

Supporting Information

***Tetra*-butylphosphonium arginine-based ionic liquid-promoted cyclization of 2-aminobenzonitrile with carbon dioxide**

State Key Laboratory and Institute of Elemento-Organic Chemistry

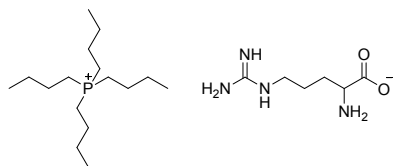
Collaborative Innovation Center of Chemical Science and Engineering (Tianjin)

Nankai University, Tianjin 300071, People's Republic of China

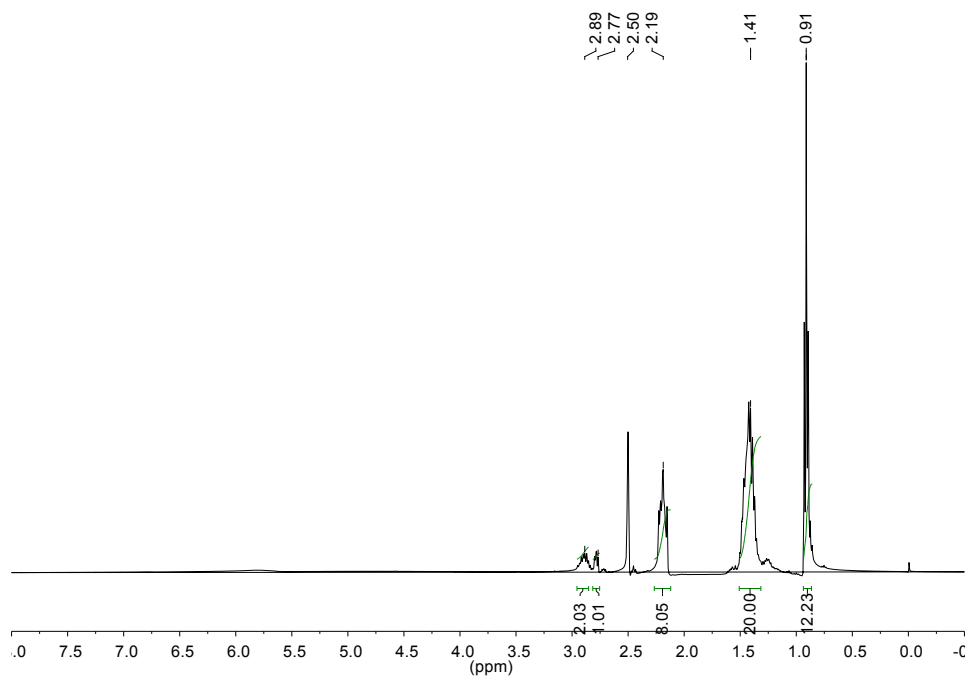
Table of Contents

	Page
1. ¹H NMR and ¹³C NMR Charts for the catalyst and products	S2
2. NMR charts for the covered catalyst	S10

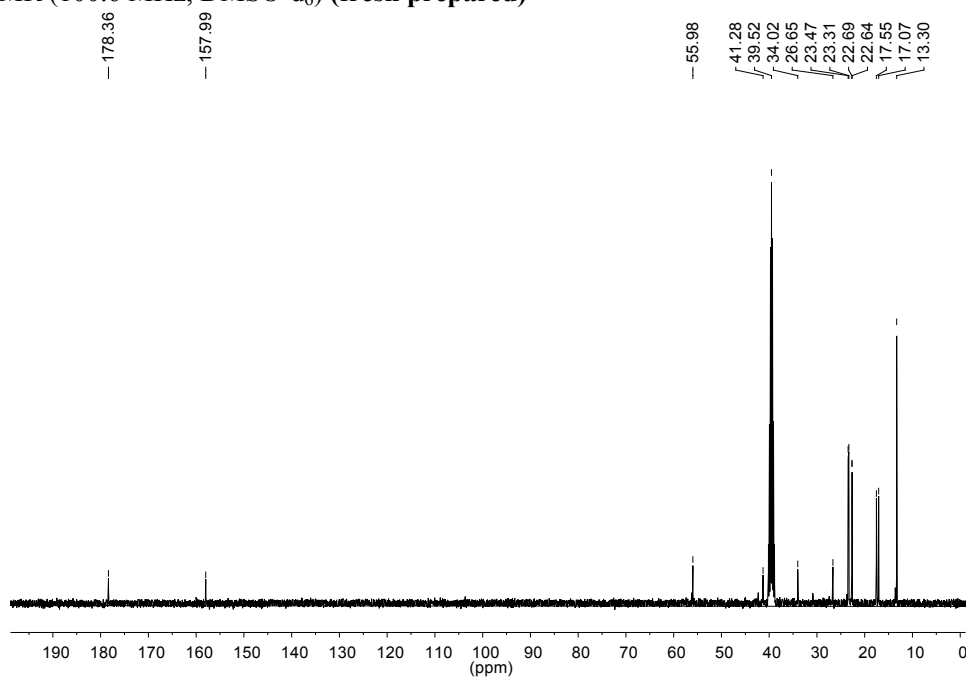
1. ^1H NMR and ^{13}C NMR Charts for the catalyst and products



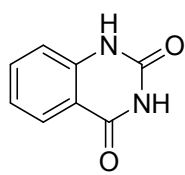
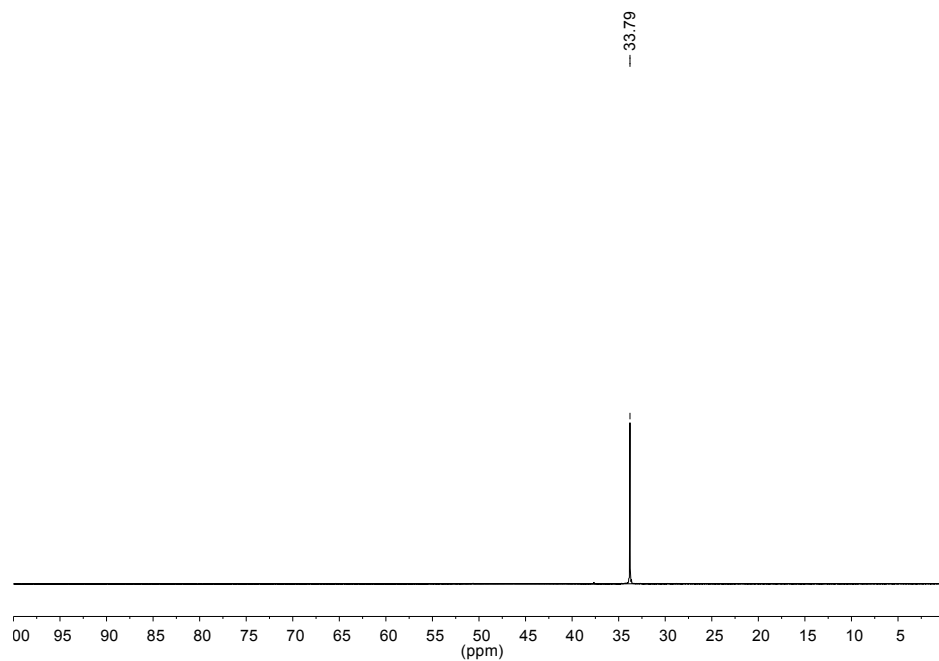
^1H NMR (400 MHz, DMSO-d_6) (fresh prepared)



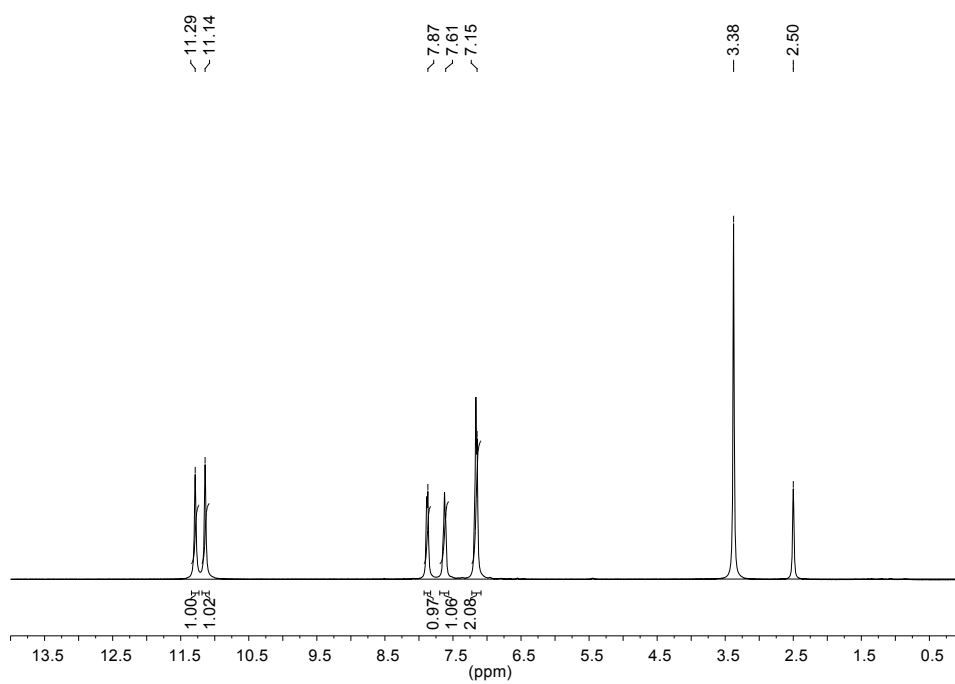
^{13}C NMR (100.6 MHz, DMSO-d_6) (fresh prepared)



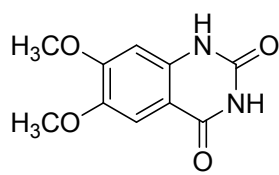
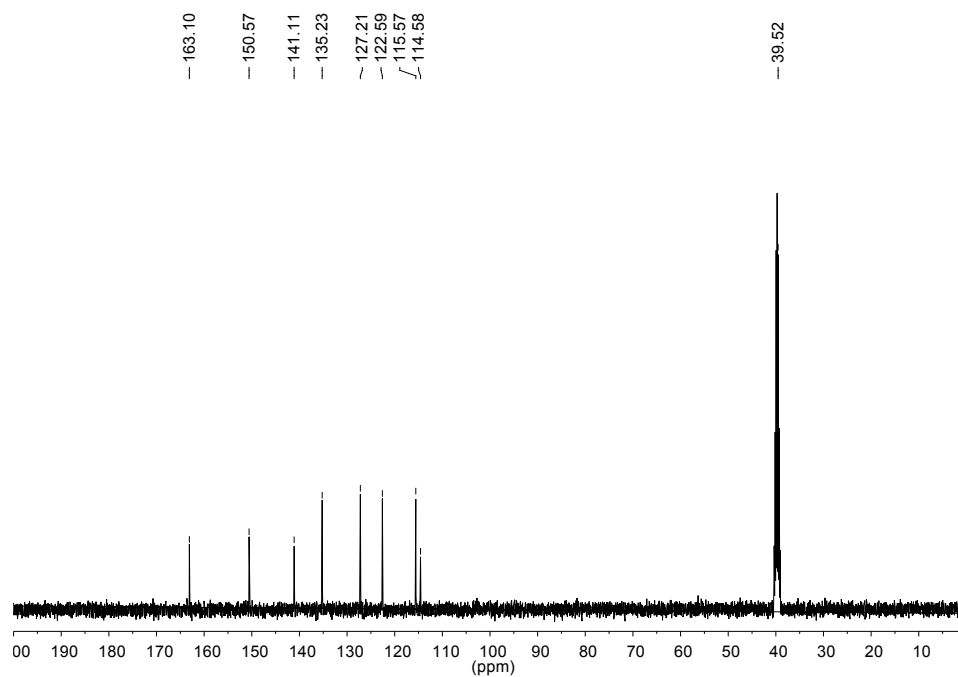
^{31}P NMR (162 MHz, DMSO- d_6) (fresh one)



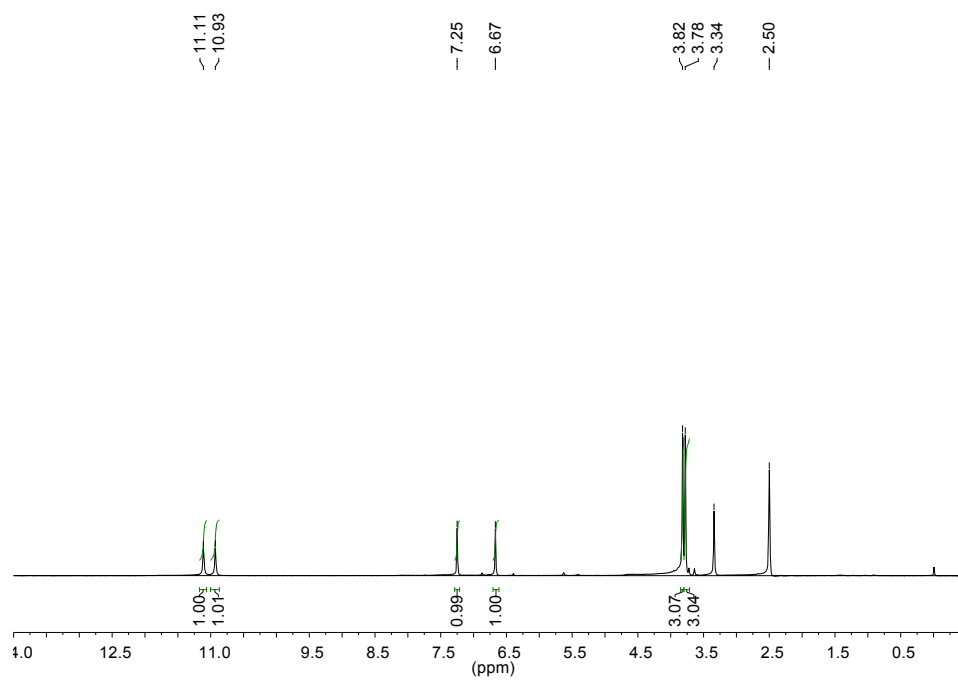
2a ^1H NMR (400 MHz, DMSO- d_6)



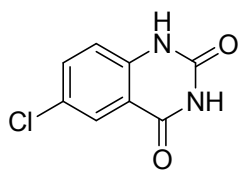
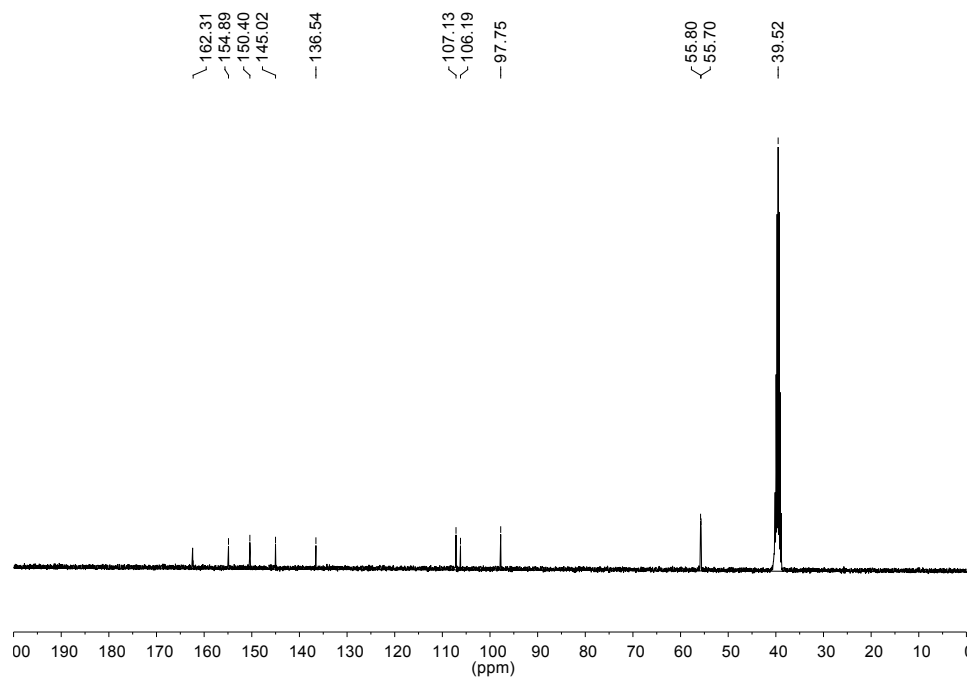
2a ^{13}C NMR (100.6 MHz, DMSO- d_6)



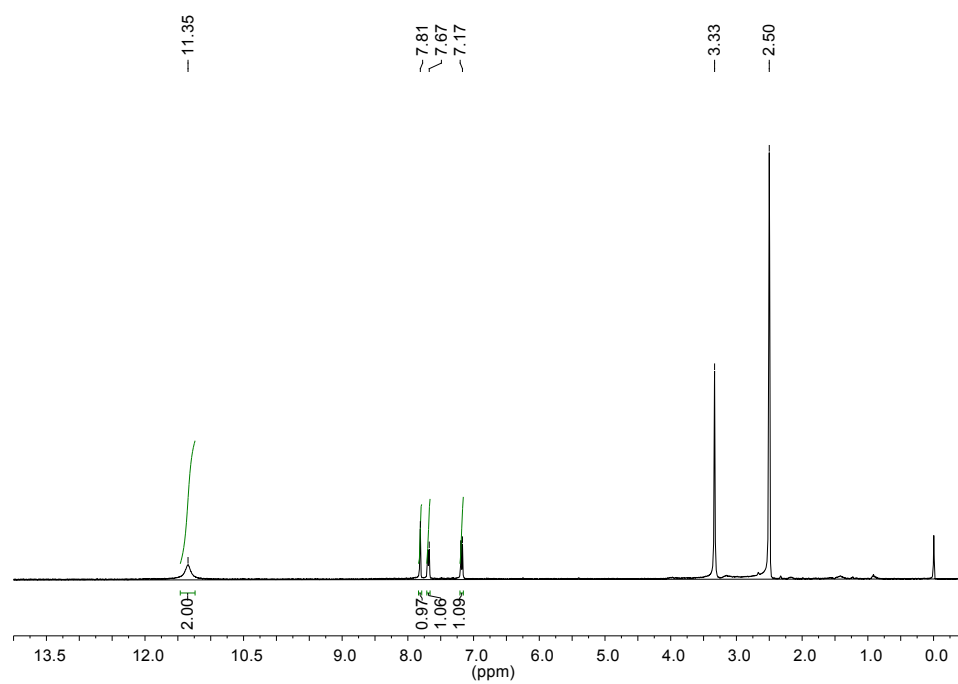
2b ^1H NMR (400 MHz, DMSO- d_6)



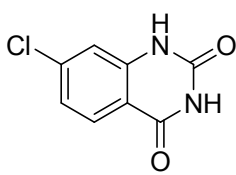
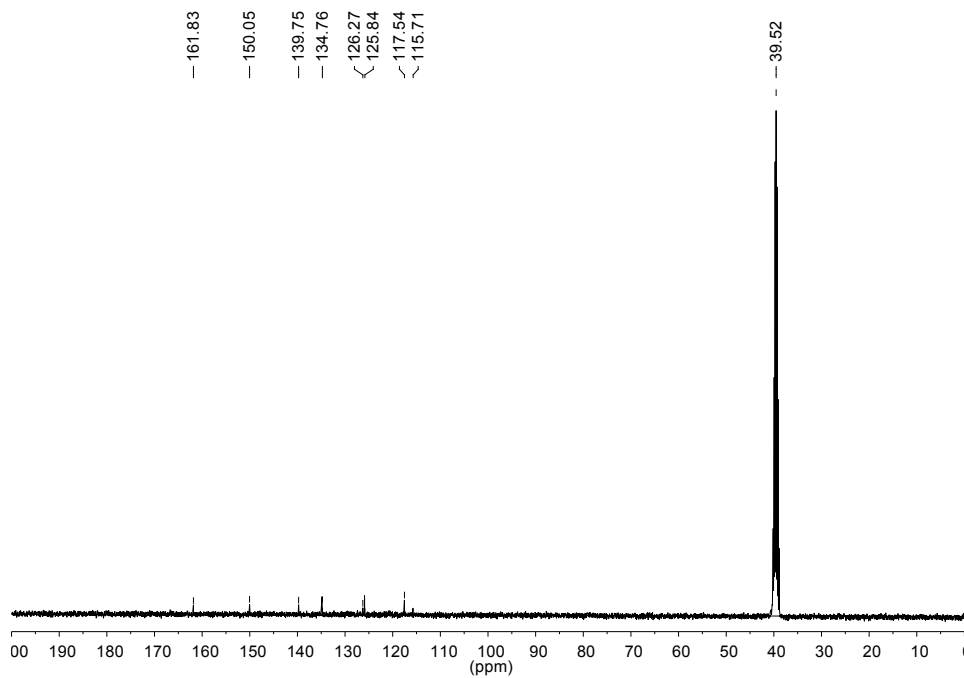
2b ^{13}C NMR (100.6 MHz, DMSO- d_6)



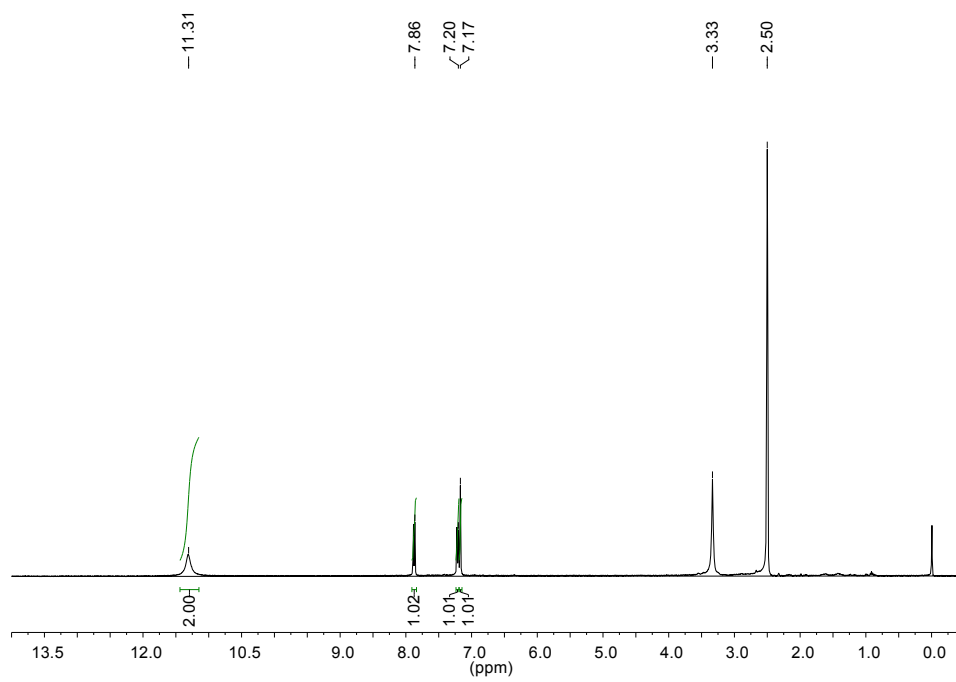
2c ^1H NMR (400 MHz, DMSO- d_6)



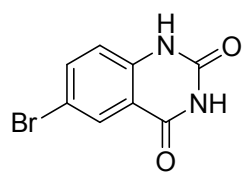
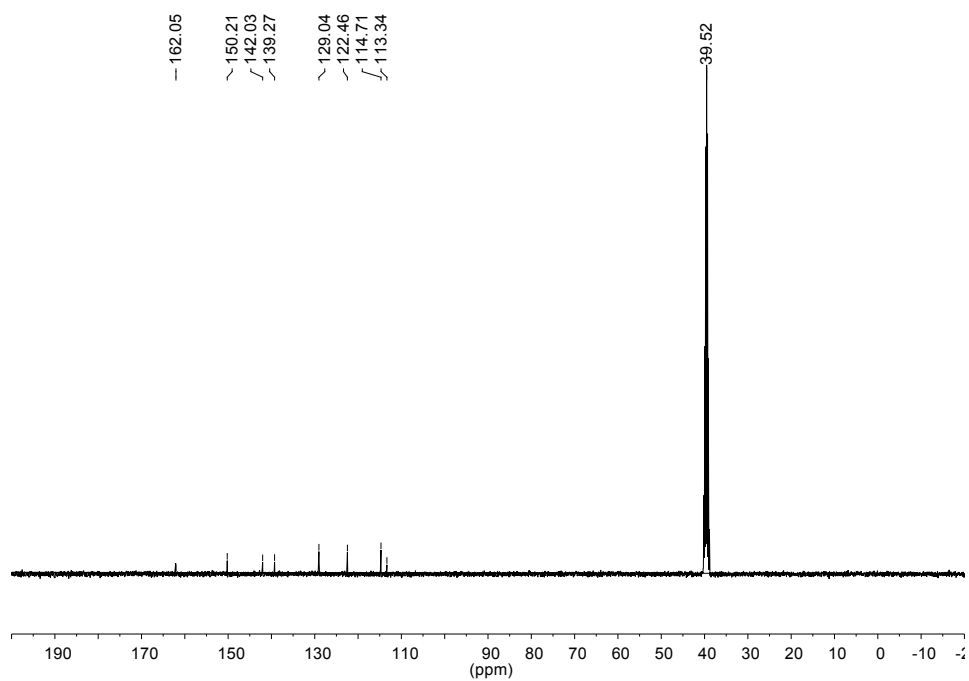
2c ^{13}C NMR (100.6 MHz, DMSO-d_6)



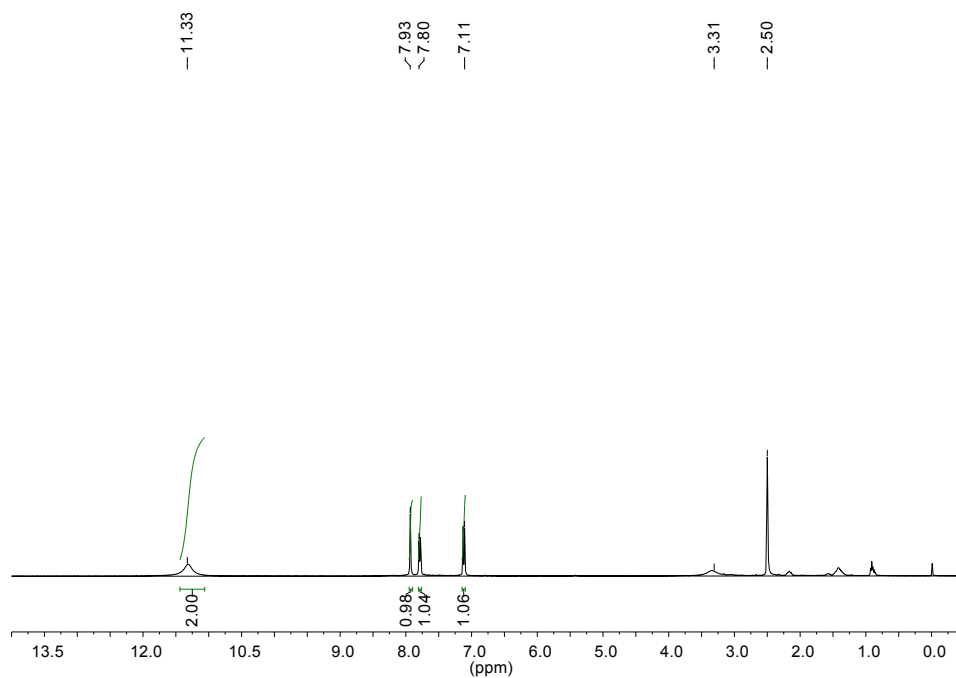
2d ^1H NMR (400 MHz, DMSO-d_6)



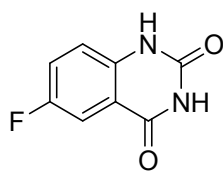
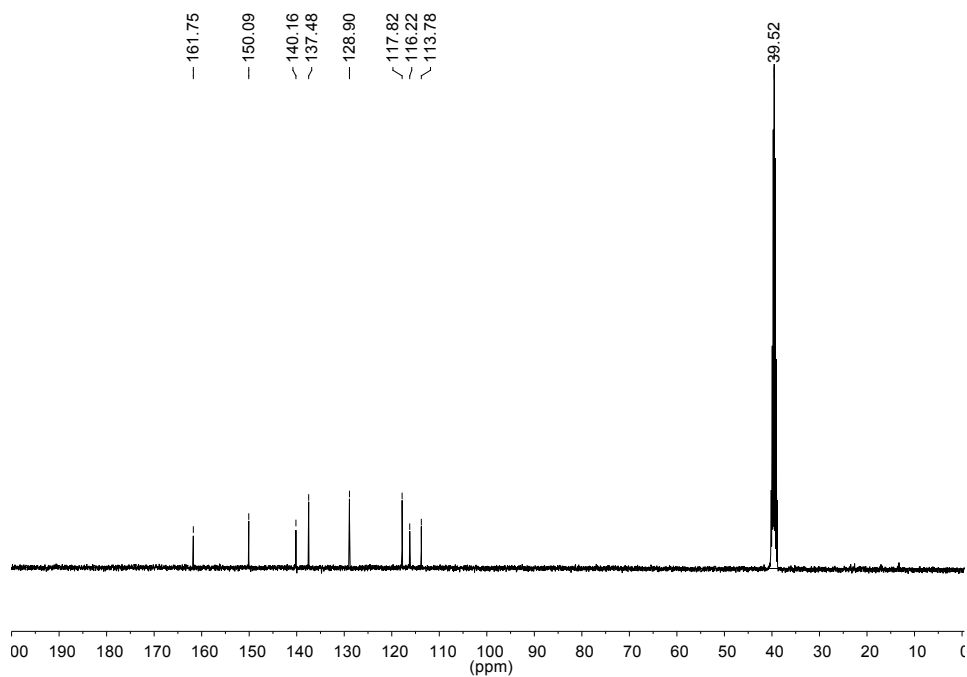
2d ^{13}C NMR (100.6 MHz, DMSO- d_6)



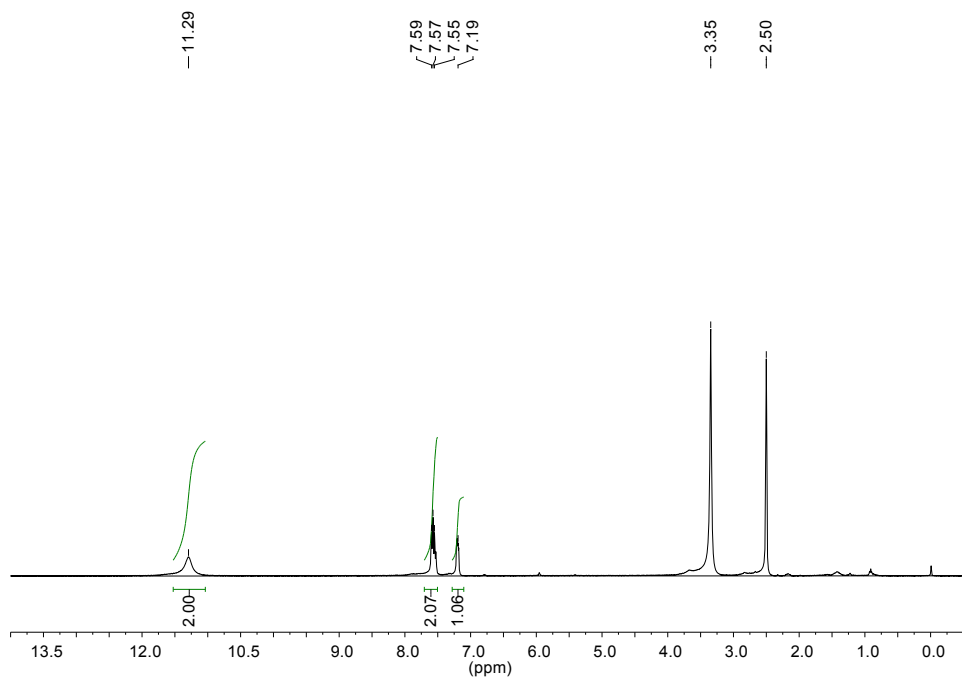
2e ^1H NMR (400 MHz, DMSO- d_6)



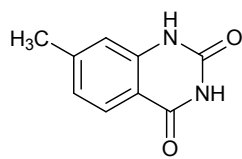
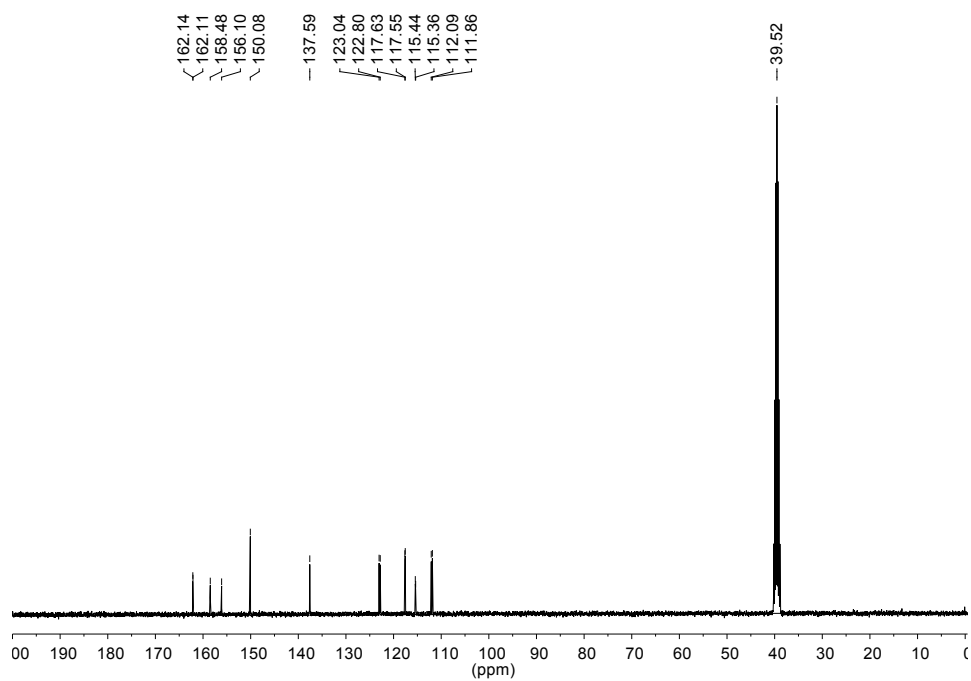
2e ^{13}C NMR (100.6 MHz, DMSO-d_6)



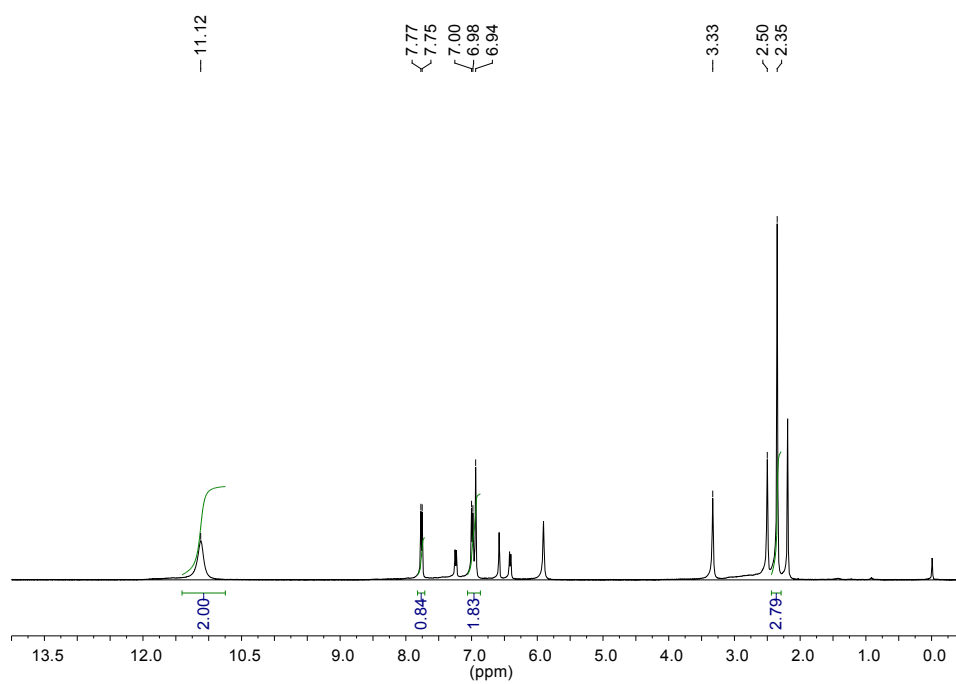
2f ^1H NMR (400 MHz, DMSO-d_6)



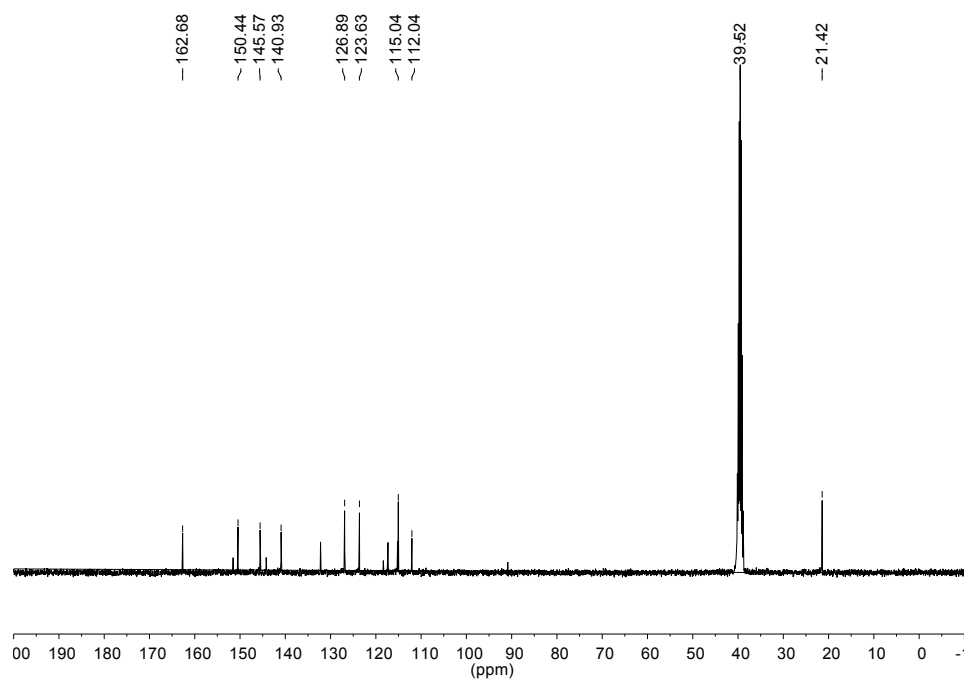
2f ^{13}C NMR (100.6 MHz, DMSO-d_6)



2g ^1H NMR (400 MHz, DMSO-d_6)

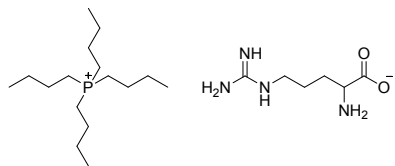


2g ^{13}C NMR (100.6 MHz, DMSO- d_6)

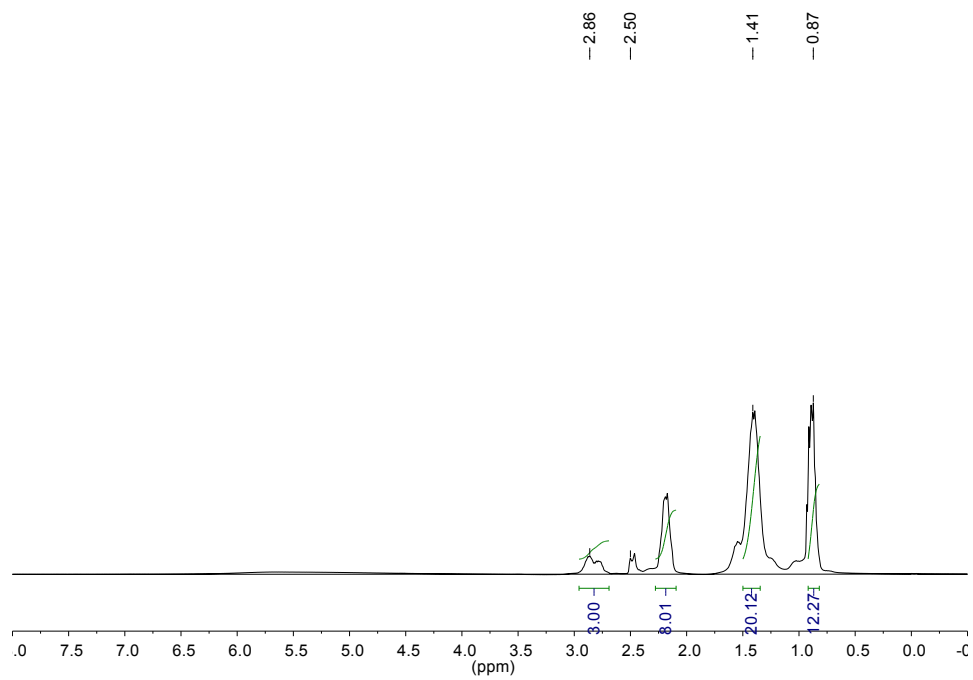


2. General procedure for catalyst recovery and NMR Charts for the recovered catalyst.

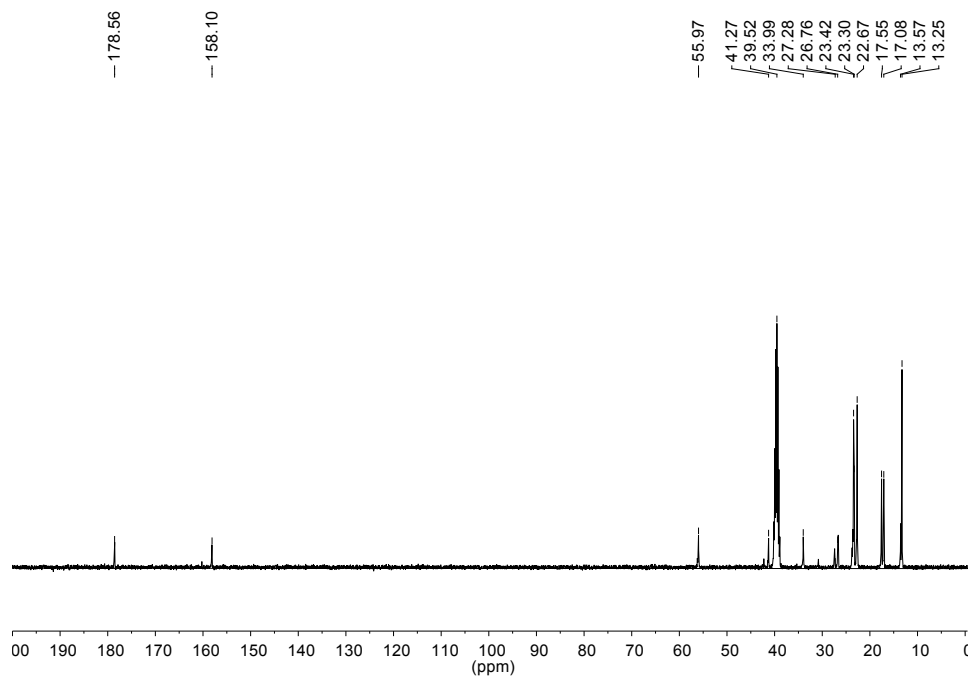
After accomplishment of the reaction, the autoclave was placed in an ice bath to allow it cooled and the inner gas was vented slowly. Water with a volume of 20 mL was poured into the mixture. The resulting precipitate was filtered and washed with water (3×20 mL) and t-BuOMe (3×20 ml), respectively. The water layer was recovered. Then the water was removed by evaporation at 50 °C and dried in vacuo for 1 day at 70 °C to give the recovered ionic liquid [TBP][Arg]. ^1H NMR and ^{13}C NMR spectra showed that the structure of the IL was not changed after reused for five times.



¹H NMR (400 MHz, DMSO-d₆) (recovered after five cycles)



¹³C NMR (100.6 MHz, DMSO-d₆) (recovered after five cycles)



^{31}P NMR (162 MHz, DMSO- d_6) (recovered catalyst)

