Pure and Al-doped ZnO obtained by the modified Pechini method applied in ethanolic transesterification of cottonseed oil

(ZnO puro e dopado com Al obtidos pelo método Pechini-modificado aplicado na transesterificação etanólica do óleo de algodão)

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Abstract

Pure zinc oxide (ZnO) and 5% Al-doped ZnO (ZNAL) were synthesized using the modified Pechini method and characterized by X-ray diffraction (XRD), X-ray fluorescence (XRF), Raman spectroscopy, infrared spectroscopy and UV-visible spectroscopy. XRF confirmed the theoretical stoichiometry, while XRD and Raman spectroscopy indicated that Al\(^{3+}\) was incorporated into the ZnO wurtzite lattice with no secondary phases, leading to a decrease in the band gap value and to a meaningful increase of the Lewis basic sites. Pure and doped ZnO were used as catalysts in the ethylic transesterification of cottonseed oil using a factorial design to determine the best synthesis conditions. Oil conversion into biodiesel was evaluated by viscosity measurements and \(^1\)H NMR spectroscopy. The results analyzed by factorial design indicated that the catalyst type and temperature were the determinant factors in the conversion indices. The highest basicity of the ZNAL lead to a significant increase of the catalytic potential, reaching a reduction of the oil viscosity next to 71% at 130 °C and greater than 85% at 200 °C.

Keywords: zinc oxide, doping, heterogeneous catalysis, biodiesel.

Resumo

Óxido de zinco puro (ZnO) e dopado com 5% de Al (ZNAL) foram sintetizados pelo método Pechini modificado e caracterizados por difração de raios X (DRX), fluorescência de raios X (FRX), espectroscopia Raman, espectroscopia no infravermelho e espectroscopia UV-visível. A análise por FRX confirmou a relação estequiométrica teórica, enquanto DRX e espectroscopia Raman indicaram que o Al\(^{3+}\) foi incorporado no retículo de wurtzita do ZnO sem fases secundárias, conduzindo a uma diminuição no valor do band gap e a um aumento significativo dos síitos básicos de Lewis. ZnO puro e dopado foram utilizados como catalisadores na transesterificação etanólica do óleo de algodão utilizando um planejamento fatorial para determinar as melhores condições de síntese. Os índices de conversão do óleo em biodiesel foram avaliados através de medidas da viscosidade e por espectroscopia RMN\(^1\)H. Os resultados analisados pelo planejamento fatorial indicaram que o tipo de catalisador e a temperatura foram os fatores determinantes nos índices de conversão. O aumento do caráter básico do catalisador ZNAL conduziu a um aumento significativo do potencial catalítico, atingindo uma redução da viscosidade do óleo próxima de 71% a 130 °C e superior a 85% a 200 °C.

Palavras-chave: óxido de zinco, dopagem, catálise heterogênea, biodiesel.

INTRODUCTION

The increased concern over the environmental impact and increasing prices of petrol products, coupled with the fact that fossil fuels are non-renewable, has prompted a continuing search for fuels that are environmentally responsible energy sources, such as biodiesel [1-3]. The most commonly employed production route for this biofuel is the transesterification of vegetable oils and animal fats using catalysts [4, 5]. Due to its reversibility, this type of reaction is strongly influenced by several factors including temperature, the size and polarity of the alcohol chain, the acidity of the oil, and the type and concentration of catalyst [6, 7]. Different solids have been used as catalysts for biodiesel synthesis and the performance of these materials is directly linked to the nature of the acidic or basic sites found in their structures. The catalytic activity of solids having Lewis basic sites is dependent upon the interaction of the alcohol used as the transesterification agent with the surface of the catalyst. Alternatively, for Lewis acid catalysts, the carbonyl groups of the fatty materials interact with the surface, leading to the formation of carbocations [6, 7].

Zinc oxide is an amphoterous oxide and has been used as a catalyst for transesterification reactions in its pure form, mixed with other oxides or salts, or as a support or matrix for other substances [8-13]. Nascimento et al. [8] compared the efficiency of ZnO with CaO, CaTiO\(_3\) and TiO\(_2\), which were all obtained by the polymeric precursor method. The
use of CaO led to a high conversion of corn oil into methyl esters at 70 °C but resulted in high levels of Ca leaching into the biodiesel. Compared with the other oxides, a greater conversion was attained with ZnO at 150 °C. Few papers report the effect of cation addition into the ZnO lattice (doping) on its catalytic activity for biodiesel synthesis. Recently, the doping of Cu into the ZnO lattice with a high conversion of waste cooking oil into methyl esters at 60 °C was reported [9].

For the ZnO-Al$_2$O$_3$ system, different papers report the catalytic activity of the ZnAl$_2$O$_4$ spinel [10, 11]. For instance, Thirunavukkarasu et al. [10] deposited Zn on gamma-Al$_2$O$_3$ and on Fe$_2$O$_3$ and, after heat treatment at 600 °C, observed the formation of ZnAl$_2$O$_4$ and ZnFe$_2$O$_4$, respectively. Both spinels are active in the methanol transesterification of vegetable oils. In other research employing ZnAl$_2$O$_4$ as a catalyst, Pugnet et al. [11] evaluated different reaction parameters in the methanolic transesterification of rapeseed oil, with a conversion of approximately 80% at 200 °C. Veiga et al. [12] obtained ZnO-Al$_2$O$_3$ catalysts with different Zn:Al ratios for the methanolic transesterification of soybean oil. Al-doped ZnO, synthesized with 25% Al, attained approximately 30% conversion at 130 °C, much lower than conversions attained with 50% Al (composed of ZnO and ZnAl$_2$O$_4$), which reached more than 80%. Another interesting paper [13] reports the use of Zn-doped Al$_2$O$_3$ (20 mol% ZnO) as a catalyst in the ethanolic and methanolic transesterification of soybean oil. The yield for the methanolic reaction at 100 °C was double that of the ethanolic reaction, which required a reaction temperature of 180 °C to reach a 78% yield. The present work is aimed at obtaining biodiesel by the ethylic transesterification of vegetable oil using ZnO and Al-doped ZnO catalysts synthesized using the modified Pechini method [14]. Despite its poor yield compared to methanol, the ethanolic route to avoid the use of petroleum-derived precursors was utilized, obtaining a truly green fuel. Moreover, Brazil is a significant producer of ethanol.

**EXPERIMENTAL**

**Catalyst synthesis and characterization**

Pure ZnO was synthesized, via the modified Pechini method, by dissolution of Zn(NO$_3$)$_2$·6H$_2$O (Aldrich) in a stirred, aqueous citric acid (Cargill) solution at 70 °C using a citric acid:metal molar ratio of 3:1. After complete dissolution, ethylene glycol (Aldrich) was added to the solution, with a citric acid:ethylene glycol mass ratio of 6:4, and the temperature was increased to 90 °C. The polymeric resin was heated at a rate of 2 °C.min$^{-1}$ and semicarbonized at 300 °C for 2 h. After heat treatment, the polymeric precursor was ground, passed through a 200 mesh (74 μm) sieve, and heat treated in an air atmosphere at 600, 700 and 800 °C for 4 h with a heating rate of 5 °C.min$^{-1}$. The same procedure was adopted for 5% Al-doped ZnO (ZNAL), but using an aqueous solution comprising Al(NO$_3$)$_3$·6H$_2$O (Aldrich) and Zn(NO$_3$)$_2$·6H$_2$O (Aldrich).

X-ray diffraction (XRD) was performed on a Shimadzu XRD 6000 instrument using K$_\alpha$(Cu) radiation (λ= 1.5406 Å), a step scan of 0.02°.s$^{-1}$ and a 2θ range of 3 to 80°. Patterns were compared to ICDD standards. X-ray fluorescence (XRF) measurements were performed on a Shimadzu XRF-1800 spectrometer using internal standards for elemental quantification. UV-visible spectra were obtained with a Shimadzu UV-2550 spectrophotometer in reflectance mode between 190 and 900 nm. Absorbance spectra were used to calculate the energy band gap according to the Wood and Tauc method [15]. Infrared spectra were obtained with a Shimadzu IRPrestige-21 spectrophotometer between 4000 and 400 cm$^{-1}$, using KBr pellets. Raman spectra were recorded on a Renishaw InVia Raman spectrometer using a 100 mW Ar laser (λ= 514 nm). Evaluation of the acidic or basic character of ZnO and ZNAL was performed by addition of 0.5 g of each catalyst into 10 mL of HCl or KOH solution (0.005 mol.L$^{-1}$) and measurement of the pH change at 6 min intervals for 1 h using a benchtop pH meter (Digimed DM22).

**Transesterification reaction**

A two level factorial design was proposed for the ethanolic transesterification of the cottonseed oil [16, 17]. The parameters of temperature, alcohol:oil ratio and catalyst type were evaluated over a 4 h reaction time. The low and high levels for each variable are presented in Table I. Experimental data were analyzed by the Statistica 8.0 software (Statistica for Windows 8.0, Statsoft, USA).

Transesterification reactions were performed in a Parr reactor, mod. 4842, with a maximum working pressure of 200 bar. Considering that this reaction proceeds via a basic catalytic mechanism, 50 g of commercial cottonseed oil were added to ethanol (Moderna) containing 3 wt% catalyst (with respect to the oil) in alcohol:oil ratios of 6:1 and 12:1. After the reaction, the suspension was centrifuged to separate the catalyst, and water was added to the liquid phase with a decantation balloon to facilitate separation of the glycerin, esters and acylglycerides. Samples were dried at 90 °C under vacuum. In order to evaluate the acidic catalytic mechanism, the reaction with the hardest conditions (200 °C and alcohol:oil ratio of 12:1) was repeated by addition of ethanol to the cottonseed oil containing 3% catalyst.

**Table I - Factorial design used in the ethanolic transesterification of cottonseed oil using temperature, alcohol:oil ratio and catalyst type as variables.**

| Variable                  | Level | Level |
|---------------------------|-------|-------|
| Temperature (°C)          |       | 130   |
| Alcohol:oil ratio         | 6:1   | 200   |
| Catalyst type             | ZnO   | ZNAL  |

[Tabela I - Planejamento fatorial para a transesterificação etanólica do óleo de algodão utilizando temperatura, razão álcool/óleo e tipo de catalisador como variáveis.]

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(ZnO or ZNAL). The products were isolated as described above. Evaluation of the viscosity of the reaction products was performed using a Julabo V18 viscometer with a Cannon Fenske glass capillary immersed in a water bath at 40 °C. Characterization of the esters and acylglycerides was performed by 1H NMR using a Varian Gemini 300BB spectrometer operating at a frequency of 200 MHz.

RESULTS AND DISCUSSION

Catalyst characterization

The XRD patterns (Fig. 1) indicate the formation of zinc oxide in ZnO and ZNAL. Increasing the heat treatment temperature resulted in increased crystallinity of the powder. All peaks were well defined, indicating a high degree of long-range order was attained and confirming that the modified-Pechini method was successful for this synthesis. All peaks were indexed according to a wurtzite hexagonal unit cell, space group P63mc, C6v (ICDD standard 89-1397). After doping, a small dislocation of the peaks to larger angles, as well as peak broadening, was observed when compared with pure ZnO (Fig. 1c), which indicates a higher degree of long-range disorder is present. The lattice parameters for ZnO were calculated, and no meaningful variation of the unit cell size was observed, as a = 3.25 Å and c = 5.20 Å for all samples. According to the XRF results, the quantitative analysis of Zn reported 99.7% purity for the pure ZnO form. For the ZNAL, quantitative analysis reported 94.6 mol% ZnO and 4.9 mol% Al2O3, suggesting satisfactory

![Figure 1: X-ray diffraction patterns of ZnO (a) and ZNAL (b) after firing at 600, 700 and 800 °C. In (c), enlargement of the XRD patterns between 30° and 38° for ZnO and ZNAL heat treated at 800 °C.](image)

![Figure 2: Raman spectra of ZnO (a) and ZNAL (b), heat treated at 600, 700 and 800 °C.](image)
Stoichiometric control was attained.

Zinc oxide with a wurtzite structure belongs to the space group $C_{4}^{6v}$, which has two formula units per unit cell, all atoms occupying 2b sites with $C_{3v}$ symmetry and the following active modes in the Raman spectrum: $A_{1} + 2B_{1} + E_{1} + 2E_{2}$. $A_{1}$ and $E_{1}$ are polar phonon modes, divided into transverse ($A_{1T}$ and $E_{1T}$) and longitudinal ($A_{1L}$ and $E_{1L}$) polar optical modes, while $E_{2}$ consists of two high- and low-frequency phonon modes ($E_{2h}$ and $E_{2l}$) [18-20]. The Raman spectra of ZnO and ZNAL are presented in Fig. 2. The bands at 101, 339, 388 and 437 cm$^{-1}$ were assigned to the wurtzite structure. The bands above 800 cm$^{-1}$ were due to the second order Raman modes [19]. An additional band was observed at 221 cm$^{-1}$, which is related to the presence of oxygen vacancies in the lattice [19-21]. These results confirm that both samples have hexagonal wurtzite structures with high degrees of short-range order, as additional bands are not observed after doping. Moreover, no meaningful changes in the spectra were observed, indicating that no symmetry breaking occurred.

The infrared spectra from 4000 to 400 cm$^{-1}$ are presented in Fig. 3. The bands assigned to tetrahedrally coordinated Zn-O or Al-O were observed between 400 and 500 cm$^{-1}$, and the bands assigned to the stretching of tetrahedrally coordinated Me-O bonds were observed between 750 and 880 cm$^{-1}$ [13, 22]. A well-defined band at 750 cm$^{-1}$ was only obtained after heat treatment at 800 °C, due to the high degree of short-range order. Bands at 3500 and 1635 cm$^{-1}$ were also observed and assigned to the OH stretching and bending, respectively, of water molecules adsorbed on the material surface [23]. Small bands were observed between 1270 and 1490 cm$^{-1}$ and were assigned to the carboxyl stretching frequencies, $\nu$(C=O) and $\nu$(C-O), respectively. These bands indicate that ester complexes were still present in the material after calcinations at 600 and 700 °C but were eliminated at 800 °C [24]. These results suggest an optimal heat treatment temperature of 800 °C for both samples.

The band gap values of the materials heat treated at 800 °C were calculated from the UV-vis absorption spectra (Fig. 4), according to the methodology described by [15], and the observed values were 3.2 eV for pure ZnO and 2.9 eV for ZNAL. Similar results were obtained by Bel-Hadj-Tahar and Mohamed [25], who observed a decrease in the band gap value when ZnO was doped with aluminum, and Mote et al. [26], who observed a decrease in the band gap value when ZnO was doped with yttrium. Bai et al. [23] evaluated the influence of Al-doping into ZnO by first-principle calculations. According to their results, band gap narrows because the conduction band undergoes a greater shift towards the low-energy region than the valence band.

Replacing a Zn$^{2+}$ ion for Al$^{3+}$ implies the need for charge compensation. According to [23, 27], this compensation is achieved through the introduction of electrons in the doped-ZnO conduction band, favored by the band gap narrowing, increasing concentration of free electrons. As
a consequence, an increase in conductivity and in sensing properties of the doped oxide have been observed [23, 27]. The charge compensation mechanism is presented in Equation A, written considering the Kröger-Vink notation [28]. This behavior may enhance the Lewis base character of the material, as electron donor defects (Al\textsuperscript{2+}Zn\textsuperscript{2+}) are formed.

\[
\text{Al}_2\text{O}_3 + \text{ZnO} \rightarrow 2\text{Al}^{2+} + 2e^- + 3\text{O}^6_-(A)
\]

It is well known that ZnO is an amphoteric material. In order to understand the prevailing acid-base character and the influence of the Al-doping, adsorption of OH\textsuperscript{−} and H\textsubscript{3}O\textsuperscript{+} species on the powder surfaces was evaluated for samples calcined at 800 °C (Fig. 5). ZnO and ZNAL samples exhibited higher basic character as indicated by the change in the pH value due to the OH\textsuperscript{−}/H\textsubscript{3}O\textsuperscript{+} adsorption/elimination. Al-doping increased the number of basic sites by more than 4 orders of magnitude relative to pure ZnO. In regard to the acidic character of the catalysts, Al-doping increased the kinetics of OH\textsuperscript{−} adsorption onto the surface.

**Ethyllic transesterification reaction**

Transesterification reactions occur in three consecutive steps through which triacylglycerides (TAG) are converted into fatty acid esters (R’COORx) and glycerol (GL). However, partial reactions can occur producing monoacylglycerides (MAG) and/or diacylglycerides (DAG) thus decreasing the overall ester content at the end of the reaction (Fig. 6)[29, 30].

Ethyllic transesterification may proceed via an acid or a basic mechanism. In the heterogeneous catalysis by the basic mechanism, the formation of an alkoxide (active species) takes place by reaction of an alcohol adsorbed on a Lewis base. In the second step, the carbonyl group of the acylglycerides gradually undergoes nucleophilic attack by the alkoxide, forming a tetrahedral intermediate. From a rearrangement of these intermediates, the formation of an ester and other alkoxy takes place which, after deprotonation of the conjugate acid, regenerates the Lewis base. This mechanism occurs on virtually all tri-, di- and monooacylglycerides are turned into alkyl esters and glycerol. In the acid mechanism the TAG adsors on the catalyst surface, with a greater acid/base interaction between the metal present in the catalyst structure and the pair of electrons from the oxygen of the carbonyl group causing an increase in the positive charge density on the carbon bonded to this oxygen, forming a carbocation which, in turn, undergoes a nucleophilic attack from the pair of electrons of the alcohol hydroxyl group, forming a tetrahedral intermediate. After elimination of the water molecule from this intermediate, a fatty ester and a DAG are formed and the catalyst is regenerated after desorption of the reaction products from the surface. MAG and glicerol are formed from similar processes [7].

As ZnO is an amphoterous solid, the type of reaction mechanism was evaluated by favoring the alcohol adsorption on the catalyst surface (basic mechanism) by mixing ethanol with the catalyst previously to the cottonseed oil addition; or favoring the TAG adsorption (acid mechanism) by adding...
the catalyst into the oil with subsequent addition of alcohol. Reactions were done at the same conditions – 200 °C for 4 h, ratio alcohol:oil of 12:1. Results are presented in Table II, which confirm that ZnO catalyzes the transesterification reaction by both mechanisms – the acid one or the basic one, with a higher efficiency for the basic mechanism, leading to a higher viscosity reduction. After Al-doping, only the basic mechanism is observed, and no conversion is attained when the reaction is performed under acid catalysis conditions. This result is in agreement with the highest basic character of ZnO and ZNAL, demonstrated in Fig. 5. For this reason, all other reactions were done with a previous mixture of the alcohol with the catalyst.

The experimental parameters of temperature, alcohol:oil ratio and catalyst type were used as variables in the factorial design to evaluate the conversions of the catalyzed transesterification reactions, as shown in Fig. 7. The significance of each variable and the possible interactions among them were evaluated by variance analysis (ANOVA) based on the Pareto graph (Fig. 7), which was plotted based on the values of the reductions in viscosities. The response surface (Fig. 8) was also used to obtain information about the experiments. According to the viscosity results (Table II), it was observed that the cottonseed oil has a kinematic viscosity of 38.4 cSt, which was reduced by 56.8 to 85.7% after the catalytic test, indicating that esters, MAG and/or DAG were formed. According to the Pareto graph (Fig. 7), the variance analysis and the response surface, the temperature produced the most significant positive effect, with the highest conversions obtained at higher temperature (200 °C). The response surface (Fig. 8) indicates that a 21.35% increase in conversion was attained when the temperature was increased from the inferior level (130 °C) to the superior level (200 °C). For the catalyst type, higher conversions were achieved when ZNAL (superior level) was used at all temperatures (Table II). A 15.6% increase in conversion was attained when changing from the inferior level (pure ZnO) to the superior level (ZNAL), as indicated by the response surface. These results confirm the importance of the highest

Table II - Kinematic viscosities after catalytic reaction: BC - basic catalysis, and AC - acid catalysis.

| Catalyst   | Temperature (°C) | Ethanol:oil ratio | Kinematic viscosity at 40 °C (cSt) | Viscosity reduction (%) |
|------------|-----------------|------------------|-----------------------------------|------------------------|
| ZnO (BC)   | 130             | 6:1              | 16.6                              | 56.8                   |
|            | 130             | 12:1             | 14.5                              | 85.5                   |
|            | 200             | 6:1              | 8.7                               | 77.3                   |
|            | 200             | 12:1             | 8.2                               | 78.6                   |
| ZnO (AC)   | 200             | 12:1             | 13.7                              | 64.3                   |
| ZNAL (BC)  | 130             | 6:1              | 11.2                              | 70.8                   |
|            | 130             | 12:1             | 9.2                               | 76.0                   |
|            | 200             | 6:1              | 7.8                               | 79.7                   |
|            | 200             | 12:1             | 5.5                               | 85.7                   |
| ZNAL (AC)  | 200             | 12:1             | 38.4                              | 0.0                    |

Figure 7: Evaluation of the ethanolic transesterification reaction of cotton seed oil via Pareto graphic of the 2³ factorial design - the vertical line defines the 95% of confidence range.

Figure 8: Response surface, considering level changes of temperature, alcohol:oil ratio, and catalyst type.

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Lewis basic character promoted by Al-doping into the ZnO lattice, indicated by Equation A and Fig. 5. Santos et al. [31] also observed a positive effect of doping in the synthesis of biodiesel. In this work, ZnAl2O4 spinel was doped with Eu3+, leading to a meaningful increase in the soybean oil conversion into biodiesel.

Each of the species involved in the transesterification reaction have characteristic signals in the 1H NMR spectra that can be utilized to demonstrate the conversion of TAG into esters. For the 1H NMR spectrum, the doublet of doublets (dd) with an integration of 4 H appearing between δ 4.0 and 4.2 ppm is assigned to the α-CH2 group of TAG and is converted into a quartet (q) with an integration of 2 H appearing between δ 4.0 and 4.2 ppm, which corresponds to the –CH2CH3 ethyl ester group [32, 33]. This difference in the spectrum profile can be observed for the cottonseed oil and the reaction products obtained using ZNAL at 200 °C (Fig. 9).

CONCLUSIONS

The Pechini method was efficient for the synthesis of zinc oxide in its pure form or doped with 5% Al3+ with good stoichiometric control. Characterization indicated that Al3+ ions were incorporated into the ZnO crystalline lattice. The addition of 5% Al3+ into zinc oxide maximizes its basic sites, driving the transesterification of cottonseed oil to a basic mechanism. The study of the experimental design indicated that adding 5% Al3+ into the ZnO lattice significantly improved its application as a catalyst for the ethyl transesterification of cottonseed oil while temperature was also a determinant factor in the conversion rates of these reactions. The reduction in kinematic viscosity values associated with the characterization of the products by 1H NMR confirmed the success of the transesterification reaction with triacylglycerides (TAG) conversion into esters.

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