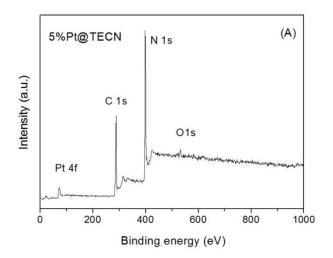
## **Electronic Supplementary Information**

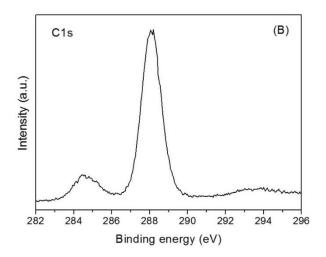
Highly selective hydrogenation of furfural to furfuryl alcohol over Pt nanoparticles supported on g-C<sub>3</sub>N<sub>4</sub> nanosheets catalysts in water

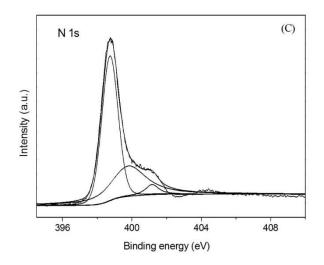
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**Figure S1** (A) XPS survey spectrum of 5%Pt@TECN catalyst, XPS spectra of (B) C1s, (C) N 1s.

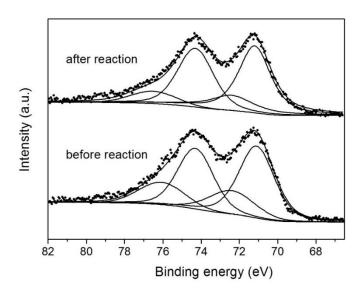


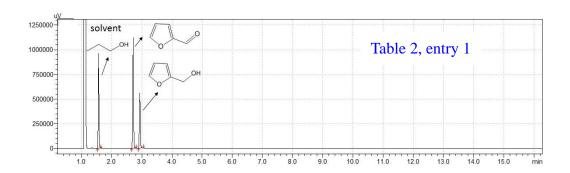
Figure S2. XPS spectra in the Pt 4f region of fresh and spent 5% Pt@TECN catalysts.

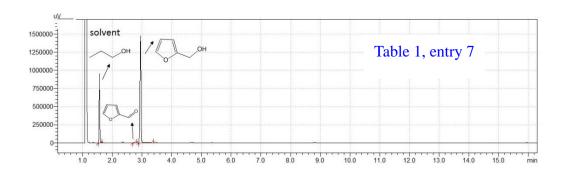
**Table S1.** Furfural hydrogenation over Pt catalysts after 1 h reaction in aqueous phase at  $100~^{\circ}\text{C}$ .

Entry	Catalyst	Conv. (%)	Furfuryl alcohol Sel. (%)
1	5% Pt@TECN	90.3	>99
2	5% Pt@C	51.1	>99

Reaction conditions: Pt catalyst (0.24 mol% Pt relative to substrate), 20 mL H<sub>2</sub>O, 0.4 mL furfural, 1.0 MPa H<sub>2</sub>, 100  $^{\circ}$ C, 1 h.

Gas-chromatographic analysis of the reaction mixture form Table 1, entry 7 and Table 2, entry 1.



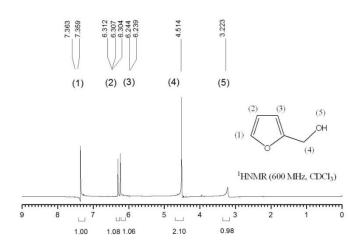


**Figure S3.** Representative GC spectrums of furfural hydrogenation in water.

## NMR analysis of the reaction mixture from Table 1, entry 7.

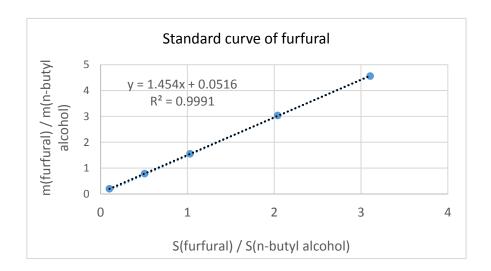
The reaction mixture was obtained by furfural hydrogenation over 5% Pt@TECN under 1MPa H<sub>2</sub> for 5 h in water (Table 1, entry 7). Then the reaction mixture was extracted with EtOAc, the organics were dried (Na<sub>2</sub>SO<sub>4</sub>), concentrated and analysed by NMR. The NMR analysis (see figure S4) confirmed that the main product of the reaction was furfuryl alcohol.

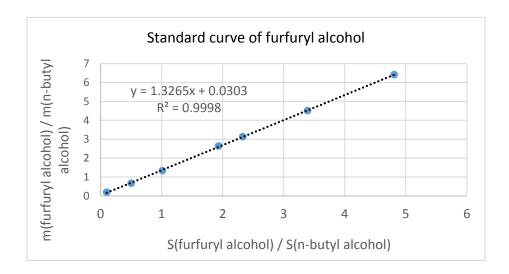
<sup>1</sup>H NMR (600 MHz, CDCl<sub>3</sub>) δ 7.36 (d, 1H), 6.30 (t, 1.08H), 6.24 (d, 1.06H), 4.51 (d, 2.1H), 3.22 (br s, 0.98H).



**Figure S4.** The <sup>1</sup>H NMR spectra of the reaction mixture.

## Quantitative GC analyses





**Figure S5.** Standard curve of furfural with n-butyl alcohol and furfuryl alcohol with n-butyl alcohol.