Supporting information

Self-assembly of hollow spherical nanocatalysts with encapsulated Pt NPs and



the effect of Ce-dipping on catalytic activity

Fig. S1 (a) the SAED pattern obtained from Pt-PVP NPs. (b) a HAADF-STEM image of the obtained Pt-PVP NPs.



Fig. S2 FT-IR spectrum of the as-synthesized (a) Pt-PVP NPs and (b) Pt@HMSiO₂ NPs.



Fig. S3 The EDX spectrum datum for the obtained (a) Pt@HMSiO₂ NPs, (b) (c) the Pt-CeO₂@HMSiO₂ NPs of different Ce content.



Fig. S4 N_2 adsorption/desorption isotherms and pore size distribution of the Pt-CeO₂@HMSiO₂ samples.



Fig. S5 TEM images of (a)Pt/SiO2 NPs, (b) Pt/SiO2 NPs calcined at 550 °C



Fig. S6 TEM image of PtCe-0.2 nanocatalyst used seven times



Fig. S7 After reaction for 8 h, TEM image of PtCe-0.2 nanocatalyst.

The propane conversion (X) was calculated by the following formulae:

propane conversion (X%)=[$(n(C_3H_8)_{in} - n(C_3H_8)_{out})$]/ $n(C_3H_8)_{in} \times 100\%$.

After reaction for 8 h, the propane conversion of PtCe-0.2 nanocatalysts decreases from 21.47% to 19.06%. Possibly, the decreasing of the catalytic activity is due to the coke deposition over the catalyst surface. ⁶²