

Supplementary material

**Production of renewable long-chained cycloalkanes from
biomass-derived furfurals and cyclic ketones**

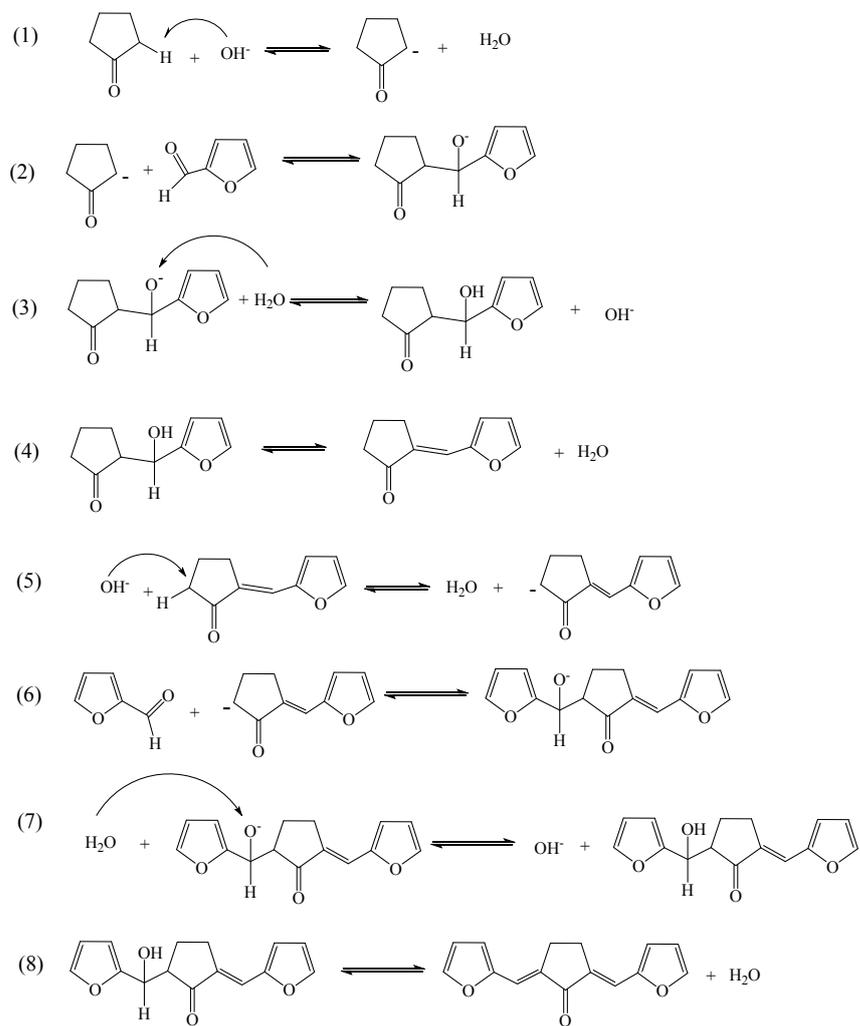
Qiyang Liu ^{a,*}, Caihong Zhang^{a,b}, Ning Shi ^{a,b}, Xinghua Zhang^{a,*}, Chenguang Wang ^a, Longlong

Ma ^a

^a *CAS Key Laboratory of Renewable Energy, Guangzhou Institute of Energy Conversion, Chinese*

Academy of Sciences, Guangzhou, 510650, P. R. China and

^b *University of Chinese Academy of Sciences, Beijing, 100049, P. R. China*



Scheme S1 Formation mechanism of FF-CP-FF by aldol condensation of CP and FF.



Fig. S1 Photographs of the condensation intermediates: (a) FF-CP-FF; (b) FF-CH-FF; (c) HMF-CP-HMF; (d) HMF-CH-HMF.

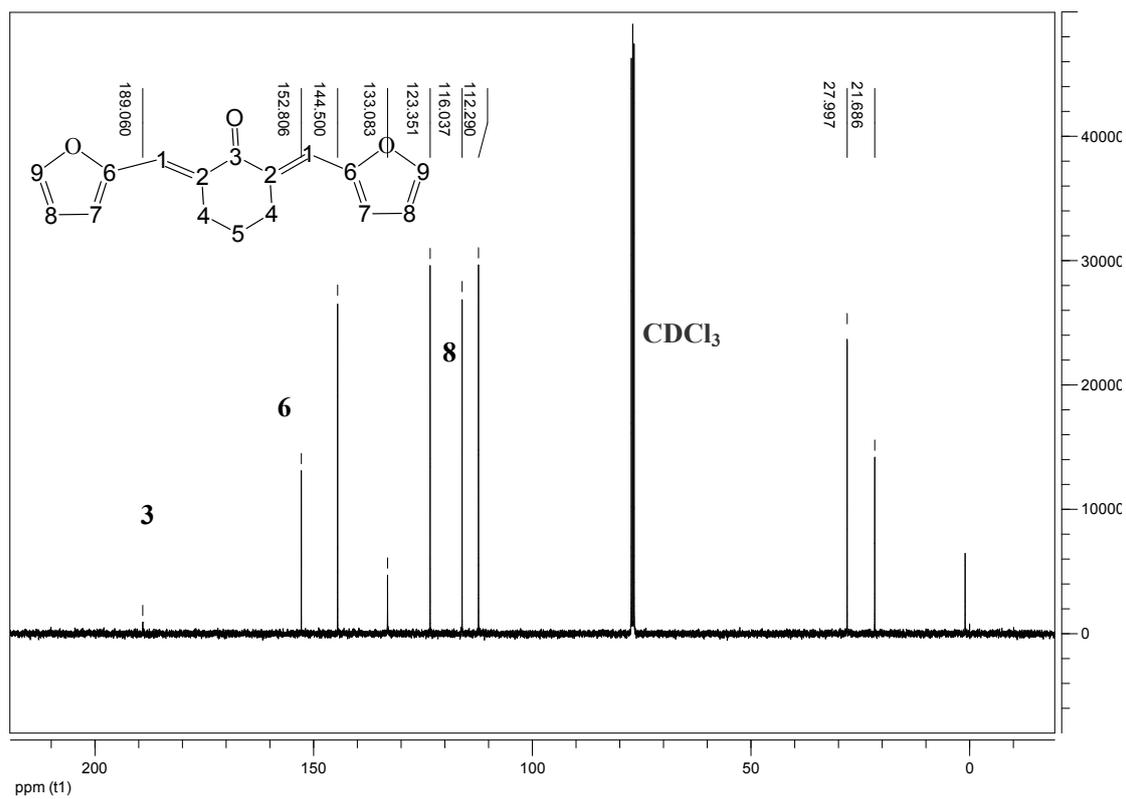


Fig. S2 The ¹³C NMR spectrum of FF-CH-FF.

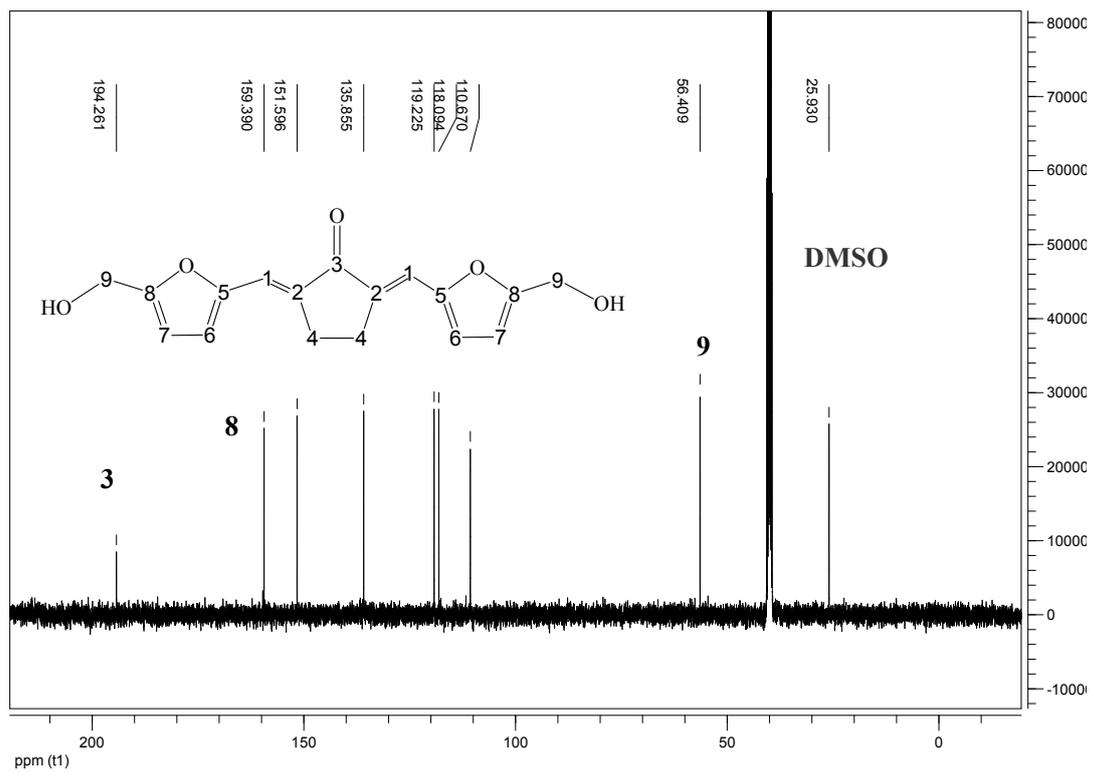


Fig. S3 The ^{13}C NMR spectrum of HMF-CP-HMF.

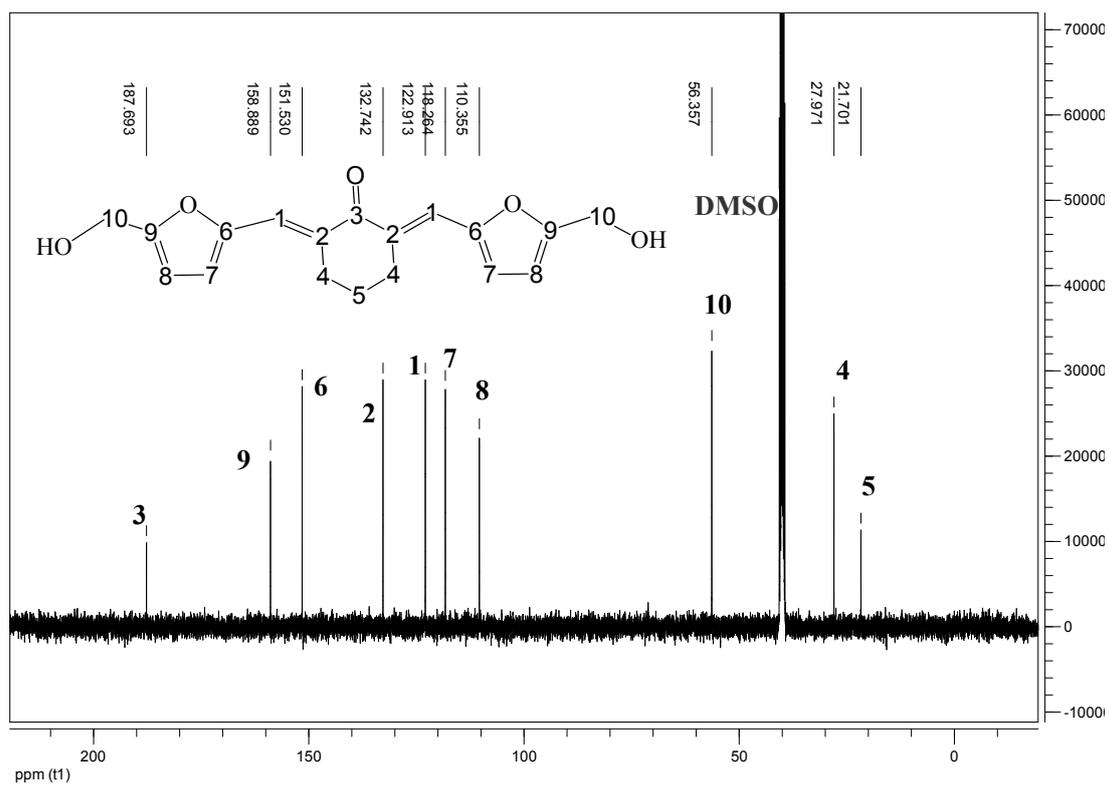


Fig. S4 The ^{13}C NMR spectrum of HMF-CH-HMF.



Fig. S5 Photograph of intermediate produced by FF-CP-FF hydrogenation at 150 °C over Pd/C.
Reaction condition: 1 mmol FF-CP-FF, 15 mL H₂O, 0.05g Pd/C, 4 MPa H₂, 3 h.

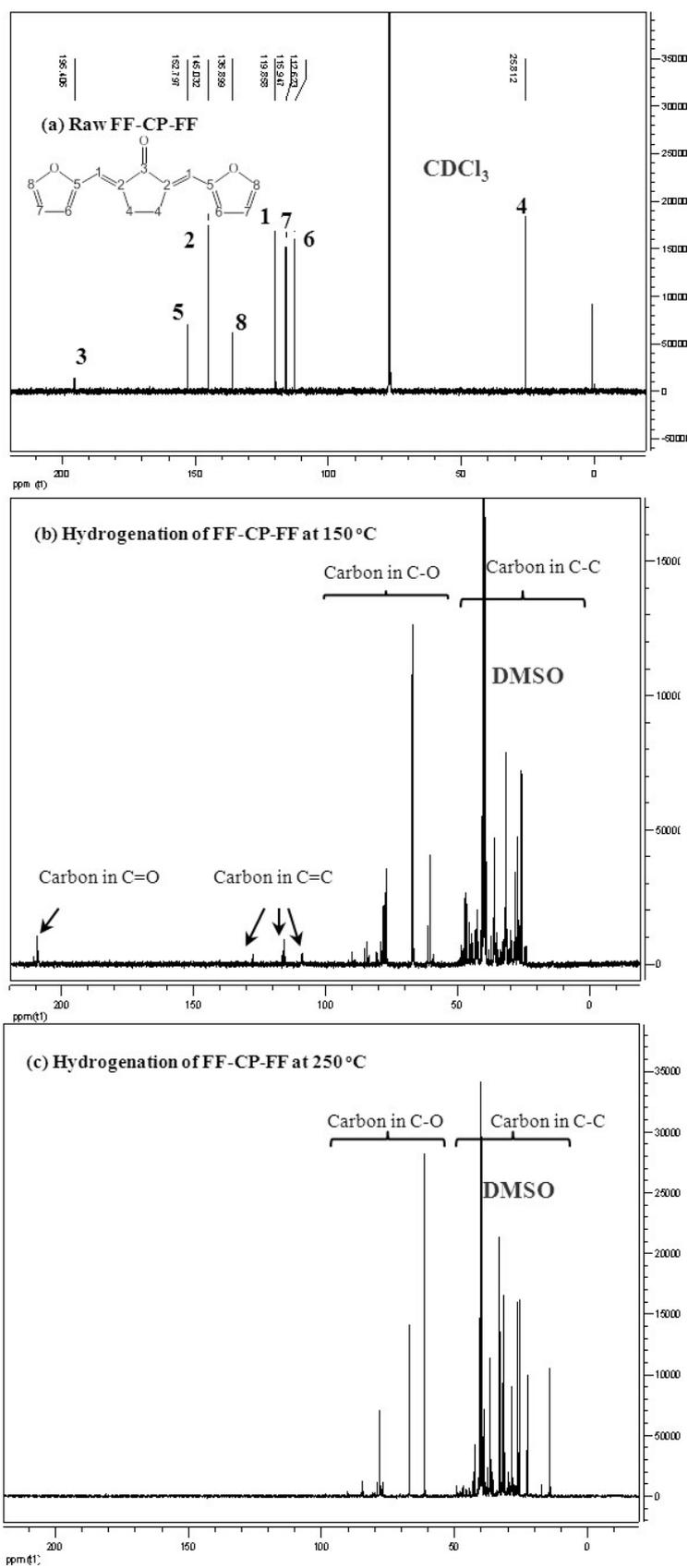


Fig. S6 The ^{13}C NMR spectra of hydrogenated products from FF-CP-FF at different temperatures. Reaction condition: 1 mmol FF-CP-FF, 15 mL H_2O , 0.05g Pd/C, 4 MPa H_2 , 3 h.

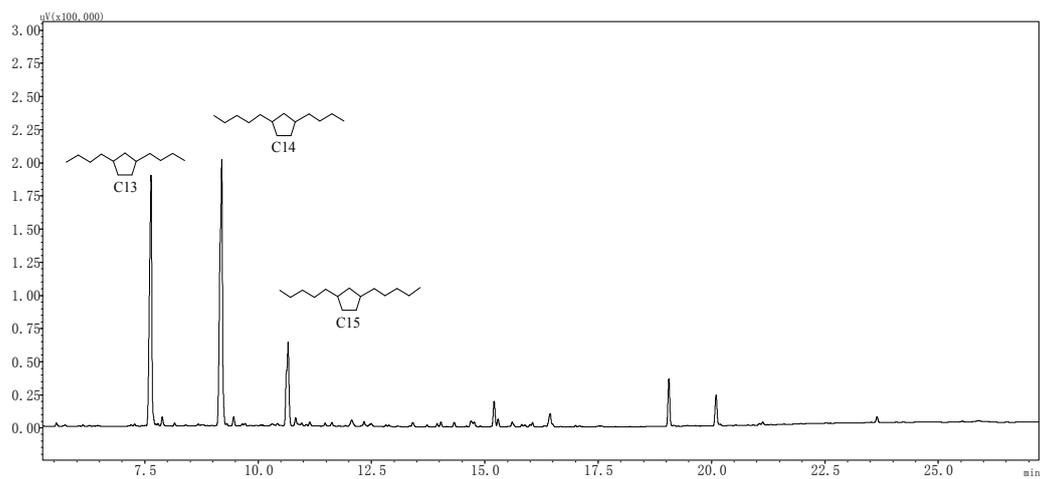


Fig. S7 Representative GC result from hydrodeoxygenation of FF-CP-FF. Reaction conditions: 1.0 mmol FF-CP-FF, 15 mL H₂O, 0.05 g Pd/C, 0.10 g ZrP, 300 °C, 3 h and 4 MPa H₂ pressure.