

Electronic Supplementary Information

A Simple '*in situ*' Co-precipitation Method for Preparation of Multifunctional CoFe₂O₄-Reduced Graphene Oxide Nanocomposites: Excellent Microwave Absorber and Highly Efficient Magnetically Separable Recyclable Photocatalyst for Dye Degradation.

Debabrata Moitra,^a Madhurya Chandel,^a Barun Kumar Ghosh,^a Raj Kumar Jani,^b Manoj Kumar Patra,^b Sampat Raj Vadera,^b and Narendra Nath Ghosh,^{*}

^aNano-materials Lab, Department of Chemistry, Birla Institute of Technology and Science, Pilani K.K. Birla Goa Campus, Goa-403726, India.

^bDefence Lab, Jodhpur 342011, India.

Corresponding author. Tel. /fax: +91 832 2580318/2557033.

E-mail address: naren70@yahoo.com (N. N. Ghosh)

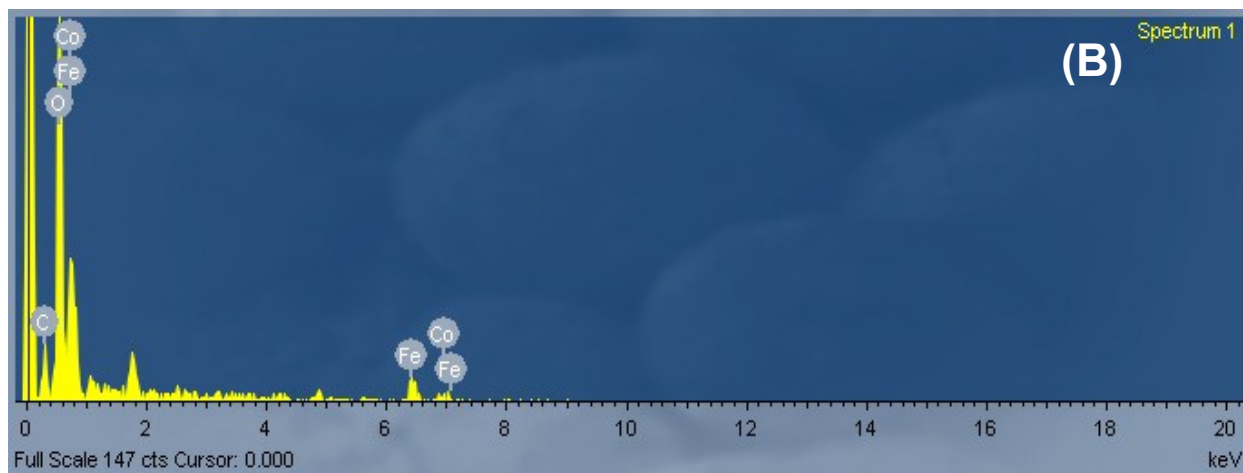
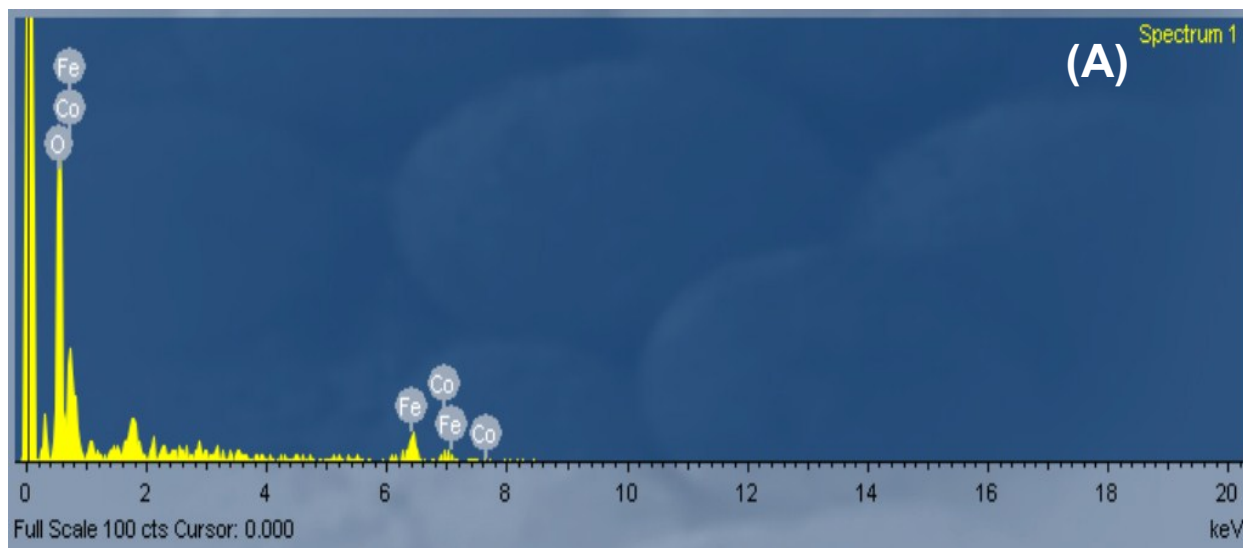


Fig S1. EDAX spectra of (A) synthesized CoFe_2O_4 nanoparticle and (B) CoFe_2O_4 -RGO nanocomposite.

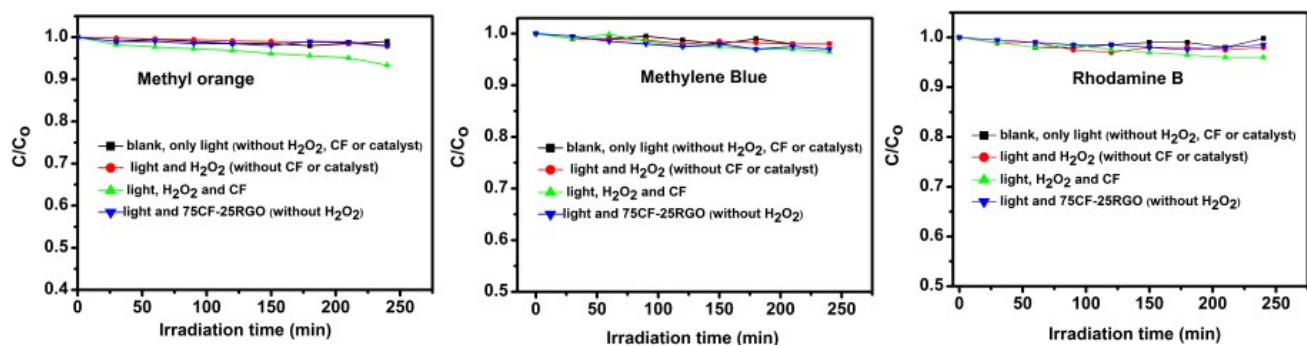


Fig S2: These C/C_0 vs. Irradiation time plots show that photocatalysis reactions of dyes do not occur in absence of CF-RGO and H_2O_2 .

Table S1: Microwave absorption properties of various ferrites and ferrite based composites

Samples	Preparation Method	Minimum Reflection loss	Corresponding Frequency (GHz)	Absorber thickness	Effective band width (RL < -10dB)	Ref.
NiFe ₂ O ₄	Hydrothermal at 180°C for 6h	-24 dB	9.5 GHz	3.5 mm	8.5 to 13 GHz	[11]
60 wt % ZnO/CoFe ₂ O ₄	Co-precipitation method 90°C	-28.3 dB	8.6 GHz	-	4.1 to 13 GHz	[12]
(Mn _{0.2} Ni _{0.4} Zn _{0.4} Fe ₂ O ₄) _x -(BaFe ₁₂ O ₁₉) _{1-x}	One pot precursor based method 800°C for 4h	-25 dB	8.2 GHz	3.5 mm	-	[28]
(NiFe ₂ O ₄) _{0.85} -(SrFe ₁₂ O ₁₉) _{0.15}	One pot precursor based method 800°C for 4h	-17 dB	8.2 GHz	3.2 mm	-	[29]
Co _{0.6} Zn _{0.4} Fe ₂ O ₄ nanofiber.	Calcination at 700°C for 2h	-16 dB	10.6 GHz	2.9 mm	7.8 to 16.8 GHz	[30]
Hollow glass microsphere/ CoFe ₂ O ₄	Co-precipitation method	-8.3 dB	18 GHz	1.5 mm	-	[31]
SrFe ₁₂ O ₁₉ / CoFe ₂ O ₄	Co-precipitation at 1200°C for 4h	-27.6 dB	10.8 GHz	-	8.2 to 9.6 GHz	[33]
N-RGO-Hematite wax composite	Refluxed in presence of PVP at 105°C for 24h	-78 dB	15.4 GHz	2 mm	11.3 to 18 GHz	[36]
ZnFe ₂ O ₄ /RGO	Hydrothermal synthesis at 180°C	-29.3 dB	16.7 GHz	1.6 mm	15.4 to 18 GHz	[47]
CoFe ₂ O ₄ hollow / Graphene composites	Vapour diffusion followed by calcination at 550°C	-18.5 dB	12.9 GHz	2 mm	11.3 to 15 GHz.	[48]
NiFe ₂ O ₄ nanorod-graphene	Solvothermal 150°C for 15h.	-29.2 dB	16.1 GHz	2.0 mm	13.6 to 18 GHz	[49]
CoFe ₂ O ₄ nanorod /Graphene	Hydrothermal at 150°C	-25.8 dB	16.1 GHz	2.0 mm	13.5 to 18 GHz.	[50]
(RGO)/CoFe ₂ O ₄ composite	One-pot hydrothermal route at 180°C for 10h	-47.9dB	12.4 GHz	2.3mm	12.4 to 17.4 GHz	[51]

Graphene-Fe ₃ O ₄	Hydrothermal at 200°C for 24h	-20 dB	16 GHz	5 mm	-	[52]
30 wt.% hollow Fe ₃ O ₄ /RGO	Solvothermal method. 200°C for 12h.	-24 dB	12.9 GHz	2.0 mm	10.8 to 15.7 GHz	[53]
RGO/CoFe ₂ O ₄	Hydrothermal at 180°C for 10h.	-47.9 dB	2.4 GHz	2.3 mm	12.4 to 17.4 GHz	[54]
5 wt. % filler RGO/MnFe ₂ O ₄ /PVDF	Hydrothermal at 140°C for 12h	-29 dB	9.2 GHz	3 mm	8 to 12.88 GHz	[55]
10 wt. % filler RGO/Co ₃ O ₄ /Poly(vinylidene fluoride)	Hydrothermal at 140°C for 12h.	-25 dB	11.6 GHz	4 mm	7.5 to 12 GHz	[56]
RGO/ Fe ₃ O ₄	One pot Co-precipitation at 80°C for 2h	-44.6 dB	6.6 GHz	3.9 mm	12.2 to 16.5 GHz	[61]
CoFe ₂ O ₄	Co-precipitation method 120°C for 12h	-55dB	9.25 GHz	2 mm	8.2 to 10.8 GHzGHz	[63]
50NZF-50RGO nanocomposite	In situ Co-precipitation method at 120°C for 12h	-19.99dB	11.58 GHz	1.8 mm	10.22 to 12.4 GHz	[68]
ZnFe ₂ O ₄ /ppy	Polymerization under 80°C for 8 h	-28.9 dB	10.8 GHz	2.7 mm	-	[70]
a) EG/PANI/ CoFe ₂ O ₄ with m _{CF} / m _{EG} / m _{PANI} of 0.8 b) CoFe ₂ O ₄	Co-precipitation method sintered at 300°C	a) -19.13 dB b)-13.58 dB	a)13.28 GHz b) 14.83 GHz	a) 0.5 mm b) 1 mm	a) 5.94GHz b)3.6 GHz.	[71]
CNT/ CoFe ₂ O ₄	Chemical Vapor Deposition	-18 dB	9 GHz	-	6.5 to 13.5 GHz	[72]
Rugby Shaped CoFe ₂ O ₄	Vapour diffusion followed by calcination at 550°C	-34.1 dB	13.4 GHz	2.5 mm	12.3 to 14.9 GHz.	[73]
CoFe ₂ O ₄ / SBA-15	Double solvent technique and incipient impregnation method	-18 dB	16 GHz	2 mm	13.5 to 18 GHz.	[74]

Polyaniline-CoFe ₂ O ₄	Citrate precursor based method at 1000°C	-21.5 dB	-	-	-	[75]
Ni _{0.5} Co _{0.5} Fe ₂ O ₄	Co-precipitation at 85°C	-18 dB	2.47 GHz	1.5 mm	-	[76]
Graphene@Fe ₃ O ₄ @SiO ₂ @NiO	Sol-Gel process with hydrothermal reaction followed by calcination at 400°C for 2h	-51.5 dB	14.6 GHz	1.8 mm	12.4-17.5 GHz	[77]
(Ni _{0.65} Zn _{0.35} Fe ₂ O ₄) _{0.85} -(BaFe ₁₂ O ₁₉) _{0.15}	One pot precursor based method 800°C for 4h	-21 dB	9.86 GHz	3.05 mm	-	[78]
Flaky graphite/cobalt zinc ferrite	Calcined at 700°C for 2h	-33.85 dB	11.7 GHz	2.5 mm.	10.3 to 13.5 GHz	[79]
BaFe ₁₂ O ₁₉ @RGO	Calcination at 900°C for 2h	-32 dB	-	3 mm.	-	[80]
BaFe _{11.92} (LaNd) _{0.04} O ₁₉ -TiO ₂ /MCNTs/wax composite	Sintering at 1200°C for 3h	-21.56 dB	11.04 GHz	2 mm	9.50 to 12.75 GHz	[81]
Fe ₃ O ₄ /C composite	Calcination at 600°C	-46 dB	12.8 GHz	2.9 mm	10.3 to 17.1GHz.	[82]
Fe ₃ O ₄ -Fe/G composite	Calcination at 420°C for 3h	-30 dB	4.72 GHz	2.0 mm	6.2 GHz	[83]
Graphene/Fe ₃ O ₄ incorporated polyaniline	Refluxing at 80°C for 5h	-26 dB	15 GHz	-	-	[84]
graphene/BaFe ₁₂ O ₁₉	Calcination at 1100°C for 1h	-58 dB	11.42 GHz.	-	-	[85]
85CF-15RGO	Co-precipitation reduction at 160°C at 16h	-31.13 dB	9.05 GHz	2.15 mm	8.2 to 10.92 GHz	This work

Table S2: Comparison of photocatalytic activity of 75CF-25RGO nanocomposite with those of the other reported RGO-Ferrite and RGO based nanocomposites.

Composite	Preparation method	Amount of catalyst	Dye concentration	Source	Time	Dye	Reference
TiO ₂ -Fe ₃ O ₄ -RGO composite	Hydrothermal at 105°C for 24h	0.01g in 40 ml	0.0535 mmol/L	Visible Light Xe lamp 500W	120 min	MB	[24]
Fe ₃ O ₄ -RGO -TiO ₂ composite	Hydrothermal at 200°C for 10h	1.5 g/L	10 mg/L	Visible light 300W UV-Vis Xenon Lamp>400nm	120 min	MB	[25]
CoFe ₂ O ₄ -TiO ₂	Hydrothermal at 150°C for 24h	20 mg in 5 ml	5 ppm aqueous solution (initial concentration)	UV light irradiation 8W UV -Hg lamp 5 sets	6 h	MB	[26]
CoFe ₂ O ₄ -RGO (10wt. %)	Hydrothermal at 180°C for 12h	20 mg in 200 ml	10 mg/L	Tungsten halogen lamp 500 W	180 min	MB	[38]
CoFe ₂ O ₄ -RGO (45 wt. %)	Ball milling for 6h at 25 sec ⁻¹	0.01g /40mL	20mg/L	Visible light Xe lamp 800W cutoff λ >420nm	180 min	MB, RhB, and MO	[39]
(3%)P25/ CoFe ₂ O ₄ /RGO	Hydrothermal at 180°C for 20h	10 mg in 30 ml	40 mg/L	Visible light Xe lamp 500W cutoff λ >420nm	240 min	MB	[40]
CoFe ₂ O ₄ -RGO (40 wt. %)	Hydrothermal at 180°C for 20h	25 mg in 100 ml	20mg/L	Visible light Xe lamp 500W cutoff λ >420nm	240 min	MB (100%) MO (94%) RhB (71%)	[41]

Composite	Preparation method	Amount of catalyst	Dye concentration	Source	Time	Dye	Reference
ZnFe ₂ O ₄ -RGO (20 wt. %)	Hydrothermal at 180°C for 12h	0.05g in 100 ml	20 mg/L	Visible light Xe lamp 500W cutoff $\lambda > 420\text{nm}$	90 min	MB	[42]
MnFe ₂ O ₄ -graphene (30 wt. %)	Hydrothermal at 180°C for 20h	0.025 g in 100 mL	20 mg/L	500 W mercury and xenon lamp cutoff $\lambda > 420\text{nm}$ Visible light	360 min	MB	[43]
RGO-MFe ₂ O ₄ (M=Zn, Co and Ni)	Hydrothermal at 200°C for 10h	20 mg in 150 ml	10 mg/L	Tungsten lamp 500 W	180 min	RhB and MB	[44]
CdFe ₂ O ₄ /Graphene	Hydrothermal at 180°C for 12h	100 mg in 100 ml	10 mg/L	Xe lamp 500W cutoff $\lambda > 420\text{nm}$ Visible light	240 min	MB	[45]
Bi ₂ Fe ₂ O ₉ /RGO (4.5 wt. %)	Hydrothermal at 95°C 36h	0.5mg/L	10mg/L	150W Xenon arc Lamp Visible light	3h	Bisphenol-A	[46]
Graphene-SnO ₂ aerosol nanocomposite	Hydrothermal at 180°C for 12h	100 mg in 200 ml	4.79×10^{-5} g/L	300W Xenon Lamp cutoff $\lambda > 400\text{nm}$ Visible light	40 min	RhB	[88]
75CF-25RGO	Co-precipitation followed by reflux at 160°C for 16h	500 mg/L	20 mg/L	100 W Reading lamp Visible light	i) 60 min ii) 75 min iii) 45 min iv) 120 min	i) MO ii) MB iii) RhB iv) Dye mixture	This work

MB= Methylene Blue, MO= Methyl Orange and RhB= Rhodamine B