Supporting Information

Copper Iodide nanoparticles supported on magnetic aminomethylpyridine functionalized cellulose: a new heterogeneous and recyclable nanomagnetic catalyst for the facile access to *N*-sulfonylamidines under solvent free conditions

Zarrin Ghasemi,* Salman Shojaei and Aziz Shahrisa

Experimental section





Figure 1: ¹H NMR spectrum (400 MHz) of compound 4a in CDCl₃.



Figure 2: Expanded (1) ¹H NMR spectrum (400 MHz) of compound 4a in CDCl₃.



Figure 3: Expanded (2) ¹H NMR spectrum (400 MHz) of compound 4a in CDCl₃.



Figure 4: ¹³C NMR spectrum (100 MHz) of compound 4a in CDCl₃.



4-Methyl-*N*-[2-phenyl-1-(piperidin-1-yl) ethylidene] benzenesulfonamide (4b):

Figure 5: ¹H NMR spectrum (400 MHz) of compound 4b in CDCl₃.



Figure 6: Expanded (1) ¹H NMR spectrum (400 MHz) of compound 4b in CDCl₃.



Figure 7: Expanded (2) ¹H NMR spectrum (400 MHz) of compound 4b in CDCl₃.



Figure 8: ¹³C NMR spectrum (100 MHz) of compound 4b in CDCl₃.



4-Methyl-*N*-[2-phenyl-1-(pyrrolidin-1-yl) ethylidene] benzenesulfonamide (4c):

Figure 9: ¹H NMR spectrum (400 MHz) of compound 4c in CDCl₃.



Figure 10: Expanded (1) ¹H NMR spectrum (400 MHz) of compound 4c in CDCl₃.



Figure 11: Expanded (2) ¹H NMR spectrum (400 MHz) of compound 4c in CDCl₃.



Figure 12: ¹³C NMR spectrum (100 MHz) of compound 4c in CDCl₃.

For previous methods for synthesis of (4a, 4b and 4c), see reference 1.



4-Methyl-*N*-[1-(4-methylpiperazin-1-yl)-2-phenylethylidene] benzenesulfonamide (4d):

Figure 13: FTIR spectrum (KBr) of compound 4d.



Figure 14: ¹H NMR spectrum (400 MHz) of compound 4d in CDCl₃.



Figure 15: Expanded (1) ¹H NMR spectrum (400 MHz) of compound 4d in CDCl₃.



Figure 16: Expanded (2) ¹H NMR spectrum (400 MHz) of compound 4d in CDCl₃.



Figure 17: ¹³C NMR spectrum (100 MHz) of compound 4d in CDCl₃.



N-[1-(4-Methylpiperazin-1-yl)-2-phenylethylidene] benzenesulfonamide (4e):

Figure 18: FTIR spectrum (KBr) of compound 4e.



Figure 19: ¹H NMR spectrum (400 MHz) of compound 4e in CDCl₃.



Figure 20: Expanded (1) ¹H NMR spectrum (400 MHz) of compound 4e in CDCl₃.



Figure 21: Expanded (2) ¹H NMR spectrum (400 MHz) of compound 4e in CDCl₃.



Figure 22: ¹³C NMR spectrum (100 MHz) of compound 4e in CDCl₃.



N-[2-phenyl-1-[4-(phenylsulfonyl) piperazin-1-yl] ethylidene] benzenesulfonamide (4f):

Figure 23: FTIR spectrum (KBr) of compound 4f.



Figure 24: ¹H NMR spectrum (400 MHz) of compound 4f in CDCl₃.



Figure 25: Expanded (1) ¹H NMR spectrum (400 MHz) of compound 4f in CDCl₃.



Figure 26: Expanded (2) ¹H NMR spectrum (400 MHz) of compound 4f in CDCl₃.



Figure 27: ¹³C NMR spectrum (100 MHz) of compound 4f in CDCl₃.



 N^{1} -Cyclohexyl- N^{1} -methyl- N^{2} -(4-methylbenzensulfonyl)-2-phenylacetamidine (4g):

Figure 28: FTIR spectrum (KBr) of compound 4g.



Figure 29: ¹H NMR spectrum (400 MHz) of compound 4g in CDCl₃.



Figure 30: Expanded (1) ¹H NMR spectrum (400 MHz) of compound 4g in CDCl₃.



Figure 31: Expanded (2) ¹H NMR spectrum (400 MHz) of compound 4g in CDCl₃.



Figure 32: Expanded (3) ¹H NMR spectrum (400 MHz) of compound 4g in CDCl₃.



Figure 33: Expanded (4) ¹H NMR spectrum (400 MHz) of compound 4g in CDCl₃.



Figure 34: ¹³C NMR spectrum (100 MHz) of compound 4g in CDCl₃.



Figure 35: Expanded (1) ¹³C NMR spectrum (100 MHz) of compound 4g in CDCl₃.



Figure 36: Expanded (2) ¹³C NMR spectrum (100 MHz) of compound 4g in CDCl₃.



Figure 37: Expanded (3) ¹³C NMR spectrum (100 MHz) of compound 4g in CDCl₃.



 N^1 -Cyclohexyl- N^1 -methyl- N^2 -(phenylsulfonyl)-2-phenylacetamidine (4h):

Figure 38: FTIR spectrum (KBr) of compound 4h.



Figure 39: ¹H NMR spectrum (400 MHz) of compound 4h in CDCl₃.



Figure 40: Expanded (1) ¹H NMR spectrum (400 MHz) of compound 4h in CDCl₃.



Figure 41: Expanded (2) ¹H NMR spectrum (400 MHz) of compound 4h in CDCl₃.



Figure 42: Expanded (3) ¹H NMR spectrum (400 MHz) of compound 4h in CDCl₃.



Figure 43: Expanded (4) ¹H NMR spectrum (400 MHz) of compound 4h in CDCl₃.



Figure 44: ¹³C NMR spectrum (400 MHz) of compound 4h in CDCl₃.



Figure 45: Expanded (1) ¹³C NMR spectrum (400 MHz) of compound 4h in CDCl₃.



Figure 46: Expanded (2) ¹³C NMR spectrum (100 MHz) of compound 4h in CDCl₃.



Figure 47: Expanded (3) ¹³C NMR spectrum (100 MHz) of compound 4h in CDCl₃.



 N^{1} -Cyclohexyl- N^{1} -methyl- N^{2} -(methylsulfonyl)-2-phenylacetamidine (4i):

Figure 48: FTIR spectrum (KBr) of compound 4i.



Figure 49: ¹H NMR spectrum (400 MHz) of compound 4i in CDCl₃.



Figure 50: Expanded (1) ¹H NMR spectrum (400 MHz) of compound 4i in CDCl₃.



Figure 51: Expanded (2) ¹H NMR spectrum (400 MHz) of compound 4i in CDCl₃.



Figure 52: Expanded (3) ¹H NMR spectrum (400 MHz) of compound 4i in CDCl₃.



Figure 53: Expanded (4) ¹H NMR spectrum (400 MHz) of compound 4i in CDCl₃.



Figure 54: ¹³C NMR spectrum (100 MHz) of compound 4i in CDCl₃.



Figure 55: Expanded (1) ¹³C NMR spectrum (400 MHz) of compound 4i in CDCl₃.



Figure 56: Expanded (2) ¹³C NMR spectrum (100 MHz) of compound 4i in CDCl₃.



Figure 57: Expanded (3) ¹³C NMR spectrum (400 MHz) of compound 4i in CDCl₃.



Figure 58: Expanded (4) ¹³C NMR spectrum (100 MHz) of compound 4i in CDCl₃.

N^1 , N^1 -Dipropyl- N^2 -tosyl-2-phenylacetamidine (4j):



Figure 59: FTIR spectrum (KBr) of compound 4j.



Figure 60: ¹H NMR spectrum (400 MHz) of compound 4j in CDCl₃.



Figure 61: Expanded (1) ¹H NMR spectrum (400 MHz) of compound 4j in CDCl₃.



Figure 62: Expanded (2) ¹H NMR spectrum (400 MHz) of compound 4j in CDCl₃.



Figure 63: Expanded (3) ¹H NMR spectrum (400 MHz) of compound 4j in CDCl₃.



Figure 64: ¹³C NMR spectrum (100 MHz) of compound 4j in CDCl₃.

For previous methods for synthesis of N^1 , N^1 -Dipropyl- N^2 -tosyl-2-phenylacetamidine (4j), see reference 2a and 2c.



 N^1 , N^1 -Dibutyl- N^2 -tosyl-2-phenylacetamidine (4k):

Figure 65: ¹H NMR spectrum (400 MHz) of compound 4k in CDCl₃.



Figure 66: Expanded (1) ¹H NMR spectrum (400 MHz) of compound 4k in CDCl₃.



Figure 67: Expanded (2) ¹H NMR spectrum (400 MHz) of compound 4k in CDCl₃.



Figure 68: Expanded (3) ¹H NMR spectrum (400 MHz) of compound 4k in CDCl₃.



Figure 69: ¹³C NMR spectrum (100 MHz) of compound 4k in CDCl₃.

For previous methods for synthesis of N^1 , N^1 -Dibutyl- N^2 -tosyl-2-phenylacetamidine (4k), see reference 2.



 N^{1} , N^{1} -dibenzyl- N^{2} -(4-methylbenzensulfonyl)-2-phenylacetamidine (4l):

Figure 70: FTIR spectrum (KBr) of compound 4I.



Figure 71: ¹H NMR spectrum (400 MHz) of compound 4l in CDCl₃.



Figure 72: Expanded ¹H NMR spectrum (400 MHz) of compound 4l in CDCl₃.



Figure 73: ¹³C NMR spectrum (100 MHz) of compound 4l in CDCl₃.



 N^{1} , N^{1} -dibenzyl- N^{2} -(phenylsulfonyl)-2-phenylacetamidine (4m):

Figure 74: FTIR spectrum (KBr) compound 4m.



Figure 75: ¹H NMR spectrum (400 MHz) of compound 4m in CDCl₃.



Figure 76: Expanded ¹H NMR spectrum (400 MHz) of compound 4m in CDCl₃.



Figure 77: ¹³C NMR spectrum (100 MHz) of compound **4m** in CDCl₃.



 N^{1} , N^{1} -Dibenzyl- N^{2} -(methylsulfonyl)-2-phenylacetamidine (4n):

Figure 78: FTIR spectrum (KBr) of compound 4n.



Figure 79: ¹H NMR spectrum (400 MHz) of compound 4n in CDCl₃.



Figure 80: Expanded (1) ¹H NMR spectrum (400 MHz) of compound 4n in CDCl₃.



Figure 81: Expanded (2) ¹H NMR spectrum (400 MHz) of compound 4n in CDCl₃.



Figure 82: ¹³C NMR spectrum (100 MHz) of compound 4n in CDCl₃.



 N^{1} , N^{1} -Dibenzyl- N^{2} -(4-methylbenzensulfonyl)-pentanamidine (40):

Figure 83: FTIR spectrum (KBr) of compound 40.



Figure 84: ¹H NMR spectrum (400 MHz) of compound 40 in CDCl₃.



Figure 85: Expanded (1) ¹H NMR spectrum (400 MHz) of compound 40 in CDCl₃.



Figure 86: Expanded (2) ¹H NMR spectrum (400 MHz) of compound 40 in CDCl₃.



Figure 87: Expanded (3) ¹H NMR spectrum (400 MHz) of compound 40 in CDCl₃.



Figure 88: ¹³C NMR spectrum (100 MHz) of compound 40 in CDCl₃.



 N^1 , N^1 -Diphenyl- N^2 -tosyl-2-phenylacetamidine (4p):

Figure 89: ¹H NMR spectrum (400 MHz) of compound 4p in CDCl₃.



Figure 90: Expanded ¹H NMR spectrum (400 MHz) of compound 4p in CDCl₃.



Figure 91: ¹³C NMR spectrum (400 MHz) of compound 4p in CDCl₃.

For previous methods for synthesis of N^1 , N^1 -Diphenyl- N^2 -tosyl-2-phenylacetamidine (4p), see reference 3.

References

[1] E. J. Yoo, I. Bae, S. H. Cho, H. Han and S. Chang, *Org. Lett.*, 2006, **8**, 1347-1350.
[2] (a) J. Kim and S. S. Stahl, *J. Org. Chem.*, 2015, **80**, 2448-2454; (b) H. E. Xinwei, S. Yongjia, H. U. Jinsong, J. U. Kai, J. Wei and W. Sufang, *Sci. China Chem.*, 2012, **55**, 214-222; (c) M. Jagadale, P. Bhange, R. Salunkhe, D. Bhange, M. Rajmane and G. Rashinkar, *Appl. Catal. A Gen.*, 2016, **511**, 95-105.

[3] T. Yang, H. Cui, C. Zhang, L. Zhang and C. -Y. Su, Inorg. Chem., 2013, 52, 9053-9059.