

Carbon Nanotube/Lithium Ferrite Nanocomposite: Magnetic and Electrochemical Optimization for Enhanced H₂O₂ Sensing

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Table 1S. The assessed crystallinity of the prepared nanomaterials

nanomaterials	Area _C (crystalline peaks)	Area (all peaks)	Crystallinity (X _C) %
CNTs	1845.24	5187.4	26.23
Li ₂ Fe ₃ O ₅ (LFO)	2093.6	5152.6	28.89
CNTs/LFO (0.5%)	2592.4	6056.7	29.97
CNTs/LFO (1%)	3117.5	6451.5	32.57
CNTs/LFO (2%)	2746.9	5913	31.71

Table 2S. The structural parameters computed for the nanoparticles that were synthesized

structure	D _v (nm) From Sherrer's analysis	δ (nm ⁻²) dislocation density	D _v (nm) From W-H analysis	(ε) Strain from W-H analysis
CNTs	2.5	0.16	2.1	1
Li ₂ Fe ₃ O ₅ (LFO)	23.73	0.0017	22.5	0.06882
CNTs/LFO (0.5%)	24.42	0.00167	23.4	0.53
CNTs/LFO (1%)	24.35	0.00168	23.5	0.51
CNTs/LFO (2%)	24.49	0.00166	23.9	0.03713

Table 3S. The magnetic parameters received from the magnetic hysteresis loop

Structure	Saturation magnetization (emu/g)	Hc (left) (G)	Hc (right) (G)	Coercivity (G)	Remanent magnetization (emu/g)	Squareness
CNTs	0.6	-691.3	1109.2	900	0.187	0.311
LFO	35.77	-16.30	27.35	21.83	3.18	0.088
CNTs/LFO (0.5%)	14.40	-21.64	40.58	31.11	1.35	0.093
CNTs/LFO (1%)	20.95	-35.21	33.73	34.47	4.5	0.214
CNTs/LFO (2%)	25.07	-17.14	78.64	47.89	2.29	0.091

Table 4S. The modified electrodes using the produced nanomaterials yielded the electrochemical parameters (CV & EIS). These values are taken from the impedimetric and voltammetric studies mentioned above.

Electrode type	I_a (μA)	I_c (μA)	$E_{\text{oxd.}}$ (V)	$E_{\text{red.}}$ (V)	ΔE_p (V)	R_s (Ω)	$R_{\text{ct (1)}}$ (Ω)	C (μF)	W (Ω)	SC (F/g)
CNTs	225.6	-238.6	0.313	0.0015	0.157	74.6	80.5	121.6	171.4	121.61802
LFO	127.04	-135.12	0.334	0.0152	0.174	33.4	1150.3	72.5	189.6	52.54385
CNTs-LFO 0.5	349.5	-357.6	0.329	-0.040	0.144	112.6	50.3	136.2	72.6	118.02871
CNT-LFO 1.0	179.01	-192.10	0.22	0.0711	0.144	214.3	205.9	112.5	94.2	88.89111
CNTs-LFO 2.0	168.6	-176.7	0.356	-0.102	0.127	391.8	700.9	95.3	168.2	83.83291

Table 5S. Comparison for different electrodes used for hydrogen peroxide (H_2O_2) detection, based on applied potential, linear range, and detection limit.

Materials	Electrode type	Buffer	pH	Applied Potential	Linear range (μM)	Detection Limit (μM)	Ref.
CNT-NiCo₂O₄	SPE	PBS	7.4	0.7	2.5-275	0.01	¹
NiCo₂O₄/RGO	GCE	PBS	7.0	-0.4	5-3000	0.41	²
MnCo₂O₃/CNTs	SPE	PBS	7.4	0.7	0.1-180	0.1	³
rGo-Pt	GCE	PBS	7.0	-0.08	0.5-3475	0.2	⁴
Co₃O₄	SPE	-	-	1.0	0.1-50	0.145	⁵
CoFe₂O₄/ CNTs	GCE	PBS	7.0	0.3	0.5-50	0.05	⁶
Pt/rGO-CNT	Paper electrode	-	-	-0.25	0.1-25	0.1	⁷
Pol(azureA)-PtNPs	SPE	PBS	7.4	0.1	0-300	0.052	⁸
Co₃O₄-rGO	GCE	-	-	-0.19	15-675	2.4	⁹
CNTs/LFO	SPE	PBS	7.4	0.7	0.01 -500	0.005	This work

Table 6S. Peroxide detection in real waste water samples using CNTs/LFO sensor.

peroxide concentration μM		RSD (%)	Recovery (%)
Added	Found		
5	5.1	0.19	102
10	9.8	0.62	98
20	19.9	1.38	99.5
30	29.7	2.96	99

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