Article
Ternary Ni-Ce-Mg-O Composites: In-Depth Optical Spectroscopy Study and Catalytic Performance in CO Oxidation
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Abstract: In the present work, ternary Ni-Ce-Mg-O composites containing various amounts of NiO and CeO₂ were synthesized via a sol-gel approach. Aqueous solutions of cerium and nickel nitrates were introduced at the stage of hydrolysis of magnesium methoxide, which allowed for avoiding the use of expensive organic precursors. It was revealed that the properties of the composites were defined by the complex interactions between NiO, CeO₂, and MgO components. In order to perform an in-depth characterization of the prepared samples, diffuse reflectance UV-vis and Raman spectroscopies were applied. According to the results of these methods, Mg²⁺ ions did not substitute Ce⁴⁺ ions in the CeO₂ lattice. However, in the case of the Ni-containing samples, approximately 2–3% of the Ce⁴⁺ ions were substituted by Ni²⁺, thus resulting in the formation of vacancies in the CeO₂. The strong interaction of NiO with MgO predictably resulted in the formation of NiₓMg₁−ₓO solid solutions. When the NiO content in the sample was 20 wt%, the composition of the formed solid solution was estimated to be Ni₀.₆₀Mg₀.₄₀O. In addition, the presence of CeO₂ affected the texture of the ternary composites, thus leading to a slight decrease in the specific surface area. The catalytic performance of the Ni-Ce-Mg-O composites was examined in the CO oxidation reaction under prompt thermal aging conditions. The choice of reaction conditions was due to a high sensitivity of the CO oxidation response toward the available metal surface area and possible metal-support interactions.

Keywords: nickel; CeO₂; MgO matrix; CO oxidation; prompt thermal aging; diffuse reflectance UV-vis spectroscopy; Raman spectroscopy

1. Introduction

One of the most industrially important applications of oxide materials is their use as catalyst support. Among such oxides, Al₂O₃ and SiO₂ attract much attention, and therefore they are used in real industrial processes more frequently. Contrarily, MgO is often considered as a model support with pronounced basic properties. The main reason for this is that Al₂O₃ and SiO₂ generally possess higher specific surface and porosity values than MgO prepared using conventional methods. However, during the last decades, MgO has been intensively studied within the composition of the catalysts for various catalytic processes. For instance, the AuAg-MgO system was effective in n-octanol oxidation [1], Ru-MgO was applicable for ammonia decomposition [2], Co/MgO showed an appropriate activity in glycerol steam reforming [3], Cu/MgO was applied in the water–gas shift reaction [4], CO oxidative coupling to dimethyl oxalate was performed over Pd/MgO [5], aerogel-prepared VOₓ/MgO system exhibited promising activity in oxidative dehydrogenation of propane [6], and Ni/MgO was used for CO₂ methanation [7,8]. In addition, the basic properties of magnesium oxide make it possible to create efficient systems for carbon dioxide capture, which can be regenerated many times without a loss in adsorption capacity [9–11].

The most attractive directions of application of MgO-based systems deal with such processes as steam reforming and carbon dioxide reforming of methane. Ni/MgO catalysts
are mostly used for these purposes [12–17]. As has been reported, such systems have excellent stability and are less inclined to coke formation. In this case, MgO plays a significant role in stabilizing the clusters of metallic nickel on the surface, thus contributing to the preservation of catalytic activity [15]. The doping of Ni/MgO catalysts with such elements as ruthenium [18], molybdenum [19], and palladium [20] have been reported to further improve the catalytic performance. The addition of cerium and zirconium oxides also positively affects the Ni/MgO catalysts, increasing their activity in both steam and dry reforming processes. Thus, Kim et al. demonstrated that such supports possessed a larger surface area and better dispersion of Ni species [21]. Khajenoori et al. [22] reported that the doping of Ni/MgO with cerium oxide allowed decreasing the number of carbon deposits formed on the Ni particles during the reaction and weakened the chemical interaction between NiO and the support, which simplified the reduction of NiO species. The optimal composition of the catalyst was found to be 10% Ni–7% CeO$_2$/MgO [23]. According to Qiujie et al. [24], the MgO-to-CeO$_2$ ratio of 1:1 allowed for achieving the highest activity, stability, and selectivity in the steam reforming of ethanol. As is known, CeO$_2$ possesses a high oxygen storage capacity that plays an important role during the catalytic process. An intensified oxygen transfer facilitates the removal of carbon deposits from the metal surface [25]. It is important to mention that once the lattice oxygen in CeO$_2$ is depleted, it can be regenerated by the atmospheric oxygen. This allows for CeO$_2$ to be used as a component of a catalytic washcoat in diesel particulate filters. The redox properties of CeO$_2$ can be enhanced by the addition of transition metals such as Cu and Ag, thus increasing the oxygen storage capacity as well as the reactivity of the surface oxygen species [26,27].

In some cases, the combination of the oxidative properties of CeO$_2$ with the basic properties of MgO provides a synergetic effect, improving the process efficiency. Thus, the introduction of CeO$_2$ into the MgO matrix allows the acceleration of the initial rate of SO$_2$ adsorption due to its oxidation by the lattice oxygen of cerium oxide to SO$_3$ [28]. Ru/CeO$_2$-MgO materials were also used for the CO$_2$ capture/methanation process. In this case, the morphology of the particles was proven to have a significant effect on the activity of CeO$_2$ [29]. Binary oxides of such composition as Ce$_{0.05}$Mg$_{0.95}$O$_{1.05}$ demonstrated excellent performance in the dry reforming of methane in the presence of H$_2$S. CO$_2$ captured on the CeO$_2$-MgO interface behaved as an oxidant in the reaction with CH$_4$ [30]. Ivanova et al. [31] showed that the introduction of CeO$_2$ into magnesium oxide can lead to the formation of (Mg, Ce)O$_2$ solid solutions along with an increase in the concentration of both strong and weak Lewis acid sites.

The properties of the ternary NiO-CeO$_2$-MgO composite systems are defined by a complex interaction between all three components. MgO can interact with NiO with the formation of Ni$_x$Mg$_{1-x}$O solid solutions [32–35]. This contributes to the stabilization of NiO species at elevated temperatures. On the other hand, the migration of Ni$^{2+}$ ions into the bulk of the MgO lattice is not a desirable process because this makes these nickel species inaccessible for the reagents and complicates their reduction. As with Ni$^{2+}$ ions, Mg$^{2+}$ ions are capable of replacing cerium ions in the CeO$_2$ lattice with the formation of oxygen vacancies [31,36]. It is obvious that the appearance of such vacancies reflects in the catalytic performance of the materials. Thus, Zhang et al. [37] emphasized the role of such vacancies in the activation of oxygen on the CeO$_2$. An increase in the number of vacancies reduced both the band gap and the activation energy, thus leading to greater specific activity. For the CO oxidation reaction over NiO/CeO$_2$ catalysts, the interactions near the NiO-CeO$_2$ interface were shown to be of great importance [36,38]. Oxygen activated on the CeO$_2$ surface reacted with CO adsorbed on NiO, which intensified the overall process. It is important to note that the surface migration of adsorbed intermediates is complicated, which emphasizes the role of the interface interactions. NiO-MgO-CeO$_2$ systems have been recently tested as oxygen carriers for the chemical looping process of CO$_2$ splitting [39].
It can be concluded from the abovementioned studies that the development of preparation methods to synthesize such ternary composites as NiO-CeO$_2$-MgO is in high demand. Therefore, the development of an effective synthetic method for the preparation of such catalysts is an important task. Various preparation approaches have been proposed to obtain multicomponent systems based on MgO, such as co-precipitation [21,31], sequential impregnation [22,23], and the solid-phase burning method [24]. Co-precipitation and impregnation methods often do not allow obtaining a uniform distribution of the active component. In addition, the surface area of commercially available MgO supports does not exceed 50–60 m$^2$/g [7,13,32]. Pure MgO and MgO-based systems with the highest surface area and pore volume can be obtained with supercritical drying and the calcination of magnesium hydroxides prepared via magnesium alkoxide hydrolysis [6]. This method also provides the possibility to introduce the second component in a form of a precursor soluble in organics.

In our previous publications, a simplified approach based on sol-gel technology to prepare MgO-containing systems was proposed [40–43]. The main idea of this approach was to introduce the second and further components in the form of an aqueous solution of suitable precursors at the stage of magnesium methoxide hydrolysis. This technique was successfully used for the preparation of a ternary Ni-Mo-Mg-O oxide system [44]. In this case, ammonium heptamolybdate was used as a water-soluble precursor of molybdenum oxide. The goal of the present work was to prepare a ternary Ni-Ce-Mg-O composite via the proposed sol-gel approach. Low-temperature nitrogen adsorption was used to determine the textural properties of the prepared samples. In order to study the possible interactions between the NiO, CeO$_2$, and MgO components of the composites in more detail, optical spectroscopic methods (Raman spectroscopy and UV–vis spectroscopy) were utilized. The catalytic performance of the Ni-Ce-Mg-O systems was examined in a model reaction of CO oxidation performed in prompt thermal aging (PTA) mode. Such conditions allowed us to follow both the low-temperature and high-temperature changes taking place during the reaction. As was mentioned above, the CO oxidation reaction was sensitive to the dispersion and structure of the active component as well as to its interaction with the support.

2. Materials and Methods

2.1. Preparation of the Composites

The sol-gel technique was used to prepare ternary composites containing NiO, CeO$_2$, and MgO. The NiO:CeO$_2$ weight ratio was varied from 1:3 to 3:1. The total loading of NiO and CeO$_2$ was 20 wt%. Two reference samples of binary composites containing 20 wt% of NiO or CeO$_2$ were prepared as well. The composition and corresponding designation of the prepared samples is presented in Table 1.

<table>
<thead>
<tr>
<th>Sample</th>
<th>NiO Loading, wt%</th>
<th>CeO$_2$ Loading, wt%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ni20</td>
<td>20</td>
<td>0</td>
</tr>
<tr>
<td>Ni15Ce5</td>
<td>15</td>
<td>5</td>
</tr>
<tr>
<td>Ni10Ce10</td>
<td>10</td>
<td>10</td>
</tr>
<tr>
<td>Ni5Ce15</td>
<td>5</td>
<td>15</td>
</tr>
<tr>
<td>Ce20</td>
<td>0</td>
<td>20</td>
</tr>
</tbody>
</table>

The preparation procedure was as follows. First, magnesium ribbon (Sigma-Aldrich, St. Louis, MO, USA) was dissolved in methanol (Avantor Performance Materials, Gliwice, Poland) with rigorous stirring at 50 °C with the formation of Mg(OCH$_3$)$_2$. In order to dissolve each 1 g of magnesium, 43 mL of methanol was used. Then, toluene was added in a 1:1 volumetric ratio to the methanol. An aqueous solution of Ni(NO$_3$)$_2$ and/or Ce(NO$_3$)$_3$ (Sigma-Aldrich, St. Louis, MO, USA) was added dropwise to the solution of magnesium methoxide. The concentrations of the salts were varied to achieve the target concentrations of NiO and CeO$_2$ in the final composites. This resulted in the immediate hydrolysis of
Mg(OCH₃)₂ and the formation of a Mg(OH)₂ gel containing Ni²⁺ and Ce³⁺ ions. The color of the gels changed from light green (for the samples containing Ni²⁺ ions only) to orange (for the samples containing Ce³⁺ ions only). The gels were stirred for 2 h and then kept overnight with mild stirring in a glass bowl to let the solvent evaporate. The samples were then dried at 150 °C for 2 h and calcined at 500 °C at a heating rate of 1 °C/min.

2.2. Characterization of the Materials

The textural characteristics of the samples were determined using low-temperature nitrogen adsorption/desorption. The isotherms were recorded at 77 K using an ASAP-2400 (Micromeritics, Norcross, GA, USA) instrument. The total pore volume (V_pore) was defined by the point of maximum adsorption. The specific surface area (A_{BET}) was calculated using the Brunauer–Emmett–Teller method.

Diffuse reflectance UV–vis spectra were recorded between 190 and 800 nm using a Varian Cary 300 UV/VIS Bio UV–vis spectrometer (Agilent Technologies, Inc., Santa Clara, CA, USA) with a Labsphere DRA-CA-3300 integrating sphere (Labsphere, North Sutton, NH, USA) and using the Spectralon® standard as reference. The samples were characterized in a naturally hydrated state under atmospheric conditions.

The UV–vis spectra were transformed into the Kubelka–Munk function (Equation (2)) [46]:

\[ F(R_\infty) = \frac{(1 - R_\infty)^2}{2R_\infty} = \frac{\alpha}{s} \]  

(2)

where R_∞ is the reflectance of infinite sample, \(\alpha\) is the absorption, and \(s\) is the scattering. The optical band gap of the samples, which characterized the \(E_g\) value, was estimated from the diffuse reflectance spectra using the expression proposed by Tauc, Davis, and Mott (Equation (3)) [47]:

\[ [F(R) \cdot h\nu]^n = A(h\nu - E_g) \]  

(3)

where \(h\) is Planck’s constant, \(\nu\) is the frequency of vibration, \(h\nu\) represents the energy per photon, function \(F(R) \sim \alpha\) is the absorption coefficient, \(E_g\) is the band gap, and \(A\) is a constant. Note that \(n = 1/2\) and \(2\) represent indirect and direct allowed transitions, respectively.

Raman spectra were registered using a Horiba Jobin Yvon LabRAM HR UV-VIS-NIR Evolution Raman spectrometer (Horiba, Tokyo, Japan) equipped with an Olympus BX41 microscope (Olympus Corp., Tokyo, Japan) and an argon laser with a wavelength of 514.5 nm. To prevent overheating of the sample, the laser power in the focal spot with a diameter of \(\sim 2\) µm was less than 0.8 mW.

2.3. Testing the Catalytic Activity

The CO oxidation reaction was used to characterize the activity of the samples. The prompt thermal aging (PTA) regime was used [48,49]. A specimen of the material (300 mg; a fraction of 0.25–0.5 mm in size) was loaded into a flow-through quartz reactor. A reaction gas mixture containing CO (0.15 vol %), oxygen (14.0 vol %), and nitrogen (balance) was fed into the reactor. The temperature was raised from 50 to 320 °C (runs 1 and 2), 600 °C (runs 3 and 4), 800 °C (runs 5–6), 900 °C (runs 7–8), 1000 °C (runs 9–10), and 500 °C (run 11) at a heating rate of 10 °C/min. The total flow rate was 334 mL/min. The reaction mixture at the outlet of the reactor was analyzed using a ULTRAMAT 6 gas analyzer (Siemens, Munich, Germany). The CO concentration was measured every two seconds. The temperature of 50% conversion of CO (T_{50}) was used as a criterion to compare the samples.
The apparent activation energy of CO oxidation was determined using the same setup. In this case, the loading of the sample was 100 mg. The conversion of CO was measured in an isothermal mode in a range of temperatures of 190–310 °C with a step of 10 °C. The normalized reaction rate \( W \) was calculated according to Equation (4):

\[
W = \left( \frac{u}{m_{\text{Ni}}} \right) \cdot X_{\text{CO}}
\]

where \( u \) is the molar flow rate of CO, \( m_{\text{Ni}} \) is the weight of Ni contained in the sample, and \( X_{\text{CO}} \) is the CO conversion value. To calculate the apparent activation energy, the Arrhenius equation was used. The CO conversion values within the range from 1% to 10% only were used for the calculation.

3. Results and Discussion

3.1. Characterization of the Composites

As was reported previously [42], magnesium oxide prepared via the sol-gel method possesses a specific surface area above 200 m\(^2\)/g. The texture characteristics of the binary and ternary samples under study are shown in Table 2. Among these composites, the largest specific surface area was observed for the Ni15Ce5 sample. Then, an increase in the concentration of cerium oxide decreased the \( A_{\text{BET}} \) value, and the lowest value of 79 m\(^2\)/g corresponded to the Ce20 sample. The pore volume contrarily increased with an increase in the CeO\(_2\) concentration and reached the maximum value of 1.13 cm\(^3\)/g in the case of the ternary Ni5Ce15 composite.

<table>
<thead>
<tr>
<th>Sample</th>
<th>( A_{\text{BET}}, \text{m}^2/\text{g} )</th>
<th>( V_{\text{pore}}, \text{cm}^3/\text{g} )</th>
<th>( \text{MgO} )</th>
<th>( \text{CeO}_2 )</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>( a, \text{Å} )</td>
<td>( D, \text{nm} )</td>
<td>( a, \text{Å} )</td>
<td>( D, \text{nm} )</td>
</tr>
<tr>
<td>Ni20</td>
<td>118</td>
<td>0.52</td>
<td>4.212</td>
<td>7.9</td>
</tr>
<tr>
<td>Ni15Ce5</td>
<td>166</td>
<td>0.62</td>
<td>4.215</td>
<td>9.4</td>
</tr>
<tr>
<td>Ni10Ce10</td>
<td>124</td>
<td>0.71</td>
<td>4.216</td>
<td>10.9</td>
</tr>
<tr>
<td>Ni5Ce15</td>
<td>114</td>
<td>1.13</td>
<td>4.215</td>
<td>10.6</td>
</tr>
<tr>
<td>Ce20</td>
<td>79</td>
<td>0.57</td>
<td>4.214</td>
<td>16.8</td>
</tr>
</tbody>
</table>

The characterization of the oxide ternary Ni-Ce-Mg-O samples via XRD analysis (Figure 1) revealed the presence of two phases of cubic-fluorite-structure CeO\(_2\) (JCPDS #34-0394) and rocksalt MgO (JCPDS #78-0430) only. The reflections attributed to CeO\(_2\) appeared at a CeO\(_2\) concentration as low as 5 wt%. This was notable because, no peaks of the second phase were observed even at a higher M\(_x\)O\(_y\) content of 15 wt% in the XRD patterns reported previously for other sol-gel-prepared M-Mg-O systems (where M is Cu, Fe, Ni, or Co) [42]. The absence of the reflections assigned to Ni species indicated the high dispersion of these species and the partial substitution of magnesium and cerium ions within the MgO and CeO\(_2\) lattices with Ni\(^{2+}\) ions. As can be seen in Table 2, the crystallite size estimated for the MgO phase decreased from 17 to 8 nm with an increase in the fraction of NiO, which agreed well with the nitrogen adsorption data. The crystallite size of the CeO\(_2\) phase ranged from 5.3 to 3.2 nm. The lattice parameter of NiO was lower than that of MgO. Therefore, the substitution with Ni\(^{2+}\) ions and the formation of Ni\(_x\)Mg\(_{1-x}\)O solid solution should have shifted the reflections toward higher angles [50]. However, in the present case, no such shift was observed, whereas an opposite effect of the expansion of the lattice parameter appeared (Table 2) [51]. The same effect could be accounted for by an increase in the CeO\(_2\) lattice parameter with a decrease in particle size (Table 2) [52,53]. The amount of Ni\(^{2+}\) in the CeO\(_2\) lattice appeared to be quite low. Slightly lower values of the lattice parameter observed for the Ce20 sample could be attributed to the shrinkage of the CeO\(_2\) unit cell due to the difference in its lattice constant (0.541 nm) with respect to that of MgO (0.412 nm) [54].
Figure 1. XRD patterns for the studied samples.

Figure 2a shows the diffuse reflection UV–vis spectra of the pure MgO support and the Ce20, Ni5Ce15, Ni10Ce10, Ni15Ce5, and Ni20 samples. The absorption spectrum in the region of 190–220 nm corresponded to low-coordinated oxygen ions on the surface of magnesium oxide (6.6 eV for O^{2−}_{5C} ions with a coordination number of 5 and 5.6 eV for O^{2−}_{4C} ions with a coordination number of 4) with a characteristic particle size of about 10 nm [55].

The charge-transfer band of Ni_{n}Mg_{1−n}O at 245 nm was resolved for two samples with a maximum NiO concentration only: Ni20 and Ni15Ce5. For the other three-component samples of this series, it was masked by an intense absorption band of CeO_{2} with a maximum of ~300 nm. The charge-transfer bands of low-coordinate ions of Ce^{3+} at ~265 nm (O^{2−} → Ce^{3+}), the charge-transfer band at ~280 nm (O^{2−} → Ce^{4+}), and also the interband transition band at ~325–350 nm (O_{2p} → Ce_{4f}) made a contribution to this band [56]. At the same time, the most interesting part of the absorption spectrum at 400–600 nm corresponded to oxygen vacancies in non-stoichiometric cerium oxide CeO_{2−x} overlapped with a d-d transition ^{3}A_{2g} → ^{3}T_{1g}(F) in Ni^{2+} at 407 nm [56,57], which made the analysis of the absorption spectra of the three-component Ni-Ce-Mg-O samples rather complex. Nevertheless, the preliminary examination of the two-component Ni20 and Ce20 samples within the series under study should allow analyzing the three-component samples.

According to the literature data, the absorption spectrum of NiO as well as the spectrum of Ni_{n}Mg_{1−n}O in a wide range of concentrations (x) is determined by the Ni^{2+}...
ions located in the octahedral surround. It consists of an intense charge-transfer band \( O^2- (2p) \rightarrow Ni^{2+}(3d) \) at 240–270 nm and unevenly widened bands at 407, 671, and 741 nm, which correspond to the d-d transitions \( ^3A_{2g} \rightarrow ^3T_{1g}(F) \), \( ^3A_{2g} \rightarrow ^1E_g \), and \( ^3A_{2g} \rightarrow ^3T_{1g}(F) \), respectively [58]. This was also observed in the case of the Ni20 sample (Figure 3a). It should be noted that the dependence of the absorption value in the region of the \( ^3A_{2g} \rightarrow ^3T_{1g}(F) \) Ni\(^{2+}\) transition at 740 nm on the nickel content of the samples (Figure 2b) was linear. Slightly smaller F(R) values for the CeO\(_2\)-containing samples corresponded to the entry of a part of Ni\(^{2+}\) into the crystal lattice of CeO\(_2\). This explains the deviation of the real composition of samples from the calculated one.

![Figure 3](image_url)

**Figure 3.** (a) Diffuse reflectance UV–vis spectra of the samples: MgO (1); Ni20 (2). (b) Dependences of \( (F(R) \cdot E)^2 \) on the photon energy \( E \) characterizing the band-gap \( E_g \) values for direct allowed transitions for the Ni20 sample. Dash line shows how the \( E_g \) value was determined.

The charge-transfer band defined the energy-gap width \( (E_g) \) for the Ni\(_x\)Mg\(_{1-x}\)O particles. The value of \( E_g \) depended on the concentration \( x \) of Ni\(^{2+}\) ions in Ni\(_x\)Mg\(_{1-x}\)O. It was experimentally determined that \( E_g \) for direct allowed transition lies in an interval ranging from 3.8 eV for NiO particles of \( \sim 6 \) nm in size to 7.8 eV for pure MgO [59]. Taking into account the narrow range of particle sizes determined via XRD (Figure 1b), it can be supposed that the dimensional effects connected to an increase in the energy-gap width with the decrease in the particle size of Ni\(_x\)Mg\(_{1-x}\)O particles were less significant than those related to the formation of solid solutions [60,61].

Based on the stoichiometry for the homogeneous Ni\(_{0.2}\)Mg\(_{0.8}\)O\(_2\) solid solution (20 wt% of NiO), the band-gap width should have been \( E_g \sim 4.68 \) eV. The experimental value \( E_g \sim 4.22 \) eV corresponded to particles with a nickel content as in Ni\(_{0.62}\)Mg\(_{0.38}\)O. This value was close to that obtained earlier for the similar Ni-Mg-O oxide system (15 wt% NiO) synthesized using the sol-gel method at pH = 1 [40]. When comparing UV–vis data with XRD results, it should be taken into account that in the case of a wide distribution of particles for Ni content, the \( E_g \) value will be determined by a portion of the particles with a concentration of Ni above average, integrally contributing enough to the absorption spectrum to form a band-gap edge.

The diffusion reflection spectrum of the Ce20 sample after subtraction of the spectrum of the MgO support is presented in Figure 4a. As can be seen, it was represented by a wide band with a maximum of \( \sim 300 \) nm. Such charge-transfer bands for low-coordinated Ce\(^{3+}\) ions as \( O^2- \rightarrow Ce^{3+} \) and \( O^2- \rightarrow Ce^{4+} \) at \( \sim 265 \) and \( \sim 280 \) nm, respectively, as well as an interband transition band \( (O2p \rightarrow Ce4f) \) at \( \sim 325 \) nm contributed to this band [56].
The valence zone of stoichiometric CeO$_2$ mainly consists of O2p states and the conduction zone consists of Ce5d states. There is a narrow band of Ce4f between them. The gap between the conduction Ce5d band and the valence band O2p is $\sim 6$ eV [62–64]. Experimental data for the band-gap width O2p $\rightarrow$ Ce4f obtained by Bremsstrahlung isochromat spectroscopy (BIS) and photoluminescence provided values of 2.6–3.39 eV [62,65]. With the formation of oxygen vacancies in the non-stoichiometric CeO$_{2-x}$, local distortion of the lattice occurs, and the Ce4f band is split into two bands: Ce4f$_{\text{filled}}$ and Ce4f$_{\text{empty}}$. Existing experimental data indicate that the Ce4f$_{\text{filled}}$ band lies in the range $\sim 2.0–2.4$ eV above the valence band [66]. The reduction limit for non-stoichiometric cerium oxide is Ce$_2$O$_3$ where all cerium ions are in the oxidation state of Ce$^{3+}$. The main difference in the zone structure of Ce$_2$O$_3$ with respect to CeO$_{2-x}$ is the fusion of the Ce4f$_{\text{empty}}$ and Ce5d bands in the conduction zone [64]. The experimental value of the energy of the absorption edge for Ce$_2$O$_3$ (Ce4f$_{\text{filled}}$ $\rightarrow$ Ce4f/Ce5d), which can be taken as an estimate of the band-gap width, is $2.4$ eV [67]. Thus, it can be expected that the formation of a non-stoichiometric cerium oxide CeO$_{2-x}$ should be accompanied by the appearance of a band at $\sim 410–420$ nm, which, upon further reduction of CeO$_{2-x}$, will form a band gap $E_g \sim 2.0–2.4$ eV.

Based on the literature, the following set of Gaussian functions converted from the energy scale to the wavelength scale in nm was used to approximate the absorption band for the Ce20 sample (Figure 4): with a maximum at 270 nm (3.66 eV and HWHM = 1.43 eV) and with a maximum at 320 nm (3.87 eV and HWHM = 0.85 eV). In addition, one more peak at 214 nm (5.79 eV and HWHM = 1.0 eV) corresponding to the edge of the band O2p $\rightarrow$ Ce5d in CeO$_2$ was used. Further, this information was considered as auxiliary since only a part of the absorption spectrum at $\lambda > 240$ nm was attractive to compare the spectra of the NiO-containing samples. This set of bands was suitable as is for approximating all the samples under consideration except for the interband transition band (O2p $\rightarrow$ Ce4f) at 320 nm, for which a small adjustment was required in the case of the NiO-containing samples.

The dependences of $(F(R) \cdot E)^2$ and $(F(R) \cdot E)^{0.5}$ on the photon energy $E$ presented in Figure 4b determined the band-gap width $E_g$ for the direct (3.4 eV) and indirect (2.85 eV) allowed transitions in the Ce20 sample. Taking into account the known experimental data for the values of the band-gap width with respect to the particle sizes of CeO$_2$ [68–70], the particle sizes for the samples under study were estimated to be $\sim 4–4.5$ nm. This value was close to the value of $\sim 5$ nm obtained from the XRD analysis (Table 2).

As can be seen for the Ce20 sample (Figure 4a), in the wavelength region above 400 nm, there were no features indicating the formation of the Ce$^{3+}$ states and the associated O-vacations that could result from the incorporation of Mg$^{2+}$ into the CeO$_2$ lattice [71,72].
According to the Raman spectra, which will be discussed in detail below, there were also no signs of the formation of oxygen vacancies. Therefore, it can be concluded that no noticeable incorporation of Mg$^{2+}$ into the CeO$_2$ lattice occurred with the synthesis method used. However, it cannot be ruled out that when synthesizing the three-component samples (Ni15Ce5, Ni10Ce10, and Ni5Ce15), part of Mg$^{2+}$ ions could enter the CeO$_2$ lattice along with Ni$^{2+}$ ions. In this study, Ce20 was used as a reference sample to investigate the incorporation of Ni$^{2+}$ into the CeO$_2$ lattice for the cases of the Ni-Ce-Mg-O samples.

To simulate the formation of a non-stoichiometric oxide CeO$_{2-x}$, the as-prepared Ce20 sample was reduced with CO at 500 °C (Figure 5a). The absorption bands that appeared at 420 nm (2.94 eV and HWHM ≈ 0.65 eV) were associated with the formation of Ce$^{3+}$ species and the associated O-vacations, and the band at 555 nm (2.23 eV and HWHM ≈ 1.35 eV) could be assigned to the surface oxygen vacancies [57]. The latter completely disappeared after oxidation at room temperature. To restore the original stoichiometry and reduce the intensity of the band at 420 nm, higher oxidation temperatures of 250–350 °C are required [73]. Similar behavior of the sample was observed during its reduction in hydrogen. With an increase in the reduction temperature and an increase in the proportion of Ce$^{3+}$ states and the associated O-vacations, the band at 420 nm formed a band gap with the width values of ≈2.0–2.4 eV [67]. According to the literature data, the surface of CeO$_2$ only is reducible at 500 °C, while a higher temperature of 650 °C is required for the reduction of bulk Ce$^{4+}$ species [74,75]. Note that the data presented for the Ce20 sample reduced with CO at 500 °C (Figure 5a) were limited due to the need to avoid texture change when characterizing the initial state of the sample [57,74]. Reduction of the sample under mild conditions only slightly decreased the band-gap values for direct (3.36 eV) and indirect (2.8 eV) allowed transitions (Figure 5b).

**Figure 5.** (a) Diffuse reflectance UV–vis spectra of the Ce20 sample reduced with CO at 500 °C after subtraction of the spectrum of the MgO support. (b) Dependences of $(F(R) \cdot E)^2$ (red) and $(F(R) \cdot E)^{0.5}$ (blue) on the photon energy E characterizing the band-gap $(E_g)$ values for direct and indirect allowed transitions for the Ce20 sample reduced with CO. Dash lines show how the $E_g$ values were determined.

In order to analyze the Ni-Ce-Mg-O samples and study the incorporation of Ni$^{2+}$ into the CeO$_2$ lattice, Figure 6a shows the diffuse reflectance UV–vis spectra of Ni5Ce15, Ni10Ce10, and Ni15Ce5 after subtraction of the spectrum of the Ni20 sample with the intensity corrected according to real absorption values in the field of transition $^3A_{2g} \rightarrow ^3T_{1g}(F)$ Ni$^{2+}$ at 740 nm. The spectrum of the pristine Ce20 sample is given for comparison. As the content of NiO in the samples increased, there was a monotonous and close to linear increase in the absorption intensity of the band at 400 nm (Figure 6b), which can be attributed to both the Ce$^{3+}$ states and the associated O-vacancies and the d-d transitions of Ni$^{2+}$, which occupied the lattice position of Ce$^{4+}$ and the associated O-vacancies. In this case, a noticeable decrease in the band-gap $(E_g)$ values for direct allowed transitions was...
observed for the sample with the maximum NiO content only. This indicated that in our case, the maximum concentration of Ni$^{2+}$ in the CeO$_2$ lattice was less than 2–3% when the effects associated with the rearrangement of the CeO$_2$ structure appeared as a change in the band-gap width \[76,77\]. Dependence of the Urbach energy ($E_U$) on the NiO content in the samples (Figure 6b) also indicated an increase in the amount of defects.

Figure 6. (a) Diffuse reflectance UV–vis spectra of the MgO support; the Ce20 sample; and the Ni5Ce15, Ni10Ce10, and Ni15Ce5 samples after subtraction of the spectrum of the Ni20 sample with corrected intensity. (b) Dependences of the absorption intensity of the d-d Ni$^{2+}$ band at 400 nm (■), the band-gap ($E_g$) values (■) for direct allowed transitions, and the Urbach energy ($E_U$) (■) on the NiO content in the samples.

Figure 7 presents a detailed curve fitting for the diffuse reflection UV–vis spectra of the Ni5Ce15 and Ni15Ce5 samples after subtraction of the spectrum of the Ni20 sample with corrected intensity. The corresponding dependences ($F(R) \cdot E$)$^2$ and ($F(R) \cdot E$)$^{0.5}$ on the photon energy $E$ characterizing the band-gap ($E_g$) values for direct and indirect allowed transitions are shown as well.

Compared to the original curve fitting set for the as-prepared Ce20 sample (Figure 4a), minor changes in the position of the interband transition band O$_2p$ → Ce4f from 320 to 313–314 nm (2.95 eV and HWHM $\sim$ 0.85 eV) were required. This can be interpreted as the appearance of a new band at 306 nm (4.06 eV and HWHM $\sim$ 0.65 eV) when the intensity of the initial band at 320 nm (3.87 eV and HWHM $\sim$ 0.85 eV) decreased. This finding correlated well with the electronic density of states (DOSs) analysis carried out by Miran and Jaf \[78\], which showed that Ni integration led to some shrinkage of Ce4f states.

The absorption band with a peak at 400 nm observed for the Ni5Ce15 sample (Figure 7a), the intensity of which increased linearly with an increase in the NiO content (Figure 6b), was well described by one broad Gaussian band at 3.1 eV with an HWHM of $\sim$1.35 eV. Only for the Ni15Ce5 sample (Figure 7c) did less intense absorption bands related to the formation of Ce$^{3+}$ states and the associated O-vacancies and the surface vacancies characteristic of the reduced state of the Ce20 sample (Figure 5a) appear.

Changes in the position of the interband transition band for the Ni5Ce15 sample appeared as a small increase in the $E_g$ values from 3.4 to 3.44 eV for direct allowed transitions (Figure 7b). As can be seen, the absorption band at 400 nm for both the Ni5Ce15 (Figure 7b) and Ni15Ce5 (Figure 7d) samples formed a band gap of $\sim$1.54–1.55 eV for indirect allowed transitions, thus indicating the formation of localized states of Ni$^{2+}$ and the associated O-vacancies between the valence and conduction bands.
In our case, when concentrations of Ni$^{2+}$ cations embedded in the CeO$_2$ lattice were small, only a detailed analysis of the structure of the absorption bands via curve fitting of the diffuse reflection UV–vis spectra allowed us to characterize the changes in structure that...
occurred. Approaches based only on the analysis of the band gap, which are commonly applied to such systems, are not informative enough.

As seen in Figure 8, the Raman spectra of the Ni-Ce-Mg-O samples were determined by the intensive lines of CeO$_2$ and NiO. Against their background, no lines corresponding to the first and second order of Raman modes for MgO nanocrystals were visible [81,82], which was due to the very low intensity of scattering in MgO.

![Figure 8](image_url)

**Figure 8.** Experimental Raman spectra of the three-component Ni-Ce-Mg-O composites and the reference two-component Ce20 and Ni20 samples.

Curve fitting of the experimental Raman spectrum of the Ce20 sample (Figure 9a) was dominated by a first-order active F$_{2g}$ band at 463 cm$^{-1}$ (HWHM $\sim$ 30 cm$^{-1}$) of CeO$_2$, which was slightly shifted with regard to single crystal ceria (465 cm$^{-1}$) [83,84]. This band was associated with the stretching of the Ce–O bond for eight-fold coordinated Ce$^{4+}$ in the cubic fluorite structure. Second-order weak Raman bands at 250, 402, and 595 cm$^{-1}$ have been assigned to 2TA and 2TO modes, and the broad band at 1178 cm$^{-1}$ has been ascribed to 2LO modes [83,85]. It should be noted that no evident shift in the position of the F$_{2g}$ mode with respect to pristine ceria and no features in the spectrum at 570–620 cm$^{-1}$ associated with the formation of oxygen vacancies [71] were seen. This indicated that in our case, the concentration of Mg$^{2+}$ impurity ions in CeO$_2$ was small, which was consistent with the data from UV–vis spectroscopy (Figure 4a).

Raman scattering of the 20%NiO-MgO samples (Figure 9b) for frequency shifts larger than 150 cm$^{-1}$ allowed us to identify the main contributions from one-phonon 1P(TO) modes at 440 cm$^{-1}$, 1P(LO) modes at 560 cm$^{-1}$, and two-phonon 2LO modes at 1100 cm$^{-1}$ for the Ni$_x$Mg$_{1-x}$O solid solutions. The two-phonon bands (2P(TO) modes at 740 cm$^{-1}$ and TO + LO modes at 925 cm$^{-1}$) were less pronounced [86,87]. In comparison with the Raman spectrum of pure NiO [88,89], the intensity of the two-magnon (2M) band at $\sim$1400 cm$^{-1}$ decreased rapidly for crystallites size smaller than 100 nm. At the same time, in the presence of the substitutional disorder as in Ni$_x$Mg$_{1-x}$O [86], the two-magnon (2M) band experienced significant broadening and disappeared completely at $x \leq 0.6$ even for polycrystalline samples.

For the presented Raman spectra of the Ni20 sample (Figure 9b), one can observe a decrease in the two-phonon band intensity relative to the one-phonon contribution and a more pronounced 1P(TO) band compared to pure NiO. This provided a possibility to estimate x in the range of $\sim$0.5–0.6 since two-phonon bands disappeared at $x \leq 0.4$ [86]. This estimate corresponded to the x value of $\sim$0.6 calculated for this sample using the band-gap ($E_g$) value obtained from the UV–vis diffuse reflection spectra.
Figure 9. Experimental Raman spectra of the powder samples of MgO-based composites: (a) Ce20; (b) Ni20; (c) Ni5Ce15; (d) Ni10Ce10; (e) Ni15Ce5. Dependence of relative intensities of lines $I_{555}/F_{2g}$ and $I_{615}/F_{2g}$ on the content of NiO in the samples is shown in (f).

For curve fitting of the experimental Raman spectrum of the Ni-Ce-Mg-O samples (Figure 9c–e), a set of Gaussian functions was used. Note that the ratios of their intensities corresponded to those for the Ce20 (Figure 9a) and Ni20 (Figure 9b) samples. This made it possible to approximate all experimental spectra with the exception of the $F_{2g}$ band of CeO$_2$. As can be seen in Figure 9c–e, with an increase in the NiO content from 5 to 10%, there was only a slight displacement and expansion of the $F_{2g}$ band of CeO$_2$ to 463 cm$^{-1}$ (HWHM $\sim$ 30 cm$^{-1}$). At the same time, the relative intensities of the emerging new lines at 555–615 cm$^{-1}$, which indicated the formation of oxygen vacancies and reduced Ce$^{3+}$ ions in CeO$_2$ [83], showed a linear ascending dependence on the total NiO content in the samples (Figure 9f). The additional band at $\sim$230 cm$^{-1}$ (HWHM $\sim$ 30 cm$^{-1}$) had a significantly smaller half-width with respect to the 2TA vibration mode of the lattice at $\sim$250 cm$^{-1}$ (HWHM $\sim$ 100 cm$^{-1}$) observed for the Ce20 sample (Figure 9a). This band is usually assigned to O–Ce longitudinal stretching of atoms in the outermost layers of CeO$_2$. 

3.2. Catalytic Performance of the Composites in the Oxidation of CO

The catalytic performance of the prepared MgO-based composites was tested in the model reaction of CO oxidation. Each sample was tested in 11 heating/cooling runs in a prompt thermal aging (PTA) regime. Such a regime considered an increase in the final temperature of each second run (as described in Section 2.3). As an example, Figure 10a shows the light-off curves of CO conversion recorded in the PTA regime for the Ni10Ce10 sample. The first curve corresponds to the initial contact of the sample with the reaction mixture. Positions of the second and third curves reflect the initial steady-state catalytic performance. Then, after aging at 600 °C, a slight reactivation effect can be observed when the curves are shifted toward lower temperatures. Further runs correspond to the aging at 800, 900, and 1000 °C. As can be seen, in these cases, the curves are shifted toward higher temperatures, thus indicating a gradual decrease in the activity of the sample.
(111) surfaces [83], which has an increased intensity for smaller ceria nanoparticles. The very close character of the Raman spectra related to NiO species in this series (Figure 9b–e) allowed us to conclude that the estimate made above for \( x \sim 0.5–0.6 \) in the composition of \( \text{Ni}_x \text{Mg}_{1-x} \text{O} \) solution solutions could be applied to all samples.

For the Ni15Ce5 sample (Figure 9e), the F\(_{2g}\) band of CeO\(_2\) was further shifted to 460 cm\(^{-1}\) and widened to HWHM \(\sim 50\) cm\(^{-1}\), which agreed with the UV-vis spectroscopy data. Along with this, in this case, the intensity of the F\(_{2g}\) band was much lower than one might expect. This may have been due to a change in the morphology of the particles for Ni15Ce5 with respect to other samples of this series accompanied, for example, by the decoration of CeO\(_2\) particles with \( \text{Ni}_x \text{Mg}_{1-x} \text{O} \) particles. Another possible reason may have been a relatively low concentration of CeO\(_2\). However, the characteristic value of the shift along with the widening of the F\(_{2g}\) band in comparison with the known experimental Raman data for Ce\(_{1-x}\)Ni\(_x\)O \([76,77]\) made it possible to estimate the concentration of Ni in CeO\(_2\) for Ni15Ce5 as \( x \sim 1–2\% \).

Thus, the characterization of the prepared composites via optical methods well confirmed the partial substitution of Ce\(^{4+}\) ions by Ni\(^{2+}\), although their concentration was about 2–3% according to various estimates. With the growth in the NiO/CeO\(_2\) ratio, the concentration of Ni\(^{2+}\) ions that substituted cerium ions grew as well. At the same time, the entry of Mg\(^{2+}\) ions into the CeO\(_2\) lattice was not found. Unlike the XRD analysis, both the UV-vis and Raman spectroscopy methods confirmed the formation of \( \text{Ni}_x \text{Mg}_{1-x} \text{O} \) solid solutions. The data presented suggest a similar interaction between NiO and MgO in all samples under study. To conclude this section, the advantages of using the detailed fitting analysis of the UV-vis and Raman spectroscopy for studying the properties of multicomponent systems have been shown. The described approaches can be used in the future for in-depth analysis of the complex interactions in various nanocrystalline oxides.

### 3.2. Catalytic Performance of the Composites in the Oxidation of CO

The catalytic performance of the prepared MgO-based composites was tested in the model reaction of CO oxidation. Each sample was tested in 11 heating/cooling runs in a prompt thermal aging (PTA) regime. Such a regime considered an increase in the final temperature of each second run (as described in Section 2.3). As an example, Figure 10a shows the light-off curves of CO conversion recorded in the PTA regime for the Ni10Ce10 sample. The first curve corresponds to the initial contact of the sample with the reaction mixture. Positions of the second and third curves reflect the initial steady-state catalytic performance. Then, after aging at 600 °C, a slight reactivation effect can be observed when the curves are shifted toward lower temperatures. Further runs correspond to the aging at 800, 900, and 1000 °C. As can be seen, in these cases, the curves are shifted toward higher temperatures, thus indicating a gradual decrease in the activity of the sample.

The comparison of the T\(_{50}\) values in 11 consecutive runs for all the samples is shown in Figure 10b. It can be clearly seen that the Ce20 sample exhibited the worst catalytic performance due to the absence of nickel species, which are responsible for catalytic activity. Oppositely, the Ni15Ce5 sample demonstrated the best performance. In general, the samples can be ranked by the T\(_{50}\) criterion as follows: Ni15Ce5 > Ni10Ce10 > Ni20 > Ni5Ce15 >> Ce20. A markedly higher specific surface area of the Ni15Ce5 sample (Table 2) may have contributed to a higher available surface area of the active component and, as a consequence, to better catalytic behavior. After aging at 600 °C, the reactivation effect mentioned above was observed for all nickel-containing samples (Figure 10b,c). Despite the noticeable deactivation of all the composite samples during the further increase in the aging temperature to 800, 900, and 1000 °C, their thermal stabilities were different. Thus, the Ni10Ce10 and Ni5Ce15 samples seemed to be more stable if compared with the Ni20 and Ni15Ce5 composites. After aging at 1000 °C, the activity row of the samples can be written as follows: Ni10Ce10 \(\approx\) Ni15Ce5 > Ni5Ce15 \(\approx\) Ni20 >> Ce20.
The comparison of the T50 values in 11 consecutive runs for all the samples is shown in Figure 10b. Comparison of T50 values at various aging temperatures for the samples under study (c). Comparison of thermal stability of the Ni15Ce5 sample with the Ni15 sample reported previously [40] (d).

Figure 10d compares the activity of the Ni15Ce5 sample with a sample containing 15 wt% NiO and 85 wt% MgO (labeled as Ni15), which was studied previously [40,90]. At the initial stages of the catalytic test, both samples behaved similarly. It is important to note that the Ni15 sample possessed a comparable value of specific surface area (154 m²/g). However, the ternary Ni15Ce5 composite sample exhibited substantially greater thermal stability after aging in the reaction mixture at temperatures of 800 °C and above. This indicated the important role of the presence of ceria within the composition of the composite material and its interaction with active species of nickel.

Table 3 summarizes the catalytic data for the Ni-containing samples under study. Despite the fact that the Ni15Ce5 sample showed the highest values of CO conversion, the specific activity decreased with an increase in the nickel content. However, the apparent activation energy values for all samples were close to each other and lay within the experimental error (Table 3, Figure 11). This may indicate a similar mechanism of the reaction. It should be emphasized, however, that the observed values of 68–73 kJ/mol were somewhat higher than the values of 50 ± 5 kJ/mol known for supported NiO-containing catalysts and catalysts based on bulk NiO [36,38,91,92]. These values refer mainly to the temperature range below 200 °C. It is likely that this was a consequence of the formation of Ni₃Mg₁₋ₓO solid solutions.

The observed effects can be explained as follows. It is natural to assume that the properties of Ni-Ce-Mg-O are defined by the complex interactions between all components of the composite system. As shown by optical spectroscopy techniques, in the studied samples, partial substitution of Ce⁴⁺ ions with Ni²⁺ ions accompanied by the formation of vacancies occurred. However, most of the nickel ions were still included in NiₓMg₁₋ₓO solid solutions of various compositions. Therefore, part of the Ni²⁺ ions appeared in the...
volume of the support, which negatively affected the catalytic performance. On the other hand, this made it possible to maintain a high dispersion of nickel species.

Table 3. Data on CO oxidation over Ni-containing composites in the catalytic experiments performed under stationary isothermal conditions (m_{cat} = 100 mg).

<table>
<thead>
<tr>
<th>Sample</th>
<th>Conversion, %</th>
<th>Specific Reaction Rate, μmol/(s·gNi)</th>
<th>Activation Energy, kJ/mol</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>250 °C</td>
<td>270 °C</td>
<td>250 °C</td>
</tr>
<tr>
<td>Ni20</td>
<td>2.8</td>
<td>5.2</td>
<td>0.62</td>
</tr>
<tr>
<td>Ni15Ce5</td>
<td>4.7</td>
<td>8.3</td>
<td>1.36</td>
</tr>
<tr>
<td>Ni10Ce10</td>
<td>4.5</td>
<td>7.8</td>
<td>1.94</td>
</tr>
<tr>
<td>Ni5Ce15</td>
<td>2.6</td>
<td>4.8</td>
<td>2.22</td>
</tr>
</tbody>
</table>

Figure 11. Arrhenius plots for the Ni-containing composites obtained in the temperature range of 190–310 °C. The experimental points of CO conversion above 1% and below 15% only were selected for the linear fitting.

Recent studies of CO oxidation on NiO/CeO$_2$ catalysts showed the decisive role of the interactions near the NiO-CeO$_2$ interface in this reaction [36,38]. Thus, oxygen activated on the CeO$_2$ surface reacted with CO adsorbed on NiO, resulting in an intensification of the process. In the ternary Ni- Ce-Mg-O composites, CeO$_2$ was present in the form of nanosized crystallites. Such a small size of CeO$_2$ crystallites provided enhanced contact between them and individual surface Ni$^{2+}$ ions and small NiO particles, which increased their activity in CO oxidation. Another factor that could influence the catalytic performance is dispersion of NiO species. Thus, an increase in Ni dispersion was observed in the systems of similar compositions upon addition of CeO$_2$ [21–23].

4. Conclusions

In this work, three-component Ni-Ce-Mg-O composites were synthesized via the sol-gel method using nitrates as precursors. The textural characteristics of the samples were found to depend strongly on their composition. The highest surface area was observed in the case of the Ni15Ce5 sample, which contained 15 wt% NiO and 5 wt% CeO$_2$. An increase in the concentration of CeO$_2$ was accompanied by the enlargement of the CeO$_2$ and MgO crystallites. Detailed investigation of the samples by means of diffuse reflectance UV–vis and Raman spectroscopies demonstrated that Mg$^{2+}$ ions did not substitute Ce ions in the CeO$_2$ lattice. However, in the Ni-containing samples, approximately 2–3% of the Ce$^{4+}$ ions were substituted by Ni$^{2+}$, thus resulting in the formation of vacancies. The interaction of NiO with MgO seemed to be much stronger in relation to its interaction with CeO$_2$. The formation of Ni$_x$Mg$_{1-x}$O solid solutions was confirmed using UV–vis spectroscopy, and the average composition was estimated as Ni$_{0.60}$Mg$_{0.40}$O. The catalytic performance of the prepared composites was examined in CO oxidation. At temperatures of 320 °C and below,
the Ni20 and Ni5Ce15 samples demonstrated the worst activity among all nickel-containing samples. In terms of the $T_50$ values, the initial activity row of the samples was as follows: Ni15Ce5 > Ni10Ce10 > Ni20 > Ni5Ce15 >> Ce20. Thermal aging of the samples at 1000 °C revealed the thermal stability of the samples, and the activity row was changed as follows: Ni10Ce10 ≈ Ni15Ce5 > Ni5Ce15 ≈ Ni20 >> Ce20. At the same time, very close values of the apparent activation energy were determined for all nickel-containing samples, which indicated a similar mechanism of CO oxidation over the composites under study. Thereby, the observed differences in the CO oxidation performance of the Ni-containing composites can be attributed to the partial substitution of Ce3+ ions by Ni2+ ions, the formation of Ni$_x$Mg$_{1-x}$O solid solutions, and the interactions on the Ni$_x$Mg$_{1-x}$O–CeO$_2$ interface.

To sum up, this work expanded the available toolkit of synthetic methods by applying the simplified sol-gel approach to the preparation of Ni-Ce-Mg-O oxide systems. In addition, the effectiveness of comparatively simple optical spectroscopy techniques such as UV-vis and Raman spectroscopy for studying complex interactions in the NiO-CeO$_2$-MgO system was also highlighted. As for the further development of this work, Ni/CoO$_2$-MgO systems are known to be effective catalysts for many industrially important processes. The performance of the Ni-Co-Mg-O systems described here will be evaluated in the dry reforming of methane, and the effect of CeO$_2$ addition on the resistance of the catalysts toward carbon formation will be studied.

**Author Contributions:** Conceptualization, A.A.V.; methodology, G.B.V. and V.O.S.; investigation, G.B.V. and V.O.S.; writing—original draft preparation, G.B.V. and V.O.S.; writing—review and editing, A.A.V.; visualization, A.A.V.; supervision, A.A.V. All authors have read and agreed to the published version of the manuscript.

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