MgO-based adsorbents for CO₂ adsorption: Influence of structural and textural properties on the CO₂ adsorption performance

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ABSTRACT

A series of MgO-based adsorbents were prepared through solution–combustion synthesis and ball-milling process. The prepared MgO-based powders were characterized using X-ray diffraction, scanning electron microscopy, N₂ physisorption measurements, and employed as potential adsorbents for CO₂ adsorption. The influence of structural and textural properties of these adsorbents over the CO₂ adsorption behaviour was also investigated. The results showed that MgO-based products prepared by solution–combustion and ball-milling processes, were highly porous, fluffy, nanocrystalline structures in nature, which are unique physico-chemical properties that significantly contribute to enhance their CO₂ adsorption. It was found that MgO synthesized by solution combustion process, using a molar ratio of urea to magnesium nitrate (2:1), and treated by ball-milling during 2.5 hr (MgO-BM2.5h), exhibited the maximum CO₂ adsorption capacity of 1.611 mmol/g at 25°C and 1 atm, mainly via chemisorption. The CO₂ adsorption behaviour on the MgO-based adsorbents was correlated to their improved specific surface area, total pore volume, pore size distribution and crystallinity. The reusability of synthesized MgO-BM2.5h was confirmed by five consecutive CO₂ adsorption–desorption times, without any significant loss of performance, that supports the potential of MgO-based adsorbent. The results confirmed that the special features of MgO prepared by solution–combustion and treated by ball-milling during 2.5 hr are favorable to be used as effective MgO-based adsorbent in post-combustion CO₂ capture technologies.

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Introduction

It is well documented that carbon dioxide (CO₂) is the most significant greenhouse gas that mainly contributes to global warming. In the past five decades, atmospheric CO₂ emissions have been increased due to the accelerated industrial growth and technological development, resulting in a potential threat for the environment and ecosystems (Shafeeyan et al., 2015). The main anthropogenic sources of CO₂ emissions are attributed to the excessive fossil materials combustion such as carbon, oil and natural gas in the energy production and in many industrial processes (Bhagiyalakshmi et al., 2011).
this context, inevitably due to the fact that fossil fuels will be the predominantly worldwide power supply, it is necessary to take urgent measures to reduce the high CO2 concentrations in the atmosphere at great scale (Ello et al., 2013). As a consequence, currently, there are different CO2 capture technologies for these purposes, including adsorption process, membrane separation, cryogenic methods, and so on. Among CO2 capture technologies, the CO2 adsorption using porous solid adsorbents has been receiving much attention because of their advantages, such as low regeneration energy requirements, no liquid waste, low cost, both pressure swing adsorption and temperature swing processes are widely used (Harrison, 2004). Indeed, also has attracted the researchers’ attention due to that can to be designed and studied a great variety of potential CO2 adsorbents of varied nature, cheap, and renewable for use in large-scale post combustion technologies. In this scenario, many solid porous materials have been widely recognized and used as potential adsorbents for CO2 adsorption, including activated carbons (Houshmand et al., 2012), zeolites (Siriwardane et al., 2005), amine compounds (Saiwana et al., 2014), organometallic networks (Liu et al., 2012), metal oxides (Busca and Lorenzelli, 1982), and hydroxalicates (Radoslaw et al., 2015), among others.

Amid these above adsorbents listed, metal oxides are promising options for suitable CO2 adsorbents and have a great potential in the future due to their easy accessibility and favorable thermodynamic properties (Kumar et al., 2015). However, the selection criteria of potential metal oxides for a large-scale application, entail their CO2 capture capacity, adsorption rate, thermal stability, regeneration heat, availability, cost, textural and structural properties (Kumar and Saxena, 2014). Specifically, the alkaline-earth metal oxide (MgO) is widely used in many technological applications as: catalysis, toxic waste remediation, additive in refractory, optically transparent ceramic windows, adsorbent for many pollutants, paint and superconductor products, among others (Jin et al., 2012). MgO is a white hygroscopic solid mineral that occurs naturally as periclase, is a source of magnesium, is a non-toxic and environmentally friendly material with higher surface reactivity and adsorption capacity, and is a suitable adsorbent for anions and cations from aqueous solution due to its favorable electrostatic attractive mechanism (Crittenden et al., 2005). In recent studies, MgO has been widely explored for CO2 capture, and has been recognized as one of the most promising adsorbents for CO2 adsorption together with CaO, because these metallic oxides combine with CO2 present in the flue gas post-combustion and form thermodynamically stable carbonates in a reversible reaction; heating the carbonates regenerates the respective oxides and thus liberates almost pure stream of CO2 that it can be directly transported to a storage site, and subsequently be used technologically (Kumar and Saxena, 2014). It has been reported that MgO can adsorb CO2 below 200°C by carbonation, and can be regenerated by calcination at relatively low temperature compared with conventionally-used CaO subjected to cyclic CO2 capture from flue gases, which makes it very desirable because it is widely accepted that there is a large scope for cost reduction and energy efficiency improvements in CO2 capture systems (Wang et al., 2011). In general, many metal oxides can qualify for CO2 capture application, however, one of the key factors in gas-solid reaction is their textural, morphological and structural properties that are essential forenhancing their CO2 adsorption. Thus, synthesis of MgO with improved specific surface area, nanocrystalline size, large total pore volume and narrow size pore distribution is of great interest as CO2 adsorbent materials. The direct relation of these MgO physico-chemical properties with the CO2 capture behaviors still not completely resolved and needs to be investigated.

On the other hand, extensive synthesis methods have been followed to obtain MgO products with porous structures and multiple morphologies, including the precipitation method (Janet et al., 2007), hydrothermal synthesis (Zhao et al., 2011), and sol-gel method (Kim et al., 2005). It is well known that the employed synthesis method and the chemical precursors to prepare powders, have a large influence on the textural and structural characteristics of materials such as porosity, specific surface area, particle size, crystallinity, between others. Particular desirable properties of compounds in their use as adsorbents are mainly extensive specific surface area that means many active sites for contaminant adsorption and a higher porous nature, which plays an important role on CO2 capture. Therefore, it is evident from the above that there is a need for a continuous improvement and development of new and potential adsorbents for gaseous, aqueous, and non-aqueous streams. Currently, the novel solution–combustion synthesis is widely used for preparing different oxide powders with high purity, high yield, large specific surface areas, and good quality mesoporous structures at short times compared to other conventional synthesis methods. Furthermore, this method is safe, simple and instantaneous for the facile fabrication of nanopowders, indeed, the solution–combustion synthesis requires simple equipment with energy saving (Tonolo et al., 2010). For this method are required high temperatures for the exothermic redox reactions and achieve the decomposition of the metal salt and the organic fuel for products formation. Therefore, extensive studies have been conducted to investigate solution–combustion synthesis of many metallic oxide powders using urea as the most commonly used chemical fuel, and some other fuels such as: starch, glycine and sorbitol (Bhatta et al., 2015; Bai et al., 2011; Devaraja et al., 2014; Granados-Correa and Bonifacio-Martínez, 2014; Mantliaka et al., 2014).

On the other hand, recent advances in nanotechnology have made much interest in preparing metallic oxide nanocompounds. The mechanical ball-milling is an extensive method to obtain nanostructured materials; this mechanical ball-milling treatment allows to activate dry solids and, especially increase its specific surface area as well as improve properties (Janusz et al., 2010). Nanopowders show substantially enhanced adsorption characteristics superior to those of the conventionally prepared powders (Liang et al., 2001). It is important to note that the gas adsorption efficiency can increase with a decrease in the MgO powder size and crystallites. In fact, the high gas adsorption efficiency of nanoparticles is also caused by their large specific surface area, and structural defects on their surface. Furthermore, it has been shown that metallic oxides doped with doping-metals such as Fe, Ni, Cd, Ce, Co among others, show significant changes in their properties with respect to original materials. In
terms of the basis of these approaches, doped metallic oxides are of research interest because of its diverse properties, which originate from its structural characteristics. In this context, MgO doped adsorbents with significant doping-metals such as Fe and Ni are here systematically studied as new and potential solid adsorbents with improved properties that can be used to separate CO₂ effectively from flue gases generated by fossil fuel combustion. Their novel properties and/or improved performance can provide an additional contribution and effort to develop an efficient CO₂ capture method.

The aim of this work was to prepare a series of MgO-based adsorbents by solution–combustion and ball-milling processes, the introduction of Fe and Ni dopants into the MgO crystal structure has also been considered as efficient approach to improve its CO₂ capture. Then, the obtained MgO materials were used comparatively as potential adsorbents for CO₂ adsorption. In this context, their CO₂ adsorption behaviors were interpreted in light of a detailed textural, morphological, and structural characterization. Indeed, the CO₂ desorption and regeneration ability of the best resulted CO₂ adsorbent was also studied.

1. Experimental

1.1. Materials

Magnesium nitrate hexahydrate (Mg(NO₃)₂·6H₂O, 99.0% purity) and urea (CH₄N₂O, 99.4% purity) were purchased from Sigma-Aldrich, St. Louis, MO. All used gases in this study were supplied by Infra México, México, with the purity) elements of analytical grade were used and obtained from Sigma-Aldrich, St. Louis, MO. All used gases in this study were supplied by Infra México, México, with the following specifications: carbon dioxide extra dry (CO₂, 99.9% purity), nitrogen (N₂, 99.9% purity), and helium (He, 99.9% purity).

1.2. Synthesis of MgO-based adsorbents

MgO-based adsorbents were synthesized via solution–combustion method and treated by ball-milling process. In order to establish the best experimental conditions for MgO formation with large specific surface area and porosity, by solution–combustion synthesis, different molar ratios of chemical precursor’s urea as chemical fuel and magnesium nitrate as oxidizer (1:1 to 2.5:1) at 800°C during 5 min were proven. For this purpose, determined quantities of urea and magnesium nitrate were placed into 50 mL porcelain crucible and dissolved with 1 mL of distilled water. The obtained homogeneous mixture was heated at 260°C during 20 min with help of electric grill for water evaporation, obtaining a consistent material with gel appearance. Then, the resulting gel was calcined for 5 min into muffle furnace at 800°C. The water effect in the MgO combustion synthesis also was studied, for this purpose, to resulting material different amounts of distilled water (0.5 to 2.5 mL) were added, prior to calcination at 800°C for 5 min. On the other hand, the MgO sample obtained by solution–combustion was ball-milling by using a stainless ball mill Spex type with a ratio mass-steel balls of 6:1 into container at different time periods (2.5 to 10.0 hr). Indeed, the MgO–Fe and MgO–Ni nanocomposites were prepared also by ball-milling at different milling time periods from 2.5 to 10.0 hr by using a ratio percent weigh MgO to metal element of 98.2 wt.%.

1.3. MgO-based adsorbents characterization

Phase composition and crystalline structures of selected prepared MgO-based adsorbents were identified by X-ray diffraction (XRD) using a SIEMENS D-5000 diffractometer, Madison, WI, with Cu-Kα radiation (λ = 1.54060 Å). The crystalline MgO structures were identified according to the Joint Committee of the Powder Diffraction Standard (JCPDS) cards. The average crystallite sizes of these MgO-based compounds were determined by means of the Sherrer’s equation, applied to the major XRD peaks (Granados-Correa et al., 2016). The morphologies and the elemental chemical composition of the MgO samples were studied by scanning electron microscopy (SEM, JEOL, Tokyo, Japan) using a JEOL-JSM-5900LV instrument equipped with an energy dispersive X-ray analysis (EDX) detector using OXFORD microprobe. Brunauer–Emmett–Teller (BET) specific surface areas (S BET), mean pore diameters (d p), and total pore volumes (V T) of the prepared MgO nanomaterials were calculated by N₂ physisorption measurements. Nitrogen adsorption–desorption isotherms were measured at 77 K, and the pore size distributions were determined using the Barrett–Joyner–Halenda (BJH) method. All N₂ physisorption measurements were made on a Micromeritics BELSORP MAX Inc. Osaka, Japan volumetric adsorption analyzer. Prior of N₂ measurements, samples were degasified in a nitrogen stream at 473 K for 3 hr.

1.4. CO₂ adsorption experiments

CO₂ adsorption experiments were carried out on MgO-based adsorbents by using a stainless reactor Parr 4592 of 50 mL capacity coupled with a temperature-controlled system. Approximately 15 mg of each material was used during each experiment. For this purpose, the samples were loaded into the reactor and degassed under vacuum at 325°C by 30 min. The equilibrium experiments were carried out at temperatures of
25, 40, 60 and 80°C and at pressures of 1, 5, 10 and 15 atm for 30 min. The amount of millimoles of CO₂ adsorbed per gram of adsorbent (mmol/g), was obtained on the basis of the weight loss of CO₂ that is adsorbed in the nanocompounds by using a TA Instruments SDT Q600 thermo gravimetric analyzer (New Castle, DE) coupled to mass spectrum analyzer TA Instruments model LLC (New Castle, DE), heating the samples at a rate of 20°C/min up to 850°C in helium atmosphere at a flow rate of 100 mL/min.

2. Results and discussion

2.1. Synthesis

It was found the optimum conditions of MgO synthesis via solution-combustion process, by using a molar ratio (2:1) of chemical precursor’s urea to magnesium nitrate, and using 1.5 mL of distilled water prior to calcination at 800°C during 5 min. In this solution-combustion synthesis, a rapid chemical process based on the explosive decomposition of urea as fuel was carried out. This decomposition provides heat for calcination and produces high temperatures since fuel-to-oxidant ratio in an oxidant-fuel mixture has a significant effect on combustion synthesis (Granados-Correa et al., 2008). Moreover, theoretically, a stoichiometric redox mixture could produce maximum energy during combustion. Then, in this work, MgO was fabricated for better textural properties by using various redox mixtures as described in the Experimental section. As a consequence, a molar ratio of urea to magnesium nitrate (2:1) was established as the best MgO solution-combustion reaction. In order to find the yield for this reaction to obtain MgO, yielding product was determined as a function of the fuel-to-oxidant ratio according to Jain and Adiga (1981). The balanced equation for the combustion reaction in this study can be written as Eq. (1):

\[
3\text{[Mg(NO₃)₂ · 6H₂O]}_\text{s} + 5\text{CH₄N₂O}_\text{s} \rightarrow 3\text{MgO}_\text{s} + 8\text{N₂}_\text{g} + 5\text{CO}_\text{2g} + 28\text{H₂O}_\text{g}
\]

In solution-combustion reaction for MgO powder formation (Eq. (1)), there is a release of N₂, CO₂ and H₂O molecules in excess, generating porous MgO with homogeneous fine particles of spherical shape. This was clearly confirmed by SEM observations. It is worthwhile to mention that in this reaction, urea acts as a chemical fuel, and H₂O has a significant role for this porous structure formation of MgO powder, due to the exothermic reaction that is generated. According to previous investigations (Cai et al., 2009a, 2009b), urea was also used as precipitating agent. In agreement to Choong-Hwan and Ji-Yeon (2010) the principles of propellants and explosives for a redox reaction between a fuel and an oxidizer, the cation (carbon, hydrogen and oxygen) acts as fuel, and the metal nitrate anion (oxygen) acts as the oxidizer. Then, in this work, to know if the mixture is rich or poor in fuel, the oxidant-fuel mixture that is usually expressed in terms of elements stoichiometric coefficient (Φₚ), was calculated using Eqs. (2) and (3) (Jain and Adiga, 1981).

\[
\Phiₚ = \frac{\sum_{\text{Cox}}(C_{\text{ox}}) \times (v)}{-(1)\sum_{\text{Cred}}(C_{\text{red}}) \times (v)}
\]

where, C₀ₓ is the coefficient of oxidizing elements in specific formula; Cᵲₓ is the coefficient of oxidizing elements in specific formula; v is the valency. Φₓ = 1 corresponds to the stoichiometric ratio, Φₓ > 1 indicates a poor mixture in fuel and Φₓ < 1 is a rich fuel mixture. The results indicated that to obtain a stoichiometric mixing of the oxidizing-reducing species (Φₓ = 1), the molar ratio of fuel-oxidizer must be 5:3 as shown in Eq. (1). However, the combustion reaction of this work was fuel rich with a molar ratio urea to magnesium nitrate (2:1) with (Φₓ = 0.598), this fuel rich can benefit the MgO formation due to the fact that more energy exists during the combustion of powders, which allowed to obtain a high yield, 99.42%, from the reaction. Therefore, this optimum performance made the process feasible by solution-combustion synthesis to successfully obtain MgO powders with increased porosity and specific surface area. The as-prepared MgO nanopowders by solution-combustion in this study were compared with another reported MgO powders from urea as fuel, and these results are in agreement with Maliyekkal et al. (2010) and Rao and Sunandana (2008), highlighting in these cases that all MgO samples prepared by this fast method, present a higher porous nature, small crystalline size, and high purity.

2.2. Characterization

Fig. 1 shows the X-ray diffraction patterns for all MgO-based adsorbents studied here. These materials only show the characteristic peaks assigned to crystalline MgO, with periclase cubic structure according to the JCPDS 01-087-0651 file. It has been reported that the ball-milling process has an important impact on the crystallinity of materials, attributed to the high pressure and high temperatures generated locally during the milling process in the ceramic powders (Granados-Correa et al., 2016). Then, an important change in crystallinity of MgO-SC sample after ball-milling process was observed. In Fig. 1c and d, Fe and Ni element phases cannot be recognized by XRD patterns.
due to their lower weight ratios (98 wt.% MgO sample: 2 wt.% Fe or Ni respectively). The average crystallite sizes of all studied MgO-based adsorbents were found to be 40.73 nm for MgO-SC sample, 8.61 nm for MgO-BM2.5h, and 10.13 and 10.49 nm for MgO-Fe and MgO-Ni samples, respectively. The discrepancy of MgO-based adsorbent crystallinity is a result of the intense shocks of milling media (steel balls) on MgO particles or due to some disaggregation of MgO-SC particles (Singh et al., 2009).

It was observed that the specific surface area for MgO-BM2.5h sample with lower average crystallite size (8.61 nm) was increased, because the solid milling favors particle fragmentation. Therefore, results revealed that there is a clear change in structure and specific surface area when the MgO-SC sample is ball-milled, which can be expected that would increase its CO2 capture. The microstructural effect of ball-milling time over MgO ball milling sample, also was studied during 5, 7.5 and 10 hr respectively, and the results indicated that for a longer ball-milling time, a decrepitation occurred, which not caused a decrease of crystallite size, but generated ultratine powders. Therefore, a short milling time of 2.5 hr is the most suitable route for producing MgO nanocrystalline adsorbent with improved structural and textural characteristics for efficient CO2 adsorption. According to Valverde et al. (2014), a reduction of metalic oxide (CaO) crystallinity by ball-milling favors diffusion and serves to promote recarbonation in accordance with the CO2 capture subject to carbonation/calcination cycles at Ca-looping, where a mechanism for CaCO3 nucleation on the CaO surface in islands has been proposed (Kumar et al., 2015). In this study, the MgO behavior is associated to that of CaO behavior. Therefore, it is widely accepted that a gas–solid carbonation reaction occurs for the MgO-CO2 reaction. According to Kumar and Saxena (2014), MgO particles are surrounded by CO2 molecules and react to form MgCO3. However, MgCO3 forms an impervious layer around unreacted MgO particles and hampers the further diffusion of CO2 molecules, as can be seen in Fig. 2, for a schematic CO2 adsorption mechanism on mesoporous MgO adsorbent.

It is recognized that when CO2 molecules make contact with the metallic oxide surface as a CO2 adsorbent, they will be in contact with two surface types: the metal oxide surface and the metal carbonate surface; when CO2 molecules make contact with the metal oxide surface, carbonation directly occurs, which is thermodynamically stable at temperature and pressure ambient conditions, while for a metal carbonate surface happens by diffusion through this layer of carbonates. It is showed that the carbonation in this stage is mainly driven by solid state diffusion, which is of practical interest. Therefore, this MgO crystal structure role is necessary in order to lead the complete carbonate conversion of MgO. Thus, a correlation between the crystallinity changes of MgO-based adsorbents with their CO2 adsorption capacities is expected because the ball-milling process causes a not very orderly atom structure in the particles, which can enhance the CO2 adsorption capacity.

Fig. 3 shows the MgO-based adsorbent morphology. The MgO-SC sample (Fig. 3a) is characterized by spherical shape particles of soft surface texture with sizes ranging from 0.5 to 5 μm. EDX analysis results indicated that MgO-SC powders are composed of Mg (62.42 at.%), and O (37.57 at.%), showing that the desired material was obtained by using solution–combustion synthesis only. The MgO-BM2.5h sample (Fig. 3b), exhibited spongy and porous aggregates of irregular shapes with rough surfaces due to the fragmentation of the dust trapped during collision of balls against the container walls during ball-milling process. In Fig. 3c and d the morphological changes of MgO-Fe and MgO-Ni samples were exhibited; in these micrographs the particles are characterized by irregular shapes with slightly rounded edges and rough textures, also clearer fine particles were observed. EDX analysis showed a heterogeneous distribution of Fe (2.12 at.%) and Ni (0.92 at.%) in MgO samples, indicating that Fe and Ni elements are built in the crystal structure of MgO-Fe and MgO-Ni samples, respectively, with well-particle dispersion which can produce reactive powders. Indeed, the sizes of particle are considered in the nanometric range. Henceforth, a narrow particle size distribution generates more specific surface area, meaning more active sites for CO2 adsorption. In order to achieve high efficiency of CO2 capture in advanced technologies based in the CO2 capture by carbonation/calcination of metallic oxides for CO2 looping cycles, the systems can adopt a process configuration that consists of using two interconnected circulating fluidized bed reactors as a carbonator and as a calciner, which allows working with higher surface speeds of gas and higher circulations of the solid adsorbent (Abanades et al., 2005). For these purposes, surface properties of MgO-based adsorbents such as specific surface area, particle size, and porosity are very critical parameters that favor the carbonation process of metallic oxides, increasing the CO2 adsorption efficiency.

The N2-physisorption measurement results of prepared MgO-based adsorbents are listed in Table 1. It was observed that the MgO-BM2.5h sample considerably increased their BET specific surface area, and total pore volume in comparison.
with MgO-SC sample from 5.71 m²/g and 0.0289 cm³/g to 15.24 m²/g and 0.1347 cm³/g respectively, and the pore size diameter was increased from 12.12 to 21.52 nm. The obtained results revealed an aggregation of particles generated by the ball-milling process, which led to the moderate pore formation. In accordance with Bhatta et al. (2015), MgO mesoporous with large BET specific surface areas and mesopore size distribution is considered a good adsorbent of CO₂ because it exhibits better selectivity.

The adsorbent pore size distribution has a significant influence on CO₂ adsorption since it can affect the transport of mass, the available pore space of CO₂ adsorbents would increase to some extent, leading to a faster diffusion of CO₂ in the pores. Fig. 4 shows the pore size distributions of MgO based adsorbents. The MgO-SC and MgO-BM2.5h samples have large pore size distributions with average pore diameters of 9.20 and 21.52 nm, respectively (Fig. 4a and b); however, the MgO–Fe and MgO–Ni samples revealed pore size distributions relatively narrow, with average pore diameters of 9.20 and 4.20 nm, respectively (Fig. 4c and d). It was observed in our results that all MgO-based adsorbents exhibited mesoporous structures for high selectivity towards CO₂ molecules of 0.33 nm of kinetic radius, enough for CO₂ molecule diffusion inside the porous of MgO-based adsorbents; in this way, intraparticle diffusion improves the CO₂ capture process, because it facilitates the access of CO₂ molecules to pore active sites in pore walls. It is known that for chemisorption, CO₂ molecule has to diffuse into the adsorbent and interacts with the internal adsorbent surface. In this context, an excellent chemical adsorbent should not only have high specific surface area for chemisorption but also have appropriate pore size distribution for facilitating the CO₂ diffusion in the adsorbent pore structure (Song et al., 2016). In our case, the MgO-BM2.5 sample with a wide pore size distribution of 21.52 nm exhibited the best CO₂ adsorption performance than the other studied MgO-based adsorbents. In fact, the

### Table 1 - Textural properties of MgO-based adsorbents synthesized by solution combustion and ball-milling processes.

<table>
<thead>
<tr>
<th>Sample</th>
<th>S\textsubscript{BET} (m²/g)</th>
<th>V\textsubscript{TP} (cm³/g)</th>
<th>d\textsubscript{p} (nm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>MgO-SC</td>
<td>5.71</td>
<td>0.0289</td>
<td>12.12</td>
</tr>
<tr>
<td>MgO-BM2.5h</td>
<td>15.24</td>
<td>0.1347</td>
<td>21.52</td>
</tr>
<tr>
<td>MgO–Fe</td>
<td>15.85</td>
<td>0.1387</td>
<td>9.20</td>
</tr>
<tr>
<td>MgO–Ni</td>
<td>30.83</td>
<td>0.1588</td>
<td>4.20</td>
</tr>
</tbody>
</table>

S\textsubscript{BET}: superficial area; V\textsubscript{TP}: total pore volume; d\textsubscript{p}: average pore diameter.

* Barrett-Joyner–Halenda (BJH) method.

Fig. 3 – SEM micrographs of MgO-based adsorbents: (a) MgO-SC, (b) MgO-BM2.5h, (c) MgO–Fe, and (d) MgO–Ni.

Fig. 4 – BJH pore size distribution of MgO-based adsorbents: (a) MgO-SC, (b) MgO-BM2.5h, (c) MgO–Fe, and (d) MgO–Ni.

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mesoporous MgO is highly basic with well-ordered porous structure to expect high CO2 adsorption.

Fig. 5 shows the N2 adsorption–desorption isotherms for the as-prepared MgO nanocompounds. All these MgO-based adsorbents showed type IV isotherms according to the International Union of Pure and Applied Chemistry (IUPAC) classification (Abd-El-Hafiz et al., 2015), in accordance with this IUPAC isotherm classification, type IV isotherms are characteristics of mesoporous materials (2 nm < pore size < 50 nm). On the other hand, MgO-SC sample (Fig. 5a) shows a type H4 hysteresis. MgO-BM (Fig. 5b), MgO-Fe (Fig. 5c) and MgO-Ni (Fig. 5d) samples showed type H3 hysteresis in 0.4–1.0 p/p0 range. Materials with type H3 and H4 hysteresis are usually solids that formed particle agglomerates and have existence of flexible pores with morphology of slit or plate type. Furthermore, the hysteresis formation begins in the middle part of the isotherms, indicative of a distribution of pore size which extends mainly to the mesopore size at 0.4 < p/p0 < 1 (Adelkhani et al., 2011). Based on these findings, the presence of mesoporous structure in the prepared MgO-based adsorbents is confirmed, as was corroborated with the BJH results.

2.3. CO2 adsorption

The CO2 adsorption results on MgO-based adsorbents at 25°C, 1 atm, and 30 min of saturation time show that for MgO-SC, MgO-BM2.5h, MgO-Fe and MgO-Ni samples, maximum adsorption capacities of 0.39, 1.61, 0.90 and 1.50 mmol of CO2 per gram of adsorbent were obtained, respectively. Fig. 6 show the thermal gravimetric analysis (TGA) curve and mass spectroscopy (MS) of CO2 adsorption on studied MgO-based adsorbents. A weight loss of 7.09% for MgO-BM2.5h sample was observed (Fig. 6b), which was higher compared than MgO-SC (1.73 wt.%), MgO–Fe (3.99 wt.%) and MgO-Ni (6.64 wt.%) samples, respectively, due to their remarkably different textural and structural characteristics that were directly correlated with their CO2 adsorption capacities. With respect to Mg-Ni and MgO–Fe performance difference, Table 1 showed that for MgO doped with Ni and Fe metals (MgO-Ni and MgO–Fe) samples, significant improvement of specific surface area, as well as the total pore volume was observed. In catalysis field, doped metal oxides are widely investigated materials and novel doping-metals such as Fe and Ni have been proposed to improve the performance of metal oxide based adsorbents. In this case, catalytic activity and selectivity behaviors are linked to dopant acidity and basicity properties (Pighini et al., 2011). In fact, it has been demonstrated that nanocomposites can exhibit the multicomponent system properties in the same material (Singh et al., 2013). It has been shown that the Ni catalyst surface acidic or basic properties can significantly affect their CO2 adsorption performance, favoring the CO2 adsorption capacity on the MgO basic sites or inhibiting the CO2 adsorption by the deposition of a solid carbonaceous residue when surfaces are in contact (Jeong et al., 2016). Moreover, to our knowledge, iron incorporated in metal oxides has an excellent redox ability to capture CO2 under high temperatures; however, no study has been reported as CO2 capture material at low and ambient temperatures. In our case, at ambient temperature, Fe powders can have the inconvenience of carbon deposition in the metallic oxide active sites during the CO2 adsorption, causing the deactivation and textural destruction of MgO nanocompounds mixed with metallic powders as Fe, consequently resulting in a decrease in active sites due to the saturation with carbon species as was observed in MgO-Fe sample (Khajenoori et al., 2015; Ross, 2005; Wang et al., 2012). Thus, it was observed that after the introduction of Ni and Fe into MgO adsorbent, both of the specific surface areas and total pore volumes increase (Table 1), benefiting their CO2 performance by increasing these important textural characteristics irrespective of their synthesis method. Additionally, the solid milling favored particle fragmentation which provides a large number of active sites, as well adding the corresponding Ni and Fe active sites. However, the corresponding average pore sizes decrease due to prolonged pore compaction caused by the ball-milling process, taking into account that MgO-Fe sample was ball-milling during 5.0 hr and MgO-Ni sample during 7.5 hr. Therefore, a significant influence of the doping element on the CO2 adsorption behavior was revealed. Moreover, even if these samples are issued from the same precursor, doped samples allowed to increase or decrease the CO2 adsorption capacity. Hence, the MgO-BM2.5h exhibited better CO2 adsorption capacity due to their morphology, textural and microstructure properties. Moreover, this MgO-BM2.5h sample showed more abundant mesoporous structure to ensure that CO2 molecules easily come into contact with the active sites that provide their extended specific surface area, improving the interaction between CO2 gas molecules with the MgO-BM2.5h mesopores. According to Li et al. (2010), the enhanced interaction at mesopores gives rise to strong adsorption of CO2, which finally improved the CO2 adsorption capacity. Similarly, Bhagiyalakshmi et al. (2010), reported that MgO powders are highly basic and can adsorb CO2 molecules in the active sites of MgO surface which are trapped into the pores by chemical reaction of MgO and CO2 to form MgCO3. Therefore, the specific surface area, total volume pore, average pore diameter, morphology and crystallinity were critically important in encouraging the CO2 adsorption capacity on these MgO-based adsorbents studied here. MgO-BM2.5h powders are highlighted to be beneficial for potential applications in CO2 adsorption under practical conditions at 25°C, 1 atm, and 30 min of saturation time.

Thermal decomposition process for the MgO-BM2.5h sample after CO2 exposure was examined using TGA and differential scanning calorimetry (DSC) techniques. The TGA/DSC curves
for CO₂ desorbed MgO-BM2.5h powders are shown in Fig. 7. It can be observed that in the DSC curve there is an endothermic peak at 99.5°C caused by the elimination of physically adsorbed water (Vu et al., 2014). At higher temperature, a second and third exothermic broad peaks from 150 to 750°C were observed, which are associated with carbonaceous material (CO₂) release. The small peak at around 370°C corresponds to weakly adsorbed CO₂ while the broad peak at around 330°C represents desorption of the strongly chemisorbed CO₂ (Bhagiyalakshmi et al., 2011). Subsequently, an endothermic peak is observed at around 800°C due to the complete MgO phase formation. From the TGA/DSC results, the release of CO₂ is due to MgCO₃ formation. This CO₂ adsorption on MgO-MM2.5h by carbonation was confirmed by FT-IR (FT: fourier transform; IR: infrared) analysis. Therefore, the TGA/DSC result indicates that CO₂ was chemisorbed on MgO-BM2.5h sample, and that at around 800°C a complete sharp regeneration of CO₂ was observed. Then, the obtained TGA/DSC results are suggested that chemisorption was mainly the mechanism that governs the CO₂ adsorption on MgO-BM2.5 adsorbent.

In order to analyze the carbonate formation, FT-IR analysis was performed. In Fig. 8, IR spectra are shown for a MgO-BM2.5h sample without further treatment (Fig. 8a), for MgO-BM2.5h sample degassed at 325°C during 30 min (Fig. 8b), and for MgO-BM2.5h sample after CO₂ adsorption. The FT-IR analysis of MgO-BM2.5h samples exhibits a typical metal oxide spectrum with significant difference between the fresh material, degasified material, and sutured material by CO₂. All spectra in Fig. 8a-c show four main adsorption bands: at 522 cm⁻¹, a remarkable adsorption band ranging from 1200 to 1700 cm⁻¹, a band at 2314 cm⁻¹, and at 3433 cm⁻¹ respectively. The bands at 522 and 3433 cm⁻¹ are assigned to the Mg–O bond and stretching vibrations of the O–H bond respectively (Devaraja et al., 2014). The IR adsorption bands of hydroxyl groups (water
surface adsorbed) at 3433 cm\(^{-1}\) are generally attributed to strong hydrogen bound OH species that are usually found in MgO even after vacuum drying (Yan et al., 2009). The band at 2314 cm\(^{-1}\) after CO\(_2\) adsorption (Fig. 8c) indicates the existence of physisorbed carbon dioxide (Ding et al., 2015). Small band at 2314 cm\(^{-1}\) also was found in Fig. 8a and b, it should be expected that this 2314 cm\(^{-1}\) band disappears in degassed MgO-BM2.5h sample (Fig. 8b), however, the 2314 cm\(^{-1}\) band for fresh material was very much smaller due to their large manipulation time for IR measurement or by hygroscopic nature of MgO. By examination of 1200 to 1700 cm\(^{-1}\) region, three different unidentate, bidentate and bicarbonate species of adsorbed CO\(_2\) were detected. According to Pighini et al. (2011), unidentate and bidentate carbonate formation requires surface basic oxygen atoms, in our study, unidentate carbonates were exhibited at 1280 and 1490 cm\(^{-1}\) adsorption bands. Bidentate carbonate stretching at 878 and 1560 cm\(^{-1}\) adsorption bands. Finally, bicarbonate species that involves surface hydroxyl groups were exhibited at 1630 and 1417 cm\(^{-1}\) adsorption bands. IR revealed the formation of MgCO\(_3\) in the adsorbent studied here. Then, based on these results, it is further demonstrated that the adsorption mechanism of the MgO-BM2.5h sample was mainly relying on chemisorption accompanying with a very small fraction of physisorption according to Song et al. (2015).

To evaluate the reuse and stability of MgO-BM2.5h-based adsorbent for practical applications, experiments of adsorption-desorption cycles on this sample were repeated five times. CO\(_2\) adsorption was carried out at 25°C, 1 atm, and 30 min of saturation time. CO\(_2\) desorption was carried out by using TGA from 0 to 850°C, with a heating ramp of 20°C/min. According to Wang et al. (2014), this study is necessary because long-time stability of an adsorbent is a critical parameter for the commercialization of a CO\(_2\) scavenger. It is well-known that the characteristic of an adsorbent in a practical application point of view is its useful life-time, because longer period of time brings to a meaningful decrease in cost of the synthesis. Fig. 9 shows the cyclical capacity of MgO-BM2.5h powders, this sample exhibited a CO\(_2\) adsorption capacity of 1.47 mmol/g at 5th cycle, equivalent to 91.3% of the obtained capacity in 1st cycle (1.61 mmol/g). Then, the cyclic CO\(_2\) adsorption–desorption showed that MgO-BM2.5h sample maintains high stability in cyclic capacity, beneficial properties for its economical and practical applications as CO\(_2\) adsorbent.

The methods reported here are advantageous because the solution–combustion synthesis is a versatile process that with only 5 min allows to synthesize MgO powders of improved textural properties, which play an important role for CO\(_2\) adsorption efficiency. Besides, the ball-milling allowed one resulting product of very fine particulates, with strain and defects forming friable agglomerates improving their textural and structural properties that enhanced CO\(_2\) adsorption efficiency on MgO powders. In summary, in this work the CO\(_2\) adsorption on MgO obtained by solution combustion and treated by ball-milling at room temperature and pressure conditions has been demonstrated, which has not been reported previously. Therefore, a better understanding of textural and structural influence properties of the as-prepared porous MgO-based adsorbents on the CO\(_2\) adsorption behavior was attained, as a contribution of alleviating the serious global warming problem.

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**Fig. 8** - FT-IR spectra of: (a) MgO-BM2.5h without further treatment, (b) MgO-BM2.5h with degassed treatment at 325°C during 30 min, and (c) CO\(_2\) adsorption on MgO-BM2.5h at 25°C and 1 atm. FT: fourier transform; IR: infrared.

**Fig. 9** - Cyclic test of CO\(_2\) adsorption on MgO-BM2.5h sample at 25°C and 1 atm.
3. Conclusions

In the present investigation, mesoporous MgO-based adsorbents were synthesized successfully by solution-combustion and ball-milling processes, likewise were mixed with Fe and Ni elements to obtain their respective nanocomposites, and their CO$_2$ adsorption behaviors were investigated. Experimental results revealed that MgO-based adsorbent obtained by solution–combustion was crystalline, highly porous with extended specific surface area. However, when this resulted MgO material was ball-milling treated during 2.5 hr (MgO-BM2.5h), their textural and structural properties were improved. The results show that this MgO-BM2.5h adsorbent was able to adsorb CO$_2$ very well at room temperature and pressure with a maximum CO$_2$ adsorption capacity of 1.611 mmol/g. After the introduction of Fe and Ni into MgO, Fe had a negligible influence on CO$_2$ adsorption behavior, in comparison with the favorable influence of Ni into MgO sample. Chemisorption was the main mechanism for CO$_2$ adsorption on mesoporous MgO-BM2.5h, accompanying with a very small fraction of physisorption which was probed by FTIR study. CO$_2$ adsorption behavior on MgO-based adsorbents was correlated with their structural, textural and structural properties as: specific surface area, total pore volume, pore size distribution and crystallinity. The recyclability of the mesoporous MgO-BM2.5h adsorbent is highly commendable, because no significant loss of performance was measured until five regeneration times. In general, the results indicate that the ball-milling process plays a crucial role in the obtaining of highly porous MgO with better textural and structural properties for enhancing the CO$_2$ adsorption capacity.

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