Supplementary Materials

Enhanced catalytic performance by copper nanoparticle**graphene based composite** Paramita Mondal,^a Arjyabaran Sinha,^b Noor Salam,^a Anupam Singha Roy,^a

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Table S1. Effect of copper source on the coupling of imidazole with iodobenzene.

Entry	Copper Source	Yield (%)
1	None	No reaction
2	Cu-metal	12
3	CuI	32
4	CuCl ₂	29
5	Cu(OAc) ₂	37
6	Cu-G composite (6 % Cu)	72
7	Cu-G composite (13 % Cu)	98
8	Cu-C composite (7 % Cu)	58
9	Cu-C composite (14 % Cu)	80

Entry	Solvent	Base	Temp. (⁰ C)	Yield (%) ^{b,c}
1	Methanol	Cs_2CO_3	reflux	22
2	CAN	Cs ₂ CO ₃	100	34
3	DMF	Cs_2CO_3	100	70
4	Toluene	Cs_2CO_3	100	12
5	DMSO	Cs ₂ CO ₃	100	98
6	DMSO	K ₂ CO ₃	100	78
7	DMSO	K ₃ PO ₄	100	45
8	DMSO	КОН	100	34
9	DMSO	KOBu ^t	100	23
10	DMSO	Na ₂ CO ₃	100	59
11	DMSO	Et ₃ N	100	09
12 ^d	DMSO	Cs_2CO_3	Room temp.	14
13	DMSO	Cs_2CO_3	60	47

Table S2. The effect of different reaction parameters on the Cu-G (13 % Cu) composite catalyzed O-arylation reaction of iodobenzene and phenol.^a

^aReaction conditions: Cu-G catalyst (0.05 g), iodobenzene (1 mmol); phenol (1 mmol); base (2 mmol); solvent (10 mL); Time (12 h) open air. ^bYield determined by GC. ^cProducts were identified by GC–MS. ^dReaction time 24 h.

Entry	Base	Solvent	Temperature (⁰ C)	Time (h)	Yield (%) ^{b,c}
1	K ₂ CO ₃	DMSO	100	7	89
2	Cs ₂ CO ₃	DMSO	100	7	94
3	K ₃ PO ₄	DMSO	100	12	78
4	Et ₃ N	DMSO	100	12	52
5	КОН	DMSO	100	12	49
6	Cs ₂ CO ₃	DMF	100	7	84
7	Cs ₂ CO ₃	NMP	100	7	78
8	Cs ₂ CO ₃	CAN	100	7	70
9	Cs ₂ CO ₃	THF	100	12	32
10	Cs ₂ CO ₃	Toluene	100	18	12
11	Cs ₂ CO ₃	DMSO	Room temp.	24	No reaction
12	Cs ₂ CO ₃	DMSO	50	12	39
13	Cs ₂ CO ₃	DMSO	70	12	58

Table S3. Screening of different solvents and bases for *N*-arylation of imidazole with iodobenzene.

^aReaction conditions: Cu-G catalyst (0.05 g), iodobenzene (1 mmol); imidazole (1.2 mmol); base (2 mmol); solvent (10 mL); ^b Yield determined by GC. ^c Products were identified by GC–MS.



Figure S2. ¹H-NMR spectrum of 4-Methoxy-diphenylether



Figure S4. ¹H-NMR spectrum of 4-Nitro-diphenylether



Figure S6. ¹H-NMR spectrum of 2-Methyl-diphenylether



Figure S8. ¹H-NMR spectrum of 1-(4-nitrophenoxy)-4-methoxybenzene



Figure S9. ¹H-NMR spectrum of 4-Cyano-4-methoxy-diphenylether





Figure S11. ¹H-NMR spectrum of 1-phenyl-1H-imidazole



Figure S12. ¹H-NMR spectrum of 1-(4-(1H-Imidazol-1-yl)phenyl)ethanone



Figure S14. ¹H-NMR spectrum of 1-(4-Methoxyphenyl)-1H-imidazole



Figure S15. ¹H-NMR spectrum of 1-(4-Methylphenyl)-1H-imidazole



Figure S16. ¹H-NMR spectrum of 1-(4-Chlorophenyl)-1H-imidazole



Figure S17. ¹H-NMR spectrum of 1-o-Tolyl-1H-imidazole



Figure S18. ¹H-NMR spectrum of 1-phenyl-1H-pyrrole



Figure S19. ¹H-NMR spectrum of 1-phenyl-1H-pyrazole



Figure S20. ¹H-NMR spectrum of 1-phenyl-1H-benzimidazole



Figure S21. ¹H-NMR spectrum of 1-(4-Methoxyphenyl)-1H-benzimidazole



Figure S22. ¹H-NMR spectrum of 1-(4-Fluorophenyl)-1H-imidazole



Figure S23. ¹H-NMR spectrum of 1-(4-Trifluoromethylphenyl)-1H-imidazole



Figure S24. ¹H-NMR spectrum of 1-(4-Methylphenyl)-1H-benzimidazole



Figure S25. ¹H-NMR spectrum of 1-(3-nitrophenyl)-1H-imidazole



Figure S26. ¹H-NMR spectrum of N-Phenylbenzamide



Figure S27. ¹H-NMR spectrum of 2-Phenylisoindoline-1, 3-dione



Figure S28. UV-visible spectra Cu-G composite after seven times reuse showing that it remains similar to as synthesized Cu-G.



Figure S29. Raman spectra of Cu-G composite after seven times reuse, showing two well documented D and G bands at 1315 and 1600 cm⁻¹ and band intensity ratio (I_D/I_G) of 2.35. Some shifting of D band is likely due to structural rearrangement of graphene.



Figure S30. Representative TEM image of Cu-G composites after seven times reuse.