

Nanosized mesoporous phosphated tin oxide as efficient solid acid catalyst

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Supporting information

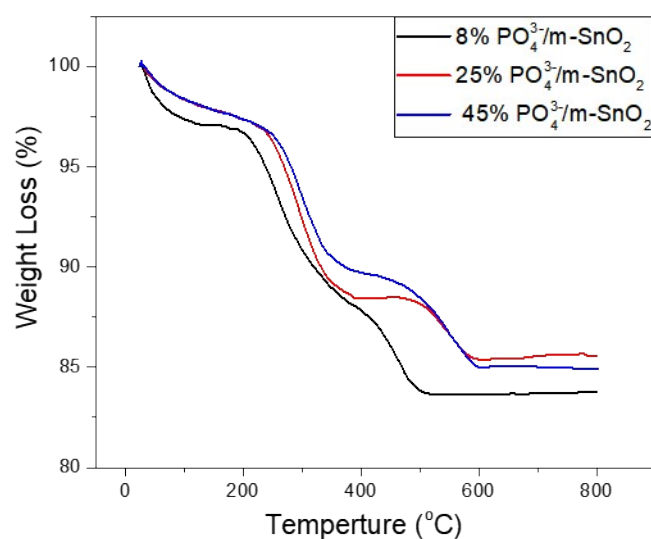


Fig. 1S: TGA curves of (a) 8%, (b) 25% (c) 45% PO₄³⁻/m-SnO₂ samples.

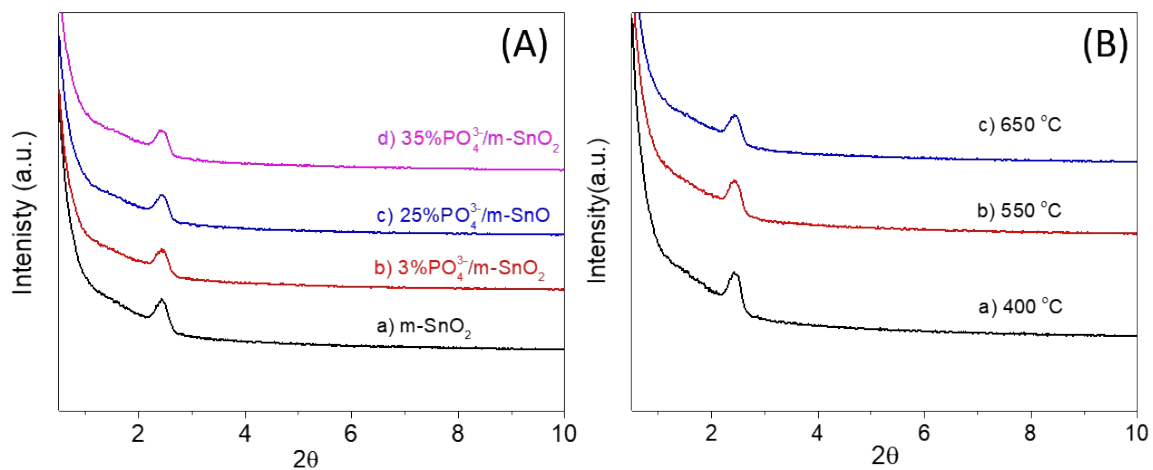


Fig.2S: (A) Low angle XRD pattern of the mesoporous (a) m-SnO₂ (550°C) and the samples PO₄³⁻/SnO₂ (400°C) at (a) m-SnO₂ (b) 3% (c) 25% (d) 35%, (B) Low angle XRD pattern of the sample 3PO₄³⁻/SnO₂ at different temperatures (a) 400°C (b) 550°C (c) 650°C.

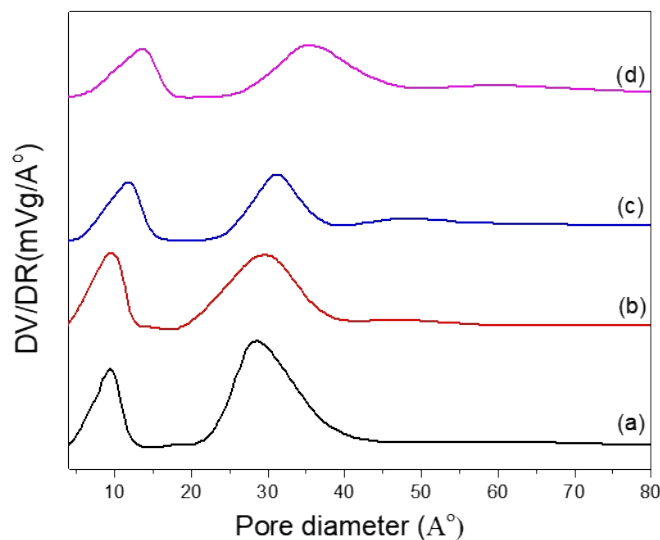


Fig.3S: Pore size distributions of 25% PO₄³⁻/m-SnO₂ samples at (a) 400°; (b) 450°; (c) 550°; (d) 650°C

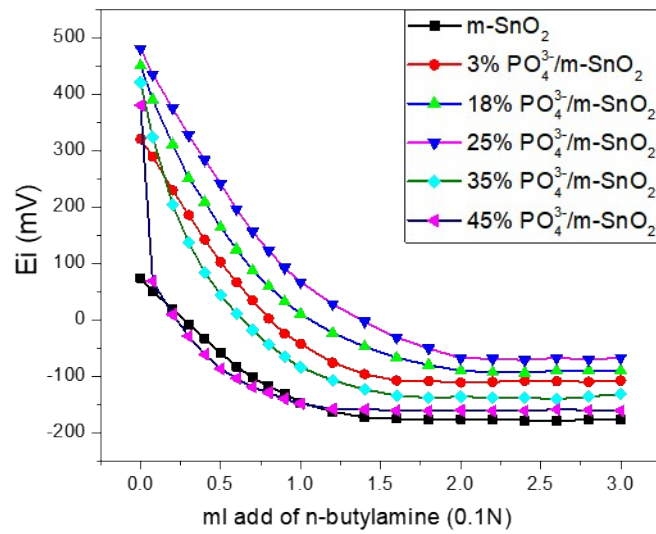


Figure 4S: Potentiometric titration of n-butylamine in acetonitrile for $\text{PO}_4^{3-}/\text{m-SnO}_2$ catalysts calcined at 400°C .

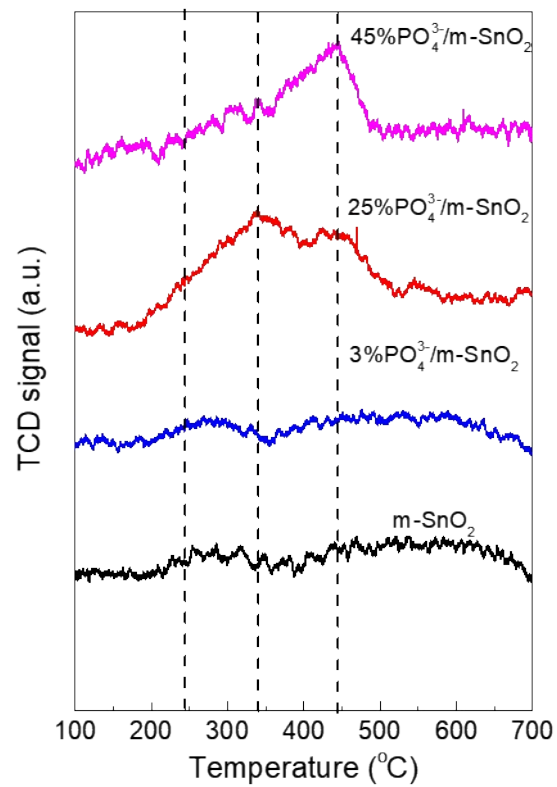


Figure 5S: NH_3 -TPD profiles of the m-SnO_2 , $3\%\text{PO}_4^{3-}/\text{m-SnO}_2$, $25\%\text{PO}_4^{3-}/\text{m-SnO}_2$ and $45\%\text{PO}_4^{3-}/\text{m-SnO}_2$ calcined at $400\text{ }^\circ\text{C}$

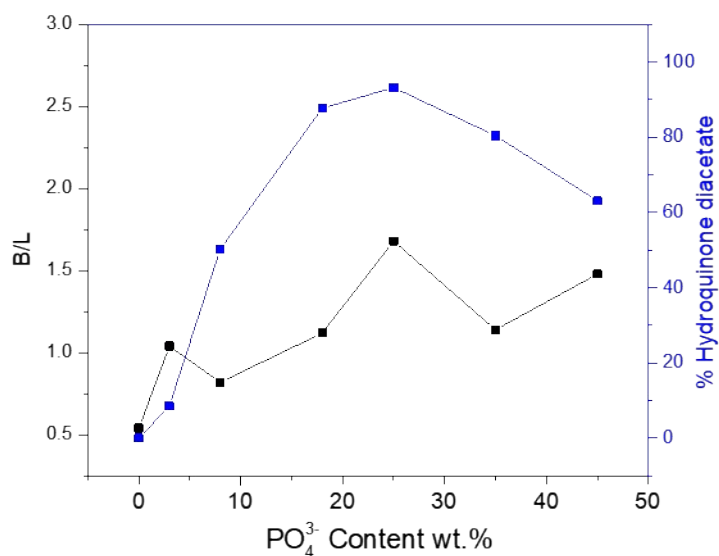


Figure 6S: Effect of PO_4^{3-} content wt. % / mSnO_2 calcined at $400\text{ }^\circ\text{C}$, on the ratio of Brönsted acid sites to Lewis acid sites and % hydroquinone diacetate.

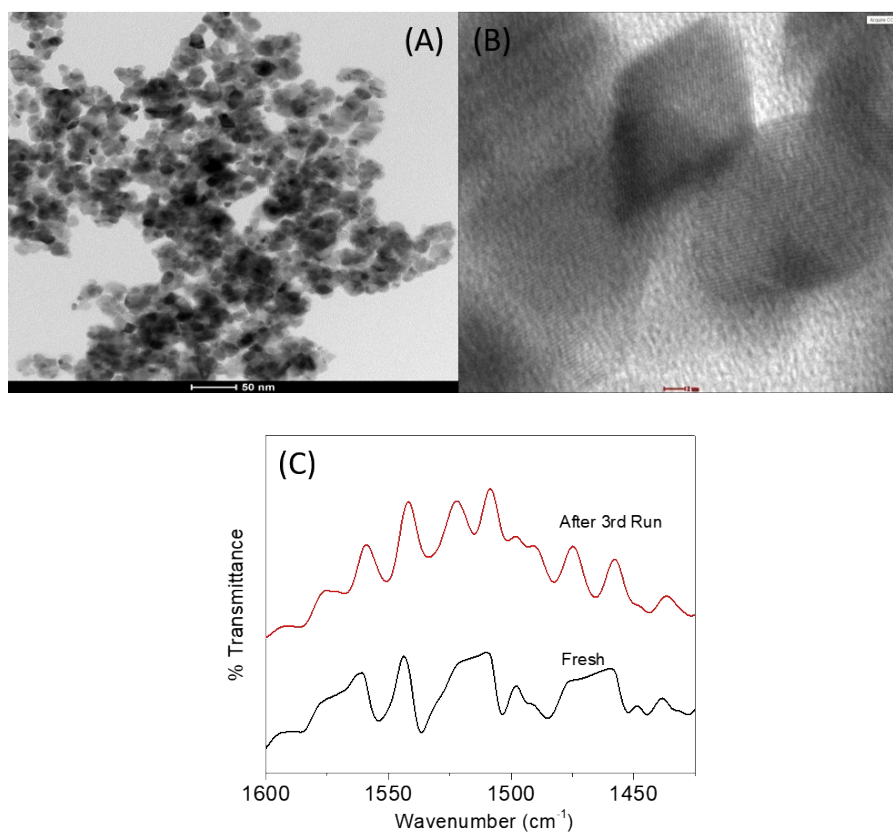


Figure 7S: (A) and (B) TEM images of $25\%\text{PO}_4^{3-}/\text{mSnO}_2$ calcined at $400\text{ }^\circ\text{C}$ after 3rd run (C) FT-IR spectra of pyridine adsorbed on $25\%\text{PO}_4^{3-}/\text{mSnO}_2$ calcined at $400\text{ }^\circ\text{C}$ after 3rd run

Table S1: comparison between solid acid catalyst and homogenous acid catalyst using 0.022 mmol of acid.

sample	Conversion %
25% PO ₄ ³⁻ /mSnO ₂	93.2
H ₂ SO ₄	82.4
HCl	64.7

mmol of Acid for the 25%PO₄³⁻/mSnO₂ has been calculated from the total number of acid sites. For the H₂SO₄ and HCl the same amount of acid concentration has been used