

## Supporting Information

Ruthenium nanoparticles supported over mesoporous TiO<sub>2</sub> as an efficient bifunctional nanocatalyst for esterification of biomass-derived levulinic acid and transfer-hydrogenation reactions

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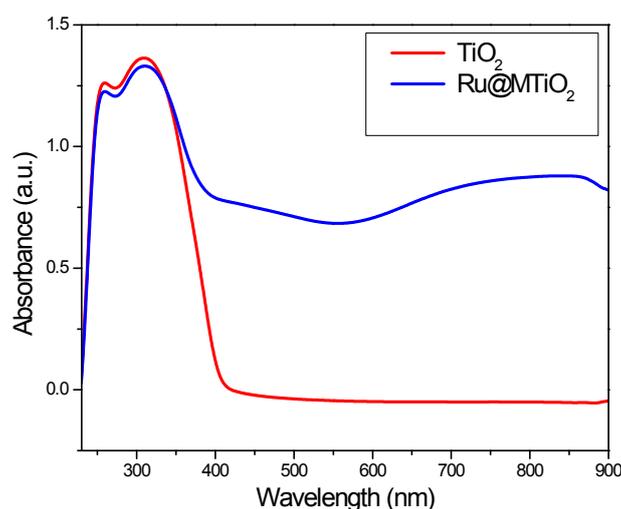
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### UV-Vis spectroscopy

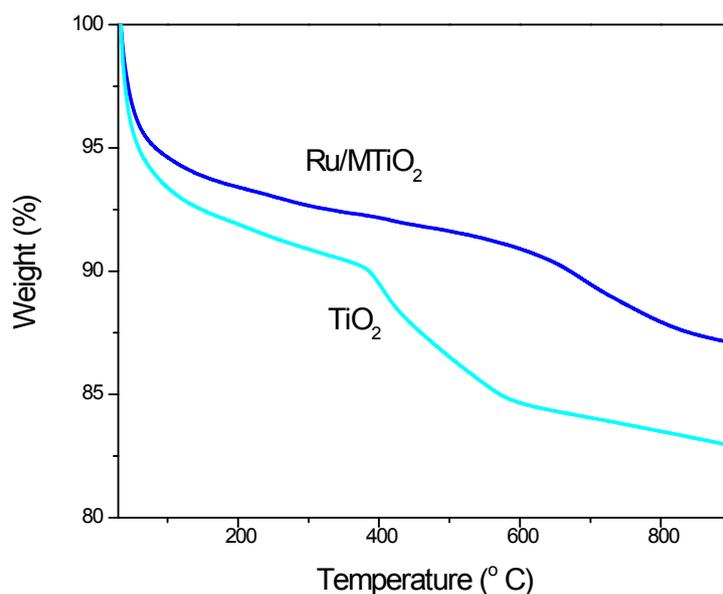
Figure 3 showed the UV-visible diffuse reflectance spectra of mesoporous TiO<sub>2</sub> and Ru@MTiO<sub>2</sub>. A broad absorption peak in the wavelength range 300-350 nm were observed for titania nanocrystal.<sup>17</sup> In case of Ru@MTiO<sub>2</sub>, it exhibited a broad absorption peak in the visible range 600-800 nm due to wide range of visible absorption.<sup>1</sup>



**Figure S1.** UV-Vis spectra of mesoporous TiO<sub>2</sub> and Ru@MTiO<sub>2</sub> nanomaterials.

## Thermogravimetric analysis (TGA)

Figure S2 showed the TGA plots of mesoporous  $\text{TiO}_2$  and  $\text{Ru@MTiO}_2$  nanoparticles under  $\text{N}_2$  flow. The TGA curve showed that *ca* 7% weight loss occurs up to *ca.* 103 °C due to desorption of physisorbed water. Decomposition of surfactant molecules starts from the temperature range of 103 °C to 376 °C and 3% weight loss observed on the surface of the nanoparticles. Again, the decomposition of the organic template starts from 376 °C to 593 °C and rapidly loses 6% of its weight. Then, weight loss of another  $\sim 1.0\%$  occurs gradually till the temperature reached upto 800 °C. In case of  $\text{Ru@MTiO}_2$  nanoparticles, same weight loss occurs up to 103 °C due to desorption of water. Only 2 % weight loss showed by  $\text{Ru@MTiO}_2$  from 103 °C to 610 °C and less than 2% weight occurs up to 800 °C, which confirmed that  $\text{Ru@MTiO}_2$  is more thermally stable than  $\text{TiO}_2$  nanoparticle.

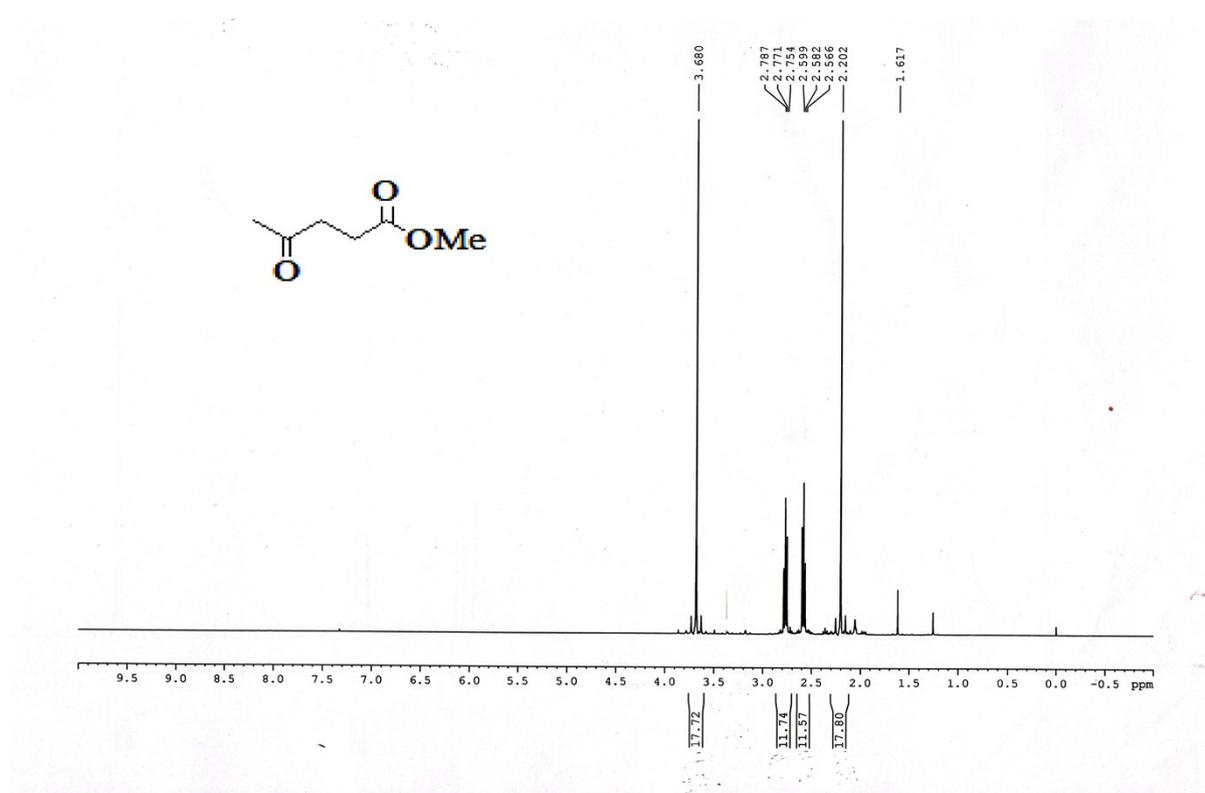


**Figure S2.** TGA plots of mesoporous  $\text{TiO}_2$  and  $\text{Ru@MTiO}_2$  nanomaterials.

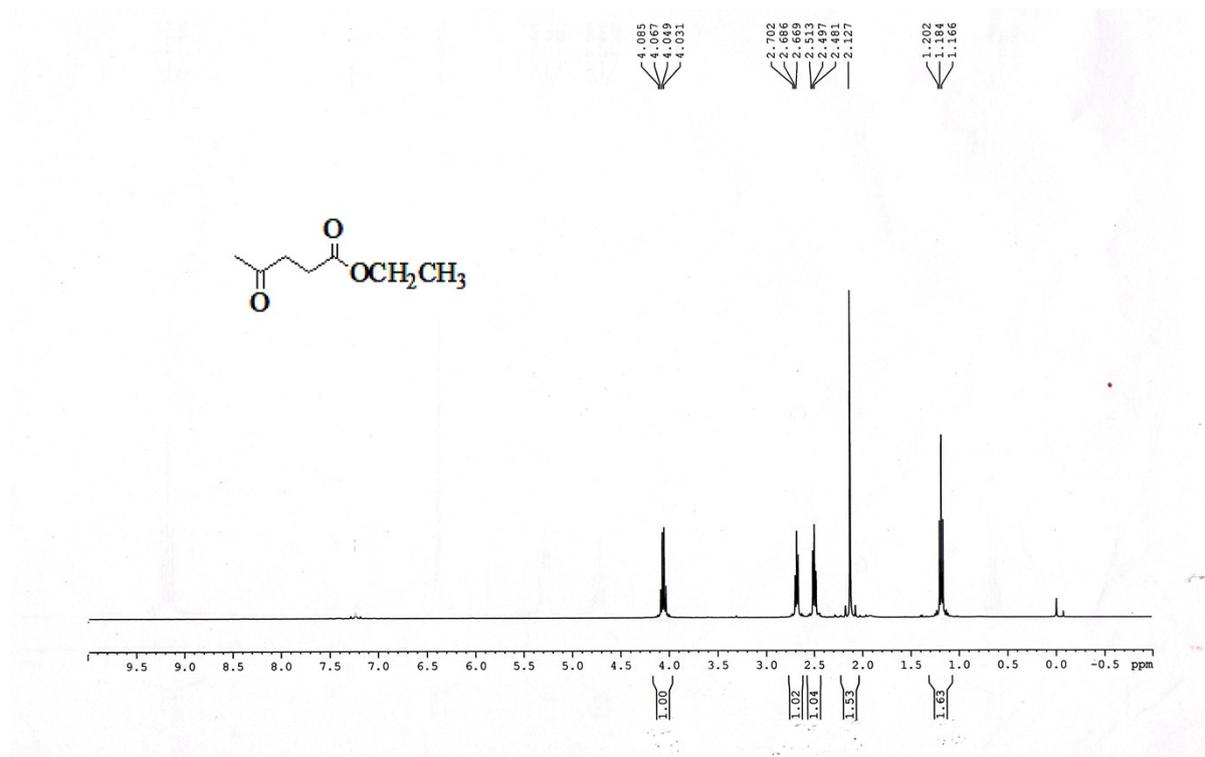
**<sup>1</sup>H NMR data of esterification reactions:**

**Methyl levulinate (Table 1, entry 1):** <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ ppm: 2.20 (s, 3H), 2.58 (t, J = 6.8 Hz, 2H), 2.77 (t, J = 6.4 Hz, 2H), 3.68 (s, 1H).

**Ethyl levulinate (Table 1, entry 2):** <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) δ ppm: 1.18 (t, J = 7.2 Hz, 3H), 2.12 (s, 3H), 2.49 (t, J = 6.4 Hz, 2H), 2.68 (t, J = 6.4 Hz, 2H), 4.03-4.08 (m, 2H).



**Figure S3.** <sup>1</sup>H NMR spectra of methyl levulinate in CDCl<sub>3</sub>.



**Figure S4.** <sup>1</sup>H NMR spectra of ethyl levulinate in CDCl<sub>3</sub>.

## References

1. S. Bang, S. Lee, T. Park, Y. Ko, S. Shin, S.-Y. Yim, H. Seo and H. Jeon, *J. Mater. Chem.*, 2012, **22**, 14141