Electronic Supporting Information

UOx@HMnO₂ Biozyme-Nanozyme Driven Electrochemical Platform for Specific Uric Acid Bioassays

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Fig S1. TEM image of SiO_2 nanoparticles and SiO_2@MnO_2 nanoparticles.



Fig S2. (a) Size distributions and (b) Zeta potential of HMnO₂ and UOx@HMnO₂.



Fig S3. EDS spectrum of UOx@HMnO2 by STEM.



Fig S4. Photographs for the chromogenic reaction of TMB catalyzed by the UOx@HMnO_2.



Fig. S5. (a) Comparison of the TMB color development of $UOx@HMnO_2$ with equal amounts of free UOx. (b) UV absorption of $UOx@HMnO_2$ vs. equal amounts of free uric acid oxidase at OD650.



Fig S6. Relationship between UA concentration and current.



Fig. S7. UOx@HMnO₂-GCE was used for the long-term detection of uric acid stability, and the vertical coordinate was calculated using the CV value on day 1 as a reference.



Fig S8. Data points and standard curves for high performance liquid chromatography-tandem mass spectrometry (UHPLC-MS/MS) tests.

Before amount (µM)	Add amount (μM)	Found amount (μM)	Recovery (%)
32.80	10	41.09	96.00
34.22	20	66.32	103.27
33.24	40	74.87	102.22

Table S1. Recoveries of UA in human urine based on the UOx@HMnO₂-GCE.