



Synergistic Design of Co₃O₄ Nanoparticles for Advanced Supercapacitor Electrodes: Hydrothermal Synthesis, Characterization, and Performance Evaluation

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Authors' contributions

This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.

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ABSTRACT

The potential of cobalt oxide (Co₃O₄) nanoparticles, synthesized via a facile hydrothermal method, as electrode materials for supercapacitors is investigated in this research. Through a comprehensive characterization approach involving X-ray diffraction (XRD), Fourier-transform infrared spectroscopy (FT-IR), scanning electron microscopy (SEM), surface wettability analysis, and Brunauer-Emmett-Teller (BET) analysis, the structural and morphological properties of the synthesized Co₃O₄ nanoparticles were thoroughly examined. XRD analysis confirmed the presence of the cubic phase of Co₃O₄, while FT-IR spectroscopy revealed the characteristic Co-O bonds. SEM imaging showcased the non-uniform aggregation of nanoparticles. BET analysis provided crucial insights into surface area and pore radius parameters. Notably, Co₃O₄ nanoparticles exhibited hydrophilic behavior. In the presence of KOH electrolyte, Co₃O₄-carbon cloth (CC) electrodes demonstrated exceptional specific capacitances (Cs) of 132 Fg⁻¹ in KOH and 79 Fg⁻¹ in Na₂SO₄, coupled with outstanding cycling stability (~75% retention after 500 cycles) in KOH, underscoring their potential as promising electrode materials for supercapacitor applications.

Keywords: Nanotechnology; Co₃O₄ nanoparticles; hydrothermal method; characterization; supercapacitor application.

1. INTRODUCTION

Nanoparticles' special qualities and wide range of uses have made them indispensable in many scientific and technical domains [1]. Exact control over the production and design of nanoparticles with specific properties has been made possible by nanotechnology, which is the manipulation and engineering of materials at the nanoscale [2]. These nanoscale structures are essential in a variety of fields, including electronics, medicine, catalysis, and energy storage, because they display unique physical, chemical, and biological characteristics [3].

Nanotechnology has completely changed the energy storage market by providing creative answers to the growing need for effective and environmentally friendly energy storage technologies [4]. Supercapacitors are one of these alternatives that has attracted a lot of attention because of its high power density, quick charging and discharging speed, and extended cycle life [5]. Supercapacitors are essential components of contemporary energy systems, finding use in anything from electric automobiles and portable devices to grid-level energy storage [6].

In an attempt to improve the efficiency of supercapacitors, scientists are investigating new electrode materials, and nanoparticles have surfaced as a serious candidate [7]. To increase the energy storage capacity and electrochemical performance of supercapacitors, nanoparticles with special properties including high surface area, variable porosity, and superior electrical conductivity are greatly desired. Scientists hope

to advance supercapacitor technology toward greater dependability and efficiency by utilizing the advantages of nanoparticles [8].

Co₃O₄ nanoparticles can be made using a variety of techniques, including as hydrothermal synthesis, chemical vapor deposition (CVD), sputtering, electrochemical processes, hydrothermal synthesis, sol-gel synthesis, and chemical bath deposition [9]. The capacity to create Co₃O₄ nanoparticles with various morphologies and structures which can greatly improve their electrochemical properties, makes hydrothermal synthesis stand out among these techniques [10].

A variety of techniques, including sol-gel, hydrothermal, and chemical vapor deposition, are commonly used in the production of nanoparticles, including Co₃O₄ [11]. Each technique has certain benefits with regard to control over size, shape, and crystallinity. In particular, hydrothermal synthesis has become more and more popular because of its ease of use, scalability, and capacity to provide nanoparticles with consistent morphology and size [12]. This process creates nanoparticles with the required characteristics by reacting precursor materials in an aqueous solution at high pressures and temperatures [12].

In order to understand the structural, morphological, and electrochemical characteristics of nanoparticles and whether or not they are suitable for a given application, characterization procedures are essential. X-ray diffraction (XRD), scanning electron microscopy

(SEM), Fourier-transform infrared spectroscopy (FTIR), and electrochemical impedance spectroscopy (EIS) are examples of common characterisation techniques. By examining the crystal structure, surface morphology, chemical makeup, and electrochemical activity of nanoparticles, researchers can better optimize synthesis procedures and create materials that are specifically suited for intended uses [13].

One interesting way to improve the performance of supercapacitors is to use Co_3O_4 nanoparticles as the electrode material [14]. Co_3O_4 nanoparticles are ideally suited for effective energy storage due to their high specific capacitance, great chemical stability, and outstanding electrical conductivity. Furthermore, the development of supercapacitors with improved energy density, power density, and cycling stability is made possible by the special qualities of Co_3O_4 in combination with the benefits of electrode designs based on nanoparticles [15]. Through the utilization of sophisticated electrode design methodologies and the synergistic impacts of nanoscale materials, researchers hope to fully realize the promise of Co_3O_4 nanoparticles for next-generation supercapacitor technologies. Comparing Co_3O_4 to other metal oxides, there are several benefits: low toxicity, economical, high chemical and thermal stability, readily available, and environmentally benign [16]. Co_3O_4 is a good option for energy storage applications because of its substantial theoretical specific capacitance (Cs) within a potential window of 0.5 V [15].

Batteries, power modules, and electrochemical capacitors (supercapacitors) are among the most practical and beneficial developments for electrochemical energy storage and processing in a variety of applications [17]. Because of their high energy thickness, excellent charge-release rate, extended cycle life (more than 5,000 000 cycles), excellent stability, and wide range of applications such as power reinforcement systems, lightweight, versatile devices, and hybrid electric vehicles supercapacitors have become increasingly commonplace in recent years [18].

This work focuses on synthesizing Co_3O_4 nanoparticles using a straightforward hydrothermal technique and examines their electrochemical characteristics in several electrolytes, including sodium sulfate (Na_2SO_4) and potassium hydroxide (KOH). One can learn

more about the applicability and potential of Co_3O_4 nanoparticles for supercapacitor applications by methodically examining their performance as electrode materials in different electrolytes.

Due to the growing global interest in energy, the rapidly expanding global economy, the decline in the use of petroleum derivatives, the growing concern over environmental issues, the growing need for convenient electronic devices, and the rising popularity of crossover vehicles, there is an urgent need for clean, sustainable, eco-friendly, high energy stockpiling and transformation innovation [19].

2. EXPERIMENTAL DETAILS

2.1 Synthesis of Co_3O_4 Powder by Hydrothermal Route

A set of procedures was followed in the hydrothermal synthesis of Co_3O_4 nanoparticles in order to create porous Co_3O_4 powder. Every reagent used in the synthesis procedure was of analytical quality. The standard protocol involved adding 1 mol of cobaltacetate $\text{Co}(\text{CH}_3\text{CO}_2)_2 \cdot 4\text{H}_2\text{O}$ to a combination of 5 ml of double-distilled water and 15 ml of methanol, then continuously swirling with a magnetic stirrer for 15 minutes. After being fully dissolved, the transparent solution was put into a 20 ml Teflon container and heated to 125°C for four hours using hydrothermal treatment. After letting the autoclave cool naturally, the finished product was gathered and thoroughly cleaned with ethanol and water to get rid of any remaining impurities. The powder that had been gathered was then heated to 60°C and then calcined for two hours at 350°C in an airtight environment [20].

2.2 Preparation of Working Electrode

A mixture of 80 weight percent Co_3O_4 electro-active material, 10 weight percent carbon black, and 10 weight percent PVDF binder (polyvinylidene difluoride) in NMP solvent (1-methyl-2-pyrrolidinone) was premixed to create the working electrode. After the combination was well mashed to guarantee homogeneity, the resultant slurry was applied to a $1 \times 1 \text{ cm}^2$ flexible carbon cloth (CC) substrate. The coated electrode's mass loading of active materials was kept constant at 41.8 mgcm^{-2} . The coated electrode was then dried at 60°C to eliminate any remaining solvent and promote the active

components' adherence to the carbon cloth substrate [21].

This method of manufacture guaranteed even dispersion of the Co_3O_4 nanoparticles in the electrode matrix, promoted electron transport, and gave the electrode mechanical stability. The electrode's overall electrochemical performance and electrical conductivity were both improved by the addition of carbon black. Furthermore, by adhering the active components to the carbon cloth substrate, PVDF binder helped to prevent their separation during electrochemical cycling [22]. The resulting Co_3O_4 -based working electrode showed superior electrical conductivity [23], effective charge transfer kinetics, and a high specific surface area, all of which are desirable properties for supercapacitor applications [24]. Due to these characteristics, the electrode can be used in high-end supercapacitor systems that provide improved long-term cycle stability and energy storage [25].

3. CHARACTERIZATION OF NANOPARTICLES

3.1 FTIR

Using infrared spectroscopy, the chemical bonding properties of materials are assessed. The FT-IR spectra of the as-prepared nanocrystalline Co_3O_4 made by hydrothermal decomposition of cobaltacetate $\text{Co}(\text{CH}_3\text{CO}_2)_2 \cdot 4\text{H}_2\text{O}$ [26]. In addition to the prominent bands indicating the presence of Co_3O_4 spinel oxide and the vibrational modes

associated with its chemical bonding, further insights can be gleaned from the infrared spectroscopic analysis of the as-prepared nanocrystalline Co_3O_4 . The observed peaks at approximately 660 cm^{-1} and 570 cm^{-1} correspond to the $\nu(\text{CoO})$ vibrational modes, which signify the stretching vibrations of the cobalt-oxygen bonds within the crystal lattice. These peaks serve as unequivocal evidence for the formation of the desired Co_3O_4 phase through the hydrothermal decomposition process employed. Furthermore, the broad absorption band centered at around 3424 cm^{-1} can be attributed to the O-H stretching vibrations of water molecules adsorbed onto the surface of the nanoparticles. This phenomenon is common in materials synthesized via wet chemical methods, such as hydrothermal synthesis, where residual water molecules often remain entrapped within the nanostructure. Similarly, the weak absorption peak at 1618 cm^{-1} corresponds to the bending vibrations of adsorbed water molecules, further confirming the presence of surface-bound water species. The detection of these water-related peaks underscores the need for thorough characterization and understanding of the surface chemistry in nanomaterial synthesis [27].

Overall, the FT-IR analysis provides valuable insights into the chemical composition and bonding properties of the Co_3O_4 nanoparticles, crucial for elucidating their structure-property relationships and informing their potential applications in various fields, including catalysis, sensing, and energy storage [26].

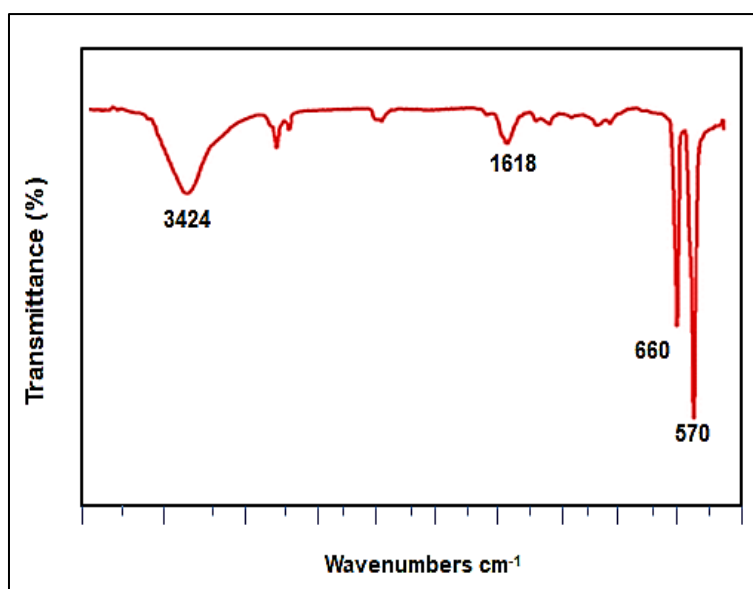


Fig. 1. FTIR spectra of Co_3O_4 nanoparticles

3.2 XRD

The XRD pattern of the synthesized Co_3O_4 nanoparticles revealed distinct diffraction peaks at specific 2θ angles, indicative of their crystalline nature and structural properties [28]. The peaks corresponding to the (110), (222), (520), (190), (400), and (422) crystallographic planes were observed at 2θ angles of 19.01° , 32.34° , 60.53° , 21.87° , 44.67° , and 56.32° , respectively. Each peak's position and intensity provide valuable information about the crystal structure and phase composition of the material [28]. The sharp, well-defined peaks suggest a high degree of crystallinity and phase purity, consistent with the formation of phase-pure Co_3O_4 . This observation is further supported by the absence of additional peaks associated with impurity phases, as verified by comparison with the standard JCPDS No. 76-1802. Phase pure Co_3O_4 has formed, as confirmed by the lack of diffraction peaks associated with CoO and other phases [29]. This is consistent with standard JCPDS No. 76-1802. Using the Debye-Scherrer equation:

$$0.9 D = (\cos \lambda) / (\beta \theta) \quad \text{eq.(1)}$$

where λ is the wavelength of Cu- α radiation (1.5418 \AA), β is the full width half-maximum (FWHM) in radians, and θ is the angle of diffraction (in radians), the average crystallite size was determined.

β (FWHM in radians) = convert from degrees to radians

θ (angle of diffraction in radians) = convert from degrees to radians

Let's say for the (110) peak at $2\theta = 19.01^\circ$:

Convert 2θ from degrees to radians: $\theta = 19.01^\circ \times \pi/180 \approx 0.332$ radians

Convert FWHM from degrees to radians: $\beta = 0.1^\circ \times \pi/180 \approx 0.00175$ radians

Now, plug these values into the equation:

$$0.9 D = (\cos \lambda) / (\beta * \theta)$$

$$D \approx (\cos 1.5418) / (0.00175 * 0.332)$$

Calculating:

$$D \approx (0.02055) / (0.000581)$$

$$D \approx 35.38 \text{ \AA (angstroms)}$$

So, for the (110) peak at $2\theta = 19.01^\circ$, the estimated average crystallite size is approximately 35.38 angstroms.

This process can be repeated for each diffraction peak observed in the XRD pattern to obtain the average crystallite size corresponding to different crystallographic planes, providing comprehensive insight into the nanostructural characteristics of the Co_3O_4 nanoparticles. The peak positions correspond to specific lattice planes within the face-centered cubic spinel structure of Co_3O_4 , as described by the $Fd\bar{3}m$ space group. The (110) peak at 19.01° , for instance, represents the interplanar spacing of the (110) crystallographic planes, while the (222) peak at 32.34° corresponds to the (222) planes, and so forth.

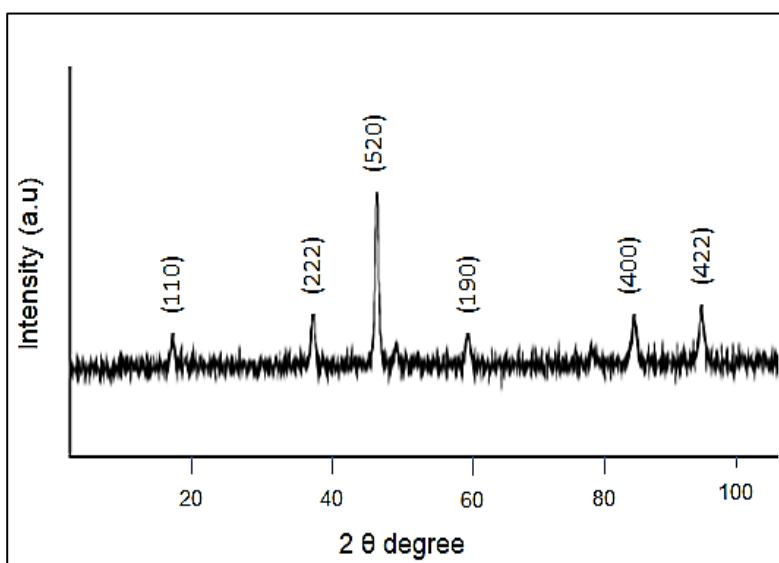


Fig. 2. XRD pattern of Co_3O_4 nanoparticles

3.3 SEM

The shape and size distribution of the Co_3O_4 nanoparticles, which were created by hydrothermally breaking down the cobalt acetate precursor, are better understood according to the SEM study [30]. The nanoparticles' asymmetrical form and surface characteristics are visible in the SEM pictures, which are a reflection of their complex growth mechanism and surface interactions during synthesis [31].

Furthermore, the Co_3O_4 nanoparticles' homogeneity and narrow size distribution, which ranges from 20 to 150 nm are shown in the SEM pictures. This uniform distribution of particle sizes is a sign of the regulated kinetics of nucleation and growth that are attained in hydrothermal settings, where temperature, pressure, and concentration of precursors are important variables that affect the final morphology of the nanostructure [32].

Furthermore, the observed sizes of the nanoparticles within the given range are very desirable for a variety of applications, such as energy storage, sensing, and catalysis, where the performance of the particles is often determined by their size. Through hydrothermal synthesis, Co_3O_4 nanoparticles' size and morphology may be accurately controlled, providing chances to modify their properties to suit particular application needs [33].

3.4 BET Characterization

BET characterization was done to look at the surface properties of the Co_3O_4 nanoparticles [34]. The CoO adsorption-desorption isotherms

for Co_3O_4 and the corresponding pore size distribution, which was investigated with the Barrett-Joyner-Halenda (BJH) method [35], are depicted in the figure. Co_3O_4 nanoparticle isotherms in the 0–1 high relative pressure (P/P0) range exhibit a hysteresis loop. The slope in the presented figure grows from 0.4 to 0.9 at high relative pressures, suggesting that the material is mesoporous.

The maximum BET surface area for Co_3O_4 nanoparticles was found to be $28.4 \text{ m}^2/\text{g}$. Within their nanocluster structure, Co_3O_4 nanoparticles have an average pore radius of 17.5 nm in the porous zone. Co_3O_4 nanoparticles' porous nanocluster form improves reaction kinetics by offering porosity, more active sites, and rapid charge-discharge capabilities. This makes them useful in applications such as supercapacitors.

3.5 Surface Wettability Test

Surface wettability was assessed to look at how the Co_3O_4 electrode and the electrolyte interacted [36]. A large contact angle ($\theta > 90^\circ$), indicating a hydrophobic surface, suggests lower wettability [37]. Conversely, a short contact angle ($\theta < 90^\circ$) indicates a hydrophilic surface and suggests more wettability. Hydrophilic materials usually show high surface energy. In this experiment, the contact angle between the water and the Co_3O_4 electrode surface was discovered to be 52° , as seen in the accompanying figure. One essential characteristic of nanocrystalline materials intended for use as supercapacitor electrodes is hydrophilicity. This result is in line with studies that looked at the hydrophilic properties of copper oxide multilayer nanosheets [38].

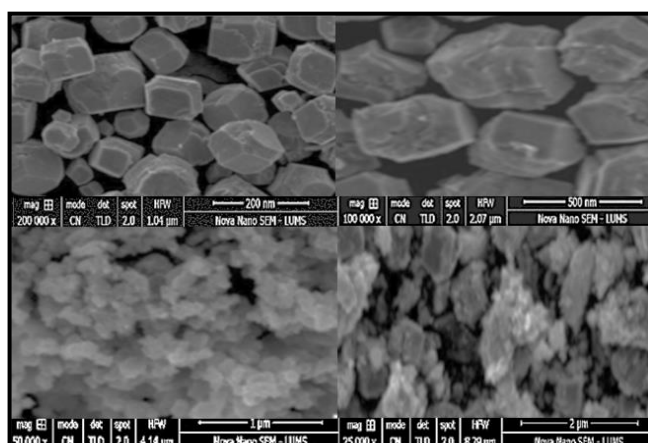


Fig. 3. SEM micrographs of Co_3O_4 nanoparticles at different resolution of 1 μm , 2 μm , 200nm and 500nm

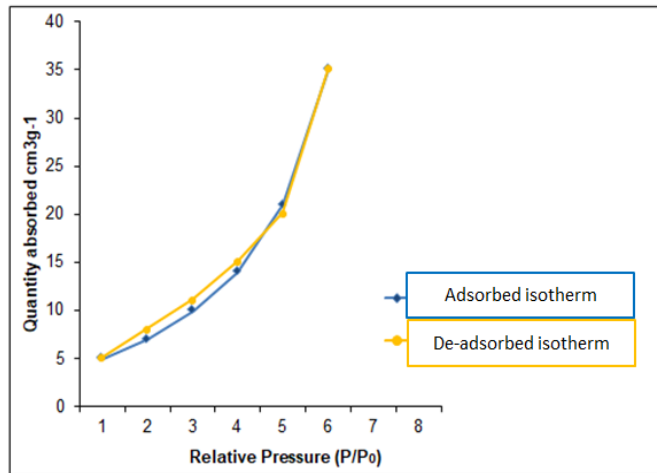


Fig. 4. CoO adsorption–desorption isotherms for Co₃O₄

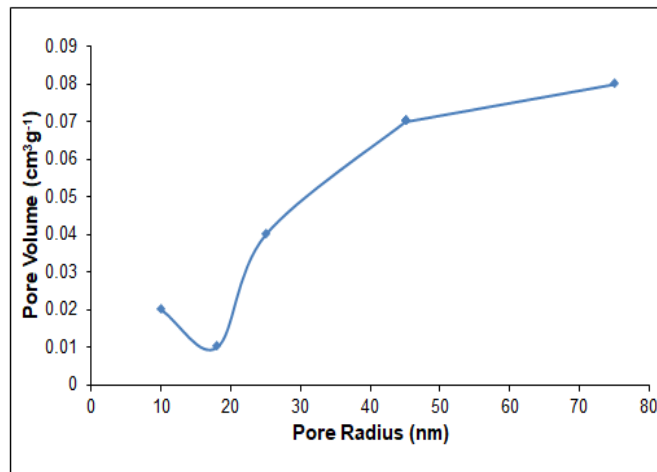


Fig. 5. Plot of pore volume vs. pore radius

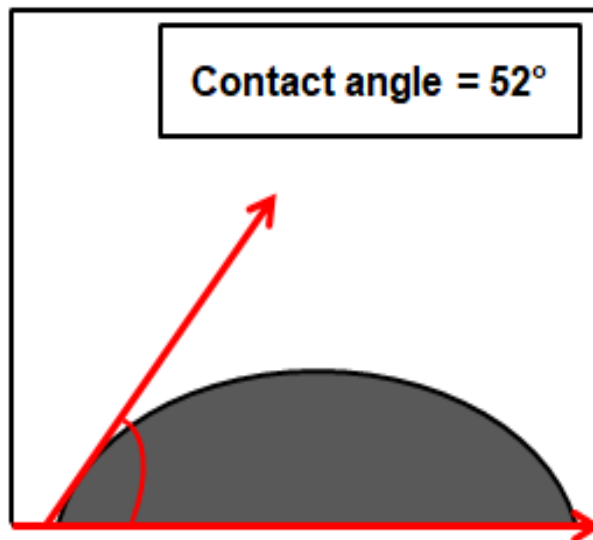


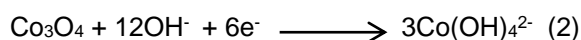
Fig. 6. Static contact angle measurement of CoO electrode

4. ELECTROCHEMICAL MEASUREMENTS

4.1 Electrochemical Measurements of Co₃O₄ Nanoparticles as Electrode

The Co₃O₄ nanoparticle electrode was electrochemically characterized using a conventional three-electrode setup. The electrode arrangement consisted of a platinum counter electrode, a SCE (Standard Calomel Electrode) reference electrode, and a Co₃O₄ nanoparticle working electrode [39]. For the measurements, one million KOH alkaline solution and one million Na₂SO₄ neutral solution were utilized.

The CV curves in KOH electrolyte at a scan rate of 5 mVs⁻¹ for both bare carbon cloth (CC) and CC modified by Co₃O₄ nanoparticles are displayed in given figure. An oxidation peak, associated with the transformation of Co₃O₄ into CoOOH, is observed at around 0.42 V. At roughly 0.27 V, the decrease peak appears as a result of the opposite reaction. When compared to the naked CC, the Co₃O₄-modified electrode exhibits a significantly higher current response, indicating better capacitance. Faradaic redox reactions can account for the charge storage mechanism of the Co₃O₄ electrode [40,41], as shown by Equations (2) and (3).



In the process of charging, Co₂O₃ absorbs electrons (6e⁻) and forms hydrated cobalt hydroxide species by reacting with hydroxide ions. In discharging, the hydrated cobalt

hydroxide species transfers electrons (6e⁻) and proceeds oxidation back to Co₃O₄, releasing hydroxide ions. In 1M KOH and 1M Na₂SO₄ electrolytes, the Co₃O₄ electrode's electrochemical performance is displayed in Fig. 7. For electrochemical performance in KOH electrolyte, the Co₃O₄ electrode outperforms Na₂SO₄. Faster rates of K⁺ ion intercalation and deintercalation are made possible by KOH's higher conductivity and ionic mobility, which makes sense [36]. The specific capacitance (Cs) values for the Co₃O₄ electrode are 145, 95, 68, 47, 44, 36, and 31 Fg⁻¹, respectively, for scan rates of 5, 10, 20, 50, 60, 80, and 100 mVs⁻¹ in 1M KOH. In 1M Na₂SO₄, the Cs values for the same scan speeds are 88, 58, 47, 34, 31, 25, and 21 Fg⁻¹. The Cs decreases with increasing scan rate, which is consistent with the results displayed in Fig. 7. Based on the cyclic voltammetry curves, the specific capacitance of the Co₃O₄ electrode was calculated using Equation (4).

$$C_s = \frac{1}{\Delta v \times m} \int_{v_{min}}^{v_{max}} I(V) dv$$

where m is the mass of active CoO deposited (g), V is the scan rate (Vs⁻¹), and Cs is the specific capacitance of the CoO electrode (Fg⁻¹). In this equation, Cs stands for the Co₃O₄ working electrode's specific capacitance (Fg⁻¹), I for current density (A), Δt for discharge time (s), Δv for potential window (mVs⁻¹), and m for the mass of active Co₃O₄ deposited. Excellent cycle stability is essential for supercapacitor applications. As shown in Fig. 8, the Co₃O₄ electrode exhibits long-term cyclic stability, retaining approximately 72% of the initial Cs after 500 cycles at a scan rate of 100 mVs⁻¹ in 1M KOH electrolyte.

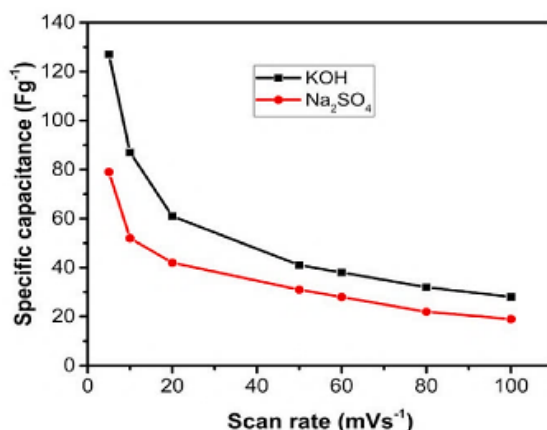


Fig. 7. Cs vs scan rates plot in KOH and Na₂SO₄ electrolytes

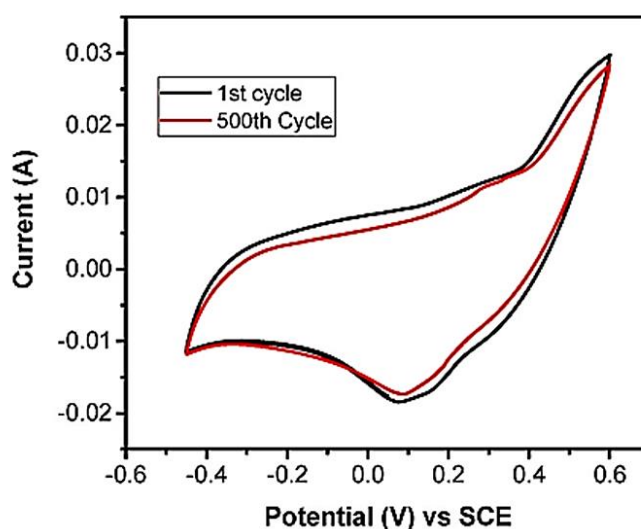


Fig. 8. Capacitance retention plot CoO-CC in KOH electrolyte

To sum up, Co_3O_4 nanoparticles show promise for use as an electrode material in alkaline electrolytes due to their greater specific capacitance values and faster charge-discharge rates when compared to neutral solutions.

5. CONCLUSION

Co_3O_4 nanoparticles synthesized by a straightforward hydrothermal process clearly show promising properties as electrode materials for supercapacitors, based on the thorough investigation reported in this study. The successful production of the cubic phase Co_3O_4 with distinctive Co-O bonds and a non-uniform aggregation of nanoparticles was confirmed by the structural and morphological investigations. While surface wettability studies revealed the hydrophilic character of the Co_3O_4 nanoparticles, which is advantageous for electrolyte interaction, the BET analysis yielded important information on the surface area and pore structure. Co_3O_4 nanoparticles on carbon cloth (CC) showed remarkable specific capacitances of 132 Fg^{-1} in KOH and 79 Fg^{-1} in Na_2SO_4 solutions when used as electrodes in supercapacitors. With almost 75% of their capacitance remaining after 500 cycles in KOH, the electrodes also demonstrated remarkable cycling stability, underscoring their durability and long-term usability. These results highlight the potential of Co_3O_4 nanoparticles as cutting-edge materials with great performance and stability as supercapacitor electrodes. The work opens the door for additional Co_3O_4 -based nanomaterial optimization and use in energy storage technologies.

COMPETING INTERESTS

Authors have declared that no competing interests exist.

REFERENCES

1. Khan I, Saeed K, Khan I. Nanoparticles: Properties, applications and toxicities. *Arabian Journal of Chemistry*. 2019;12(7): 908-931.
2. Goesmann H, Feldmann C. Nanoparticulate functional materials. *Angewandte Chemie International Edition*. 2010;49(8):1362-1395.
3. Ealia SAM, Saravanakumar MP. A review on the classification, characterisation, synthesis of nanoparticles and their application. In *IOP conference series: Materials Science and Engineering*. IOP Publishing; 2017.
4. Heiligttag FJ, Niederberger M. The fascinating world of nanoparticle research. *Materials Today*. 2013;16(7-8):262-271.
5. Yan J, et al. Recent advances in design and fabrication of electrochemical supercapacitors with high energy densities. *Advanced Energy Materials*. 2014;4(4): 1300816.
6. Zuo W, et al. Battery-super capacitor hybrid devices: Recent progress and future prospects. *Advanced Science*. 2017;4(7): 1600539.
7. Muzaffar A, et al. A review on recent advances in hybrid super capacitors: Design, fabrication and applications.

- Renewable and Sustainable Energy Reviews. 2019;101:123-145.
8. Yaseen M, et al. A review of super capacitors: Materials design, modification, and applications. *Energies*. 2021;14(22):7779.
 9. Wang X, et al. Recent advance in Co₃O₄ and Co₃O₄-containing electrode materials for high-performance super capacitors. *Molecules*. 2020;25(2):269.
 10. Kumar S, et al. Metal oxides for energy applications, in *Colloidal Metal Oxide Nanoparticles*. Elsevier. 2020;471-504.
 11. Pang H, et al. Synthesis of functional nanomaterials for electrochemical energy storage. Springer; 2020.
 12. Kokila G, Mallikarjunaswamy C, Ranganatha VL. A review on synthesis and applications of versatile nanomaterials. *Inorganic and Nano-Metal Chemistry*. 2022;1-30.
 13. Salem SS, et al. A comprehensive review of nanomaterials: Types, synthesis, characterization, and applications. *Biointerface Res. Appl. Chem*. 2022;13(1):41.
 14. Abouali S, et al. Electrospun carbon nanofibers with in situ encapsulated Co₃O₄ nanoparticles as electrodes for high-performance super capacitors. *ACS Applied Materials and Interfaces*. 2015;7(24):13503-13511.
 15. Meng T, et al. Co₃O₄ nanorods with self-assembled nanoparticles in queue for supercapacitor. *Electrochimica Acta*. 2015;180:104-111.
 16. Liu F, et al. Facile synthesis of ultrafine cobalt oxide nanoparticles for high-performance supercapacitors. *Journal of Colloid and Interface Science*. 2017;505:796-804.
 17. Kim BK, et al. Electrochemical super capacitors for energy storage and conversion. *Handbook of Clean Energy Systems*. 2015;1-25.
 18. Zhao J, Burke AF. Electrochemical capacitors: Materials, technologies and performance. *Energy Storage Materials*. 2021;36:31-55.
 19. Gupta P, et al. Recent developments and research avenues for polymers in electric vehicles. *The Chemical Record*. 2022;22(11):e202200186.
 20. Nassar MY. Size-controlled synthesis of CoCO₃ and Co₃O₄ nanoparticles by free-surfactant hydrothermal method. *Materials Letters*. 2013;94:112-115.
 21. Xu J, et al. Preparation and electrochemical capacitance of cobalt oxide (Co₃O₄) nanotubes as super capacitor material. *Electrochimica Acta*. 2010;56(2):732-736.
 22. Aghazadeh M, et al. High performance Pseudocapacitor electrode material fabricated by pulse electro-synthesized cobalt oxide nanostructures. *International Journal of Electrochemical Science*. 2016;11(12):11016-11027.
 23. Umar A, et al. Perforated Co₃O₄ nanosheets as high-performing supercapacitor material. *Electrochimica Acta*. 2021;389:138661.
 24. Saikia BK, et al. A brief review on super capacitor energy storage devices and utilization of natural carbon resources as their electrode materials. *Fuel*. 2020;282:118796.
 25. Biswas S, Chowdhury A. Organic super capacitors as the next generation energy storage device: Emergence, opportunity, and challenges. *Chem Phys Chem*. 2023;24(3):e202200567.
 26. Kaviyarasu K, Raja A, Devarajan PA. Structural elucidation and spectral characterizations of Co₃O₄ nano flakes. *Spectrochimica Acta Part A: Molecular and Biomolecular Spectroscopy*. 2013;114:586-591.
 27. Yang YP, et al. Preparation and electrochemical performance of nanosized Co₃O₄ via hydrothermal method. *Transactions of Nonferrous Metals Society of China*. 2007;17(6):1334-1338.
 28. Deori K, et al. Morphology controlled synthesis of nano porous Co₃O₄ nanostructures and their charge storage characteristics in super capacitors. *ACS Applied Materials and Interfaces*. 2013;5(21):10665-10672.
 29. Jamil S, Janjua MRSA, Khan SR. Synthesis and structural investigation of polyhedron Co₃O₄ nanoparticles: Catalytic application and as fuel additive. *Materials Chemistry and Physics*. 2018;216:82-92.
 30. Andrade-Sanchez M, et al. Temperature and pH effect on reaction mechanism and particle size of nanostructured Co₃O₄ thin films obtained by sol-gel/dip-coating. *Materials Research Express*. 2021;8(2):025015.
 31. El-Shamy OA, Deyab MA. The most popular and effective synthesis processes for Co₃O₄ nanoparticles and their benefit in preventing corrosion. *Zeitschrift für*

- Physikalische Chemie. 2023;237(3):333-350.
32. Kumarage GW, Comini E. Low-dimensional nanostructures based on cobalt oxide (Co₃O₄) in chemical-gas sensing. Chemosensors. 2021;9(8):197.
 33. Wang S, et al. One-dimensional porous Co₃O₄ rectangular rods for enhanced acetone gas sensing properties. Sensors and Actuators B: Chemical. 2019;297:126746.
 34. Zheng MB, et al. Preparation of mesoporous Co₃O₄ nanoparticles via solid-liquid route and effects of calcination temperature and textural parameters on their electrochemical capacitive behaviors. The Journal of Physical Chemistry C. 2022;113(9):3887-3894.
 35. Pudukudy M, et al. Facile synthesis of bimodal mesoporous spinel Co₃O₄ nanomaterials and their structural properties. Superlattices and Microstructures. 2019;64:15-26.
 36. Jadhav SL, et al. Influence of deposition current and different electrolytes on charge storage performance of Co₃O₄ electrode material. Journal of Physics and Chemistry of Solids. 2023;180:111422.
 37. Rajeshkhanna G, Umeshbabu E, Rao GR. Charge storage, electrocatalytic and sensing activities of nest-like nanostructured Co₃O₄. Journal of Colloid and Interface Science. 2017;487:20-30.
 38. Zhao L, et al. Electrolyte-wettability issues and challenges of electrode materials in electrochemical energy storage, energy conversion, and beyond. Advanced Science. 2023;10(17):2300283.
 39. Adhikari H. Synthesis and Electrochemical Performance of Hydrothermally Synthesized Co₃O₄ Nanostructured Particles; 2020.
 40. Adekunle AS. Electrochemical and electrocatalytic properties of carbon nanotubes integrated with selected metal and metal oxide nanoparticles. University of Pretoria; 2019.
 41. Yang W, Peng D, Kimura H, Zhang X, Sun X, Pashameah RA, Alzahrani E, Wang B, Guo Z, Du W, Hou C. Honeycomb-like nitrogen-doped porous carbon decorated with Co₃O₄ nanoparticles for superior electrochemical performance pseudo-capacitive lithium storage and super capacitors. Advanced Composites and Hybrid Materials. 2022;5(4):3146-57.

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