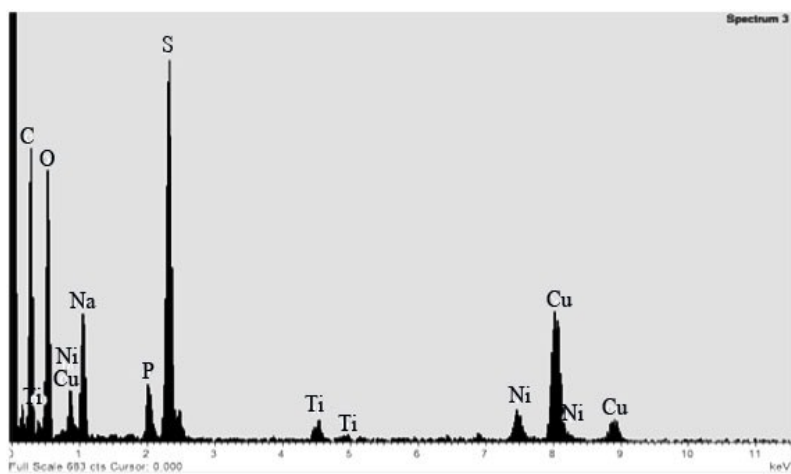
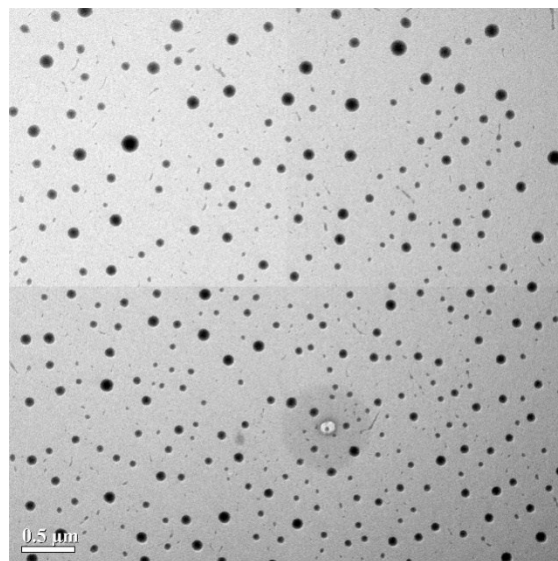
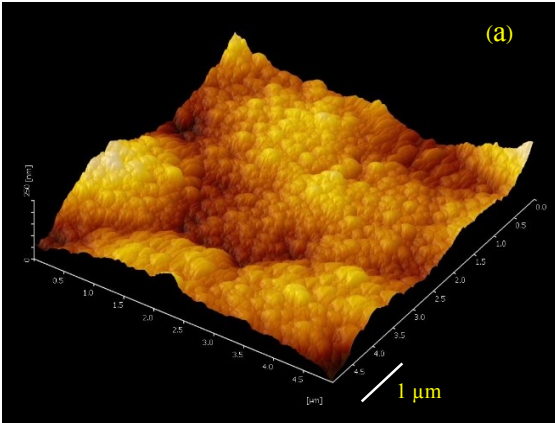
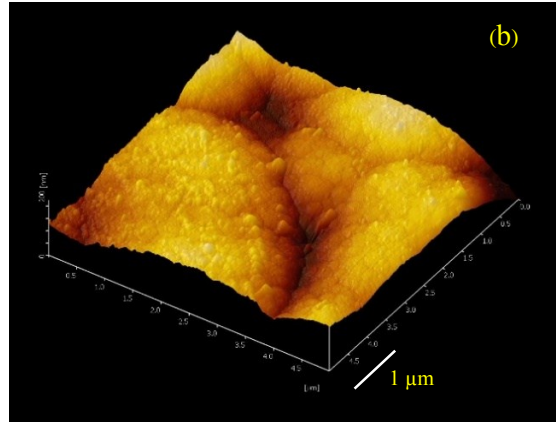


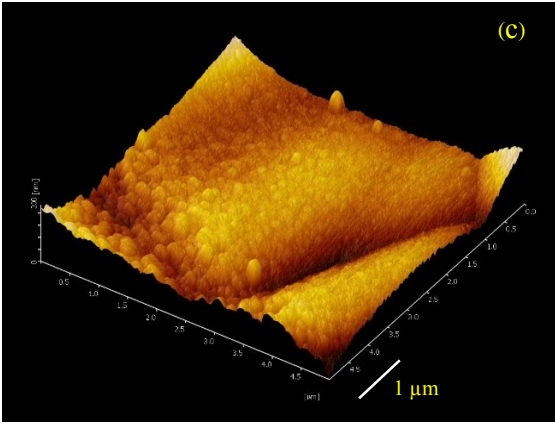
Fig. S3 SEM images of TiO_2 sol in Ni-P electroless bath.



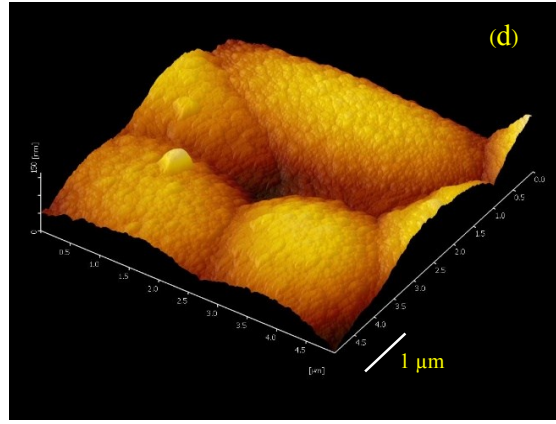
Ra = 22.52 nm
 $S = 2.574 \times 10^7 \text{ nm}^2$



Ra = 22.59 nm
 $S = 2.544 \times 10^7 \text{ nm}^2$



Ra = 19.02 nm
 $S = 2.646 \times 10^7 \text{ nm}^2$



Ra = 19.96 nm
 $S = 2.522 \times 10^7 \text{ nm}^2$

Fig. S4 AFM images indicating the surface area (S) of (a) Ni-P-TiO₂ (2 g/L of TiO₂) coating, (b) Ni-P/Ni-P (0 g/L of TiO₂) coating, (c) Ni-P/Ni-P-TiO₂ (2 g/L of TiO₂) coating and (d) Ni-P/Ni-P-TiO₂ (4 g/L of TiO₂) coating. (SPA-400 atomic force microscope (Seiko Instruments, Inc., Japan), using non-contact mode).

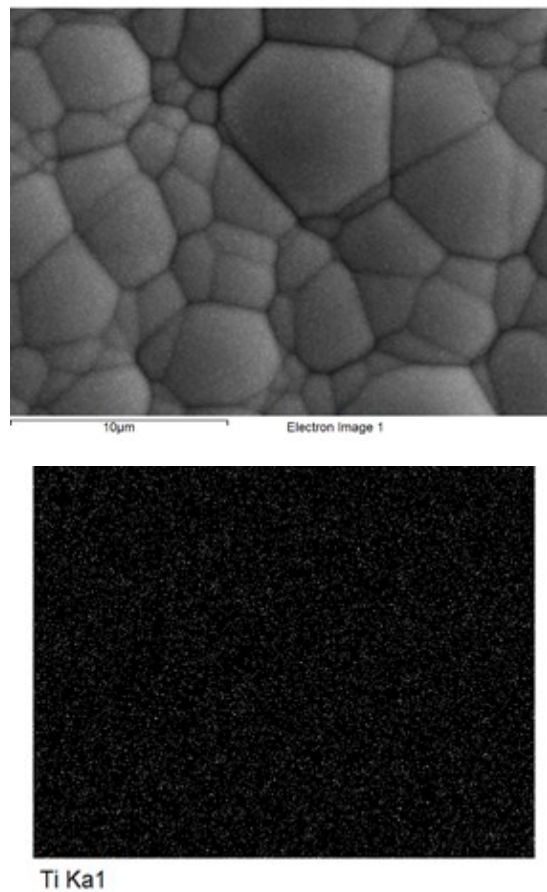


Fig. S5 An SEM image of Ni-P-TiO₂ (top) and the surface mapping indicates the distribution of Ti on the coated surface (bottom).

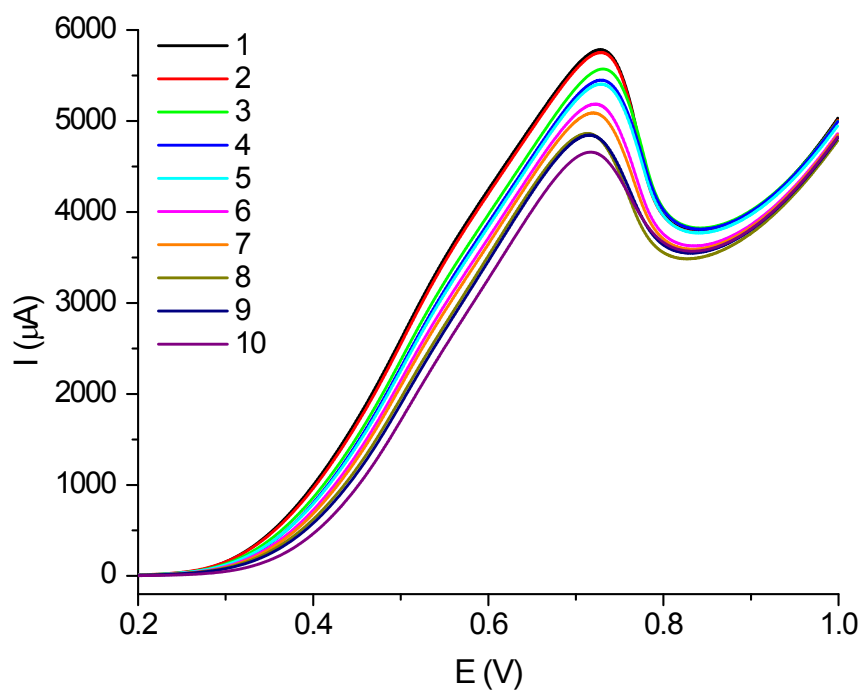


Fig. S6 Reproducibility of Ni-P/Ni-P-TiO₂ electrode for 10 consecutive detection of sorbitol.

Table S1 Stability of electrode for the detection of different compounds after storage for 7 days.

Electrode	% of current signal compared to an original current signal				
	Methanol	Ethanol	Isopropanol	Sorbitol	Glucose
Ni-P	85.1 ± 13	84.4 ± 25	83.0 ± 11	85.0 ± 16	83.2 ± 12
Ni-P-TiO ₂ (2 g/L of TiO ₂)	89.0 ± 3.9	90.1 ± 2.6	89.0 ± 3.5	91.0 ± 4.1	83.3 ± 1.9
Ni-P/Ni-P-TiO ₂ (2 g/L of TiO ₂)	91.5 ± 1.4	93.7 ± 1.2	93.6 ± 1.6	96.4 ± 1.1	89.5 ± 3.4