

Supporting Information

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Microwave-assisted preparation of graphene quantum dots immobilized nanosilica as an efficient heterogeneous nanocatalyst for the synthesis of xanthenes

Somayeh Pahlavan Moghanlo and Hassan Valizadeh^{1*}

Department of Chemistry, Faculty of Sciences, Azarbaijan Shahid Madani University, Tabriz, Iran

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*Corresponding author: Tel: +98-41-34327520, Fax: +98-4134327522; P.O. Box 53714-161; E-mail address: hvalizadeh2@yahoo.com

S.1. Spectral and analytical data of selected compounds

14-Phenyl-14H-dibenzo [a, j] xanthene (2a): Yield 94%, mp. 188-190 °C. FTIR (KBr, cm⁻¹): 3072 (Ar-H stretch), 2922 (C-H stretch), 1618 (C=C stretch). ¹H NMR (CDCl₃, 300 MHz): δ= 8.11 (d, 2H, J = 8.2 Hz), 7.77 (t, 4H, J = 9.3 Hz), 7.58-7.38 (m, 8H), 7.00 (t, 2H, J = 7.5, 7.2 Hz), 7.00 (t, 1H, J = 7.1 Hz) and 6.47 (s, 1H). ¹³C NMR (CDCl₃, 75 MHz): δ= 147.25, 142.90, 130.42, 130.87, 126.70, 126.76, 126.38, 127.20, 125.56, 126.11, 123.10, 120.63, 116.28, 115.38 and 37.42 ppm.

14-(4-Nitrophenyl)-14H-dibenzo [a, j] xanthene (2b): Yield 98%, mp. 314-316 °C. FTIR (KBr, cm⁻¹): 3068 (Ar-H stretch), 2924 (C-H stretch), 1635 (C=C stretch). ¹H NMR (CDCl₃, 300 MHz): δ= 8.00 (d, 2H, J = 8.2 Hz), 7.91 (d, 2H, J = 8.5 Hz), 7.5-7.38 (m, 4H), 7.56-7.40 (m, 8H) and 6.20 (s, 1H). ¹³C NMR (CDCl₃, 75 MHz): δ= 140.18, 127.39, 126.18, 127.10, 126.87, 126.02, 125.98, 123.10, 122.88, 120.84, 120.45, 115.25, 113.48 and 33.37 ppm.

14-(4-Cyanophenyl)-14H-dibenzo [a, j] xanthene (2c): Yield 98%, mp. 293-294 °C. FTIR (KBr, cm⁻¹): 3072 (Ar-H stretch), 2921 (C-H stretch), 2220 (C≡N stretch), 1618 (C=C stretch). ¹H NMR (CDCl₃, 300 MHz): δ= 8.25 (d, 2H, J = 8.4 Hz), 7.80 (m, 4H), 7.61-7.39 (m, 10H) and 6.52 (s, 1H). ¹³C NMR (CDCl₃, 75 MHz): δ= 150.00, 147.87, 131.80, 130.51, 130.30, 128.90, 129.59, 128.29, 126.68, 124.12, 122.47, 117.91, 117.38, 115.75, 109.31 and 36.76 ppm.

14-(4-Trifluoromethylphenyl)-14H-dibenzo [a, j] xanthene (2d): Yield 98%, mp. 260-261 °C. FTIR (KBr, cm⁻¹): 3055 (Ar-H stretch), 2959 (C-H stretch), 1616 (C=C stretch). ¹H NMR (CDCl₃, 300 MHz): δ= 8.28 (d, 2H, J = 8.1 Hz), 7.83-7.77 (m, 4H), 7.60-7.38 (m, 10H) and 6.52 (s, 1H). ¹³C NMR (CDCl₃, 75 MHz): δ= 148.81, 146.56, 129.22, 129.00, 128.09, 127.51, 126.08, 126.89, 125.31, 123.52, 123.48, 122.78, 121.19, 116.08, 114.75 and 35.18 ppm.

14-(3-Nitrophenyl)-14H-dibenzo [a, j] xanthene (2e): Yield 95%, mp. 212-214 °C. FTIR (KBr, cm⁻¹): 3080 (Ar-H stretch), 2924 (C-H stretch), 1620 (C=C stretch), 1527 (N=O stretch), 1455 (C-H bend), 1342 (N=O bend). ¹H NMR (CDCl₃, 300 MHz) δ= 8.39 (s, 1H), 8.28 (d, 2H, J = 8.1 Hz), 7.85-7.78 (m, 6H), 7.61 (t, 2H, J = 7.3, 7.6 Hz), 7.51-7.39 (m, 4H), 7.29-7.25 (m, 1H) and 6.58 (s, 1H). ¹³C NMR (CDCl₃, 75 MHz) δ= 148.11, 147.56, 146.32, 133.69, 130.40, 128.87, 128.91, 128.38, 126.64, 123.96, 120.45, 121.38, 121.08, 117.48, 115.24 and 37.07 ppm.

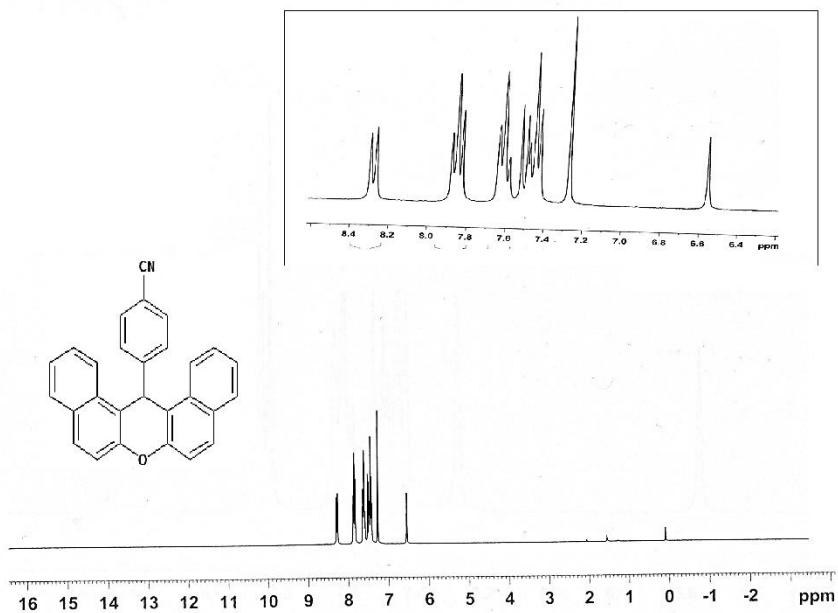


Figure S1. 400 MHz ^1H -NMR spectrum of compound **2c**

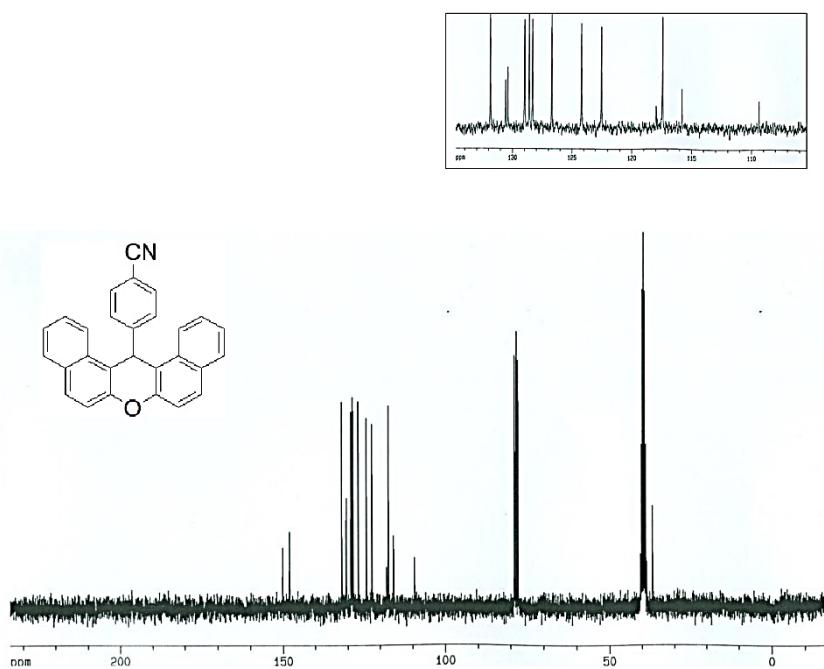


Figure S2. 100 MHz ^{13}C -NMR spectra of compound **2c**

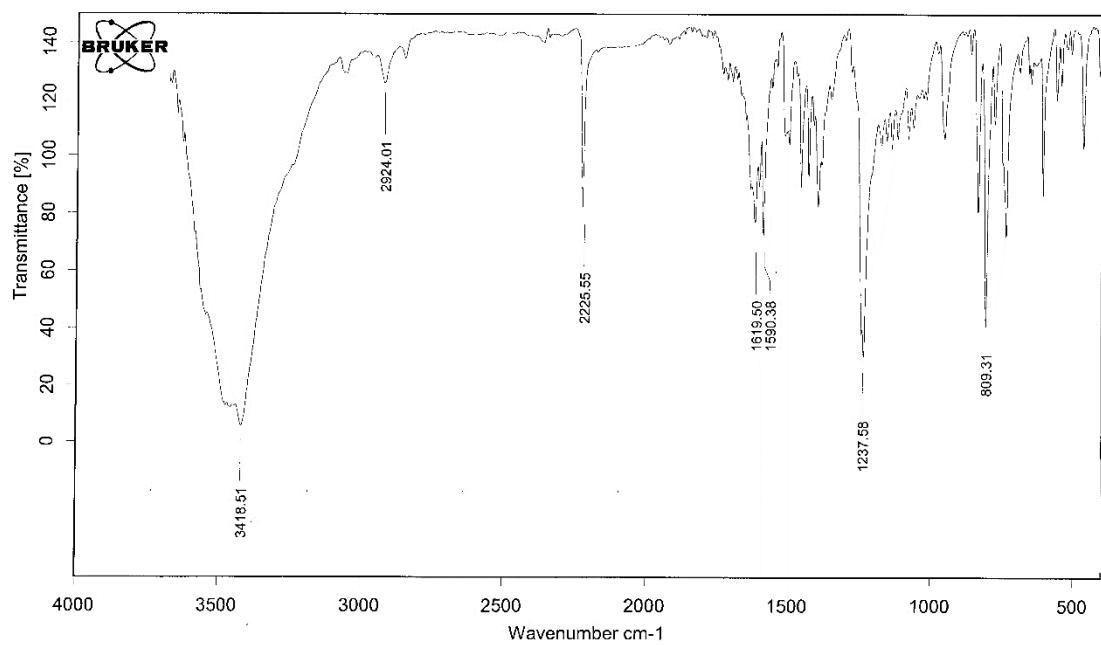


Figure S3. FT-IR spectrum of compound **2c**

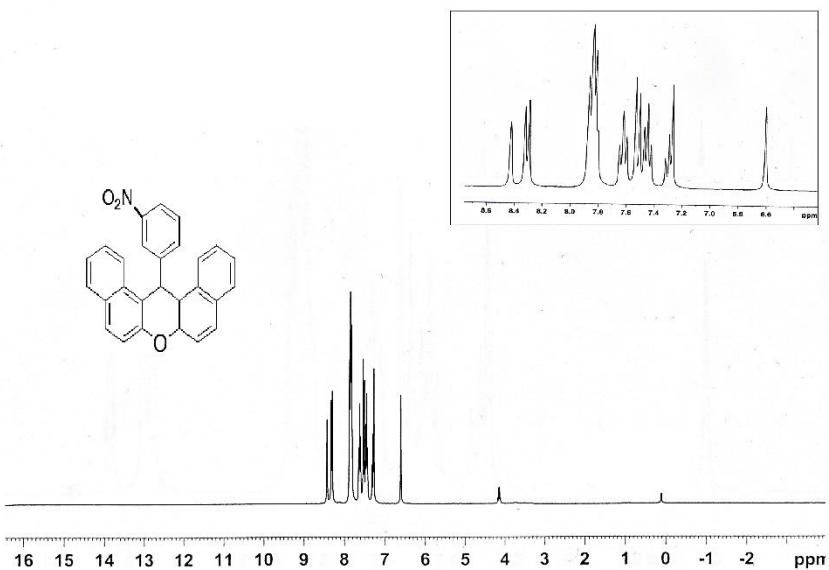


Figure S4. 400 MHz ^1H -NMR spectrum of **2e**

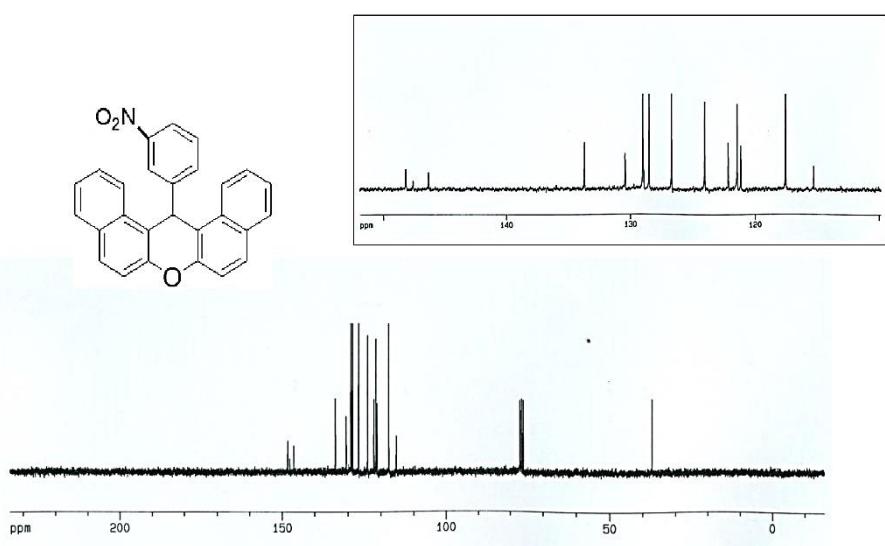


Figure S5. 100 MHz ^{13}C -NMR spectrum of **2e**

Figure S6. FT-IR spectrum of compound **2e**

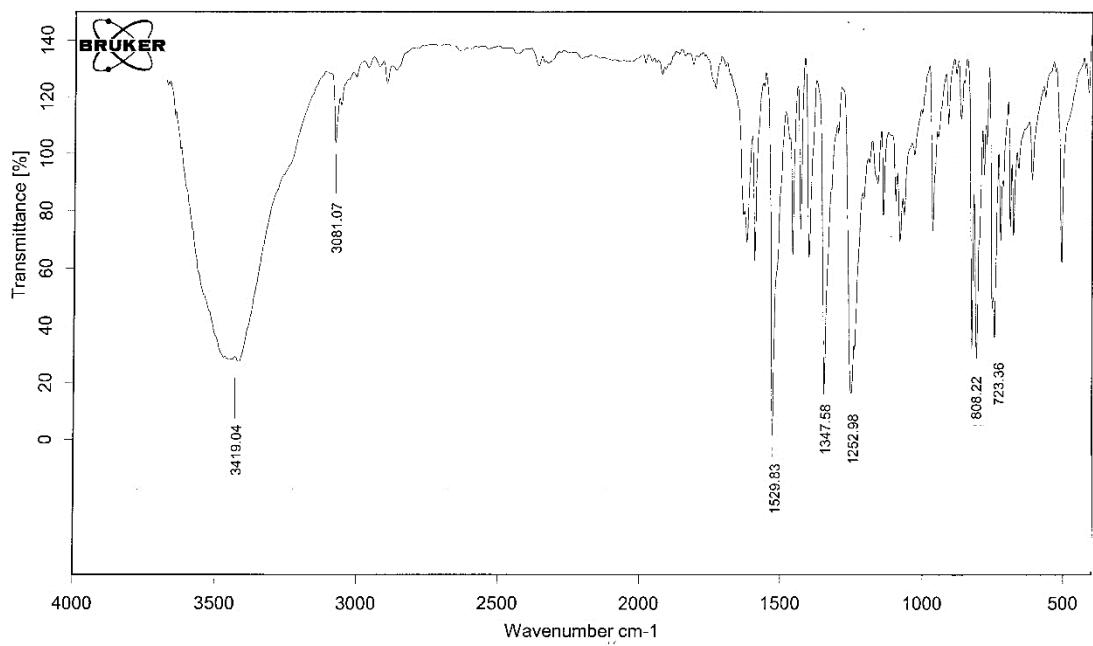


Figure S6. FT-IR spectrum of compound **2e**