ELECTROCATALYSIS OF COBALT DOPED CeO$_2$/rGO NANOCOMPOSITE FOR OXIDATION OF METHANOL AND FORMIC ACID

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ABSTRACT

In this study, a novel cobalt-doped cerium oxide (CeO$_2$) on reduced graphene oxide (rGO) substrate was successfully prepared through a hydrothermal method. The comprehensive characterization of the synthesized nanocomposite involved X-ray diffraction studies (XRD), Raman spectrum analysis, High Resolution Transmission Electron microscopy (HRTEM), and Field Emission Electron Microscopy (FE-SEM). The effective surface area of CeO$_2$/rGO and Co-CeO$_2$/rGO was determined using the Randles-Sevcik equation, revealing values of 4.02X10$^{-5}$ cm$^2$ and 6.35X10$^{-5}$ cm$^2$, respectively. Cyclic Voltammetry (CV) was employed to investigate the electrochemical reversibility behavior, demonstrating promising results for Co-CeO$_2$/rGO nanocomposite. Electrochemical impedance spectroscopy (EIS) further highlighted low charge transfer resistance (Rct) and enhanced double-layer capacitance ($C$$_{dl}$). Notably, the synthesized nanocomposite exhibited excellent electrocatalytic activity for methanol and formic acid oxidation in an acidic medium. Overall, this work provides valuable insights into the synthesis characterization, and electrochemical performance of Co-CeO$_2$/rGO nanocomposite, showcasing their potential applications in energy-related processes.

Keywords: rGO, Co-CeO$_2$/rGO, Electrocatalysis, Methanol, Formic Acid Oxidation.

INTRODUCTION

Graphene’s structural features make it a focal point in contemporary science and technology$^1$, particularly in various energy storage applications due to its unique physicochemical properties.$^2$ Economical large-scale production of graphene is achievable through the chemical or thermal reduction of graphene oxide.$^3$ Functionalized graphene, obtained by reducing graphene oxide$^4$, serves as a substrate for nanocomposite preparation.$^5$ Transition metal oxides have garnered significant attention in the field of storage devices due to their properties, such as their structural, mechanical, electronic, and pseudo-capacitance characteristics. These properties arise from their variable oxidation states, a quality achievable through the use of carbon-based materials.$^6,7$ Noble transition metal oxides such as RuO$_2$ and IrO$_2$ are costly, have lower abundance and poor stability prone to show catalytic poisoning other non-noble metal oxides such as NiO, MnO$_2$, Co$_3$O$_4$, CeO$_2$, Cr$_2$O$_3$, Fe$_2$O$_3$ have been in use as they are cheaper and environmentally friendly.$^8$-10 CeO$_2$, with its distinct structural properties, has emerged as a supporting material in electrochemical devices. Outstanding results have attracted researchers to explore the synthesis of transition rare earth metal oxide doped with graphene composites.$^{15}$ Hence Co-CeO$_2$/rGO is selected and there are various well-known methods to synthesize, among them Hydrothermal is an efficient method where the nanomaterial is synthesized with morphologically controlled growth.$^{16,18}$ With this importance, we have prepared Co-CeO$_2$/rGO. Supercapacitors have two main energy storage mechanisms: pseudocapacitor and electric double-layer capacitor.$^{19,20}$ The first one employs precious metals such as platinum (Pt), gold (Au), and silver (Ag), while the second approach utilizes carbon-based materials combined with transition metal oxides. Mixed transition metal oxides have garnered significant attention in the latter method.$^{21,23,69}$
cells have a growing interest as a future energy source. This is due to suitable fuel feed and easy handling even at low temperatures. Ethanol and methanol as fuel, have a potential feature such as reactivity at low temperatures and easy handling compared to all organic reagents. Methanol and formic acid are more advantageous than ethanol concerning selectivity to carbon dioxide formation during electrochemical oxidation. The goal is to create a remarkably effective and long-lasting catalyst for the oxygen reduction reaction (ORR), serving as a cost-effective alternative to platinum-based catalysts in both traditional fuel cells and direct methanol fuel cells (DMFCs). Therefore, in this present paper, we have prepared cobalt-doped CeO$_2$/rGO nanocomposite using the hydrothermal method and confirmed by XRD, Raman, SEM, and TEM techniques. The obtained material was used to study the oxidation of methanol and formic acid.

**EXPERIMENTAL**

**Material and Methods**

Ammonium cerium nitrate, reduced graphene oxide, cobalt (II) chloride hexahydrate, sodium hydroxide as fuel, quaternary ammonium surfactant hexacetyltrimethylammoniumbromide (CTAB), and other chemicals like KCl, KOH, C$_6$H$_5$FeK$_3$, C$_6$FeK$_3$Na, methanol, HCOOH, and H$_2$SO$_4$ have been ordered from a reputed supplier of analytical research-grade chemicals. They have been supplied in their purest form.

**Synthesis of Co-CeO$_2$/rGO**

0.04g of graphene oxide (GO) was dissolved in 30mL of 2M caustic soda solution for 60 minutes using ultrasonication. Subsequently, 8mL of 0.05M ammonium cerium nitrate (NH$_4$)$_2$Ce(NO$_3$)$_6$ solution was stirred using a magnetic stirrer for 30 minutes. To maintain a consistent cobalt (Co) dopant percentage, a fixed proportion of 400mL of CoCl$_2$6H$_2$O was added. The entire mixture was then transferred to a 100mL hydrothermal bomb enclosed in a stainless steel container. The reaction took place in a hot air oven at a temperature of 180 °C for 24 hr. Once the reaction was allowed to cool to laboratory temperature, the resulting nanocomposite was isolated through centrifugation and subsequently dried at 60 °C for a duration of 12 hours.

**Preparation of Working Electrode for Electrochemical Studies**

The process of achieving a homogenous carbon paste involved manually blending a mixture of synthesized nanocomposite, graphite, and silicone oil using a mortar and pestle for approximately 30 minutes. Further, it's filled in a Teflon electrode with copper wire at the other end which is sharpened to remove polymer for electrical contact. This cylindrical Teflon electrode is used to study cyclic voltammetry (CV) by applying a potential range between 10 to 100 mV/s. To study the fuel cell application and Electrochemical Impedance (EIS), a Nickel mesh electrode is used which is prepared by making a blend of nanocomposite, graphite, and PTFE (Polytetrafluoroethylene) binder using mortar and pestle to get a homogenous blend which is pasted on Nickel mesh, for electrical contact copper wire is inserted the electrode is dried at room temperature for an about 24 hr. and used as a working electrode.

**RESULTS AND DISCUSSION**

**XRD**

The crystallography and phases of Co-rGO/CeO$_2$ synthesized by the hydrothermal method were characterized using XRD. The XRD pattern of rGO, CeO$_2$, rGO/CeO$_2$ and Co-rGO/CeO$_2$ is shown in the Fig.-1. A diffraction peak of rGO is observed at 26.2°, 42°, and 54° 20 angles. For CeO$_2$, the diffraction peaks are observed at 29°, 33°, 47°, 56°, 59°, and 69° with respect to reflection planes of (111), (200), (220), (311), (222), and (400) which is confirmed with standard JCPDS Card No-34-0394. For rGO/CeO$_2$, position of the peak remains the same but with intensity changed. Due to the insertion of Co in the CeO$_2$ space lattice, there is a displacement in the diffraction peak to a higher 20 with higher intensity indicating that the shift in higher 20 is due to the replacement of Ce with Co, in the crystal structure. The avg. particle of synthesized nanocomposite was calculated using Debye-Scherer Eq. (1).

$$D = \frac{kl}{\beta \cos \theta}$$

The measurement considered the FWHM of the extremely strong diffraction. The crystalline sizes of rGO, rGO/CeO$_2$, and Co-rGO/CeO$_2$ were found to be 9.2, 7.66, and 5.78 nm respectively.

**Raman Spectroscopy**

The Raman spectra of rGO/CeO$_2$ and Co-rGO/CeO$_2$ are displayed in Fig.-2. The D and G bands at 1364 and 1598 cm$^{-1}$ which are characteristics of GO, are shown in the graph. However, the graph depicts a shifted band at 1352 cm$^{-1}$ and 1579.3 cm$^{-1}$ which are commonly labeled as D and G bands which is due to the reduction of GO to Rgo. The sp$^3$ disordered carbon atoms give the d band whereas the G band ascends mainly from the oscillation of
basic sp² hybridized carbon in graphite. The shift at 462.5 cm⁻¹ shown by the composites also reflects the symmetrical stretching vibration mode of Cerium and oxygen.

There is a high chemical influence between rGO and CeO₂ reflected in the Raman shift. It indicates the homogeneous wrapping of rGO sheets by CeO₂ on its surface. A downward shift is also seen for Co-doped CeO₂ indicating the excellent imperfection created due to oxygen vacancy. Such a shift is mainly due to a decrease of Ce⁴⁺ to Ce³⁺ enhancing the O₂ vacancies and balancing the total charge. From the literature survey it’s clear that the ID/IG value was found to be much higher than GO as in rGO which is due to smaller sp² domains due reduction of GO. I_D/I_G value for CeO₂/rGO is 0.8584 and for Co-CeO₂/rGO is 0.8518. It’s clear that as metal is doped I_D/I_G value decreases.

FE-SEM images were used to characterize the analysis and morphology of the built materials, as shown in Fig.-3. The prepared rGO depicts a flame-like sheet structure in Fig.-3 (a), with a few layers stacked on top of one another. The sheets appear to be between 50 and 100 nm thick and a few microns long. The nano balls and nano-rods-like structure were obtained for CeO₂ NPs as depicted in Fig.-3 (b). Some nano-ball shape particles (spherical) are formed and some of the particles are almost agglomerated. The flame layer sheet image of rGO edges and CeO₂ nanoparticles overlapped with each other is shown in Fig.-3(c). The majority of the molecules are aggregated, and metal oxide with distinctive structures was used cobalt oxide to anchor the surface of rGO layers in Fig.-3(d).

HR-TEM images of Co-doped rGO/CeO₂ were obtained by the hydrothermal method as displayed in Fig.-4. From Fig.-4 (a), displays a typical TEM micrograph image of rGO formed by a flakes-like structure and multiple-nano-layer sheets with a size of 10-100 nm and CeO₂ NPs overlapped with each other, and the majority of the molecules are aggregated with cobalt. The distance between two symmetric diffraction sites can be used to calculate the d-spacing value. The d-spacing value was found to be 3.12 nm Fig.-4 (e). The SEAD patterns of Co-doped rGO/CeO₂ NCs obtained bright spots with ring-like patterns displayed in Fig.-4 (b and d) indicating the crystalline properties of rGO/CeO₂.

The CeO₂, CeO₂/rGO, and Co-CeO₂/rGO were analyzed in a 5 mM [K₃/K₄Fe(CN)₆] + 0.5M KCl electrolyte by increasing the scan rate within the range of 10 to 100 mV/s is depicted in Fig.-5. A steady increase in current is observed, which also indicates the reversibility of the redox process.
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![Image of FE-SEM and HR-TEM images]

**Fig.-3**: FE-SEM of (a) rGO (b) CeO$_2$ (c) CeO$_2$/rGO (d) Co-CeO$_2$/rGO at Different Magnifications

**Fig.-4**: HR-TEM images of Co-rGO/CeO$_2$ (a) 100 nm (c) 500 nm (e) d-spacing, and (b, d) SAED Pattern of NCs

This also reveals the reproducibility and thermodynamic stability which is supported by the linear regression value as shown in Fig.-5(a) to Fig.-5(c). It also shows that the above process is controlled by adsorption. The following mechanism depicts the rapid reversible Faradaic redox reaction which is electrochemically possible.

$$[Fe(CN)_6]^{4-} \leftrightarrow [Fe(CN)_6]^{3-} + e^- \quad (2)$$

The above reaction is also responsible for charging, where the conversion of Fe$^{2+}$ to Fe$^{3+}$ and during discharging, the conversion Fe$^{3+}$ to Fe$^{2+}$ is a Faradaic feasible redox reaction. From the above explanation, it’s clear that by comparing the current due to doping of Co to CeO$_2$/rGO, the surface area is increased as well as ORR in cyclic voltammetry as shown in Table-1. The following Table-2 reveals the effective surface area of the electrode, which is calculated using the Randles-Sevecik equation.

$$I_{pa} = (2.69 \times 10^5) \times n^{2/3} \times A_{eff} \times D^{1/2} \times v^{1/2} \times C \quad (3)$$

Where D is diffusion coefficient, $C_o$ is bulk concentration, $A_{eff}$ is effective surface area, and $v$ is scan rate respectively.

<table>
<thead>
<tr>
<th>Electrode</th>
<th>$I_{pa}$ (A) $\times 10^{-3}$</th>
<th>$I_{pc}$ (A) $\times 10^{-3}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>CeO$_2$</td>
<td>2.058</td>
<td>-1.805</td>
</tr>
<tr>
<td>CeO$_2$/rGO</td>
<td>3.067</td>
<td>-3.869</td>
</tr>
<tr>
<td>Co-CeO$_2$/rGO</td>
<td>4.504</td>
<td>-4.742</td>
</tr>
</tbody>
</table>

It is clear that cerium and cobalt are synergic with rGO as indicated in Table-2. When comparing the data above; there is a consistent rise in the current when compared to CeO$_2$. In the case of NCs, there is a flipping of quick redox couple between Ce$^{3+}$ and Ce$^{4+}$ and vice versa, which is due to the enhanced surface area.
Table-2: Effective Surface Area of Various Electrodes

<table>
<thead>
<tr>
<th>Electrode</th>
<th>Effective surface area $\times 10^6$ cm$^2$</th>
</tr>
</thead>
<tbody>
<tr>
<td>CeO$_2$</td>
<td>3.36</td>
</tr>
<tr>
<td>CeO$_2$/rGO</td>
<td>4.02</td>
</tr>
<tr>
<td>Co-CeO$_2$/rGO</td>
<td>7.35</td>
</tr>
</tbody>
</table>

Fig.-5: CV Images of (a) CeO$_2$ (b) CeO$_2$/rGO (c) Co-CeO$_2$/rGO (d) Comparative of Reversibility

**Electrocatalysis of Methanol Oxidation**

The cobalt-doped CeO$_2$/rGO exhibits a remarkable enhancement in the redox peak, indicating better electrocatalytic activity. The addition of 0.2M CH$_3$OH + 0.5M KOH was also performed during these experiments. Figure-6 illustrates the CV data acquired within the potential window of -1 to 1 V, employing a scan speed of 50 mV/s. Notably, a redox peak is observed at different potential ranges for the three materials: -0.1349 to 0.1568V for CeO$_2$, -0.1142 to 0.0677V for CeO$_2$/rGO, and -0.1207 to 0.0755V for Co-CeO$_2$/rGO. This redox peak indicates an enhanced anodic current density when methyl alcohol is added to the supporting electrolyte. Furthermore, the appearance of the redox peak reveals the oxidation of methanol specifically at the CeO$_2$/rGO interface. This observation suggests that the presence of Co-CeO$_2$/rGO enhances the electrocatalytic activity during the oxidation of methanol in comparison to CeO$_2$ and CeO$_2$/rGO. The CV response of Co-CeO$_2$/rGO at different concentrations of methanol (0.025M to 1M) demonstrates that with an increase in the concentration of methanol, the oxidation of methanol intensifies, indicating more methanol molecules available for electrochemical reaction leading to more electrons being transferred and consequently higher current. It also suggests kinetically more favorable where there is increased chances of successful collision between methanol molecules and the catalyst on the anode surface.

Fig.-6: CV of (a) CeO$_2$ (b) CeO$_2$/rGO (c) Co-CeO$_2$/rGO (d) Comparative Redox System

This leads to the development of intermediates and products that start blocking the active sites on the electrocatalyst, thereby hindering further adsorption of methanol. As a consequence, the oxidation reaction gradually decreases, and the peak in the cyclic voltammogram remains relatively constant, as shown in Fig.-7.
This behavior shows that the electrocatalyst is affected by the increasing concentration of methanol, ultimately limiting its catalytic activity for methanol oxidation. Additionally, during the backward redox reaction, the methyl alcohol will undergo deoxidation once more.\textsuperscript{57} The negative onset potential also indicates more electrocatalytic activity of the material.\textsuperscript{58,59}

**Electro Catalysis for Formic Acid Oxidation**

An identical experiment was conducted to assess the oxidation of formic acid. From Fig.8, it’s clear that CeO\textsubscript{2}, CeO\textsubscript{2}/rGO, and Co-CeO\textsubscript{2}/rGO also show formic acid oxidation similar to methanol oxidation but with a slightly lower anodic potential\textsuperscript{60-62}. EIS typically acknowledged as the most efficient technique for evaluating electron transport at the electrolyte-electrode interface and determining the ionic conductivity of both the electrode and electrolyte during the O\textsubscript{2} evolution route, stands out as the preferred method.\textsuperscript{63}

In Fig.9, a pronounced semicircle representing the charge transfer resistance in the high-freq., band is shown together with a slope representing the impedance seen in the low-freq., range. The appearance of a depressed curve in the high-frequency region of the spectra indicates a charge transfer mechanism, signifying the least charge...
The decrease in $R_{ct}$ as frequency increases may indicate that the system is becoming more capacitive at higher frequencies. Capacitors have lower impedance at higher frequencies, which is reflected in the Nyquist plot as a decrease in the real part of impedance as indicated in Table-3. Based on this observation, it is evident that the synthesized nanocomposite performs effectively as an electrode. The data indicates $R_s$, $R_{ct}$, and $C_{dl}$ related to active materials catalytic properties, obtained by fitting with experimental data using the equivalence circuit. There is a decrease in $R_{ct}$ and an increase in $C_{dl}$ indicating the surface catalytic activity.

<table>
<thead>
<tr>
<th>Electrode</th>
<th>$R_s$(Ω)</th>
<th>$R_{ct}$(Ω)</th>
<th>$C_{dl}$/F</th>
</tr>
</thead>
<tbody>
<tr>
<td>rGO</td>
<td>71.28</td>
<td>278.2</td>
<td>$8.12 \times 10^{-9}$</td>
</tr>
<tr>
<td>CeO$_2$/rGO</td>
<td>49.25</td>
<td>132.7</td>
<td>$6.133 \times 10^{-8}$</td>
</tr>
<tr>
<td>Co-CeO$_2$/rGO</td>
<td>29.51</td>
<td>108.8</td>
<td>$3.301 \times 10^{-7}$</td>
</tr>
</tbody>
</table>

**CONCLUSION**

The hydrothermal technique was used to prepare the cobalt-doped CeO$_2$/rGO NCs. Because of the increased surface area, the redox performance of cobalt-doped CeO$_2$/rGO was significantly improved. The nanocomposite exhibited enhanced electocatalytic performance during the oxidation of formic acid and methanol. The EIS demonstrated that the bigger the capacitance, the lower the charge transfer resistance. Based on the findings, Co-CeO$_2$/rGO material may be a better electrocatalyst for methanol and formic acid in acidic media for fuel cell applications.

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**CONFLICT OF INTERESTS**

The authors declare that there is no conflict of interest.

**AUTHOR CONTRIBUTIONS**

All the authors contributed significantly to this manuscript, participated in reviewing/editing and approved the final draft for publication. The research profile of the authors can be verified from their ORCID ids, given below:

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