



PAPER

Oxidase-mimicking activity of perovskite $\text{LaMnO}_{3+\delta}$ nanofibers and their application for colorimetric sensing

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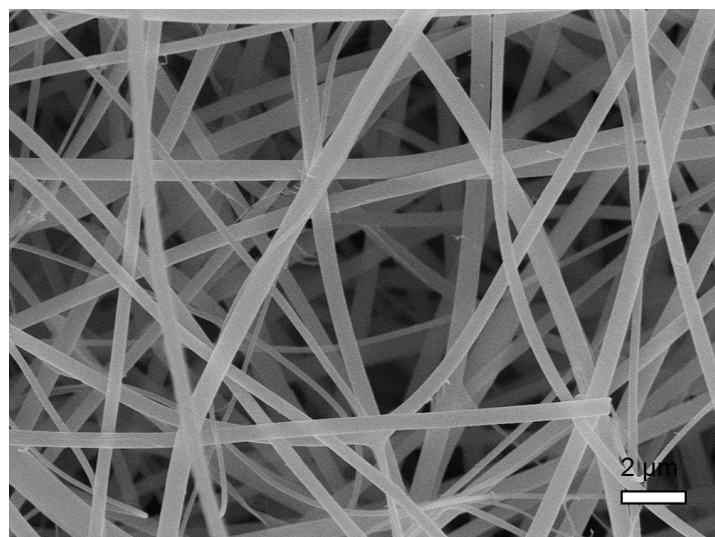


Fig. S1. SEM image of the electrospun PVP/La(NO₃)₃/Mn(Ac)₂ nanofibers.

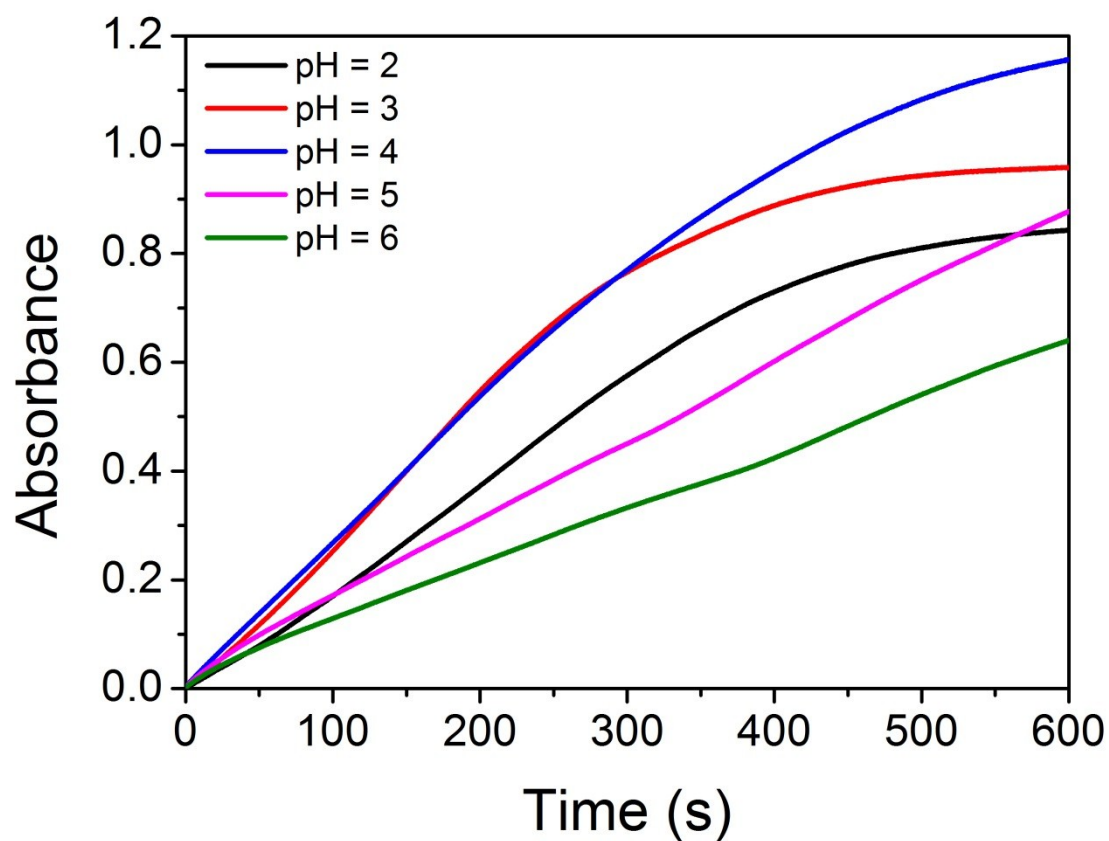


Fig. S2. The oxidase-like catalytic activity of the catalyst in acetate buffer solution with diverse pH values.

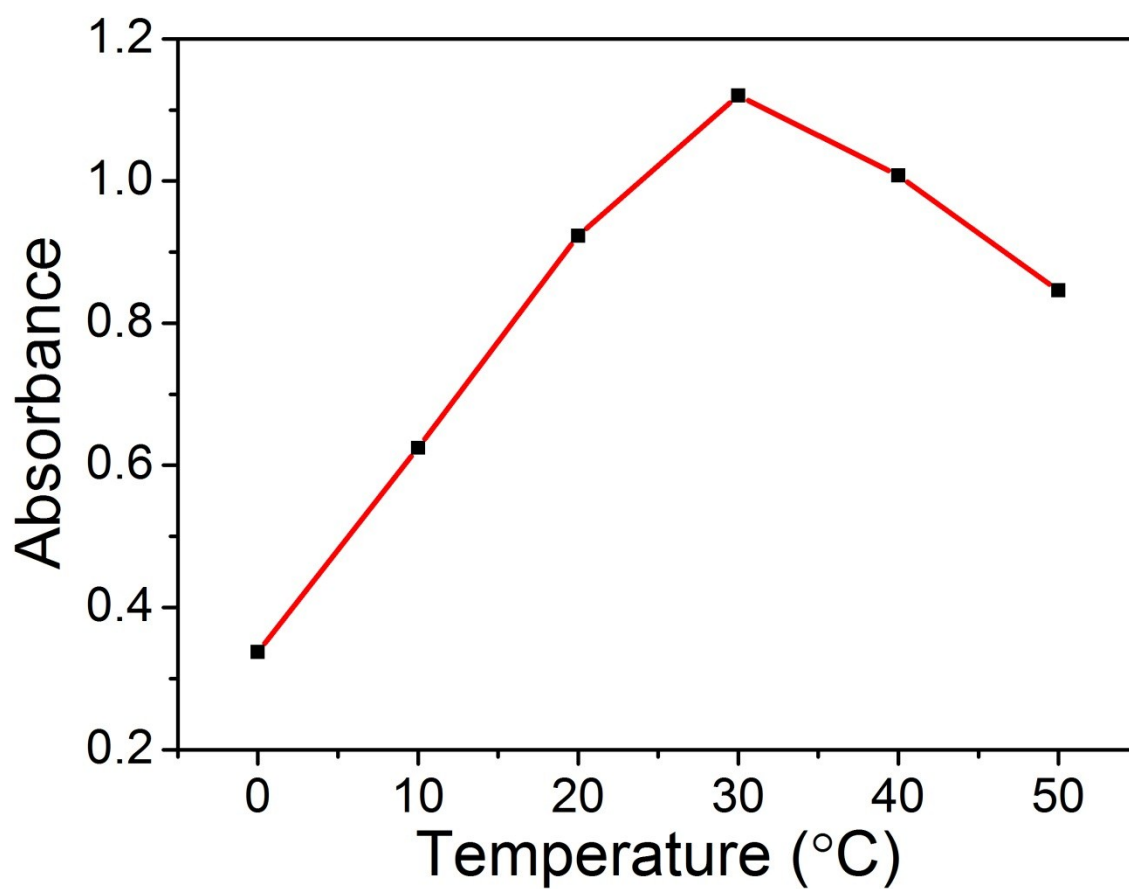


Fig. S3. Dependence of the oxidase-like activity of perovskite $\text{LaMnO}_{3+\delta}$ nanofibers on the temperature.

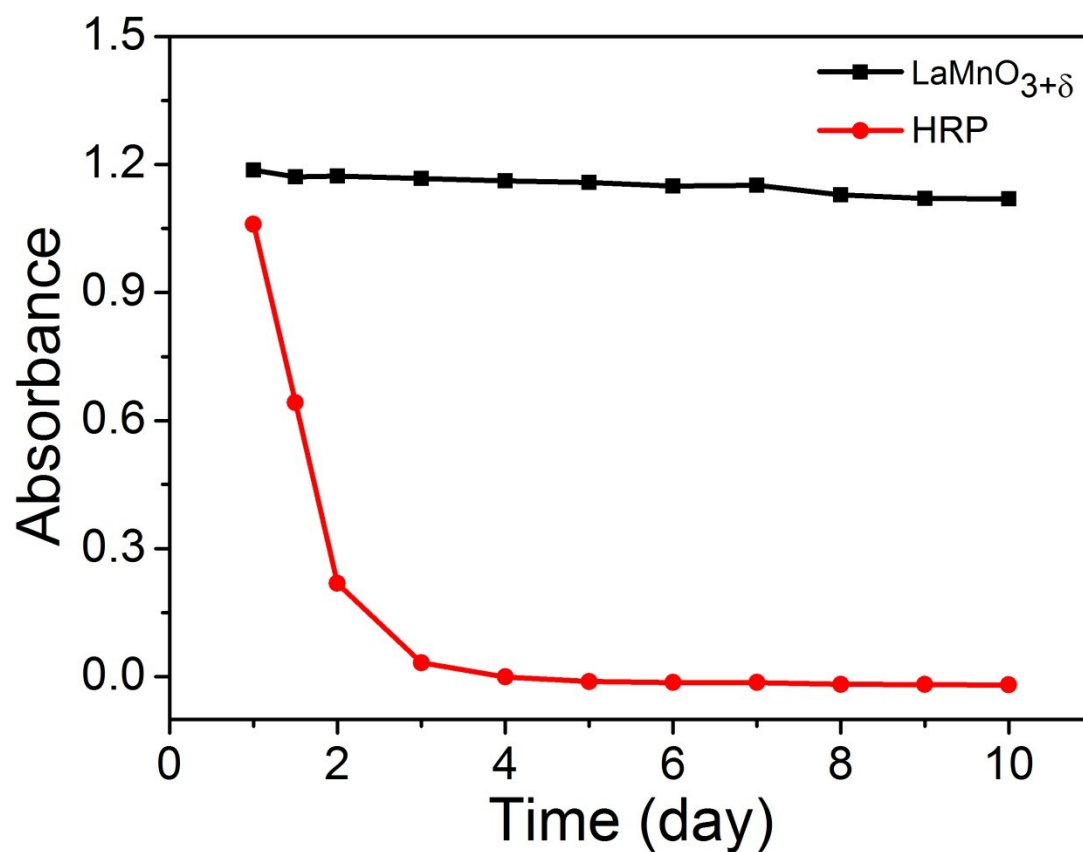


Fig. S4. Long-term stability of LaMnO_{3+δ} nanofibers and HRP for enzyme mimicking. The upper system contains 20 $\mu\text{g}/\text{mL}$ of LaMnO_{3+δ} nanofibers and 0.1 mM TMB in an acetate buffer solution (pH = 4) and the bottom system contains 40 ng/mL of HRP, 0.1 mM TMB and 65 mM H₂O₂ in an acetate buffer solution (pH = 4).

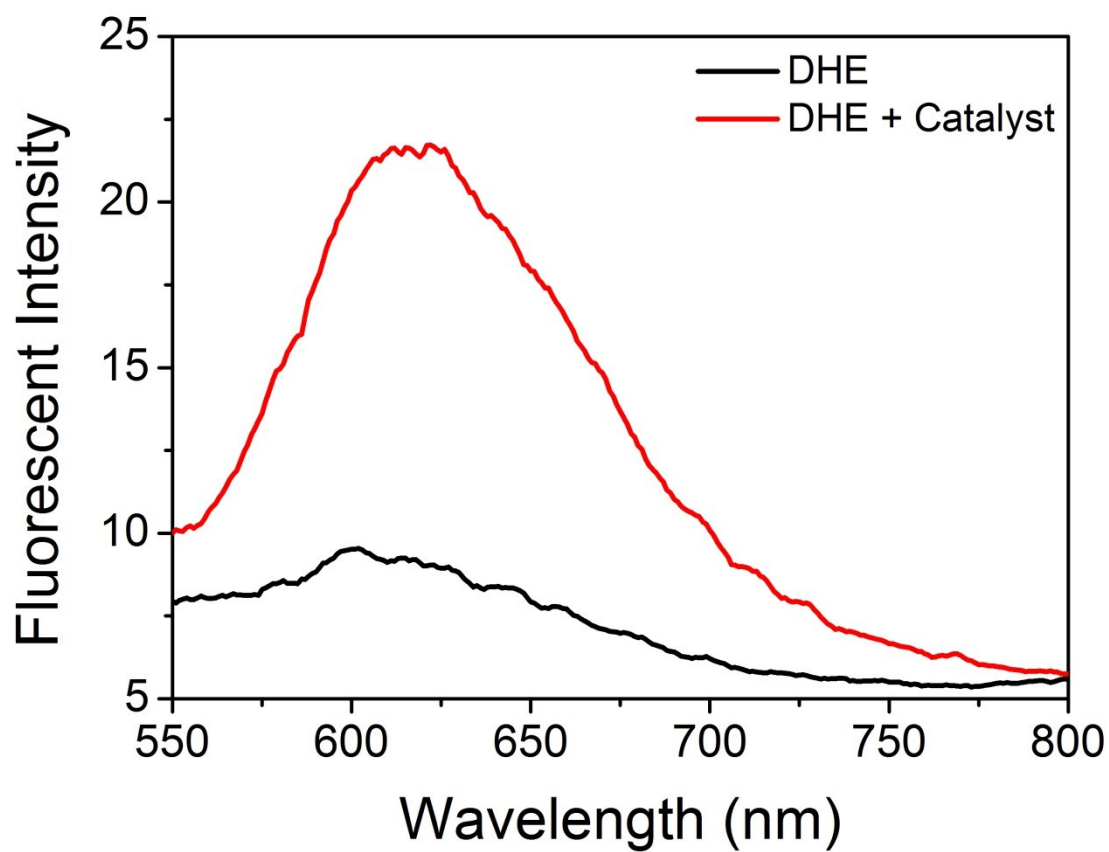


Fig. S5. The fluorescent spectrum of the specific detection of ROS using DHE as fluorescence probes. $\lambda_{\text{ex}} = 500$ nm.

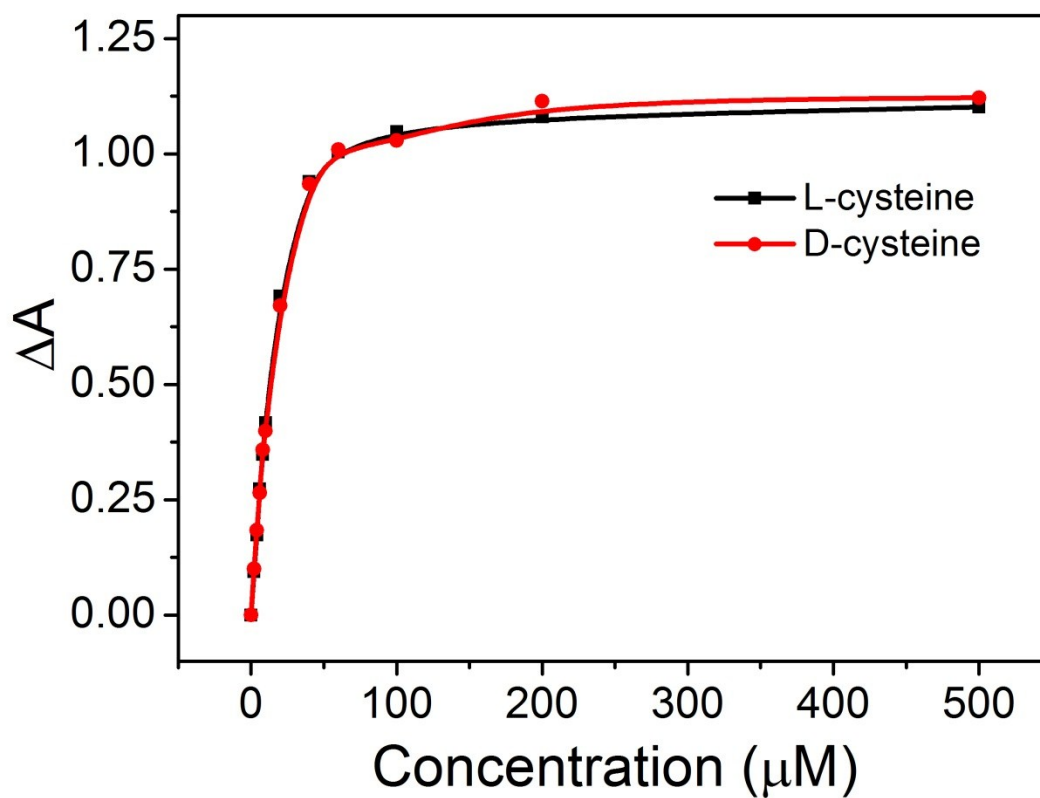


Fig. S6. Comparison of the absorbance evolution at 651 nm of the oxidation of TMB after 10 min with the existence of L-cysteine or D-cysteine at different concentrations.

Table S1. Determination of L-cysteine concentration in the serum sample.

| Serum sample (150-folds diluted) | Without spiking (μM) \pm SD (n=3) | L-cysteine spiking (μM) | Inhibitor measured (μM) \pm SD (n=3) | Recovery (%) |
|--|---|---|--|--------------|
| | | 4 | 8.209 \pm 0.057 | 101.3 |
| Sample | 4.155 \pm 0.236 | 6 | 10.047 \pm 0.237 | 98.2 |
| | | 8 | 12.171 \pm 0.258 | 100.2 |