Supporting Information

Conducting Polymer nanofibers supported Pt Alloys: Unprecedented Materials for MethanolOxidation with Enhanced ElectrocatalyticPerformance and Stability

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Fig. S1 XRD pattern of Ppy nanofibers.



Fig.S2 Transmission electron micrograph of (a,b) Pd/Ppynanocomposites at two different magnification, (c) HRTEM of Pd/Ppy, (d, e) Au/Ppyat two different, higher magnification, (f) HRTEM of Au/Ppy.



Fig.S3 Transmission electron micrograph of (a, b) $Pt_{49}Pd_{51}/Ppy$ nanocompositeat two different magnification,(c)HRTEM of $Pt_{49}Pd_{51}/Ppy$, (d, e) $Pt_{25}Pd_{75}/Ppy$ nanocompositeat two different, higher magnification, (f) HRTEM of $Pt_{25}Pd_{75}/Ppy$.



Fig.S4 Magnified XPS spectrum, (a) C1s, (b) Cl 2p and (c) N 1s in Pt/Ppynanocomposites. (d)
(b) XPS pattern for the synthesized nanocomposites, Pt/Ppy, Pt₆₆Pd₃₄/Ppy, Pt₄₉Pd₅₁/Ppy, Pt₂₅Pd₇₅/Ppy.



Fig.S5 Cyclic voltammetry curves for MOR on Pt₆₆Pd₃₄/Ppynanocompositeat different scan rate variation, (b) plot of current density at different scan rate.



Fig S6 Effect of mass of polymer on Pt based catalysts for electro oxidation of methanol.

Table S1 Comparison of the electrochemical performance of Pt₆₆Pd₃₄/Ppy catalysts for the oxidation of methanol at different temperature. The main characteristics measured from cyclic voltammograms associated with the electrocatalytic oxidation of 1 M MeOH in 0.1 M KOH. The working electrode was a glassy carbon disc modified with the Pt nanostructures. The reference electrode was an Ag/AgCl electrode. The current density is referred to the geometric area of the glassy carbon support.

Temperature (°C)	$j_{\rm f}$ (mA. cm ⁻²)	j_b (mA. cm ⁻²)	j₁⁄jь
20	0.70	.008	87.5
30	11	4.3	2.5
40	17.6	8.3	2.1
50	25	17	1.4