

A facile preparation method for efficiency a novel LaNiO₃/SrCeO₃ (p-n type) heterojunction catalyst in photocatalytic activities, bactericidal assessment and dopamine detection

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ABSTRACT

The wide occurrence of medicines and textiles pollutants in water bodies, as well as their by-products, has raised concerns about the impact of damage to the environment and human health. In this study, the photocatalytic effect of Lanthanum nickelate (LaNiO₃), strontium cerate (SrCeO₃), and LaNiO₃/SrCeO₃ nanocomposite were studied using the co-precipitation method at room temperature. Various techniques were used to analyze the structural, morphological, optical properties of these synthesized nanoparticles. The results indicate uniform growth of LaNiO₃ nanoparticles on the surface of SrCeO₃ nanoparticles in sphere-shaped forms which confirms the formation of a p-n type junction catalyst. The energy bandgap of SrCeO₃, LaNiO₃ and LaNiO₃/SrCeO₃ was found to be 2.79, 2.06, and 1.65 eV, respectively. The photocatalytic activity was studied by degradation of Methylene Blue dye under visible light. As a result, reusability test shows the stability of catalysts for long-term period. The LaNiO₃/SrCeO₃ composites depict the effective degradation for MB dye was 93.5% at 100 min. The rate constant (K, min⁻¹) of SrCeO₃, LaNiO₃, and LaNiO₃/SrCeO₃ was 0.0050, 0.0088, and 0.0156, respectively. The antibacterial action of LaNiO₃/SrCeO₃ versus *Klebsiella pneumoniae* and *Bacillus cereus* observed by using the antimicrobial test. The peroxidase activity of LaNiO₃/SrCeO₃ was performed for colorimetric detection of dopamine. The linear range of the method is 1-200 nM via the detection limit of 3.48 μM. The LaNiO₃/SrCeO₃ has high potential in analysis of dopamine.

1. Introduction

Water pollution has arisen as a key concern for the scientific community in recent years, with fast-rising businesses and populations, necessitating an intensive and valid response. Colored dyes used in the textile industry, like methylene blue are harmful to aquatic life and people who suffer from skin diseases, and lung infections [1–3]. As a

result, a novel low-cost way of eliminating these dyes before releasing them into the environment is required [4]. There are several ways to decompose these organic pollutants in wastewater, but advanced oxidation processes are still a fairly new technique [5–7]. The best methods for reducing pollutants in wastewater systems is photocatalytic degradation with semiconductor photocatalysts [8–11]. Previous research has shown that heterogeneous photocatalysts are particularly

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effective in removing a wide impurities from our wastewater system [12–16]. Various metal oxides, such as TiO₂, Fe₂O₃, ZnO, ZrO₂, Nb₂O₅, V₂O₅, and WO₃, have been employed as excellent catalysts for photocatalytic water treatment [17–20]. Hybrid semiconductor materials produced by p-n-type heterostructures are an extremely effective way of removing organic contaminants [21]. Hybrid semiconductor materials such as TiO₂-SnO₂ [22], CuO-ZnO [23], Mn₂CuO₄/CdO [14], SnFe₂O₄/BiFeO₃ [24] are widely used for photocatalysis process. The use of rare earth metal oxide such as Lanthanum nickelate (LaNiO₃), and strontium cerate (SrCeO₃) with band gaps at ~2.0 eV and 3.0 eV, is the new technique in photocatalysis system [25–27]. These minerals are used frequently to enhance chemical resistance, luminescence, and mechanical strength and have exciting applications in the industrial and technological fields [25]. Because of its crystal structure, and high oxygen storage capacity, SrCeO₃ has received a lot of interest in the photocatalytic degradation of wastewater pollutants [27]. Because the activities and selection of pure SrCeO₃ catalysts are generally increased by modifying the morphology and particle size. The catalytic oxidation can be modified by the LaNiO₃ [26]. The LaNiO₃, which has remarkable properties and is a prospective material in many industries [26].

The important hormone in human body is dopamine (DA). The presence of DA is necessary to maintain the normal activity of the body. Abnormal amount of DA will bad effect on human health and nervous system diseases, such as schizophrenia, and Parkinson's disease were conducted [28]. Thereupon, the quantitative DA detection method is essential for disease diagnosis. The colorimetric technique has many benefits such as facile operation and low cost, and this great for detection of DA [14].

In this study, the synthesize LaNiO₃/SrCeO₃ heterostructures was conducted by a precipitation technique. The physiochemical, optical, and structural aspects of LaNiO₃/SrCeO₃ nanostructures were investigated and the photocatalytic degradation capacity was conducted by employing the MB as a pollutant. The reusability analysis of the LaNiO₃/SrCeO₃ nanocomposites was conducted to investigate the stability of catalysts for usage. The antibacterial properties of the LaNiO₃/SrCeO₃ was conducted against *Klebsiella pneumoniae* and *Bacillus cereus*. The peroxidase like activity of the LaNiO₃/SrCeO₃ nanocomposites was studied in the colorimetric detection of DA. The detection limit was calculated.

2. Experimental

2.1. Chemicals

Materials and reagents are provided in Support Information Text S1.

2.2. Fabrication of LaNiO₃/SrCeO₃ Nanocomposites

Nanocomposite (p-type LaNiO₃/n-type SrCeO₃) catalysts were fabricated by using the co-precipitation method in two steps. In first solution, 0.3 M of lanthanum nitrate hexahydrate, 0.3 M of Nickel nitrate hexahydrate and 0.5 M of sodium hydroxide was dissolved in 150 mL distilled water, which was stirred with a magnetic stirrer for about 50°C for 3 hours. Similarly, in second solution, 0.3 M of cerium nitrate hexahydrate and 0.3 M of strontium nitrate were dissolving in 150 mL distilled water, which was stirred with a magnetic stirrer for about 50°C for 3 hours and 0.5 M of sodium hydroxide solution was added drop by drop until the pH of the solution rises to pH-10. The precipitate was washed and dried at 90°C for 8 h.

To make a suspension, the LaNiO₃ and SrCeO₃ samples were mixed with ethanol as a liquid solvent. The LaNiO₃ powder was dispersed in 100 ml ethanol for 2 hours using a magnetic stirrer. Then, in a similar ratio of (1:1) SrCeO₃ powder was added to the suspension and mixed until a homogeneous mixture was formed. After evaporate the solvent, the product was aged at 150°C for 6 h, and calcined at 500°C for 3 h.

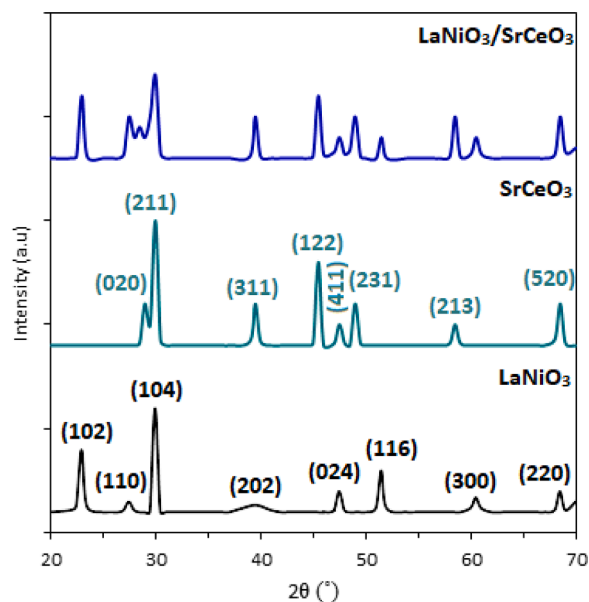


Fig. 1. XRD Patterns of synthesis LaNiO₃, SrCeO₃ and LaNiO₃/SrCeO₃ nanocomposite.

2.3. Characterizations technique

The crystallite size and type of the prepared nanomaterials was conducted by using the PANalytical diffractometer. The surface morphology was examined by FE-SEM (Mira3TESCAN). The absorbance properties and band gap of the prepared sample was recorded by UV-vis spectrophotometry (Agilent carry). The photoluminescence properties was recorded spectrophotometer (Shimadzu RF-5301). The chemical states of elements was recorded by X-ray photoelectron spectroscopy (XPS, VG ESCALab220i- XL). The electrochemical impedance (EIS) was studied by SP-240 Potentiostat.

2.4. Photocatalytic test

The photocatalysis test under UV light (450 W) was investigated using MB dye degradation at room temperature. MB degradation was studied in various exposure time intervals. The spectrophotometer is specifically developed for UV irradiation to analyze catalytic performance. The 30 mL of MB (10 mg/L) was used to this test. The distance of UV lamp from mixture was 13.3 cm. The equilibrium reaction was found after 30 min in dark condition. The LaNiO₃, SrCeO₃, and LaNiO₃/SrCeO₃ catalyst were used to this test. The reusability test was conducted by fifth cycle times. The photocatalysis mechanism was studied by using the scavenging test in the presence of silver nitrate, EDTA, vitamin C, and ethanol for quenching the e⁻, h⁺, •O₂⁻, and •OH.

2.5. Antibacterial test

The antimicrobial activity of the LaNiO₃/SrCeO₃ was determined by calculating the % inhibition of *Klebsiella pneumoniae* and *Bacillus cereus*. Here, the bacterial culture was inoculated and incubated at 37°C for 18 h. The culture was washed with saline water. The 5 mL of the nano-samples concentration (1, 15, 30, 60 mg) was added to 5 mL of bacterial cultures and shacked for 4 h. The OD of the solutions were recorded and the inhibition percentage was determined as the previous study [15].

2.6. Dopamine detection

The LaNiO₃/SrCeO₃ nanocomposites was applied as probe to

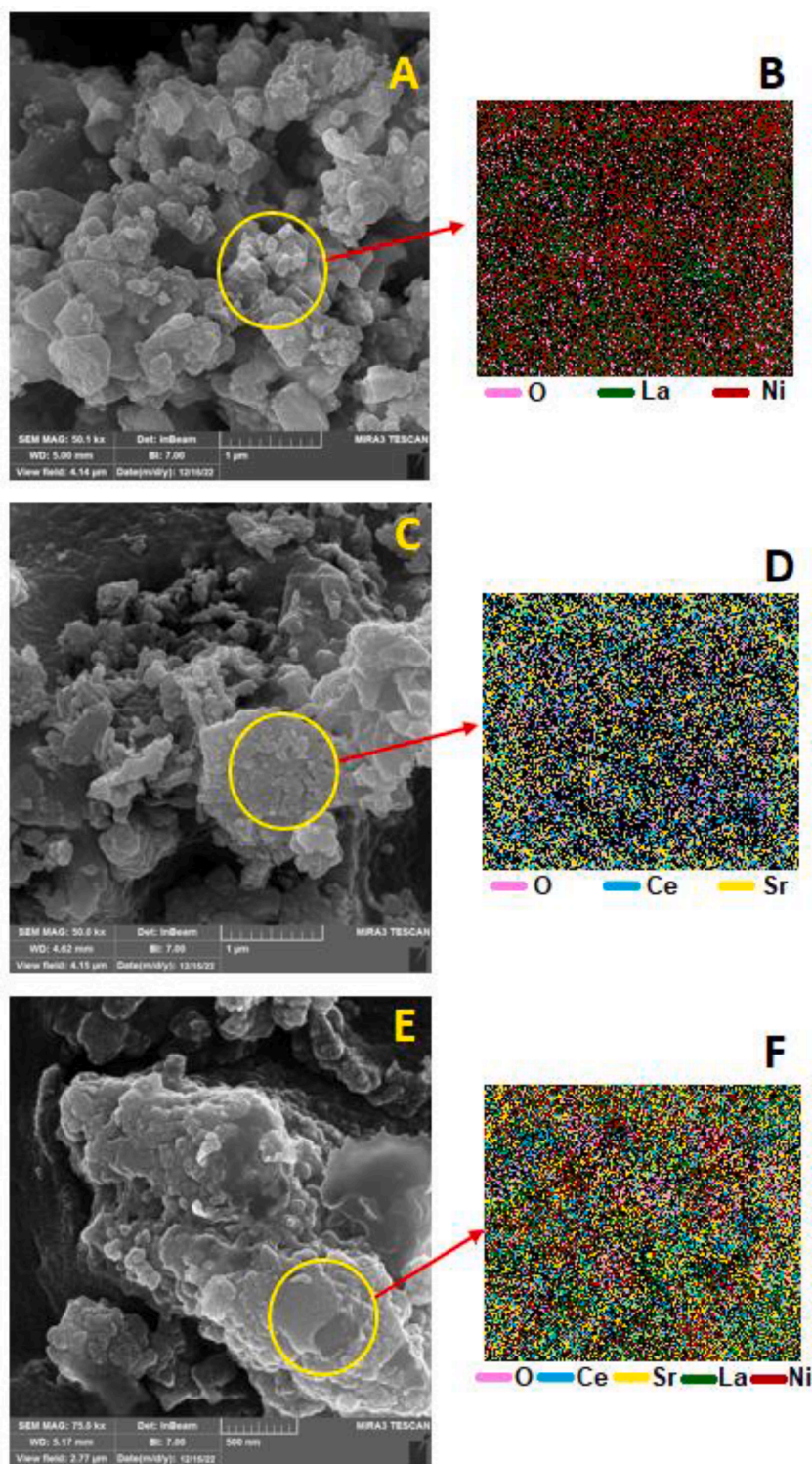


Fig. 2. FE-SEM and elemental mapping of (A, B) LaNiO_3 , (C, D) SrCeO_3 , (E, F) $\text{LaNiO}_3/\text{SrCeO}_3$ nanocomposite.

colorimetric detection of DA by using the $\text{H}_2\text{O}_2/\text{TMB}$ in peroxidase activity. The interaction absorbance was spectrophotometrically monitored by using UV-visible (Agilent carry) spectroscopy. The reaction was conducted by the 50 μL of $\text{H}_2\text{O}_2/\text{TMB}$ solution (1.0 mM to 1.0 mM), 5 mg/L of the $\text{LaNiO}_3/\text{SrCeO}_3$ were added to buffer solution. The selectivity of DA detection was studied by mixing the 30 μL of DA solutions and the interface compounds. The reaction was stirred at 40°C for 30 min. The DA was detected from serum, and urine. The limit of detection for this method was calculated.

3. Results and discussion

3.1. Characterization

The X-ray (XRD) diffraction patterns of LaNiO_3 , SrCeO_3 , and $\text{LaNiO}_3/\text{SrCeO}_3$ nanocomposites are illustrated in Fig. 1. For LaNiO_3 , the diffraction peaks relates with the planes (102), (110), (104), (202), (024), (116), (300), and (220) from hexagonal crystal structure (ICDD 79-2451) [26]. The diffraction peaks for SrCeO_3 are represented by the

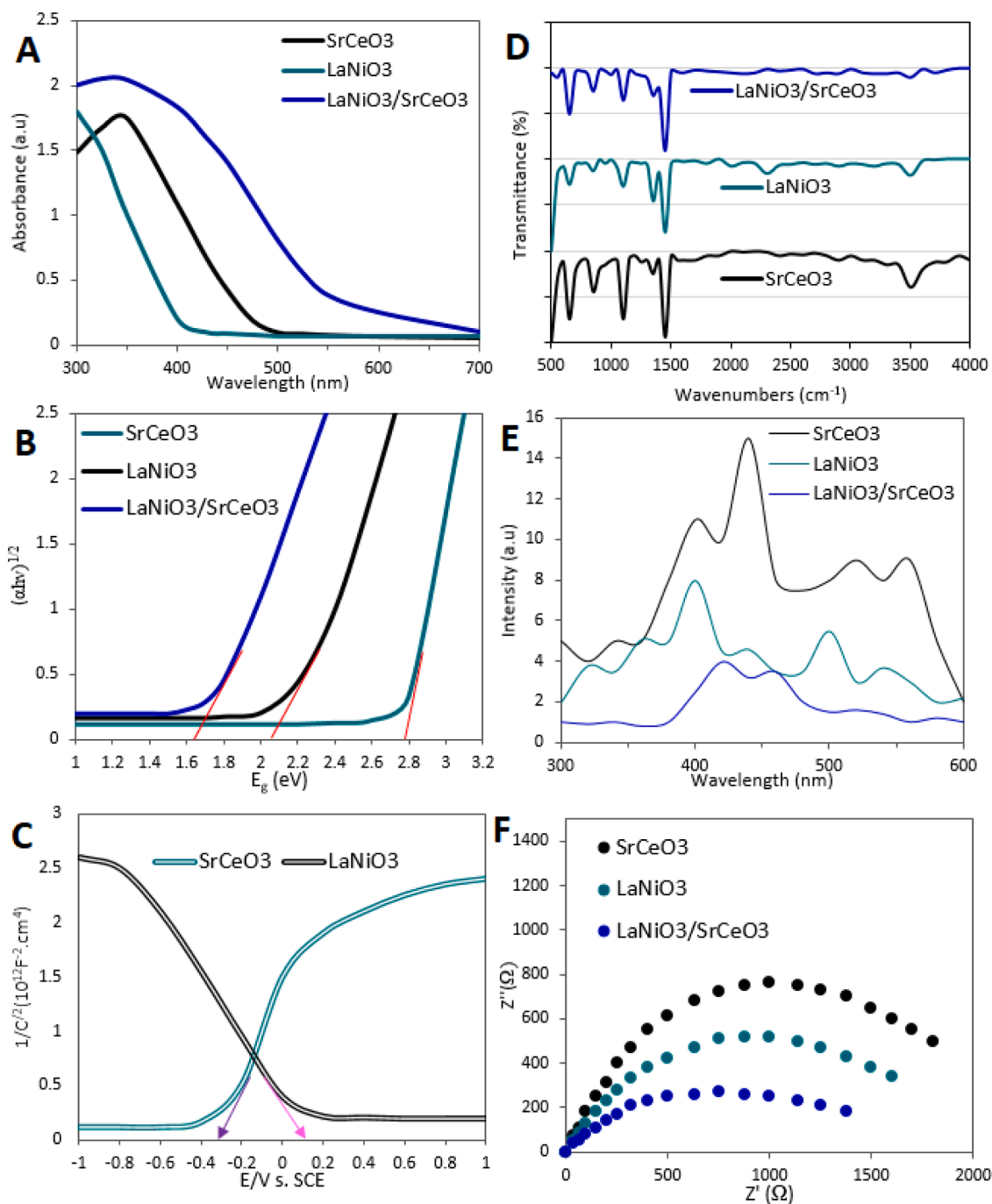


Fig. 3. (A) The absorbance vs wavelength, (B) energy band gap, (C) Mott-Schottky test, (D) FTIR spectra, (E) Photoluminescence spectra, and (F) EIS spectra for LaNiO₃, SrCeO₃ and LaNiO₃/SrCeO₃ nanocomposite

Miller indices (020), (211), (311), (122), (411), (231), (213), and (520) from orthorhombic crystal (ICDD 81-26) [27]. The XRD curve of LaNiO₃/SrCeO₃ nanocomposites confirmed the presence of LaNiO₃ and SrCeO₃ nanoparticles in composite. The result shows that the crystalline phase of the LaNiO₃/SrCeO₃ nanocomposite without any impurities. Using the formula, the average crystallite size was estimated from the Debye-Scherrer equation [15]. The average crystallite size of LaNiO₃, SrCeO₃, and LaNiO₃/SrCeO₃ nanocomposites is 18.3, 25.4, and 32.3 nm. Fig. 2 (A, B) depicts the FESEM images of LaNiO₃ nanoparticles with elemental mapping. It can be seen the spherical shape of LaNiO₃ nanoparticles with smooth surfaces. The elemental mapping shows the presence of La, Ni, and O. Fig. 2 (C, D) shows the irregular morphology of SrCeO₃ with the elemental mapping from Sr, Ce, and O. As shown in

Fig. 2 (E, F), the surface morphology of the LaNiO₃/SrCeO₃ nanocomposite was spherical, with irregular shapes of agglomerated particles. As a result of the elemental mapping of the LaNiO₃/SrCeO₃ depicts the presence of La, Ni, Sr, Ce, and O in composites.

The optical properties of produced LaNiO₃, SrCeO₃, and LaNiO₃/SrCeO₃ materials were investigated UV-vis spectroscopy. The absorption measurements were performed to investigate the band gap level. The optical absorbance of SrCeO₃ was about 400 nm, and increase for LaNiO₃, and LaNiO₃/SrCeO₃ nanocomposites. The intensity of LaNiO₃/SrCeO₃ was higher than SrCeO₃, and LaNiO₃ in the UV region (Fig. 2A). Fig. 2B depicts the energy bandgap (E_g) from the Tauc relation expression [29–37]. As shown, the energy bandgap of SrCeO₃, LaNiO₃ and LaNiO₃/SrCeO₃ was found to be 2.79, 2.06, and 1.65 eV,

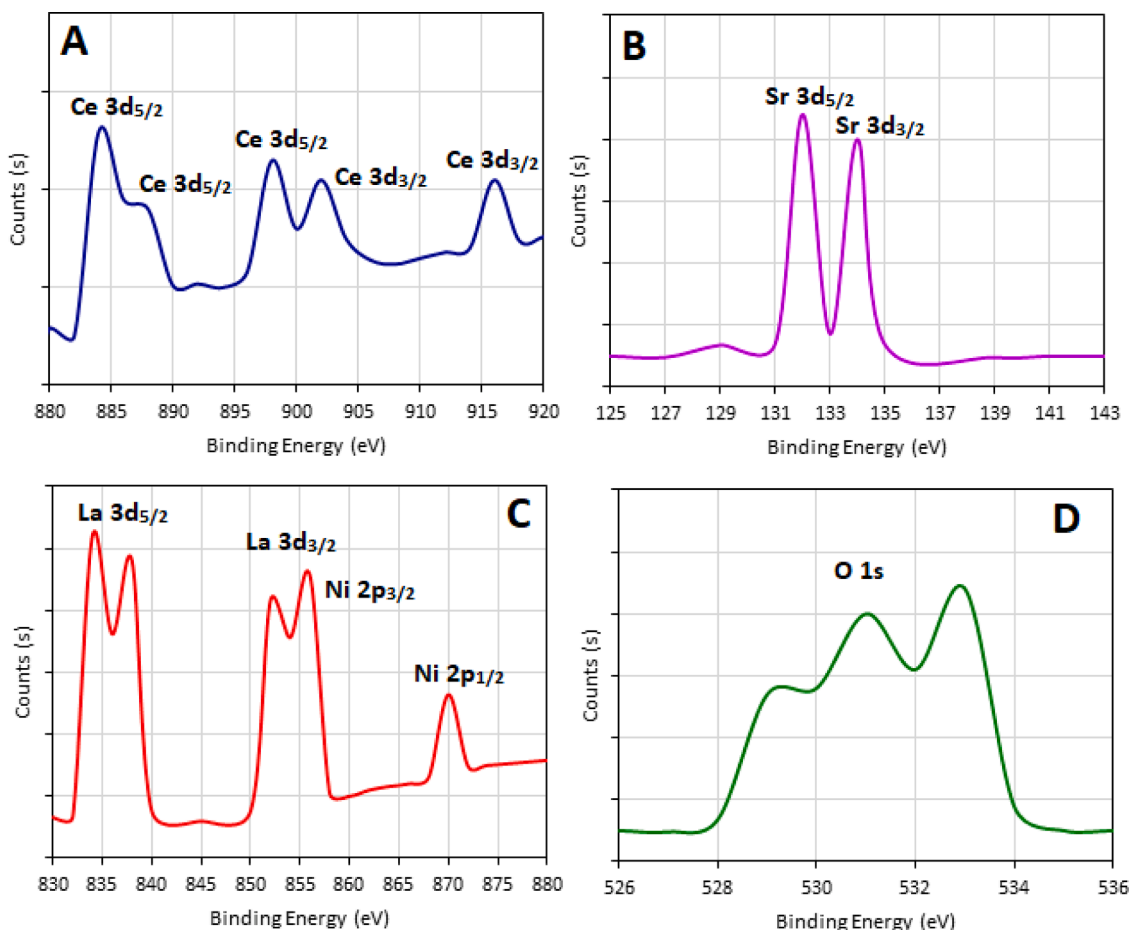


Fig. 4. XPS de-convoluted of (A) Ce 3d, (B) Sr 3d (C), La 3d, Ni 2p, and (D) O 1s from the LaNiO₃/SrCeO₃ nanocomposite.

respectively. The Mott-Schottly test was utilized to checking the flat band potential. As the previous studies, the p-type catalyst such as LaNiO₃, which that shows the activity in high wavelength compared to UV range via lower band gap. The n-type SrCeO₃ catalyst with wide band gap was activated in UV range [38–40]. The flat band potential of LaNiO₃ and SrCeO₃ was investigated and illustrated in the Fig. 2C. As can be observed that the positive slope of n-type SrCeO₃ and negative slope of p-type LaNiO₃ displayed the flat band potential properties [41–43]. Flat band potential values for SrCeO₃ and LaNiO₃ are -0.35 V and +0.13 V, respectively. FTIR spectra were used to examine the major functional groups existing in pure LaNiO₃, SrCeO₃, and LaNiO₃/SrCeO₃. Fig. 3D displays LaNiO₃ with a weak absorption band at 3606 cm⁻¹ in the IR region due to the presence of an O-H stretching vibration from water molecules [26]. Strong broad peaks around 637 cm⁻¹ are related to metal-oxygen stretching vibrations. The sharp peak of about 1463 cm⁻¹ and the weak peak of around 1384 cm⁻¹ can be the asymmetric stretching of CO₃²⁻ functional groups. Furthermore, The C-H stretching is denoted by the peak at 1556.36 cm⁻¹ and the C-O band is represented by the peaks at 1062 cm⁻¹ and 854 cm⁻¹. As a result of FTIR spectra of LaNiO₃/SrCeO₃, indicating that the SrCeO₃ particles were coated with LaNiO₃ [27].

Photoluminescence (PL) and electrochemical impedance spectroscopy (EIS) were applied to investigate the rate of excitation of e⁻/h⁺ pair recombination for synthesized LaNiO₃, SrCeO₃, and LaNiO₃/SrCeO₃. PL measurement is the most effective method for studying the photo induced charge carrier properties of semiconductor materials. The photoluminescence of LaNiO₃, SrCeO₃, and LaNiO₃/SrCeO₃ were investigated. The LaNiO₃, SrCeO₃, and LaNiO₃/SrCeO₃ samples present substantial PL intensity signals and are represented in in Fig. 3E. The SrCeO₃ sample exhibited the highest intensity. As displayed, the

LaNiO₃/SrCeO₃ nanocomposites has much lower PL intensity than the pure LaNiO₃ and SrCeO₃ samples, displaying that bonding with the LaNiO₃ layer on SrCeO₃ may minimize surface defects for charge recombination at an interface [48].

The electrochemical impedance spectroscopy (EIS) was studied to study charge transport carriers for the prepared samples. The EIS spectra were evaluated using the Nyquist plot was displayed in Fig. 3F. A lower arc radius, as is widely known, suggests less interference for electron-hole transmission, meaning that the performance has greater charge carrier separation [35]. The EIS result, demonstrates that the arc radius of the samples is ordered: SrCeO₃ > LaNiO₃ > and LaNiO₃/SrCeO₃ NC has the smallest arc radius, which indicates it has the lowest charge transfer impedance [47]. At low PL and EIS intensity exhibits that, fabricated hybrid LaNiO₃/SrCeO₃ has a lower recombination rate for induced photo-generated charge carriers, resulting the improved photocatalytic degradation efficiency [35]. XPS analysis was studied for information about the chemical and electronic state. XPS analysis of LaNiO₃/SrCeO₃ was conducted. Fig. 4 shows the high resolution XPS scan for the presence of lanthanum, nickel, strontium, cerium, and oxygen in composite. For Ce 3d_{5/2} and Ce 3d_{3/2} XPS spectrum (Fig. 4A), the de-convoluted peaks at 883.2, 889.4 and 897.9, 902.5 and 915.9 eV were suggested that the 4+ (IV) oxidation state of cerium in SrCeO₃ [27]. The Sr 3d spectrum (Fig. 4B) displays the two strongest peaks at 132.3 and 133.9 eV, which related to the Sr 3d_{5/2} and Sr 3d_{3/2} core-level spin-orbits of Sr²⁺ oxidation state, respectively [27]. The La 3d and Ni 2p XPS spectra was depicted in Fig. 4C. It can be seen that the La 3d_{3/2} peak overlap Ni 2p_{3/2} peak, and located at 855.4 eV [26]. The La 3d_{5/2} peak was observed at 834–837 eV. The Ni 2p_{1/2} peak was observed at 870.01 eV. The presence of La 3d and Ni 2p was observed, due to LaNiO₃ [26]. The O 1s can be de-convoluted into three binding energy peaks

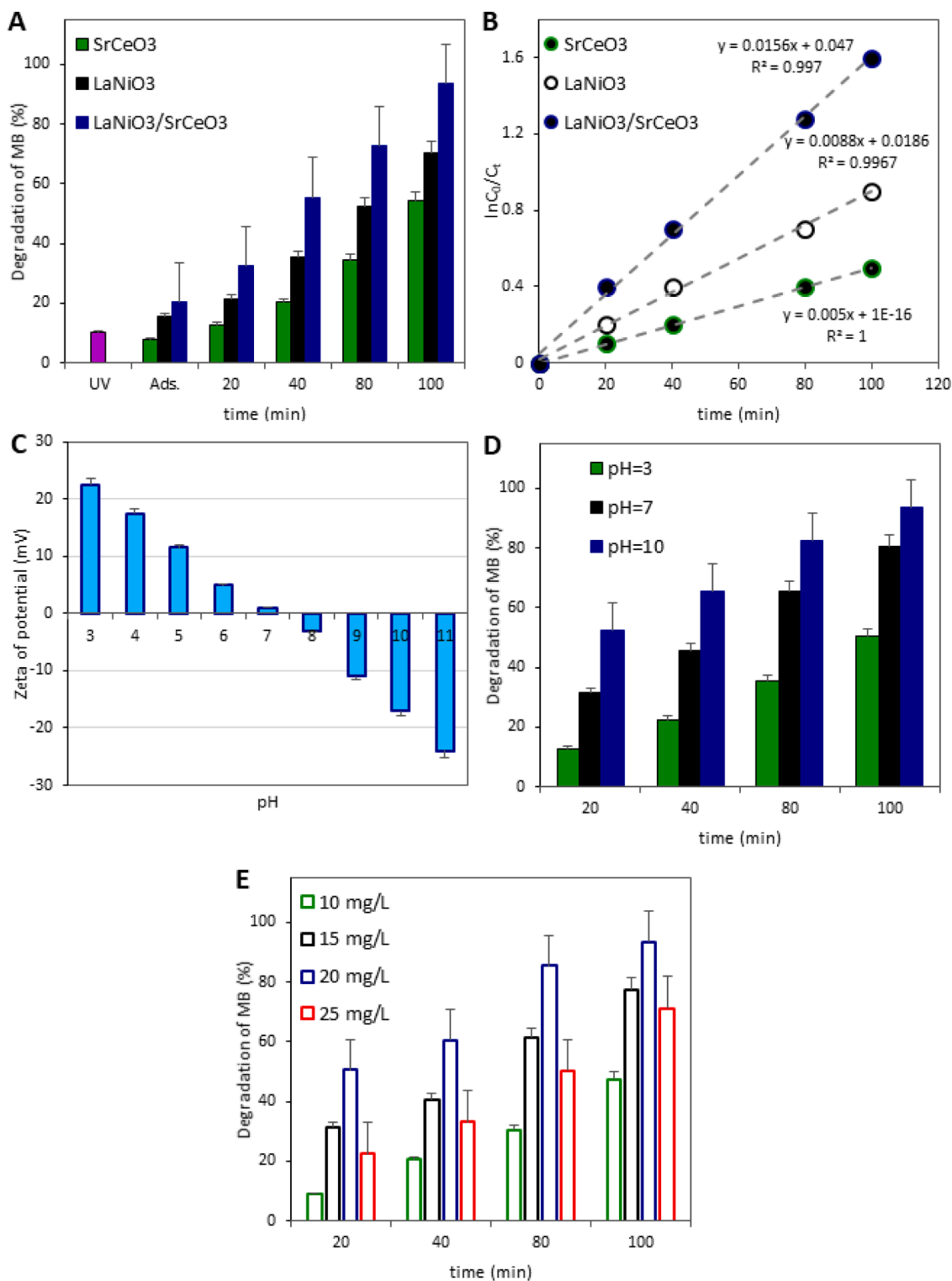


Fig. 5. (A) Degradation of MB dye for UV-light by different catalysts, (B) Photocatalytic reaction kinetics, (C) zeta potential of LaNiO₃/SrCeO₃ nanocomposite, (D) effect of pH on MB degradation, and (E) effect of catalyst dosage on MB degradation.

centered at 529.3, 531.5 and 532.8 eV were ascribed to the lattice oxygen, O-Sr, O-Ni, O-Ce, and O-La [26] (Fig. 4D).

3.2. Photocatalytic activity

The photocatalytic efficiency of the LaNiO₃, SrCeO₃, and LaNiO₃/SrCeO₃ nanocomposites were investigated by degradation of MB with UV irradiation. Fig. 5A shows the degradation of MB in different

condition. It can be seen that the MB was degraded about 10% under UV light without any catalyst. The adsorption capacities of LaNiO₃, SrCeO₃, and LaNiO₃/SrCeO₃ were studied via removal of MB in dark condition. After 30 min, the adsorption capacities of SrCeO₃, LaNiO₃, and LaNiO₃/SrCeO₃ were found 8.31, 15.50, and 20.32%, respectively. The photocatalytic capacities of the prepared catalysts was investigated in different time (20-100 mins). As can be observed, the photocatalysis activity of LaNiO₃/SrCeO₃ was 93.50 % at 100 min, and it's higher than

Table 1
Comparison of photocatalytic degradation of MB.

Catalyst	Light source	Degradation	References
CeO ₂ /SnO ₂	UV	80 %	[54]
CeO ₂ /CuO	Visible	70 %	[55]
TiO ₂ /La ₂ O ₃	UV	95 %	[56]
SnO ₂ /TiO ₂	Visible	90 %	[57]
LaNiO ₃ /SrCeO ₃	UV	93.5 %	This work

the activity of SrCeO₃ (54.35%), and LaNiO₃ (70.51%). The low band gap of LaNiO₃/SrCeO₃, and lower the PL intensity and arc radius can be present the substantial photocatalysis response for degradation of MB under UV light irradiation [44–53].

For different reaction times, Fig. 5B depicts the reaction kinetics of MB degradation by SrCeO₃, LaNiO₃, and LaNiO₃/SrCeO₃. The rate constant (K, min⁻¹) for photocatalytic degradation reactions for all catalysts was found from Fig. 5B. The rate constant (K, min⁻¹) of SrCeO₃, LaNiO₃, and LaNiO₃/SrCeO₃ was 0.0050, 0.0088, and 0.0156, respectively. It can be seen that the rate reaction of LaNiO₃/SrCeO₃ was 1.77, and 3.12 times higher than the LaNiO₃, and SrCeO₃ nanoparticles.

The pH is an important role in the photo-degradation of pollution. Fig. 5C shows the effect of pH on degradation of MB under different times. Based on the results, the effectiveness of MB degradation under UV irradiation utilizing LaNiO₃/SrCeO₃ nanocomposite were 93.5% for alkaline media (pH=10), 80.51% for neutral media (pH=7), and 50.35% for acidic media (pH=3). The surface charges of LaNiO₃/SrCeO₃ at different pHs was investigated and displayed in Fig. 5D. The isoelectric point for LaNiO₃/SrCeO₃ is 7.0. The LaNiO₃/SrCeO₃ has negative charge at the higher isoelectric point. At acidic medium, the LaNiO₃/SrCeO₃ surface has positive charge, and reducing response due to the positive charge of the MB dye [54–56]. The stronger electrostatic interaction with MB and LaNiO₃/SrCeO₃ was done at basic medium. To evaluation of the catalyst dose effect on photocatalytic degradation, different dose of LaNiO₃/SrCeO₃ nanocomposites was added to the MB solution (Fig. 5E). The most effective catalyst dosage was 20 mg/L (93.5 %). The lower catalyst dose leads to less photon absorption by the LaNiO₃/SrCeO₃ nanocomposite, which results in decreasing the catalytic performance. At high catalyst concentrations (25 mg/L), the surface area of the catalyst decreases due to particle agglomeration [57].

Table 1 illustrates the comparison this study to a previous hybrid photocatalyst for the degradation of various organic pollutants. The synthesized LaNiO₃/SrCeO₃ nanocomposite displayed remarkable degradation efficiency for MB dye via exposed to UV-light. Catalyst regeneration in photocatalytic systems is a significant appearance of photocatalysis efficiency for the wastewater purification process [58–62]. After recycling the catalyst, the LaNiO₃/SrCeO₃ were applied for the degradation of fresh MB solution. This reaction was repeated for fifth cycle. Fig. 6A shows the recycling performance of catalytic efficiency. It can be seen that the photocatalysis capacity of the LaNiO₃/SrCeO₃ was negligibly decreased after 5th cycle time (93.5% to 92.51%). The XRD of LaNiO₃/SrCeO₃ was studied after 5th cycle time and are showed in Fig. 6B. The XRD analysis shows the stability of crystal phase after fifth photo-degradation reaction, and confirms the appreciable photostability of p-n type LaNiO₃/SrCeO₃ nanocomposite as a photocatalyst.

To determine the role of reactive species in the MB degradation, a scavenging analysis was conducted. Scavenging investigations paralleled the MB photo-degradation process. To detect such species, four scavengers, such as vitamin C (VC), ethanol (Et), Ethylene diamine tetra acetic acid (EDTA), and silver nitrate (SN) was added to the reaction to scavenging superoxide anions (•O₂⁻) radicals, hydroxyl radicals (•OH), holes (h⁺), and electron (e⁻), respectively (Fig. 6C). The presence of VC and Et reduces the MB decomposition (%). It can be seen that the presence of EDTA and SN can be decreased the photocatalysis amount, due to the •OH, and •O₂⁻ generated by using the h⁺, and e⁻,

respectively, which that showed the •OH, and •O₂⁻ play important role in the MB degradation by LaNiO₃/SrCeO₃ nanocomposites [63–65].

The electron spin resonance (ESR) spectra was applied to evaluation of degradation mechanism as depicted in Fig. 6D, The high signals with intensities were observed for the DMPO-•O₂⁻ and DMPO-•OH under UV irradiation. The signal intensity of DMPO-•OH was approximately same to the DMPO-•O₂⁻, which that showed the •OH and •O₂⁻ are keys for MB degradation process by LaNiO₃/SrCeO₃ catalyst [66].

The diagrammatic mechanism for MB decomposition with LaNiO₃/SrCeO₃ nanocomposite is presented in Fig. 6E. The Mott-Schottly test shows the conduction band of SrCeO₃ and LaNiO₃ are -0.35 V and +0.13 eV, respectively. The band gap of SrCeO₃ and LaNiO₃ was calculated from kubelka-munk plot, and are 2.79, and 2.06 eV, respectively. The valence band was calculated as below:

$$E_{VB} = E_{CB} + E_g \quad (1)$$

By using the equation, the valence band of SrCeO₃ and LaNiO₃ are 2.44, and 2.19 eV, respectively. When a LaNiO₃/SrCeO₃ semiconductor absorbs a photon, electron and hole were generated. Based on the energy diagram, the h⁺ can be transferred from the VB of n-type SrCeO₃ to the VB of p-type LaNiO₃. The excited electrons can be transferred from the CB of p-type LaNiO₃ to the CB of n-type SrCeO₃. Thus, the p-n heterojunction LaNiO₃/SrCeO₃ has better photocatalytic response than the LaNiO₃ and SrCeO₃ [21]. Moreover, the VB potential of LaNiO₃ and SrCeO₃ are positive than the E₀ of •OH/H₂O (1.99 eV) [14], indicating that the h⁺ on the surface of LaNiO₃ can be oxidize H₂O into •OH. In addition, the CB potential of LaNiO₃ is near to SrCeO₃ and are approximately negative to reduce the O₂ to the •O₂⁻ (E₀=-0.33 eV) [15]. Therefore, the role of •OH for degradation of MB was stronger than •O₂⁻ by using the p-n heterojunction LaNiO₃/SrCeO₃ nanocomposites under UV light irradiation, the presence of •OH and •O₂⁻ in system, can be MB degraded to CO₂ and H₂O.

3.3. Antibacterial activities

The antibacterial efficiency of SrCeO₃, LaNiO₃ and LaNiO₃/SrCeO₃ was investigated against *Klebsiella pneumoniae* and *Bacillus cereus* and are revealed in Fig. S1. The antibacterial performance was investigated via various nano samples concentration. The inhibition percentage of nano samples on *K. pneumoniae* and *B. cereus* enhances with raise in the nano samples concentration. The inhibition percentage of LaNiO₃/SrCeO₃ was effective than SrCeO₃ and LaNiO₃ nanoparticles, due to the LaNiO₃/SrCeO₃ can be penetrates in the bacteria cell membrane, and inhibiting the formation of nucleic acid [67]. The production of ROS enhances the inhibition percentage of the nano LaNiO₃/SrCeO₃ material.

3.4. Colorimetric DA detection

In this work, the peroxidase like activity of the synthesized LaNiO₃/SrCeO₃ nanocomposites was investigated for detection of dopamine (DA) in the presence of TMB and H₂O₂. Fig. 7A depicts the absorbance spectra of different samples. It can be seen that the LaNiO₃/SrCeO₃ as probe without interaction with DA, has high absorbance in compared to other samples. The peroxidase activity of LaNiO₃/SrCeO₃ was great response in interaction with DA. It is noteworthy, the LaNiO₃/SrCeO₃ probe shows great response in interaction with DA in the presence of other compound such as urea, glutamic acid, arginine, valine, and lysine. Fig. 7B shows the effect of DA concentration on the peroxidase activity of the LaNiO₃/SrCeO₃ probe, which that showed the response of DA detection was enhanced in the lowest DA concentration. The LOD equation is [Y = 0.2947 X + 0.178, R² = 0.9868], and calculated from linear range of 1-200 nM [13–15]. The LOD was 3.48 nM (S/N = 10). The selectivity response of DA detection was studied by LaNiO₃/SrCeO₃ probe in the presence of other substance such as Urea, Glu A, Arg, Val, and Lys, and illustrated in Fig. 7C. The result depicts that the LaNiO₃/SrCeO₃ detect the DA via other compound without any

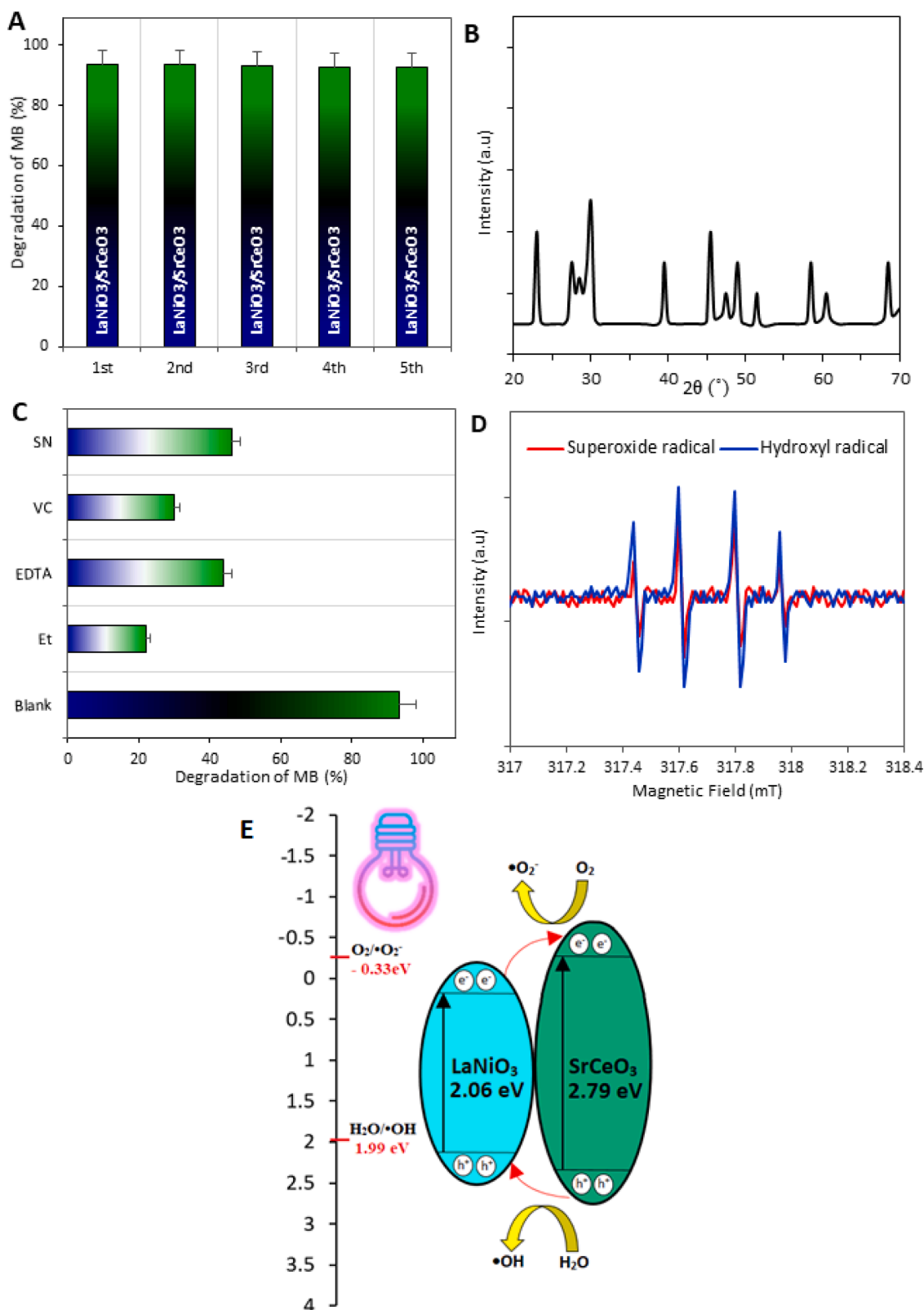


Fig. 6. (A) reusability test, (B) XRD of $\text{LaNiO}_3/\text{SrCeO}_3$ nanocomposite after 5th degradation of MB, (C) scavenging test, (D) DMPO spin-trapping ESR spectra of $\text{LaNiO}_3/\text{SrCeO}_3$, and (E) The possible photocatalytic mechanism.

noises. The accuracy of the prepared method was evaluated by the standard addition technique. After adding known concentration of DA (10, and 20 nM) into 1.0 mL of urine or serum, the probe was used to concentration of DA in two samples. Fig. 7D depicts the DA detection

from urine and serum. It can be seen, DA detected as 9.8 and 19.9 nM in serum, and 10.2 and 21.2 nM in urine. The recovery rate of this probe is 98-106%.

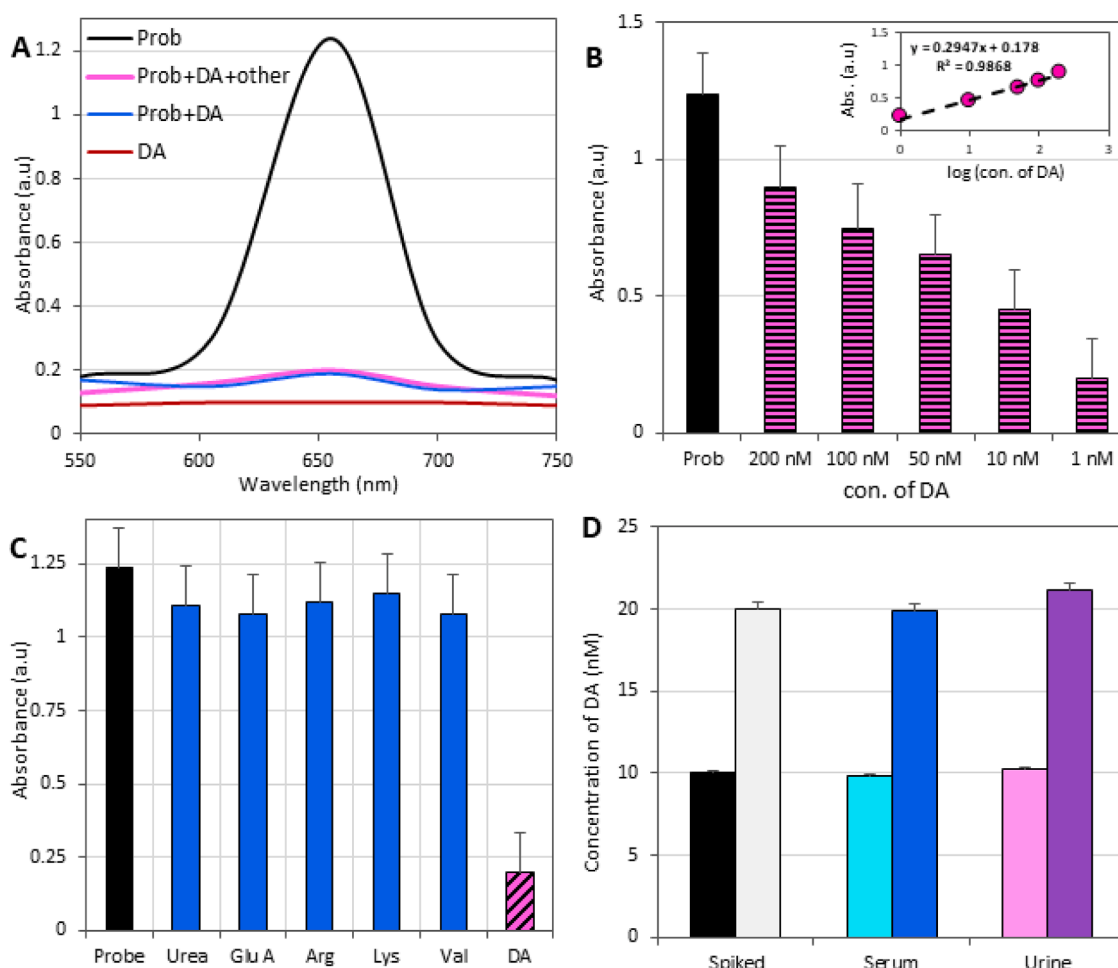


Fig. 7. (A) The peroxidase activity of samples in UV–vis spectra, (B) the effect of DA concentration (inset: calibration curve), (C) the selectivity DA detection, and (D) the DA detection from serum and urine.

4. Conclusion

In conclusion, a LaNiO₃/SrCeO₃ nanocomposites was prepared using a co-precipitation method, and the structural and optical properties were studied using various techniques. The XRD results the formation of a LaNiO₃/SrCeO₃ nanocomposites. The energy band gap of prepared nanocomposite was studied using UV-visible spectroscopy and shows that the combining of LaNiO₃ nanoparticles to SrCeO₃ can be decrease the energy band gap of LaNiO₃/SrCeO₃. The investigation confirms that the degradation of MB for LaNiO₃/SrCeO₃ nanocomposite was 93.5%, which is higher when compared to the pure LaNiO₃ and SrCeO₃ nanoparticles. The rate reaction of LaNiO₃/SrCeO₃ was 1.77, and 3.12 times higher than the LaNiO₃, and SrCeO₃ nanoparticles. Furthermore, the role of •OH for degradation of MB was stronger than •O₂ by using the p-n heterojunction LaNiO₃/SrCeO₃ nanocomposites under UV light irradiation. The inhibition percentage of LaNiO₃/SrCeO₃ versus *Klebsiella pneumoniae* and *Bacillus cereus* was effective than SrCeO₃ and LaNiO₃ nanoparticles. The as-synthesized LaNiO₃/SrCeO₃ probe was used to dopamine detection. The probe depicts the superior analysis parameters. The LOD value of LaNiO₃/SrCeO₃ probe for DA detection was 3.48 nM.

CRedit authorship contribution statement

Ying Chen: Methodology, Investigation, Formal analysis. **Ali Jihad:** Formal analysis, Methodology, Investigation, Validation. **Fadhil Husam:** Formal analysis, Methodology, Investigation. **Salah Hassan Zain Al-Abdeen:** Formal analysis, Methodology, Investigation. **Jamal**

Mohammad Hussein: Formal analysis, Methodology, Investigation. **Zainab Hussein Adhab:** Formal analysis, Methodology, Investigation. **Zahraa Hamzaa Abd Alzahraa:** Formal analysis, Methodology, Investigation. **Irfan Ahmad:** Formal analysis, Methodology, Investigation, Validation, Resources. **Leila Fatolahi:** Investigation, Validation, Software, Data curation. **Baadal Jushi Janani:** Writing – review & editing, Methodology, Investigation, Validation, Software, Project administration, Conceptualization, Supervision.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Data availability

The data that has been used is confidential.

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