Electronic Supplementary Information

for

Multi-functional MnO₂ nanomaterials for photo-activated applications by a plasma-assisted fabrication route

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§ S-1. Characterization

§ S-1.1 X-ray photoelectron spectroscopy (XPS)

The surface C1s signal (Fig. S1) could be fitted by means of four distinct components. Bands (I) and (II), centered at BE = 284.8 eV and 286.3 eV, respectively, were attributed to adventitious carbon contamination and to surface C-O moieties.\textsuperscript{1, 2} The two components located at 289.0 eV (III) and 292.6 eV (IV) were mainly ascribed to CF and CF\textsubscript{2} residuals from the used Mn precursor.\textsuperscript{3} The surface C1s photopeak was reduced to noise level after Ar\textsuperscript{+} erosion for 5 min, indicating thus that carbon presence was limited to the sample surface and highlighting the purity of the obtained systems.

![Figure S1. Surface C1s photopeak for a manganese oxide specimen grown at 200 °C.](image-url)
§ S-1.2 Energy dispersive X-ray spectroscopy (EDXS)

EDXS compositional analyses were carried out both in plane and in cross-sectional mode. Fig. S2a reports a representative spectrum, in which O, Mn and F signals could be clearly discerned. Na, Si and Sn signals originate from the growth substrate. Cross-sectional analyses (Fig. S2b) evidenced an almost parallel distribution of O, Mn and F throughout the deposit, suggesting their common chemical origin and demonstrating a uniform in-depth fluorine distribution.

Figure S2. (a) EDXS spectrum for a manganese oxide sample deposited at 200 °C. (b) Cross-sectional EDXS O Kα, Mn Kα and F Kα line-scans for the same specimen.
§ S-1.3 Atomic force microscopy (AFM)

![Graph showing RMS roughness vs. temperature]

**Figure S3.** Evolution of the root mean square (RMS) roughness obtained by AFM measurements as a function of the adopted deposition temperature.
§ S-1.4 Kelvin probe (KP) and valence band XPS results

Figure S4. (a) Absolute Fermi energy obtained by KP. (b) Valence band onset with respect to the Fermi level as determined by XPS valence band measurements. In the inset, experimental data for the sample grown at 200° C are provided as an example.
§ 5-1.5 Photocatalytic tests

The photocatalytic activity mechanism leading to the degradation of the target dye can be described as follows:4-9

\[
\text{MnO}_2 + h\nu \rightarrow \text{MnO}_2(e^-) + h^+ \quad (S1)
\]

\[
h^+ + \text{H}_2\text{O} \rightarrow \text{HO}^* \quad (S2)
\]

\[
e^- + \text{O}_2 \rightarrow \text{O}_2^{*-} \quad (S3)
\]

\[
\text{O}_2^{*-} + \text{H}_2\text{O} \rightarrow \text{HO}_2^* + \text{HO}^* \quad (S4)
\]

\[
\text{O}_2^{*-}/ \text{HO}^* / \text{HO}_2^* + \text{organic pollutant} \rightarrow \text{degradation products} \quad (S5)
\]

Upon irradiation of the catalyst, holes ($h^+$) and electrons ($e^-$) are simultaneously generated in its valence and conduction bands, respectively [equation (S1)]. Subsequently, holes reaction with water molecules results in the formation of hydroxyl radicals [equation (S2)]. In turn, photogenerated electrons can be scavenged by dissolved O$_2$, yielding superoxide radical anions and generate HO$_2^*$ and HO$^*$ species [equations (S3) and (S4)]. Finally, the target organic pollutant can be effectively oxidized by the photogenerated radicals, generating mineralized products (CO$_2$, H$_2$O,...) [equation (S5)].

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<th>$k$ (min$^{-1}$)</th>
<th>$t_{1/2}$ (min)</th>
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<tbody>
<tr>
<td><strong>UV irradiation</strong></td>
<td>(0.025±0.001)</td>
<td>28</td>
</tr>
<tr>
<td><strong>Vis irradiation</strong></td>
<td>(0.022±0.002)</td>
<td>31</td>
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**Table S1.** Comparison of first-order reaction rate constants and half-life times for the photobleaching of Plasmocorinth B solutions under UV and Vis illumination for a $\beta$-MnO$_2$ specimen deposited at 200 °C.
Figure S5. Schematic representation of the band structures of $\beta$-MnO$_2$ and cubic Mn$_2$O$_3$\textsuperscript{10,11}.

Figure S6. Different UV photodegradation cycles of Plasmocorinth B for a MnO$_2$ sample fabricated at a growth temperature of 200 °C.
References


