Preparation and Characterization of LaCr$_{0.99}$Fe$_{0.01}$O$_3$ Nanomaterial

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Abstract. Perovskite compounds show great potential in a variety of applications due to their diverse and unique properties and can be modified. Nanomaterial LaCr$_{0.99}$Fe$_{0.01}$O$_3$ has been prepared using sol-gel and freeze drying methods. The nanomaterial was then calcined at 700ºC for 10 hours and characterized by the analysis techniques of X-Ray Diffraction (XRD), Fourier Transform Infrared (FTIR), Diffuse Reflectance UV-Vis Spectroscopy (DRS UV-Vis), and Transmission Electron Microscopy (TEM). The result of XRD analysis showed that two main crystal phases were formed from the ABO$_3$ perovskite structure, namely LaCrO$_3$ and LaFeO$_3$ which were superimposed and the crystalline phase size based on the Scherrer method was 23.5 nm. Furthermore, the TEM analysis result shows that there are various forms and hollows. Then, the FTIR spectrum states that there is an interaction between transition metals, and the bandgap energy is 2.71 eV.

Keywords: perovskite, nanomaterial, hollow, bandgap energy

1. Introduction
Perovskite material is well known as a material with many benefits [1–4], and each year researchers publish more than 2000 articles in various applications [5] related to perovskite material. The uniqueness of perovskite compounds, especially the type of oxide, is due to the ease of controlling the valence of the cations at A and B sites through partial substitution as well as non-stoichiometry control through cationic or anionic vacancies in their oxide structure based on an electroneutralite point of view [6]. This control leads to the existence of various cationic oxidation states which have characteristics as catalysts [7-9], ferroelectricity [10, 11], piezoelectricity [12, 13], pyroelectricity [14, 15], thermoelectricity [16], magnetism [17, 18], superconductivity [19, 20], sensors [21-23], optics [24, 25], and fuel cells [26] so that the application of oxide-type perovskite materials has proven to be very wide [27].

Oxide-type perovskites can be prepared from transition metal oxides or their salts using a variety of preparation methods such as hydrothermal [28-30], co-precipitation [31-33], and sol-gel [34, 35]. In general, the perovskite structure is ABO$_3$ although the cations at sites A and B can be modified such as A$_1$A$_2$BO$_3$ or AB$_1$B$_2$O$_3$. It can even be more complex A$_1$A$_2$B$_1$B$_2$O$_3$ and the structural model can be presented in Figure 1 below.
The success of preparing perovskite compounds depends also on the selection of cations for sites A and B, which generally follow the following rules,

\[ t = \frac{(r_A + r_O)}{\sqrt{2(r_B + r_O)}} \]

Where \( t \) is the Goldschmidt tolerance factor, \( r_A, r_B, \) and \( r_O \) are the radii of A, B, and oxygen ions, respectively. The value of the Goldschmidt tolerance factor must be in the range from 0.75 to 1.00 in order to obtain the perovskite structure [36]. Several combinations of A- and B-site cations can form a stable perovskite-like structure. Additionally, A- and B-site cations, as well as oxygen anions can be partially substituted by other suitable elements [37, 38].

Based on the unique characteristics and flexibility of its structure [39, 40], where the A cation and the B cation that fill the octahedral and tetrahedral positions can exchange even be replaced by other inorganic cations or organic molecules. So that the modified perovskite material provides the desired characteristics for its application [41, 42]. One of the most commonly known perovskite oxide types is lanthanum chromite, \( \text{LaCrO}_3 \) with a wide range of applications such as electronics, opto-electronics, energy storage, thermal catalysis, sensors and photocatalysts [43-46].

In the field of photocatalysis, one of the characteristics of the \( \text{LaCrO}_3 \) perovskite as a reference is the ease of promoting electrons from the valence band (HOMO) to the conductance band (LUMO). The smaller the energy required to move electrons from the valence band to the conductance band, the greater the chance that the electrons will initiate the target reaction. Based on the obtained references, the perovskite gap energy of \( \text{LaCrO}_3 \) and its modification is in the \( \leq 3.0 \) eV range [47, 48] so that the perovskite of this type of oxide can function as a photocatalyst because it is semiconductor [49].

On this occasion, we reported the preparation and characteristics of perovskite oxide types of lanthanum chromites doped with Fe cation, \( \text{LaCr}_{0.99}\text{Fe}_{0.01}\text{O}_3 \) using the sol gel method and its possible application in photocatalytic reaction by analyzing the material using X-Ray Diffraction, Fourier Transform Infrared, Diffuse Reflectance UV-Vis Spectroscopy, and Transmission Electron Microscopy.

2. Experimental

2.1. Material
The materials used in this study include \( \text{La(NO}_3)_2 \cdot 6 \text{H}_2\text{O} \) (Merck, 99%), \( \text{Cr(NO}_3)_2 \cdot 9 \text{H}_2\text{O} \) (Merck, 99%), \( \text{Fe(NO}_3)_2 \cdot 9 \text{H}_2\text{O} \) (Merck, 99%), \( \text{NH}_3 \) (Merck, 99%), pectin and distilled water.
2.2. Instruments
The instruments used for characterization include X-Ray Diffractometer (XRD) model PW 1710 with Cu-Kα radiation for structure identification, Fourier Transform Infrared (FTIR) type Shimadzu Prestige-21 for functional group identification, Diffuse Reflectance UV-Vis Spectroscopy (Agilent Cary 60) to determine the band-gap energy, and Transmission Electron Microscopy (TEM) JEOL JEM 1400 for identifying grain size distribution and crystallite morphology.

2.3. Preparation of \( \text{LaCr}_{0.99}\text{Fe}_{0.01}\text{O}_3 \)
The \( \text{LaCr}_{0.99}\text{Fe}_{0.01}\text{O}_3 \) nanocatalyst was prepared by the sol-gel method. The procedure was described in the previous work [50].

2.4. Characterization of \( \text{LaCr}_{0.99}\text{Fe}_{0.01}\text{O}_3 \)

2.4.1. X-Ray Diffraction Analysis. The X-ray diffraction pattern \( \text{LaCr}_{0.99}\text{Fe}_{0.01}\text{O}_3 \) was recorded at \( 2\theta = 10-90^\circ \) using a Philips diffractometer model PW 1710 Cu-Kα radiation. Phase identification was carried out by matching the diffractogram of the sample to the standard diffractogram using the JCPDF published by ICCD PDF. The particle size was determined by the Scherrer method [51].

2.4.2. FTIR Analysis. After heating at 90°C, the samples were analyzed using FTIR spectroscopy. Spectroscopic analysis is performed by grinding the sample with KBr and scanning it at a wave number 4000-400 cm\(^{-1}\) [52].

2.4.3. UV-Vis DR Spectroscopy Analysis. Determination of the \( \text{LaCr}_{0.99}\text{Fe}_{0.01}\text{O}_3 \) band-gap energy, a number of samples were analyzed using UV-Vis Diffuse Reflectance Spectroscopy and scanned at a wavelength of 200-800 nm [53].

2.4.4. Transmission Electron Microscope Analysis. To evaluate the surface morphology, the samples were characterized using TEM. The analysis was conducted on polished and thermally etched samples with different magnifications [54].

3. Results and Discussion

3.1. Structural Analysis of X-Ray Diffractogram
Based on the diffractogram shown in Figure 2, the diffraction pattern of \( \text{LaCr}_{0.99}\text{Fe}_{0.01}\text{O}_3 \) was compared with the standard using the JCPDF program. The results showed that two main crystalline phases were formed which is superimposed with each other, namely \( \text{LaCrO}_3 \) (PDF-24-1016) and \( \text{LaFeO}_3 \) (PDF-37-1493) with the highest intensity at \( 2\theta = 32^\circ \). The crystal size of the sample is determined by the following Scherrer equation:

\[
D = \frac{K\lambda}{\beta\cos\theta}
\]

Where \( K \) is the crystallite form factor, \( \lambda \) is the X-ray wavelength (1.5406 nm for CuKα), \( \beta \) is the width of the diffraction peaks, and \( \theta \) is the Bragg angle. The average size of the crystalline phase of \( \text{LaCr}_{0.99}\text{Fe}_{0.01}\text{O}_3 \) was 23.5 nm.
3.2. Analysis of FTIR Spectrum

Figure 3 shows the FTIR spectra of the LaCr\textsubscript{0.99}Fe\textsubscript{0.01}O\textsubscript{3} nanomaterial prepared at a calcination temperature of 700ºC in the wave number range 4000-400 cm\textsuperscript{-1}. The FTIR spectrum of LaCr\textsubscript{0.99}Fe\textsubscript{0.01}O\textsubscript{3} is shown in Figure 4. The broad band observed at 3445 cm\textsuperscript{-1} and 1636 cm\textsuperscript{-1} corresponds to the stretching vibration and bending vibration of O-H groups respectively. The symmetric stretching of carboxyl C=O bond was not observed at 2356 cm\textsuperscript{-1} due to the adsorption of atmospheric carbon dioxide, as the measurement was recorded with pretreatment. The weaker intense peak as a shoulder observed at around 500 cm\textsuperscript{-1} due to the presence of stretching vibration of Fe-O confirms the octahedral FeO\textsubscript{6} present in the LaCr\textsubscript{0.99}Fe\textsubscript{0.01}O\textsubscript{3} [55]. At the wave number 601 cm\textsuperscript{-1}, it shows the presence of Fe-O stretching vibrations [56] and La-O-La bending vibrations [57]. The presence of Cr-O and O-Cr-O stretching vibrations at 884 cm\textsuperscript{-1} absorption [58].
3.3. Analysis of DRS Spectrum

Optical absorption bands were detected on perovskite LaCr$_{0.99}$Fe$_{0.01}$O$_3$ through the % reflectance and absorbance spectra of the wavelength in Figure 4 above, where the four basins in Figure 4A at wavelengths of about 380, 450, 620, and 740 nm show absorption peaks in Figure 4B. The peaks indicate that there is an interaction between the 3d Fe$^{3+}$ orbital and the 3d Cr$^{3+}$ orbitals with the 2p O$^{2-}$ orbitals, respectively, which are shown at the 380 and 450 nm wavelengths of absorption, respectively. Most probably weak absorption bands at 630 and 740 nm belongs to d-d transition in Fe$^{3+}$ and Cr$^{3+}$ ion in an octahedral symmetry [58], which is influenced by replacing Cr with Fe in LaCr$_{0.99}$Fe$_{0.01}$O$_3$ perovskite structure. Intra-atomic d-d transitions are of much lower intensity comparison to the inter-atomic transitions at region of fundamental absorption edge, which are mainly determining the color of iron-rich oxides [59].

![Figure 4](image_url)

**Figure 4.** UV-Vis DR Spectra of LaCr$_{1-x}$Fe$_x$O$_3$

Figure 4C shows the bandgap energy obtained graphically through extrapolation when \( (F(R)\cdot h\nu)^2 \) to bandgap energy \( (E_g) \) on the X-axis of 2.71 eV. Order 2 in the equation \( (F(R)\cdot h\nu)^2 \) denotes an electronic transition that occurs indirectly. Based on the \( E_g \) value obtained, it can be said that the LaCr$_{0.99}$Fe$_{0.01}$O$_3$ nanomaterial can function as a photocatalyst.

3.4. Morphological Analysis

Characterization of the samples using TEM produced the micrographs as presented in Figure 5 below. In general, there are several structural shapes identified such as cubic (brown arrow), circular (yellow arrow) and pipe (blue arrow). If focusing on perovskite structure, it displays, the existence of LaCr$_{0.99}$Fe$_{0.01}$O$_3$ crystalline phase as a unit cell since LaCrO$_3$ and LaFeO$_3$ is superimposed. However, in a certain area, the presence of a bigger perovskite structure can be observed, which confirms the existence of layers (red arrow) and hollow cubes (green arrow).
Although various forms of perovskite appear to exist, the resulting sizes are mostly in the nano category. It seems that further analysis is needed regarding the hallow form. Based on the 50 nm scale (right-micrograph), it can be said that the grains are 50-100 nm in size. Few grains appeared to be in the planar connection so that they seem to be greater than 100 nm, and a few appear as circular with a size less than 50 nm. In addition, there was a tube with a diameter of 98 nm and its length of 163.3 nm. Overall result proved that the material prepared is nano size and hollow. The size result is smaller than that of LaCrO$_3$ modified by Fe and Co which is 168 nm [60].

4. Conclusion

Based on the analysis of LaCr$_{0.99}$Fe$_{0.01}$O$_3$ material characteristics, it can be concluded that the prepared materials have the perovskite structure consisting of LaFeO$_3$ and LaCrO$_3$ which is superimposed and formed into layers or piles that stick together. In fact, the structure formed is various. It has nanopipes, nanocubes, nanocircular and hollow structures. So that the perovskites are nano hollow material. Furthermore, since its band-gap energy is around 3.0 eV, it is implied that this material could be used in photonic applications.

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