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# Extraction of silica from sugarcane bagasse ash and its utilization in zeolite 4A synthesis for CO<sub>2</sub> adsorption

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Sugarcane bagasse ash (SCBA) is a solid waste containing a high amount of silica (SiO<sub>2</sub>) and is suitable to utilize as a silica source for synthesizing zeolite NaA. SCBA is typically calcined at high temperatures before silica extraction. The method is not environmentally friendly because it consumes energy and produces CO<sub>2</sub>. This work demonstrates an alternative extraction method of SiO<sub>2</sub> from SCBA by treating it with hydrochloric (HCl) and sodium hydroxide (NaOH) solution. The obtained mixture was separated by paper filter No. 1 (P) and a combination of paper filter and syringe filter (PS). The solution was neutralized by HCl solution, producing silica (SiO<sub>2</sub>-P and SiO<sub>2</sub>-PS) with a purity of 98 wt%. Both SiO<sub>2</sub> samples and SCBA were utilized to synthesize zeolite NaA for CO<sub>2</sub> adsorption. The CO<sub>2</sub> adsorption capacities of NaA-P and NaA-PS were 4.30 and 4.10 mmol g<sub>adsorbent</sub><sup>-1</sup>, in the same range as commercial NaA. The capacity is influenced by the total basicity of zeolite. The CO<sub>2</sub> adsorption behavior of all samples correlates well with the Toth model. The CO<sub>2</sub> adsorption kinetics agrees well with the pseudo-second-order kinetic model. Overall, this work shows the successful extraction of silica *via* using a direct NaOH solution, yielding high-purity silica sufficient for synthesizing zeolite NaA, a promising adsorbent of CO<sub>2</sub>.

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## 1. Introduction

Fossil fuels continue to dominate as the primary source of global energy consumption, and this reliance is expected to persist for several more decades. The use of fossil fuels generates carbon dioxide (CO<sub>2</sub>),<sup>1</sup> the main greenhouse gas. The atmospheric CO<sub>2</sub> concentration has been reported to reach a crucial threshold of 425 ppm.<sup>2</sup> Consequently, several countries have taken action to reduce CO<sub>2</sub> emissions through various approaches, including carbon capture and storage (CCS).<sup>3</sup> Carbon capture is generally employed in large-scale resources releasing a high concentration of CO<sub>2</sub>, such as flue gas in coal plants and industries.<sup>4</sup> Given the rising levels of CO<sub>2</sub>, it remains imperative to persistently advance methods to reduce CO<sub>2</sub> emissions.

Techniques for removing CO<sub>2</sub> include adsorption, cryogenic distillation, microalgae bio-fixation, and membrane separation.<sup>2</sup> The most common method is adsorption on solids, which is simple and cost-effective, with the possibility of regenerating

the adsorbent.<sup>5</sup> Zeolites are among the effective adsorbents due to their large surface area, high thermal stability, and defined pore structure.<sup>6,7</sup> Those properties could allow efficient CO<sub>2</sub> adsorption at high temperatures, beneficial for flue gas utilization in coal plants and industries. Besides, the pore structure of zeolite could serve as shape-selective adsorption of CO<sub>2</sub>.

Zeolites have an aluminosilicate framework comprising tetrahedral TO<sub>4</sub> units (T = Si or Al) joined by shared oxygen atoms.<sup>8</sup> The silicon-to-aluminum ratio (Si/Al) strongly influences the zeolite structure and adsorption capacities.<sup>9</sup> There are reports that zeolites with low Si/Al ratios, such as NaX and NaA have high CO<sub>2</sub> adsorption capacity due to their high basicity.<sup>7,9,10</sup> It was reported that NaA and NaX synthesized from rice husk silica have CO<sub>2</sub> adsorption capacity of 97.7 and 107.8 mg-CO<sub>2</sub> per g-zeolite, respectively.<sup>10</sup> Despite the lower CO<sub>2</sub> adsorption capacity, NaA is a promising adsorbent due to its high CO<sub>2</sub> adsorption capacity, simple synthesis method with low energy consumption from a short crystallization time.<sup>10,11</sup> Siriwardane *et al.*<sup>7</sup> suggested that the presence of more cations (Na<sup>+</sup>) of zeolite promotes the basicity of the material, resulting in the greater adsorption capacity of CO<sub>2</sub>. The capacity could be improved by increasing the sodium content in the synthesis gel. Overall, it is worth focusing on the improvement of CO<sub>2</sub> adsorption on NaA zeolite by synthesizing with a higher molar ratio of Na content in the synthesis gel.

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Some downsides in the zeolite synthesis are the energy cost and non-environmentally friendly silica sources, such as silicon alkoxides, silicic acid, and fumed silica.<sup>12,13</sup> Thus, the utilization of silica from biomass waste for zeolite synthesis is considered a cheaper route due to the value-added waste and its alternative application.<sup>14</sup> There are several biomass wastes with high silica content such as rice husk ash,<sup>8</sup> cogon grass weed,<sup>15</sup> and sugarcane bagasse ash (SCBA) that have been used in the zeolite synthesis.<sup>16</sup> Accordingly, Thailand is one of the top world sugar producers, generating a large amount of sugarcane bagasse.<sup>17,18</sup> The sugarcane bagasse is further utilized as a fuel in cogeneration to generate steam and electricity. Consequently, the SCBA remains the final waste with low commercial value in the sugar production chain.<sup>18</sup> SCBA waste consists of ~13 wt% silica content;<sup>19</sup> it has potential as a silica source for the synthesis of NaA zeolite.

To obtain zeolite with high purity, the silica source in the synthesis gel should have high purity.<sup>20</sup> Subsequently, SCBA waste must be treated to improve the silica purity before use in the synthesis gel of zeolite. The residual carbon in SCBA could be removed by calcination in air at high temperatures, *i.e.*, from 600 °C to 1000 °C for several hours followed by a treatment in an acid solution to remove metal impurities.<sup>21,22</sup> The calcination at high temperatures is energy-consuming and produces CO<sub>2</sub> to the environment. Thus, the process could be considered non-environmentally friendly.

Herein, we demonstrate a selectively extracted silica from SCBA by direct treatment with NaOH solution in the form of sodium silicate solution (Na<sub>2</sub>SiO<sub>3</sub>), followed by precipitation of SiO<sub>2</sub> in mild HCl solution.<sup>23</sup> The product is directly employed as a silica source for the synthesis of zeolite NaA. Moreover, the residual carbon particles are separated from the Na<sub>2</sub>SiO<sub>3</sub> solution by filtration with filter paper (P) followed by a syringe filter (PS). The remaining residual carbon particles might be useful in other applications such as fertilizer and/or adsorbent of pollutants without any loss of benefits.<sup>18</sup>

The goals of this study are to synthesize the NaA zeolites using silica from SCBA by treatment with NaOH solution and to investigate their CO<sub>2</sub> adsorption capacity, compared with commercial NaA zeolite. Moreover, the physicochemical properties, thermal stability, morphology, basicity, adsorption model, and kinetics of NaA zeolites using silica from SCBA, are discussed and compared to that of commercial-NaA zeolite.

## 2. Results and discussion

Fig. 1a–c demonstrates the color of as-received SCBA, SiO<sub>2</sub>-P, and SiO<sub>2</sub>-PS. As-received SCBA is dark powder due to unburned carbons, similar to the appearance reported in the literature.<sup>19</sup> The SiO<sub>2</sub>-P and SiO<sub>2</sub>-PS samples are gray and white powders, respectively, implying the normal color of silica. The color difference in those samples implies their distinct purities.<sup>23</sup> The yields of silica powder calculated according to eqn (6) for SiO<sub>2</sub>-P and SiO<sub>2</sub>-PS were 20.4 wt% and 19.6 wt%, respectively. This result suggests that the two-step filtrations might eliminate impurities from NaSiO<sub>3</sub> solution resulting in the whiter powder for SiO<sub>2</sub>-PS.

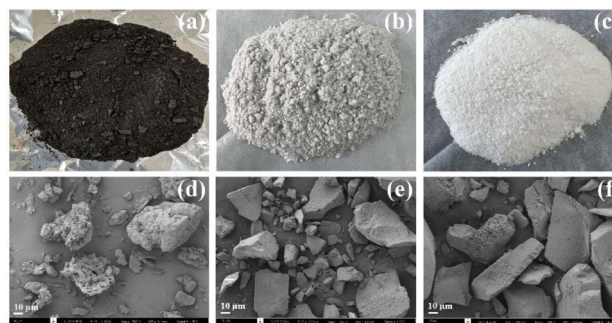


Fig. 1 Photographs of powder and SEM images of as-received SCBA (a and d), SiO<sub>2</sub>-P (b and e), and SiO<sub>2</sub>-PS (c and f).

Fig. 1d–f reveals the morphologies of as-received SCBA, SiO<sub>2</sub>-P, and SiO<sub>2</sub>-PS from SEM. The as-received SCBA has morphology of irregular shapes with a rough surface of particles containing various sizes of fibers.<sup>19</sup> After the extraction process, SiO<sub>2</sub>-P, and SiO<sub>2</sub>-PS gave the aggregation of dense particles and smooth surfaces. These results are consistent with the morphology of commercial SiO<sub>2</sub> reported in the literature.<sup>24</sup>

Fig. 2a demonstrates the functional groups of raw-SCBA, SiO<sub>2</sub>-P, and SiO<sub>2</sub>-PS, and the band assignment is shown in Table 1. The as-received SCBA exhibits mainly the vibration spectra of Si–O–Si, and carbonate species,<sup>25</sup> suggesting that silicon oxide and metal carbonate are the main components. In contrast, SiO<sub>2</sub>-P and SiO<sub>2</sub>-PS only show the peaks corresponding to Si–O–Si, indicating that HCl solution removes metal impurities and NaOH selectively extract silicon oxide from the as-received SCBA.<sup>23</sup> The additional bands at 968 cm<sup>-1</sup> and 1636 cm<sup>-1</sup> were attributed to the vibrations of Si–OH and water adsorbed on the surface SiO<sub>2</sub>.

Fig. 2b displays thermograms showing the thermal decomposition of as-received SCBA, SiO<sub>2</sub>-P, and SiO<sub>2</sub>-PS from room temperature to 800 °C under air zero. The thermal decomposition of as-received SCBA occurs in three ranges: 50–200 °C, 350–500 °C, and 640–700 °C. Those weight losses were attributed to the removal of adsorbed water, decomposition of organic compounds,<sup>26,27</sup> and decomposition of CaCO<sub>3</sub>, respectively.<sup>28</sup> The result is in good agreement with the report from Castaldelli *et al.*<sup>26</sup> that Ca species is one of the second major components remaining in raw SCBA from the sugar industry in Brazil. The thermograms of SiO<sub>2</sub>-P, and SiO<sub>2</sub>-PS gave only two main regions in the temperature ranges of 80–150 °C, and 150–400 °C contributing to the water desorption and unburned carbon. One should notice that more weight loss in the range of 150–400 °C was pronounced in SiO<sub>2</sub>-P implying some trace amount of organic matter remaining in the sample.

Fig. 3 exhibits the phases of as-received SCBA, SiO<sub>2</sub>-P, and SiO<sub>2</sub>-PS determined by XRD. The raw SCBA provided the crucial peaks of quartz (SiO<sub>2</sub>), calcite (CaCO<sub>3</sub>), and hematite (Fe<sub>2</sub>O<sub>3</sub>), respectively. The quartz gave essential peaks at 20.8° and 26.5°. Besides, the calcite displayed character peaks at 23.1°, 29.3°, 39.5°, 43.2°, 47.3°, and 48.4°, respectively<sup>30</sup> and hematite exhibited the crucial peak at 36.1°. The result is not much surprising because these phases have been reported as the



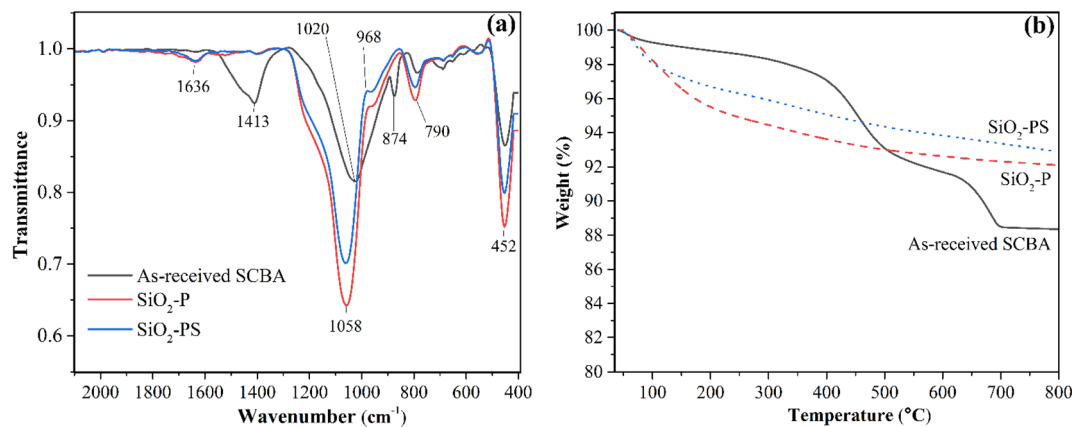


Fig. 2 FTIR spectra (a) and TGA curves (b) of as-received SCBA, SiO<sub>2</sub>-P, and SiO<sub>2</sub>-PS.

Table 1 Bands assignment of FTIR for raw-SCBA, SiO<sub>2</sub>-P, and SiO<sub>2</sub>-PS

Wavenumber (cm <sup>-1</sup> )	Assignment	Reference
<b>As-received SCBA</b>		
452	Bending vibration of Si–O–Si	25
788	Symmetric vibration of Si–O–Si	
874	Out-of-plane angular deformation of group CO <sub>3</sub> <sup>2-</sup> vibration	
1020	Asymmetric stretching of Si–O–Si	
1413	Asymmetric vibration of carbonate species	
<b>SiO<sub>2</sub>-P</b>		
452	Bending vibration of Si–O–Si	25 and 31
790	Symmetric vibration of Si–O–Si	
968	Vibration of Si–OH	
1058	Asymmetric stretching of Si–O–Si	
1636	Vibration of water adsorbed on the surface	
<b>SiO<sub>2</sub>-PS</b>		
452	Bending vibration of Si–O–Si	25 and 31
790	Symmetric vibration of Si–O–Si	
967	Vibration of Si–OH	
1060	Asymmetric stretching of Si–O–Si	
1636	Vibration of water adsorbed on the surface	

major components of SCBA.<sup>16</sup> Moreover, the presence of the calcite phase aligns well with the results from FTIR and TGA. In terms of SiO<sub>2</sub>-P, and SiO<sub>2</sub>-PS, only a broad peak at 22° attributed to the characteristic peak of amorphous silica was observed. The result implies that both samples contain a high purity of amorphous silica (SiO<sub>2</sub>). This assumption correlates well with the result from elemental compositions from XRF as demonstrated in Table 2. Moreover, the obtained silica in the amorphous phase could be beneficial for zeolite synthesis because it can be dissolved easily in NaOH solution to produce sodium silicate (Na<sub>2</sub>SiO<sub>3</sub>).<sup>15</sup>

Fig. 4a demonstrates FTIR spectra of NaA-SCBA, NaA-P, NaA-PS, and commercial-NaA. All samples exhibit five crucial peaks of the zeolite NaA framework. The peaks at 462 cm<sup>-1</sup> and 545 cm<sup>-1</sup> were attributed to the internal tetrahedral vibration of O–T–O bending when T is Si or Al and, the external vibration of double four-rings. Besides, the peaks at 662 cm<sup>-1</sup> and 972 cm<sup>-1</sup>

corresponded to the internal vibration of Si–O–T symmetric stretching and asymmetric O–H scissors-like vibration of adsorbed water.<sup>32</sup> Moreover, NaA-SCBA, NaA-P, and NaA-PS samples had similar FTIR patterns to commercial-NaA. This result suggests that the zeolite synthesized using silica from SCBA has essentially zeolite NaA type.

The thermographs of NaA-SCBA, NaA-P, NaA-PS, and commercial-NaA are shown in Fig. 4b. The thermal behavior of physisorbed water on zeolite implies the character of each zeolite type.<sup>33</sup> NaA-SCBA shows weight loss in four main regions: 100–200 °C, 200–400 °C, 400–500 °C and 600–700 °C, attributing to the desorption of physisorbed water, water in the zeolite structure, volatile organic compounds, and the decomposition of CaCO<sub>3</sub>, respectively.<sup>26,27,34</sup> The character weight losses of organic compounds and CaCO<sub>3</sub> were observed probably due to the direct use of silica from as-received SCBA containing high impurities. Other three zeolites including NaA-P, NaA-PS, and



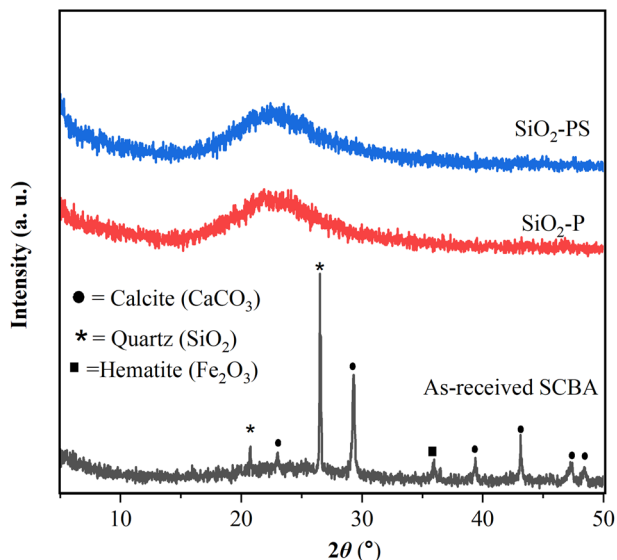


Fig. 3 XRD patterns of as-received SCBA, SiO<sub>2</sub>-P, and SiO<sub>2</sub>-PS.

Table 2 Elemental compositions determined by XRF of as-received SCBA, SiO<sub>2</sub>-P, and SiO<sub>2</sub>-PS<sup>a</sup>

Sample	Component (wt%)					
	SiO <sub>2</sub>	Al <sub>2</sub> O <sub>3</sub>	CaO	Fe <sub>2</sub> O <sub>3</sub>	MnO <sub>2</sub>	TiO <sub>2</sub>
As-received SCBA	77.2	6.9	12.3	2.7	0.5	0.4
SiO <sub>2</sub> -P	98.1	1.9	n.d.	n.d.	n.d.	n.d.
SiO <sub>2</sub> -PS	97.9	2.1	n.d.	n.d.	n.d.	n.d.

<sup>a</sup> n.d.: not detected.

commercial-NaA exhibited similar weight loss in two ranges: 100–200 °C and 200–400 °C attributed to the desorption of physisorbed water and water in the zeolite structure, respectively.<sup>34</sup> The percentage weight losses were 21.3%, 18.8%, and 18.9% for NaA-P, NaA-PS, and commercial-NaA, respectively. The thermal behavior of physically adsorbed water, similar to commercial-NaA could imply that the synthesis of NaA zeolite is successful by using SiO<sub>2</sub>-P and SiO<sub>2</sub>-PS. The more weight loss

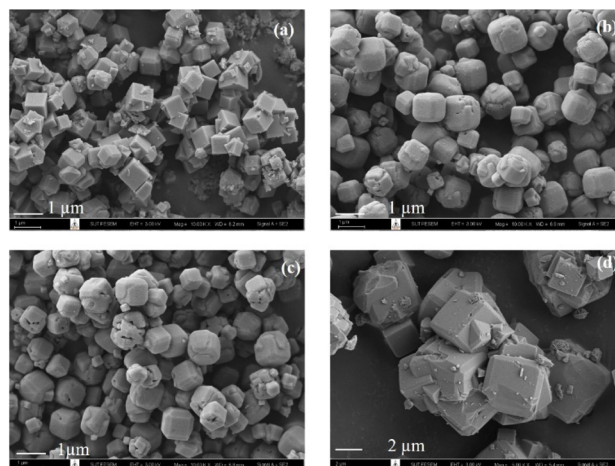


Fig. 5 SEM images of NaA-SCBA (a), NaA-P (b), NaA-PS (c), and commercial-NaA (d).

from NaA-P was probably due to the remaining carbon matter from the silica source (see Fig. 2b).

Fig. 4c demonstrates XRD patterns of NaA-SCBA, NaA-P, NaA-PS, and commercial-NaA. It is noticeable that the NaA-SCBA gave mixed characteristic peaks of zeolite NaA (5A) according to JCPDS #39-0222, and quartz.<sup>16</sup> Meanwhile, NaA-P, NaA-PS, and commercial-NaA gave essentially characteristic peaks of pure zeolite NaA (4A).<sup>10</sup> The substitution of sodium ions from the initial gel with calcium ions from the raw SCBA results in the generation of pore diameter of 5 Å (zeolite 5A) instead of 4.1 Å (zeolite 4A). This evidence confirms the success in the synthesis of pure zeolite NaA (4A) from silica extracted from SCBA by both SiO<sub>2</sub>-P and SiO<sub>2</sub>-PS approaches.

Fig. 5a–d displays the morphologies of NaA-SCBA, NaA-P, NaA-PS, and commercial-NaA. The NaA-SCBA (Fig. 5a) provided mainly cubic morphology with a sharp edge, a characteristic shape of zeolite 5A,<sup>35</sup> and mixed with small particles. The presence of small particles is probably due to the remaining quartz phase suggested by the XRD result. Moreover, the dominant cubic morphology with average particle sizes around 800 nm was mainly observed. The NaA-P and NaA-PS (Fig. 5b and c) showed similar morphologies of a cubic shape and curvy

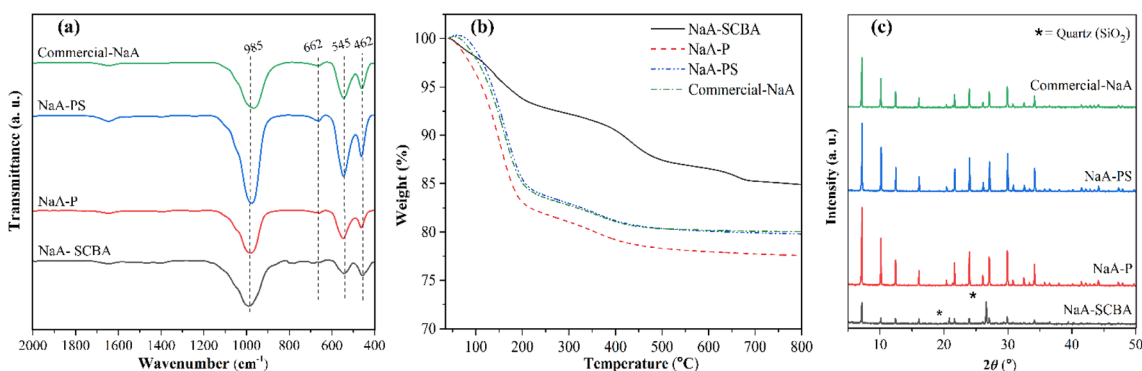


Fig. 4 FTIR spectra (a), TGA curves (b), and XRD patterns (c) of NaA-SCBA, NaA-P, NaA-PS, and commercial-NaA.



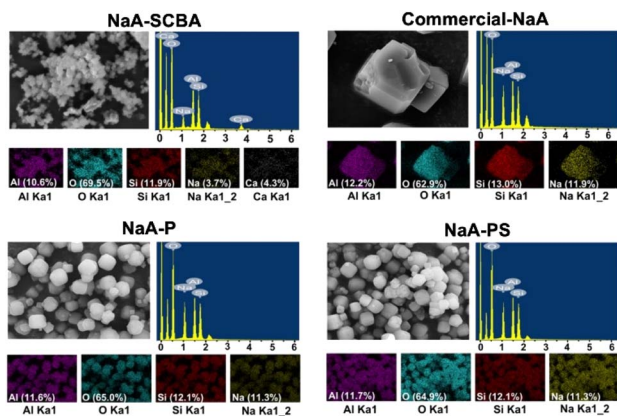


Fig. 6 SEM-EDS mapping of NaA-SCBA, commercial-NaA, NaA-P, and NaA-PS.

edge crystals with an average particle size of around 0.9  $\mu\text{m}$ . The morphology was identical to zeolite 4A in the literature.<sup>36</sup> The difference is that the commercial-NaA has a larger particle, *i.e.*, around 4  $\mu\text{m}$ . The observation correlates well with the morphology of commercial-NaA (4A) reported in the literature.<sup>34</sup>

Fig. 6 demonstrates SEM-EDS mapping of NaA-SCBA, NaA-P, NaA-PS, and commercial-NaA. The NaA-SCBA revealed the dispersion of Al, Si, O, Na, and Ca throughout the particles of zeolite. The presence of Ca in the zeolite structure could be caused by the ion exchange with  $\text{Na}^+$  resulting in the formation of zeolite 5A, consistent with the result from XRD.<sup>34</sup> The hypothesis also correlates well with the small amount of Na (3.7%) observed in the zeolite, in contrast to the other three zeolites which contain a higher amount of Na (11%). Besides, NaA-P, NaA-PS, and commercial-NaA exhibited the suitable

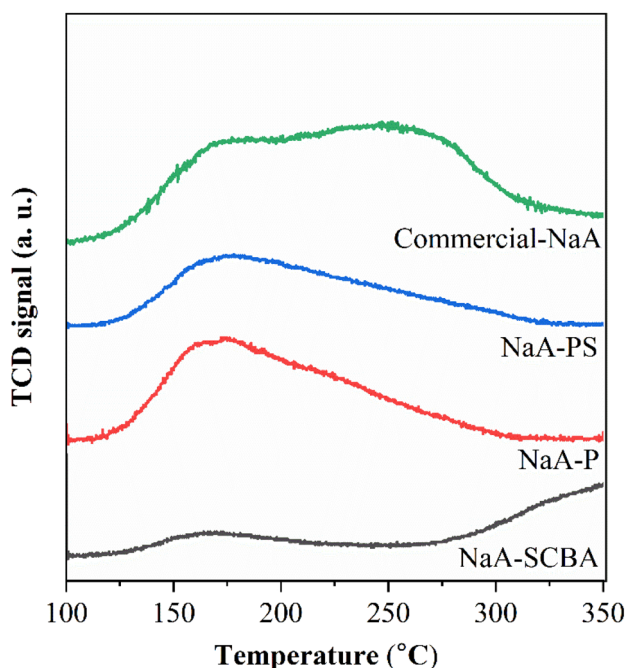


Fig. 7  $\text{CO}_2$ -TPD profiles of NaA-SCBA, NaA-P, NaA-PS, and commercial-NaA.

Table 3 Basicity and Si/Al ratios of NaA-SCBA, NaA-P, NaA-PS, and commercial-NaA

Sample	Basicity <sup>a</sup> ( $\text{mmolCO}_2 \text{ g}_{\text{adsorbent}}^{-1}$ )	Si/Al (atomic ratio) <sup>b</sup>	Na/Si (atomic ratio) <sup>b</sup>
NaA-SCBA	0.029	2.64	0.11
NaA-P	0.094	1.30	0.47
NaA-PS	0.064	1.32	0.42
Commercial-NaA	0.117	1.30	0.63

<sup>a</sup> Calculated from the desorption peak in the temperature range from 100  $^\circ\text{C}$  to 350  $^\circ\text{C}$ . <sup>b</sup> Obtained from ED-XRF result.

elements including Al, Si, O, and Na with the Si/Al ratio of about 1 of zeolite NaA.<sup>37</sup>

Fig. 7 manifests  $\text{CO}_2$ -TPD profiles of NaA-SCBA, NaA-P, NaA-PS, and commercial-NaA, and the amount of basicity in each sample is summarized in Table 3.  $\text{CO}_2$ -TPD was employed to evaluate the strength of the basic sites available in zeolites for  $\text{CO}_2$  adsorption.<sup>38</sup> It was observed that all samples had different desorption peaks and intensities implying the presence of their distinct basic strengths and basicities. The amount of basicity was in the order of commercial-NaA > NaA-P > NaA-PS > NaA-SCBA. The difference in basicity could be the consequence of the distinct values of Si/Al and Na/Si ratio in each sample. As known, the framework of zeolite is a building of tetrahedral  $\text{TO}_4$  units ( $\text{T} = \text{Si}$  or  $\text{Al}$ ) bonded together by sharing oxygen atoms. The zeolite framework will present the net negative charge that requires Na in the positive form ( $\text{Na}^+$ ) as a charge balancing.<sup>37</sup> Either a net negative charge or positive Na ( $\text{Na}^+$ ) could be an active site for  $\text{CO}_2$  adsorption.<sup>10</sup> Thus, zeolite consisting of low Si/Al and high Na/Si ratios could be beneficial for  $\text{CO}_2$  adsorption. As a result, the Si/Al and Na/Si ratios of zeolites were correlated well with the amount of basicity as demonstrated in Table 3. Moreover, Gouveia *et al.*<sup>4</sup> have suggested that the total amount of basic sites in the adsorbent has a stronger influence on  $\text{CO}_2$  adsorption than that of basic strength. The presence of the  $[\text{AlO}_4]^{5-}$  tetrahedron in zeolite (NaA) results in a negative charge, which is balanced by the positively charged  $\text{Na}^+$  cation. This cation could serve as the adsorption center for  $\text{CO}_2$  in zeolites. Therefore, the concentration of  $\text{Na}^+$  cation could play a crucial role in governing the adsorption process, consistent with the increase of Na/Si ratio in Table 3. This assumption agrees well with the report in the literature<sup>10,39</sup> that the positively charged  $\text{Na}^+$  cations in zeolite NaX and NaA are the main adsorption sites for  $\text{CO}_2$ . Thus, the improvement of the Na/Si ratio in zeolites to enhance adsorption capacity could be done in several ways, for example, by changing the gel composition and optimizing the crystallization conditions.<sup>36</sup>

Fig. 8 exhibits  $\text{CO}_2$  adsorption isotherms of NaA-SCBA, NaA-P, NaA-PS, and commercial-NaA performed at 25  $^\circ\text{C}$  and the pressure range of 0.0–1.0 bar. It was noticed that the adsorption behavior in all samples showed type I isotherm according to the IUPAC classification suggesting a physical adsorption character in all samples.<sup>10</sup> The sorption isotherm has a sharp curve at the low pressure because of the abundant adsorption sites followed by the gradient decreased and became constant at high



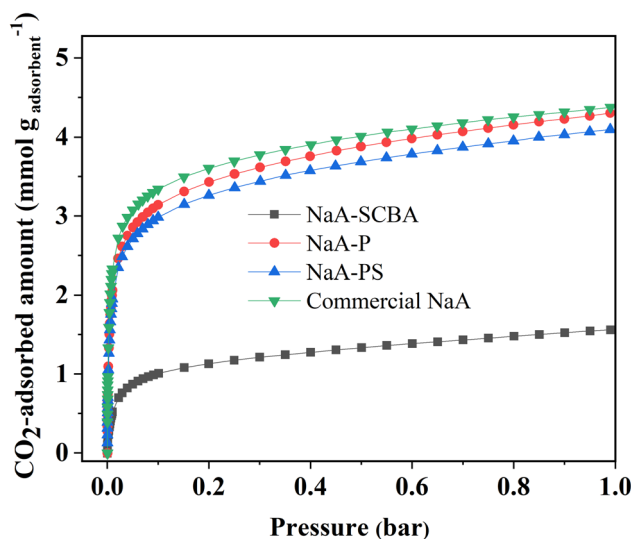


Fig. 8 CO<sub>2</sub> adsorption isotherms of NaA-SCBA, NaA-P, NaA-PS, and commercial-NaA performed at 25 °C and the pressure range of 0.0–1.0 bar.

pressure. Then, the adsorption behavior was almost unchanged at the pressure close to 1 bar.

The fundamental interactions between zeolites and CO<sub>2</sub> molecules are mostly electrostatic force. The zeolite adsorbs CO<sub>2</sub> molecules due to the strong quadrupolar interactions between adsorbate molecules and the persistent electric field created by the charge-balancing cation of zeolites. The type of the cations (Na<sup>+</sup>) and their location in the zeolite pore reflect the material's permanent electric field. Therefore, CO<sub>2</sub> uptake is dramatically increased at the beginning due to the available cation in the zeolite framework. At high relative pressure, there is no availability of unoccupied cations, thus the adsorption curve became almost constant.<sup>7,40</sup>

Furthermore, the adsorption capacity of CO<sub>2</sub> was in the following: commercial-NaA ~ NaA-P > NaA-PS > NaA-SCBA. This order is in good agreement with the amount of total basicity obtained from CO<sub>2</sub>-TPD. Moreover, this evidence is consistent with the report by Gouveia *et al.*<sup>4</sup> that the basicity has a strong influence on CO<sub>2</sub> adsorption in zeolitic materials. One should rationalize that zeolite purity is one of the important factors enhancing CO<sub>2</sub> adsorption.<sup>41</sup> This assumption correlates well with the report by Madhu *et al.*<sup>10</sup> that zeolite NaA with a higher purity synthesized from kaolinite provides a greater CO<sub>2</sub> adsorption capacity. Besides, the CO<sub>2</sub> adsorption capacity of NaA-P had almost similar to that of commercial NaA (4.30 vs. 4.38 mmol g<sup>-1</sup> in Table 5) indicating a promising strategy process and adsorbent due to the use of silica from SCBA waste.

Moreover, an equilibrium isotherm is crucial information delineated through adsorption isotherm data, delineating specific parameters that elucidate the surface characteristics and affinity of the adsorbent for the intended adsorbate. There are three models employed in this work to predict the adsorption isotherm including Langmuir, Freundlich, and Toth models.

The Langmuir adsorption model assumes a monolayer adsorption on a surface with a finite number of identical sites. It proposes that adsorption occurs through the formation of a saturated monolayer, and once all sites are occupied, no further adsorption can take place as demonstrated in equation (eqn (1)).

$$\frac{C_e}{q_e} = \frac{1}{K_L q_m} + \frac{1}{q_m} C_e \quad (1)$$

where,  $q_m$  (mg g<sup>-1</sup>) represents the theoretical maximum adsorption capacity,  $K_L$  (L mg<sup>-1</sup>) signifies the Langmuir isotherm constant,  $C_e$  denotes the equilibrium concentration (mg L<sup>-1</sup>).

The Freundlich adsorption model is an empirical equation used to describe non-ideal adsorption on heterogeneous surfaces. It assumes that the surface contains a range of sites with different energies of adsorption as shown in eqn (2).

$$\ln q_e = \frac{1}{n} \ln C_e + \ln K_f \quad (2)$$

where,  $K_f$  (L mg<sup>-1</sup>) and  $n$  represent the Freundlich isotherm constant and heterogeneity factor, respectively.

The Toth model was derived from the Langmuir model, monolayer adsorption isotherm with two parameters of the affinity of adsorbate molecule to the adsorption areas ( $K_f$ ) and the concentration of adsorbed molecules ( $q_m$ ), to rectify the deviation discrepancies between experimental data and the calculated value at equilibrium. In the case of the Toth equation, three variables were considered with the additional  $n$  symbol as shown in eqn (3).

$$\frac{q}{q_m} = \frac{\alpha_T C_e}{[1 + (\alpha_T C_e)^n]^{1/n}} \quad (3)$$

where  $\alpha_T$  and  $C_e$  could be attributed to  $K_L$  and equilibrium concentration, respectively, and  $n$  is the values correlated with the heterogeneity.<sup>42</sup>

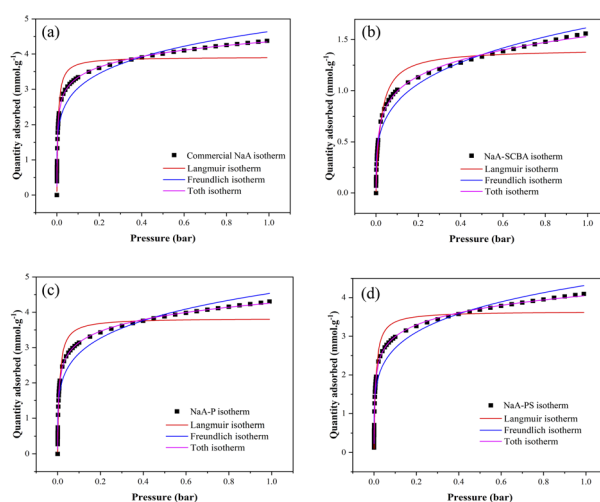


Fig. 9 CO<sub>2</sub> adsorption isotherms of commercial-NaA (a), NaA-SCBA (b), NaA-P (c), and NaA-PS (d) fitted with Langmuir, Freundlich, and Toth models.



Table 4 CO<sub>2</sub> adsorption isotherm properties of commercial-NaA, NaA-SCBA, NaA-P, and NaA-PS

Samples	Correlation coefficients ( $R^2$ )			$n$ value from the Toth model
	Langmuir isotherm	Freundlich isotherm	Toth isotherm	
Commercial-NaA	0.94969	0.94030	0.99882	0.89205
NaA-SCBA	0.96252	0.97310	0.99855	0.82932
NaA-P	0.95420	0.95021	0.99952	0.87752
NaA-PS	0.95380	0.94983	0.99960	0.87762

Fig. 9 reveals the behavior of the CO<sub>2</sub> adsorption isotherm, examined by applying the principles of the Langmuir, Freundlich, and Toth models. The correlation coefficient ( $R^2$ ) values of all samples obtained from each model are shown in Table 4. It was observed that the behavior of the CO<sub>2</sub> adsorption in all samples correlated well with the Toth isotherm with  $R^2$  of over 0.998, rather than Langmuir and Freundlich. A higher  $R^2$  value could indicate better applicability and reliability of a given model. These results also agree well with several reports in the literature<sup>43–45</sup> that the CO<sub>2</sub> adsorption behavior of zeolitic materials such as 13NaX and ZSM-5 follows the Toth model.

This behavior could be defined as an interaction between the adsorbed substances, which results in a slight growth after the plateau.<sup>46</sup> The isotherms of these NaA samples are properly

fitted by the Toth model, and it is marked that larger Al atom quantities lead to more heterogeneity, which could enhance the ability of CO<sub>2</sub> adsorption.<sup>47</sup> The highest  $n$  value and the Na/Si ratio, which could be assumed as the amount of Al, were obtained in the commercial NaA with 0.89205 and 0.63, respectively, while the lowest numbers were 0.82932 and 0.11 that were received from NaA-SCBA. Wang and coworkers<sup>47</sup> also reported with a heterogeneous interpretation that 3.30 mmol g<sup>-1</sup> of CO<sub>2</sub>/H<sub>2</sub>O adsorption was obtained when silicalite-1 was utilized as an adsorbent, followed by significantly increasing up to 3.99 mmol g<sup>-1</sup> when Al was incorporated into the synthetic preparation of ZSM-5. Except for the Na content, the tendency of CO<sub>2</sub> uptake in this work is also in line with the  $n$  values from the Toth model.

Table 5 Comparison of CO<sub>2</sub> adsorption capacity from this work to current works in the literature

Sample	Condition pretreatment	CO <sub>2</sub> adsorption capacity		Reference
		(mg g <sup>-1</sup> )	(mmol g <sup>-1</sup> )	
Zeolite NaA-SCBA	300 °C, 3 h under vacuum	68.64	1.56	This work
Zeolite NaA-P	300 °C, 3 h under vacuum	189.2	4.30	This work
Zeolite NaA-PS	300 °C, 3 h under vacuum	180.4	4.10	This work
Commercial-NaA	300 °C, 3 h under vacuum	192.7	4.38	This work
Zeolite Y	No treatment	96.0	2.18	4
Zeolite 13X <sup>a</sup>	100 °C, 1 h	154.0	3.50	57
Hydrophobic-coated zeolite 13X (HZ)	ND	149.6	3.40	58
Zelite NaX from rich husk silica	300 °C, 3 h	107.8	2.45	10
Zeolite NaA from rice husk silica	300 °C, 3 h	97.7	2.20	10
Zeolite LTA	300 °C, 3 h	135.1	3.07	51 and 59
Zeolite 4A	200 °C, 12 h under vacuum	149.2	3.39	60
Zeolite 5A	150 °C, 12 h under vacuum	161.9	3.68	61
N-rGO-ZnO	150 °C, 3 h	156.2	3.55	62
Nitrogen-doped hierarchically constructed interconnected porous carbon nanofibers derived from polyaniline (PANI)	200 °C, 2 h under vacuum	176.4	4.01	63
Activated carbon	105 °C, 10 min	72.3	1.64	64
HAC-0.5 (activated carbon)	300 °C, 3 h under vacuum	174.7	3.97	65
Chitosan-derived porous carbon (activated carbon)	150 °C, 2 h under vacuum	169.8	3.86	66
N-Doped polypyrrole-based porous carbons	200 °C and 1 bar	171.6	3.9	67
Rich nitrogen-doped ordered mesoporous carbon	200 °C, 8 h under vacuum	140.8	3.2	68

<sup>a</sup> CO<sub>2</sub> adsorption was measured at 25 °C and 1 bar in all samples, except zeolite 13X which was tested at 35 °C and 1 bar.



Besides, even if NaA-SCBA had a poor Na/Si ratio, the heterogeneity character remained high, probably as a result of interferences that raised the  $n$  value. Note that the Toth adsorption isotherm was frequently seen once activated carbon was employed as a material to remove  $\text{CO}_2$ .<sup>42,46,48</sup> Zeolite 13X initially provided a Sips  $\text{CO}_2/\text{N}_2$  isotherm,<sup>34</sup> then turned to the Toth model when 5% carbon black was included.<sup>43</sup>

Table 5 displays the comparison of the  $\text{CO}_2$  adsorption capacity of the present work to the relevant works including zeolitic, carbon, and N-dropped carbon-based materials in the literature. It was noticeable that the NaA-P provided the best  $\text{CO}_2$  adsorption capacity than other adsorbents reported in the literature despite they were tested in almost similar conditions. In terms of the zeolitic material category, the pretreatment step also plays an important role in  $\text{CO}_2$  adsorption.<sup>41,49</sup> Majchrzak-Kuceba and Nowak<sup>49</sup> have revealed that the pretreatment temperature of zeolite NaA strongly affects the  $\text{CO}_2$  adsorption capacity. Likewise, the  $\text{CO}_2$  adsorption capacity enhances with the increase in pretreatment temperature from 100 °C to 500 °C. This finding indicates that the activation temperature of the zeolite is a critical factor for the  $\text{CO}_2$  sorption process, as higher activation temperatures lead to dehydration and the creation of additional adsorption sites.

Compared to the result in the present work (Table 5), the  $\text{CO}_2$  adsorption capacity of NaA-P is the largest of other adsorbents

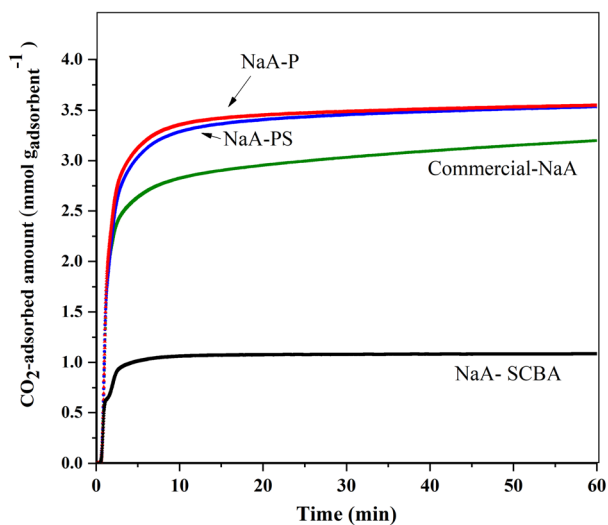


Fig. 10  $\text{CO}_2$  adsorption capacity of zeolite samples as a function of time measured at 25 °C.

could be caused by both the higher pretreatment temperature, 300 °C for 3 h under vacuum. When compared with other adsorbents such as some carbon and/or N-dropped carbon-based materials reported in the literature (Table 5), NaA-P also had the highest adsorption capacity of  $\text{CO}_2$ . The consequence might be caused by the greater basic density of NaA-P providing the impressed  $\text{CO}_2$  adsorption capacity. This result indicates the advantage of the NaA-P using silica from SCBA waste with the simplicity in the synthesis. This remark agrees well with the report in the literature<sup>1,50</sup> that zeolite NaX 13 and NaA (4A) give greater adsorption capacity of  $\text{CO}_2$  than activated carbon (AC). Moreover, zeolite NaA has more selective to  $\text{CO}_2$  adsorption than zeolite NaY and NaX.<sup>51</sup> This high selectivity of zeolite A is beneficial because  $\text{CO}_2$  needs to be separated from flue gas containing nitrogen, water, and oxygen.

Overall, this finding confirms that the NaA-P synthesized from SCBA silica is a promising adsorbent for  $\text{CO}_2$  adsorption because it provides high adsorption capacity and simplicity in the synthesis with mild conditions. Besides, the utilization of silica source material derived from SCBA is considered a cost-effective and eco-friendly approach for production, with the notable benefit for large-scale production.

Fig. 10 demonstrates  $\text{CO}_2$  adsorption kinetics of the zeolite samples as a function of time measured at 25 °C and determined by the gravimetric method. The  $\text{CO}_2$  adsorption in all samples was initially rapid before reaching equilibrium. Similar behaviors were reported on zeolite NaX and 5A in the literature.<sup>52</sup> The adsorption capacity of NaA-P and NaA-PS were nearly the same and higher than commercial NaA and NaA-SCBA.

Moreover, the adsorption kinetics of  $\text{CO}_2$  was investigated and analyzed according to the pseudo-first and pseudo-second order models reported in the literature on various zeolites.<sup>52,53</sup> The pseudo-first-order model (eqn (4)) assumes that the concentration of one reactant is much higher than the concentration of the other reactants<sup>52</sup>

$$\ln(q_e - q_t) = \ln q_e - k_1 t \quad (4)$$

where  $q_t$  ( $\text{mmol g}^{-1}$ ) is the amount of adsorbate adsorbed at the time of  $t$  (minute),  $q_e$  ( $\text{mmol g}^{-1}$ ) is the adsorption capacity at equilibrium and  $k_1$  ( $\text{min}^{-1}$ ) is the rate constant of the pseudo-first-order.

The pseudo-second-order model (eqn (5)) assumes that the availability of active sites on the adsorbent surface is always proportional to the adsorption capacity.<sup>52</sup>

Table 6 Pseudo-first and pseudo-second order kinetics model parameters of  $\text{CO}_2$  adsorption on zeolites

Adsorbent	Pseudo-first order			Pseudo-second order		
	$q_{e(\text{cal})}$ ( $\text{mmol g}^{-1}$ )	$K_1$ ( $\text{min}^{-1}$ )	$R^2$	$q_{e(\text{cal})}$ ( $\text{mmol g}^{-1}$ )	$K_2$ ( $\text{min}^{-1}$ )	$R^2$
Commercial NaA	3.41	0.0663	0.8693	3.50	0.0035	0.9984
NaA P	3.83	0.0769	0.9101	3.92	0.0049	0.9990
NaA-PS	3.76	0.0793	0.8985	3.85	0.0055	0.9983
NaA-SCBA	1.31	0.0849	0.8536	1.32	0.0453	0.9998



$$\frac{t}{1/q_t} = \frac{1}{k_2 q_e^2} + \frac{1}{q_e} t \quad (5)$$

where  $k_2$  ( $\text{min}^{-1}$ ) is the rate constant of the pseudo-second-order.

The kinetics parameters, correlation coefficient ( $R^2$ ) and adsorption capacity of all samples are summarized in Table 6. All samples correlated well with the pseudo-second-order kinetics model. The results are consistent with the adsorption kinetic model of  $\text{CO}_2$  on various zeolites including 13X, 5A, and 4A.<sup>52,53</sup>

One should note that the capacity of dynamic  $\text{CO}_2$  adsorption in commercial NaA is lower than those in NaA-P and NaA-PS, the opposite trend from the result of static adsorption isotherm measured by Autosorb IQ3 in Fig. 8. This phenomenon could be the consequence of the larger particle size in commercial NaA as shown in Fig. 5. It is reported in the literature<sup>54,55</sup> that the smaller particle size in nano-sized CHA zeolite improved the  $\text{CO}_2$  adsorption capacity when compared with the micron-sized zeolite due to the minimized diffusion resistance of  $\text{CO}_2$ .

## 3. Experimental

### 3.1 Chemicals and materials

Sugarcane bagasse ash was obtained from Mitr Phol Sugar industry, Chiyaphum Province, Thailand. Chemicals used for silica extraction and zeolite synthesis were hydrochloric acid (37%, ANAPURE®), sodium hydroxide (97% wt, Carlo Erba), sodium aluminate (50–56%  $\text{Al}_2\text{O}_3$ , 40–45%  $\text{Na}_2\text{O}$ , Sigma-Aldrich). Commercial NaA zeolite was purchased from Sigma-Aldrich (CAS number: 1318-02-1). Chemicals employed for  $\text{CO}_2$  adsorption and  $\text{CO}_2$  temperature-programmed desorption ( $\text{CO}_2$ -TPD) were  $\text{CO}_2$  gas with a purity of 99.99% and  $\text{CO}_2$  gas with 10% in He, respectively.

### 3.2 Silica extraction from sugarcane bagasse ash waste

Silica extract extraction from sugarcane bagasse ash waste was done following the method adapted from the literature.<sup>23</sup> Firstly, raw SCBA (20.00 g) was ground and sieved with particle sizes of 100  $\mu\text{m}$  and subsequently treated in HCl solution (1.0 M) with the SCBA and HCl solution ratio of 1 : 6 W/V at 30 °C for 2 h to remove some metal impurities. Then, the acid-treated SCBA was separated by centrifugation and dried at 100 °C overnight. Afterward, the silica extraction of acid-treated SCBA was done by NaOH solution treatment (1.0 M) with the ratio of acid-treated and NaOH solution = 1 : 8 W/V at 90 °C for 2 h. Then, the sodium silicate ( $\text{Na}_2\text{SiO}_3$ ) solution was separated by the paper filter for particles smaller than 11  $\mu\text{m}$  (no. 1) followed by a syringe filter for that than 0.45  $\mu\text{m}$ . Finally,  $\text{SiO}_2$  powder was obtained by precipitation of hydrated  $\text{Na}_2\text{SiO}_3$  in a mild HCl solution (1.0 M), separated using centrifugation, and dried at 100 °C overnight. The obtained silica powders from paper filter no. 1 were named  $\text{SiO}_2$ -P and that from paper filter no. 1 followed by syringe filter was named  $\text{SiO}_2$ -PS. The yield of obtained  $\text{SiO}_2$  powder was calculated according to eqn (6).

$$Y_{\text{SiO}_2} = \frac{W_{\text{SiO}_2}}{W_{\text{acid-SCBA}}} \times 100 \quad (6)$$

where  $Y_{\text{SiO}_2}$  is yield of silica powder while  $W_{\text{SiO}_2}$  and  $W_{\text{acid-SCBA}}$  are the weight of silica powder from precipitation of sodium silicate and the weight of dried acid leaching of as-received SCBA, respectively.

### 3.3 Synthesis of zeolite 4A using silica extracted from sugarcane bagasse ash waste

Zeolite 4A samples were synthesized directly from silica of as-received SBCA,  $\text{SiO}_2$ -P, and  $\text{SiO}_2$ -PS with an overall molar gel ratio of  $6\text{Na}_2\text{O} : \text{Al}_2\text{O}_3 : \text{SiO}_2 : 150\text{H}_2\text{O}$  modified from the literature.<sup>56</sup> NaOH (3.07 g) was dissolved in distilled water (22.52 g), and the NaOH solution was divided equally. The first part of the NaOH solution was mixed with sodium aluminate (1.59 g). Another part was added to the PP bottle and extracted silica from SBCA (2.00 g).

The first part was added to the PP bottle, followed by adding 5.00 g of SBCA or 2.00 g of extracted silica from SBCA ( $\text{SiO}_2$ -P and  $\text{SiO}_2$ -PS). The mixture was heated at 90 °C for 3 h and agitated magnetically. The second part was mixed with sodium aluminate (1.59 g) and stirred until clear. After that, the sodium aluminate mixture was immediately added to the sodium silicate mixture. The mixture was crystallized under hydrothermal conditions in an oven at 80 °C for 2.5 h. After cooling down, the mixture was separated using a centrifuge (4000 rpm for 10 minutes), washed with DI water several times until the washing water was neutral, and dried in an oven at 100 °C overnight. The obtained solid products were denoted as NaA-SCBA, NaA-P, and NaA-PS.

### 3.4 Characterization of silica from sugarcane bagasse ash waste

Morphology of zeolite particles was studied by scanning electron microscopy (SEM, Carl Zeiss, Auriga® series, Germany) with an accelerating voltage of 30 kV. The samples were coated with gold by sputtering. Also, elemental compositions were examined using a JEOL JSM 7800F field-emission scanning electron microscopy (FE-SEM) and scanning electron microscopy (SEM) with energy-dispersive X-ray spectroscopy (EDS). The phases of the zeolite samples were determined by X-ray diffraction (XRD, Bruker D8 ADVANCE, Germany) with Cu  $K\alpha$  radiation. Elemental compositions of samples were determined by X-ray fluorescence spectroscopy (ED-XRF, Horiba 5200).

Fourier-transform infrared spectroscopy (FT-IR) using a Bruker Tensor 27 in Attenuated Total Reflectance (ATR) was used to detect the functional groups of SCBA and NaA zeolites. Moreover, their thermal decompositions were studied by using thermogravimetric analysis (TGA, Mettler TGA/DSC1). Each sample (10 mg) was heated from room temperature to 800 °C at a rate of 10 °C  $\text{min}^{-1}$  under an air zero with a flow rate of 50  $\text{mL min}^{-1}$ .

$\text{CO}_2$  temperature-programmed desorption ( $\text{CO}_2$ -TPD) on a BELSORP-mini<sup>-1</sup> was used to assess the basicity of each adsorbent. Each sample (0.050 g) was packed in a quartz U-tube reactor before being processed at 500 °C for 1 h under a He flow rate of 50  $\text{mL min}^{-1}$  and cooled to 70 °C under He flows. The system was then filled with  $\text{CO}_2$  (10%) diluted in He at a total



flow rate of 50 mL min<sup>-1</sup> for 30 min to allow for adsorption. Then, the temperature was raised to 100 °C and the system was purged for 1 h to eliminate the physisorbed CO<sub>2</sub>. Finally, the temperature was raised from 100 to 350 °C at a rate of 10 °C min<sup>-1</sup>. The amount of CO<sub>2</sub> absorbed in mmol of CO<sub>2</sub> was measured online by a TCD detector compared with the peak area of the known amount of CO<sub>2</sub> from the calibration curve of the injection loop. The basicity in mmol<sub>CO<sub>2</sub></sub> g<sub>adsorbent</sub><sup>-1</sup> was calculated following the eqn (7).

$$\text{Basicity (mmol}_{\text{CO}_2} \text{ g}^{-1}) = \frac{\text{CO}_2 \text{ adsorbed (mmol) of sample}}{\text{sample weight (g)}} \quad (7)$$

### 3.5 CO<sub>2</sub> adsorption isotherms of zeolite NaA synthesized from SCBA waste

The CO<sub>2</sub> adsorption was tested following the method in the literature.<sup>10</sup> Briefly, each sample was degassed under vacuum at 300 °C for 3 h to remove physisorbed water, and then the adsorption process was achieved at 25 °C and the pressure range of 0.0–1.0 bar by using a Quantachrome Autosorb IQ3 analyzer. After that, the CO<sub>2</sub>-adsorbed amount was plotted against pressure to compare the adsorption isotherm. The CO<sub>2</sub> adsorption capacity (mmol) of each adsorbent was calculated at the saturated point of 1 bar and 25 °C by changing the adsorbed volume of CO<sub>2</sub> to mmol, respectively. The CO<sub>2</sub> adsorption capacity (mmol g<sub>adsorbent</sub><sup>-1</sup>) was calculated according to the eqn (8).

$$\text{CO}_2 \text{ adsorption capacity (mmol g}_{\text{adsorbent}}^{-1}) = \frac{\text{volume adsorbed of CO}_2(\text{cm}^3) \text{ in sample}}{\text{volume adsorbed of 1 mol}_{\text{CO}_2@25^\circ\text{C}} (\text{cm}^3 \text{ mol}^{-1}) \times \text{sample weight (g)}} \quad (8)$$

### 3.6 CO<sub>2</sub> adsorption kinetics of zeolite NaA synthesized from SCBA waste

The CO<sub>2</sub> adsorption kinetics was measured by Gravimetric measurement adapted from the literature.<sup>69</sup> Before the adsorption measurement, each sample (20 mg) was degassed at 300 °C for 1 h at the flow rate of 30 mL min<sup>-1</sup> in N<sub>2</sub>. The adsorption process was performed at temperatures of 25 °C with a pressure of 1 bar. The flow rate of pure CO<sub>2</sub> (99.99%) during the adsorption process was maintained at 30 mL min<sup>-1</sup> for 1 h to ensure that the adsorption process reached equilibrium. Changes in mass were recorded by using thermogravimetric analysis (TGA) on Mettler Toledo TGA/DSC 3+. The CO<sub>2</sub> adsorption capacity (mmol g<sub>adsorbent</sub><sup>-1</sup>) was calculated following eqn (9).

$$\text{CO}_2 \text{ adsorption capacity (mmol g}_{\text{adsorbent}}^{-1}) = \frac{M_t - M_0}{44 (\text{g mol}^{-1}) \times M_0} \quad (9)$$

where  $M_t$  and  $M_0$  are the change in mass of the sample at time  $t$  (min) and the mass of the sample after pretreatment (before CO<sub>2</sub> adsorption).

## 4. Conclusions

The development of an alternative silica extraction from sugarcane bagasse ash (SCBA) waste and utilization in NaA zeolite synthesis was accomplished by etching in NaOH solution to produce Na<sub>2</sub>SiO<sub>3</sub>. Then, the Na<sub>2</sub>SiO<sub>3</sub> solution was separated either by paper filter No. 1 only or paper filter No. 1 and a syringe filter (0.45 μm). SiO<sub>2</sub> powder was obtained by the precipitation of those filtrates in HCl solution (1.0 M). The SiO<sub>2</sub> purities obtained from the paper filter (SiO<sub>2</sub>-P, 98.1 wt%) and paper filter followed by syringe filter (SiO<sub>2</sub>-PS, 97.9 wt%) were significantly improved when compared with that from the as-received SCBA (77.2 wt%). Pure zeolite NaA (4A) was successfully synthesized from both SiO<sub>2</sub>-P and SiO<sub>2</sub>-PS. Meanwhile, mixed phases between zeolite NaA (5A) and quartz were obtained from the synthesis using SiO<sub>2</sub>-raw SCBA. From adsorption studies, the CO<sub>2</sub> uptake was in the following order: commercial-NaA ~ NaA-P > NaA-PS > NaA-SCBA, with respect to the amount of total basicity suggested by CO<sub>2</sub>-TPD. The CO<sub>2</sub> adsorption behavior of all samples fits well with the Toth model. The CO<sub>2</sub> adsorption kinetics correlates with the pseudo-second-order model. Moreover, the CO<sub>2</sub> adsorption capacity of zeolite NaA-P in the present work was the largest of the reported values in the relevant literature. The research confirmed that the method to extract silica from SCBA was practical, affordable,

and environmentally benign, and might be beneficial for large-scale manufacturing processes in the future.

## Author contributions

Chalermpan Keawkumay: conceptualization, methodology, investigation, formal analysis, data curation, writing-original draft preparation, and writing-review and editing. Panot Krukkratoke: methodology, formal analysis, investigation, data curation, writing-original draft preparation, and writing-review and editing. Saran Youngjan; formal analysis and investigation. Nattawut Osakoo: visualization, validation, methodology, investigation, formal analysis, data curation, writing-original draft preparation, and writing-review and editing. Krittanun Deekamwong: methodology and writing-review and editing. Pongtanawat Khemthong: formal analysis and investigation. Jakkapop Phanthasri: formal analysis and investigation. Sanchai Prayoonpokarach: conceptualization, supervision, validation, and writing-review and editing. Jatuporn Wittayakun:



conceptualization, supervision, validation, and writing-review and editing. All authors have read and agreed to the published version of the manuscript.

## Conflicts of interest

There are no conflicts to declare.

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