

Applicable anode based on Co_3O_4 – SrCO_3 heterostructure nanorods-incorporated CNFs with low-onset potential for DUFCS

Zafar Khan Ghouri¹ · Khaled Elsaid² · Saeed Al-Meer¹ · Nasser A. M. Barakat^{3,4}

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Abstract Besides the high-current density, lower onset potential of urea electrooxidation is key parameter which influences the direct urea fuel cell performance. In the present article, low-onset potential has been reported for nickel-free (NF) electrocatalyst in urea electrooxidation. The nickel-free electrocatalyst: Co_3O_4 – SrCO_3 heterostructure nanorods-incorporated carbon nanofibers (CNFs) were synthesized by electrospinning technique, followed by calcination of electrospun mat composed of strontium acetate, cobalt acetate, and poly(vinyl alcohol) sol–gel in inert environment at 750 °C. Physiochemical characterizations confirmed the formation of Co_3O_4 – SrCO_3 heterostructure nanorods-incorporated CNFs. The electrochemical activity of resultant nickel-free electrocatalyst toward the electrooxidation of urea in alkaline medium is evaluated using cyclic voltammetry measurements (CV). Co_3O_4 – SrCO_3 heterostructure nanorods-incorporated

CNFs reveals high-current density of 21.33 mA/cm² at low-fuel concentration. Notably, the low-onset potential has been observed, showing a good application prospect in direct urea fuel cells.

Keywords Electrospinning · Urea · DUFC · Onset potential · Electrocatalyst · Current density

Introduction

The ultimate rising requirement for energy and environmental concerns has driven the search for new resources to produce eco-friendly energy (Ghouri et al. 2015). Foregoing research demonstrated that direct liquid fuel cells (DLFCs) are a competitive alternative resource of green energy (Radenahmad et al. 2016). Among all types of direct liquid fuel cells, direct urea fuel cell (DUFCS) is a strong candidate working simultaneously for resolving energy and environmental problems (Xu et al. 2016). Besides urea fuel cell, other liquid fuel cells, such as methanol, ethanol, and formic acid fuel cells, need a supply of fuels from process industry by chemical or biological production, which requires strong storage and transportation infrastructure. For instance, direct urea fuel cells have many advantages compared to the above-mentioned liquid fuel cells, such as urea, which are not expensive, available in solid granules and, thus, easy to transport and store (Lan and Tao 2011). More interestingly, urea contains 10 wt% hydrogen with high-energy density (16.9 MJ l⁻¹) (Lan and Tao 2011; Xu et al. 2014; Lan et al. 2010). Moreover, urea is the main nitrogen-containing component in the urine/liquid waste (2–2.5 wt%) (Lan et al. 2010; Kojima et al. 2006; Boggs et al. 2009). The total theoretical cell potential of urea oxidation (0.37 V vs SHE) is much lower

✉ Zafar Khan Ghouri
zafarkhan.ghouri@qu.edu.qa;
zafarkhanghouri@hotmail.com

✉ Saeed Al-Meer
salmeer@qu.edu.qa

✉ Nasser A. M. Barakat
nasser@jbnu.ac.kr

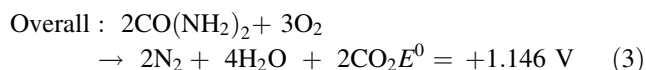
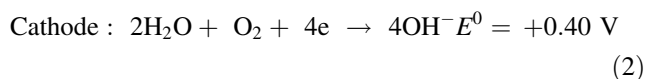
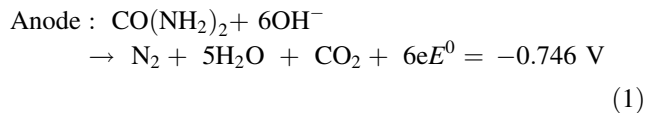
¹ Central Laboratories Unit, Qatar University, PO Box 2713, Doha, Qatar

² Chemical Engineering Program, Texas A&M University at Qatar, PO 23874, Doha, Qatar

³ Department of Organic Materials and Fiber Engineering, Chonbuk National University, Jeonju 54896, Republic of Korea

⁴ Department of Chemical Engineering, Minia University, El-Minia, Egypt

than that of water electrolysis (1.23 V vs SHE), providing a cheaper source of hydrogen production (Boggs et al. 2009; King and Botte 2011). Therefore, it has been reported that either hydrogen or electricity can be generated from urea-rich aqueous solution, according to following reactions (King and Botte 2011; Wang et al. 2013).



Unfortunately, the use of costly noble metals as an anode material not only limits their possible application for urea oxidation in industrial-scale, but also exhibited low-current densities with high over potentials (Boggs et al. 2009). Among various non-precious metals, nickel-based catalyst was found to be very successful, but the transformation of $E_{\text{Ni}^{2+}}^{2+}/E_{\text{Ni}^{3+}}^{3+}$ proceeds at a high potential that will directly affect the current density and energy efficiency of urea fuel cells (Xu et al. 2016; Chen et al. 2015; Guo et al. 2015). Hence, it is desirable to synthesize nickel-free (NF) catalysts that can oxidize urea efficiently at lower over potential. Recently, cobalt-based electrocatalysts have been studied significantly. In particular, tricobalt tetraoxide (Co_3O_4), a magnetic p-type semiconductor, is a versatile inorganic transition metal oxide that has received extensive attention recently for the potential applications in many fields, including fuel cells, supercapacitors, lithium ion batteries, photocatalysis, and electrochemical sensors (Hu et al. 2008; Shahid et al. 2014; Jiao and Frei 2009; Li et al. 2005, 2008; Kim et al. 2014). In addition, cobalt-based nanostructures as electrocatalysts supported on carbon nanofibers have appealed great attention due to its distinct characteristics over the other structures (Ghouri et al. 2015; Ghouri et al. 2016). Inspired by cobalt's catalytic activity toward electrode reactions and potential applications of SrCO_3 in material chemistry, we propose to develop a bimetallic catalyst, composed of cobalt oxide (Co_3O_4) and strontium carbonate (SrCO_3), which could potentially oxidize urea efficiently at lower onset potential and thus support to improve the overall efficiency of direct urea-based fuel cells. In this context, we develop a simple strategy to fabricate Co_3O_4 - SrCO_3 heterostructure nanorods supported by carbon nanofibers (CNFs) as efficient nickel-free electrocatalysts via electrospinning.

Experimental

Catalyst synthesis, characterizations, and electrochemical measurements

In a typical synthesis, 0.3 g of strontium acetate (99.9%) and 0.7 g of cobalt(II) acetate tetrahydrate (98%) salts were dissolved in 3 mL of deionized water, 15 g of PVA (10 wt%) solution was added and stirred at 50 °C for 6 h, and the resultant solution was electrospun at 22 kV. Finally, after conventional drying, nanofiber mat was annealed at 750 °C for 6 h in inert environment.

The detailed physicochemical characterizations were done by field-emission scanning electron microscopy (FESEM), Raman spectroscopy, Fourier transform infrared spectroscopy (FTIR), X-ray diffraction (XRD), and thermal gravimetric analysis (TGA) measurements. Electrochemical measurements were carried out using 2 mg of active material on an electrochemical workstation (VersaSTAT 4, USA) with a three-electrode cell at room temperature.

Results and discussion

The XRD pattern of synthesized Co_3O_4 - SrCO_3 heterostructure nanorods-incorporated CNFs are shown in Fig. 1. The apparent diffraction peaks observed at 2θ values of 25.20°, 35.02°, 44.10°, 47.83°, and 50.37° indexing as the (111), (200), (221), (212), and (231) reflection planes, respectively, conformed the formation of pure orthorhombic strontianite (JCPDS no.: 84-0418). The additional diffraction peaks at 2θ values of 31.40°, 36.90°, 44.70°, 55.70°, and 77.38° corresponding to (220), (311),

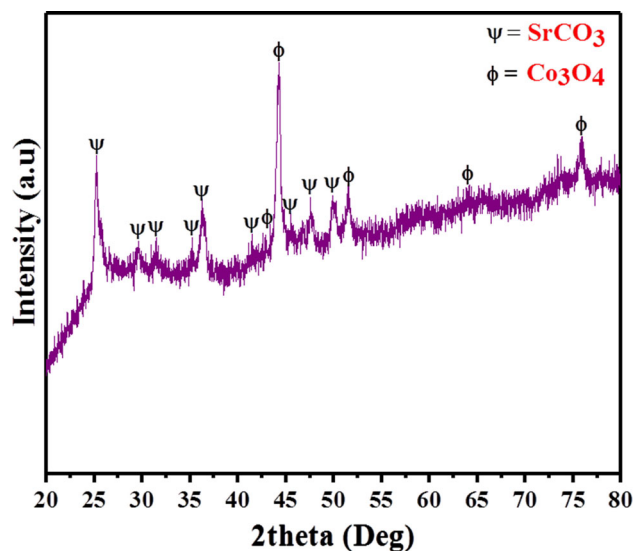


Fig. 1 XRD pattern for the obtained Co_3O_4 - SrCO_3 heterostructure nanorods-incorporated CNFs after calcination at 750 °C for 6 h in inert environment

(400), (422), and (533) reflection planes, respectively, of the cubic spinel crystalline structure of Co_3O_4 (JCPDS no.: 42-1467).

The phases of synthesized Co_3O_4 - SrCO_3 heterostructure nanorods-incorporated CNFs are additionally confirmed by Raman spectroscopy. A typical Raman spectrum presented in Fig. 2 exhibits two set of active bands at 478 and 520 cm^{-1} corresponding to the E_g and F_{2g} symmetry modes of Co_3O_4 , respectively (Alvarez et al. 2012). The weak band located at around 608 cm^{-1} is also accounted for F_{2g} mode of Co_3O_4 (Cheng et al. 1998). The two high-intensity Raman modes at 181 (caused by in-plane bending) and 700 cm^{-1} [comes from librational motion of the $(\text{CO}_3)^{2-}$] are attributed to the vibration bands of SrCO_3 (Divya 2016). In addition, two significant bands were also observed at 1359 and 1595 cm^{-1} , which are assigned to the D- and G-bands of carbon nanofibers, respectively.

FTIR spectra (Fig. 3) of synthesized Co_3O_4 - SrCO_3 heterostructure nanorods-incorporated CNFs were recorded over the 400 – 4000 cm^{-1} wavenumbers. Commonly, the free-planar CO_3^{2-} complexes have D_{3h} symmetry and the absorption bands were produced by the vibrations in CO_3^{2-} complex ions at 400 – 1800 cm^{-1} wavenumbers. The two strong absorption bands (located at around 1463 and 855 cm^{-1}) and one weak absorption band (located at around 1064 cm^{-1}) for SrCO_3 are assigned to be out-of-plane and in-plane bending vibrations (Choi and Huh 2009; Davar et al. 2011). In addition, further two absorption bands are located at 580 and 664 cm^{-1} , indicating the presence of typical stretching vibrations of Co–O group in Co_3O_4 (Shahabuddin et al. 2016). Moreover, the absorption

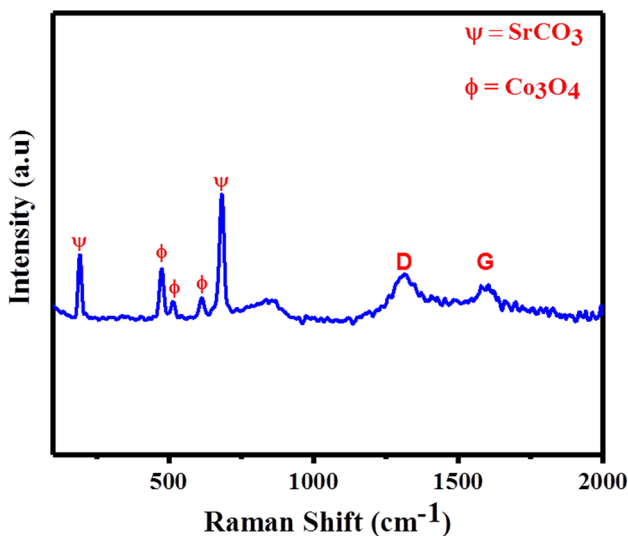


Fig. 2 Raman pattern for the obtained Co_3O_4 - SrCO_3 heterostructure nanorods-incorporated. CNFs after calcination at $750\text{ }^\circ\text{C}$ for 6 h in inert environment

peak at around 1670 cm^{-1} corresponds to the O–H group in the water molecule (Xu et al. 2016).

Thermal behavior was measured from room temperature to $750\text{ }^\circ\text{C}$ by TGA analysis. As shown from the thermo-graph (Fig. 4), initial weight loss occurred at below $200\text{ }^\circ\text{C}$, due to the removal of physically adsorbed water. The second major weight loss was observed between 200 and $590\text{ }^\circ\text{C}$, due to simultaneous dehydration of structural water and thermal decomposition of acetate ions, and final weight loss that was observed between 590 and $750\text{ }^\circ\text{C}$ can

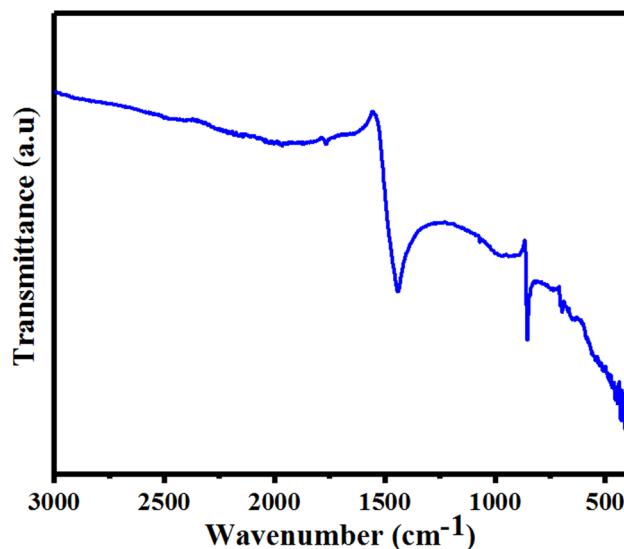


Fig. 3 FTIR pattern for the obtained Co_3O_4 - SrCO_3 heterostructure nanorods-incorporated. CNFs after calcination at $750\text{ }^\circ\text{C}$ for 6 h in inert environment

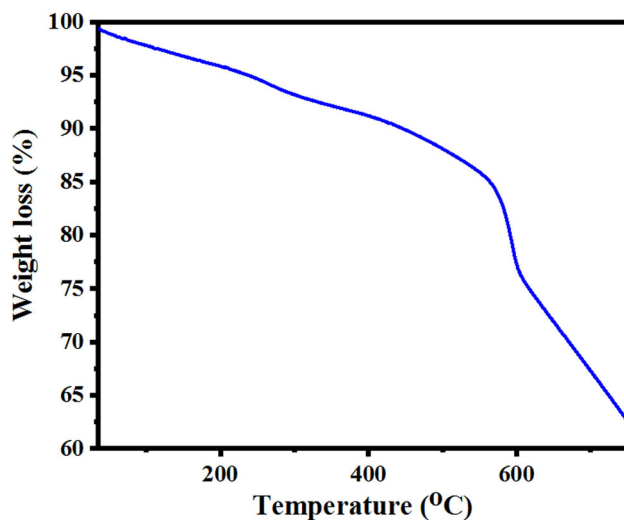


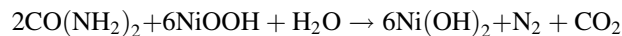
Fig. 4 TGA pattern for the obtained Co_3O_4 - SrCO_3 heterostructure nanorods-incorporated. CNFs after calcination at $750\text{ }^\circ\text{C}$ for 6 h in inert environment

be accounted for the decomposition of polymeric chain and phase transformation.

The morphology of synthesized $\text{Co}_3\text{O}_4\text{-SrCO}_3$ heterostructure nanorods-incorporated CNFs was analyzed by field-emission scanning electron microscope (FESEM). Figure 5a–c shows the FESEM images of the $\text{Co}_3\text{O}_4\text{-SrCO}_3$ heterostructure nanorods-incorporated CNFs at different magnifications. As shown in low-magnification image (Fig. 5a), various dense nanoparticle clusters were observed on the surface of carbon nanofibers. Interestingly, it can be observed that the $\text{Co}_3\text{O}_4\text{-SrCO}_3$ nanoparticles were composed of nanorods with a thickness of ≈ 50 nm and average length about 150–200 nm, which were grown up uniformly on the skeleton of carbon nanofibers (Fig. 5b). However, as shown in Fig. 5c, not only nanorods but also nanoballs were observed; it is possible that some nanorods may be vertically aligned and the X-ray beam parallel to those vertically aligned nanorods may result in the observation of nanoballs.

Elemental profile was determined from EDX line scan analysis (Fig. 6) along the selected line. The results confirm that all the carbon, oxygen, strontium, and cobalt signals are dominantly distributed along at the surface of nanofibers.

Considering the preceding information and the fact that nickel-based catalysts have low electrocatalytic activity, it can be electrochemically activated by consecutive cyclic voltammetry sweeping in an alkaline environment for creating NiOOH active layer on the surface, which is mainly responsible for worthy electrochemical activity. According to the following equation, NiOOH ionic layer is responsible for electrooxidation process in direct urea fuel cells (Wang et al. 2014).



Therefore, it is rational to carry out multiple cyclic voltammetry sweeping to investigate the electrochemical activity of the non-nickel-based electrocatalyst. Figure 7 shows the cyclic voltammogram (CV) of $\text{Co}_3\text{O}_4\text{-SrCO}_3$ heterostructure nanorods-incorporated CNFs electrode in the potential range from -200 to 1000 mV/s (vs. Ag/AgCl) at a scan rate of 50 mV/s in 1.0 M KOH. As shown, current density gradually increases with the number of potential sweeps at low voltage (0.4 V), due to the introduction of OH^- ions into the skeleton of catalyst; this suggests that the adsorption–desorption processes are not reversible. Interestingly, no oxidation/reduction peaks are observed, proposing that the performance of catalyst is independent

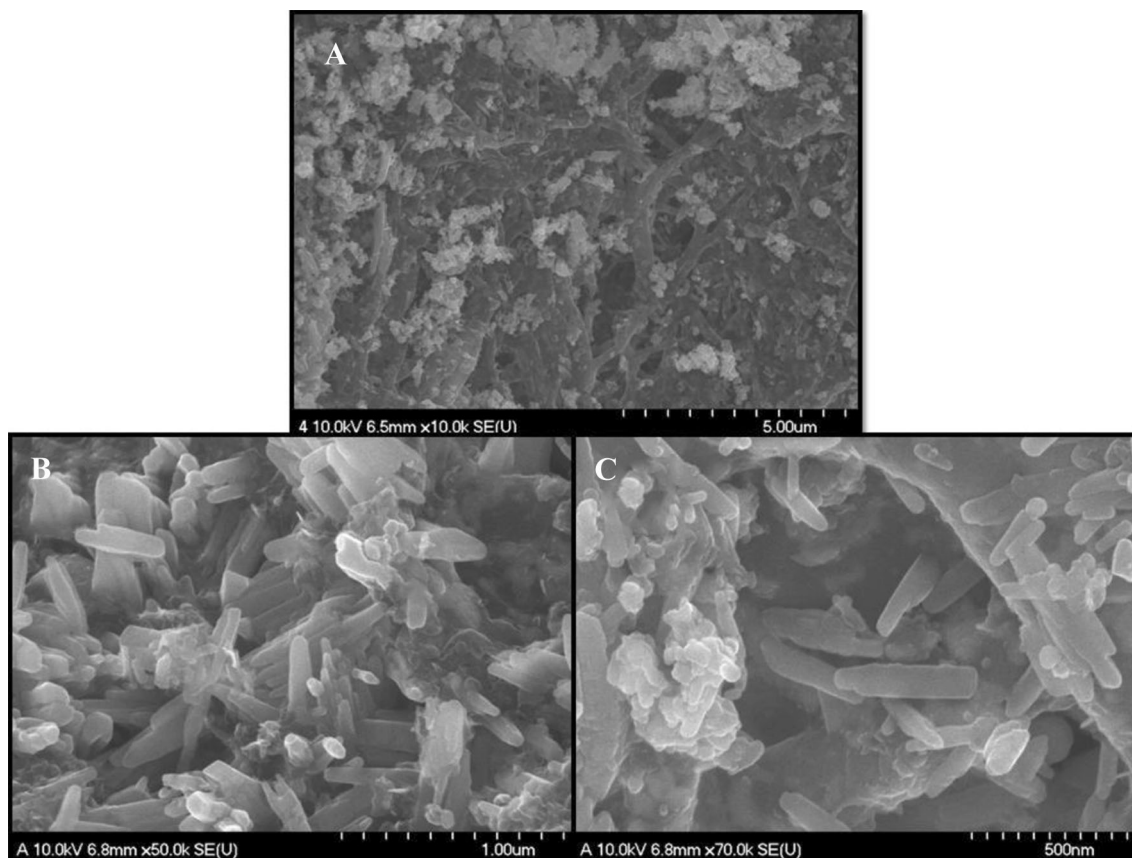


Fig. 5 a–c FESEM image of $\text{Co}_3\text{O}_4\text{-SrCO}_3$ heterostructure nanorods-incorporated CNFs after calcination at 750 °C for 6 h in inert environment

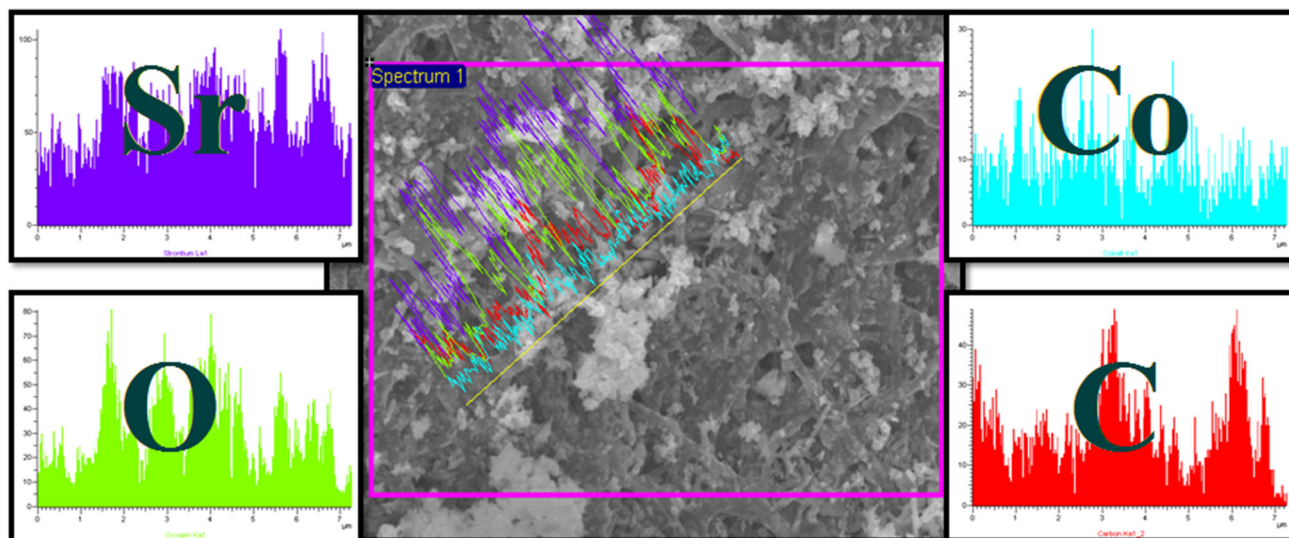


Fig. 6 Line EDX for the obtained $\text{Co}_3\text{O}_4\text{-SrCO}_3$ heterostructure nanorods-incorporated CNFs after calcination at 750°C for 6 h in inert environment

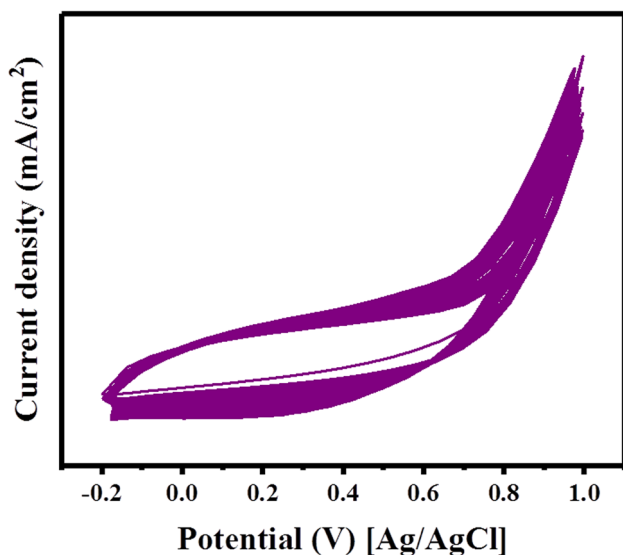


Fig. 7 Cyclic voltammograms for the obtained $\text{Co}_3\text{O}_4\text{-SrCO}_3$ heterostructure nanorods-incorporated CNFs after calcination at 750°C for 6 h in inert environment at scan rate of 50 mV/s at room temperature

of the formation of active ionic layer; this revealed a good catalytic activity of $\text{Co}_3\text{O}_4\text{-SrCO}_3$ heterostructure nanorods-incorporated CNFs electrode.

Figure 8 displays the cyclic voltammograms (CVs) of $\text{Co}_3\text{O}_4\text{-SrCO}_3$ heterostructure nanorods-incorporated CNFs electrodes in a solution of urea ranging from 0.1 to 3.0 +1.0 M KOH. As shown, polarization was started between -2.0 and 1.0 V under a sweep rate of 10 mV/s vs. Ag/AgCl. As revealed in high magnification voltammo-

gram (Fig. 8b), the presence of urea in the alkaline solution has strong influence on anodic current density. However, the observed low-current density at high urea concentration (1.0, 2.0, and 3.0) is understandable due to kinetics limitations of electrode material. Interestingly, no significant difference was observed among the maximum current density obtained from 0.5 and 0.33 M urea concentration that may be due to a diffusion controlled process. Regardless of high-current density ($29.5@0.33$ M urea), the onset potential at which the urea electrooxidation actually starts is considered as a key factor to influence the direct urea fuel cell performance. As shown in Fig. 8c, the minimum onset potential observed was 0.5 V vs. Ag/AgCl, which is very low compared to non-nickel-based catalysts. This could be caused by the exposed active sites of the Co^{3+} ions, which would act as donor-acceptor reduction sites on the surface of $\text{Co}_3\text{O}_4\text{-SrCO}_3$ nanostructures. In addition, Co^{3+} ions can produce surface electronic states by Jahn–Teller effect in alkaline media, which can be predicted as the active sites for the alkaline electrocatalytic reactions (Unwin and Compton 1989). Moreover, these results could not only be accredited to the unique architecture of electrode which would supply more active sites for urea electrooxidation, but incorporation of SrCO_3 in cobalt oxide increases the structural defects of cobalt oxide which lead to providing convenient electroactive pathway for rapid ionic transformation on the surface of electrode during urea electrooxidation. Probably, the conductive network of carbon with strongly attached nano-metallic counterparts not only adsorbs greater number of urea

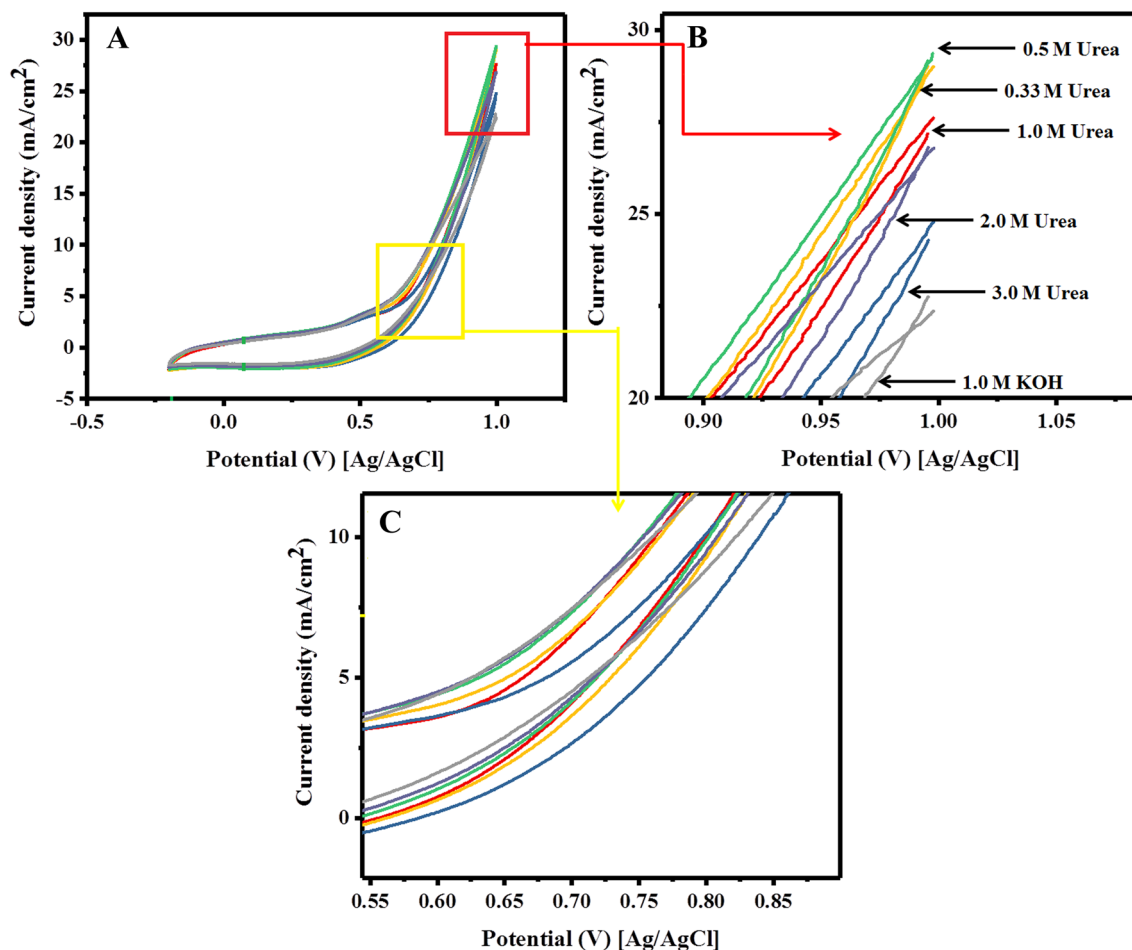


Fig. 8 **a** Typical Cyclic voltammograms for the obtained Co_3O_4 – SrCO_3 heterostructure nanorods-incorporated CNFs in different concentrations of urea (+1 M KOH), **b** high magnification to show

the influence of urea concentration on current density, and **c** high magnification to show the onset potential at scan rate of 50 mV/s at room temperature

molecules but is capable of electrooxidation of urea at low-onset potential with high-current density.

Conclusions

In conclusion, we have successfully synthesized nickel-free electrocatalyst based on Co_3O_4 – SrCO_3 heterostructure nanorods-incorporated CNFs, by a simple electrospinning technique followed by a thermal treatment in inert environment at 750 °C. The resultant Co_3O_4 – SrCO_3 heterostructure nanorods-incorporated CNFs exhibit high-current density at low-fuel concentration. Remarkably, because of good conductive network of carbon nanofibers with high dispersion of metallic counterpart on the surface of carbon nanofibers support, the nickel-free electrode was capable of lowering the onset potential toward urea electrooxidation, representing hopeful applications in direct urea fuel cells anodes.

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