



Cite this: *RSC Adv.*, 2019, 9, 28916

Natural rubber latex foam with particulate fillers for carbon dioxide adsorption and regeneration

Krittaya Panploo,^a Benjapon Chalermssinuwana ^{ab} and Sirilux Poompradub ^{abc}

To reduce the carbon dioxide (CO₂) concentration in the atmosphere, natural rubber (NR) was developed as a rubber foam for CO₂ adsorption. Although the CO₂ adsorption capacity of the NR latex foam produced by mixing with a cake mixer (CM) was higher than that produced with an overhead stirrer (OS), both capacity values were still low. To improve the CO₂ adsorption capacity, the use of unmodified and (3-aminopropyl) triethoxysilane-modified silica particles as fillers in the CM rubber foam matrix was examined. The highest CO₂ adsorption capacity, from a mixed gas flow rate of 100 mL min⁻¹ at ambient temperature and pressure, was obtained with the CM foam filled with 5 parts by weight per hundred parts of rubber filled with modified silica particles (4.08 mg g⁻¹). The CO₂ adsorption capacity of this foam was approximately 1.11- and 2.87-fold higher than that of the CM foam filled with unmodified silica particles (3.69 mg g⁻¹) and unfilled CM rubber (1.42 mg g⁻¹), respectively. Morphological analysis supported that the cell size and number of pores per cell of the NR latex foam, which were higher in the CM foams than the OS foams, were important factors for evaluating CO₂ adsorption. In addition to physisorption between CM and CO₂, chemisorption between the modified silica particles and CO₂ increased the CO₂ adsorption capacity.

Received 2nd August 2019
 Accepted 6th September 2019

DOI: 10.1039/c9ra06000f

rsc.li/rsc-advances

1. Introduction

The atmospheric carbon dioxide (CO₂) concentration has increased continuously over the last century and has become a worldwide issue, being a major contributor to global climate change. Recently, CO₂ capture and storage methods have been considered as a key option for the mitigation of the impact of increasing CO₂;¹⁻³ these approaches include absorption,^{4,5} adsorption,⁶⁻⁸ cryogenic methods,^{9,10} membrane separation,¹⁰⁻¹² and CO₂ reduction and utilization.^{13,14} Among these, the removal of CO₂ from CO₂-containing streams in industrial processes is considered to be effective. Solid sorbents are usually used, such as metal oxides (calcium oxide, magnesium oxide, and lithium zirconate); however, these sorbents require high temperature for both CO₂ adsorption and regeneration of the adsorbent.¹⁵⁻¹⁷ However, the possibility to capture CO₂ directly from the atmosphere is of outstanding importance for the future development of this technology. Therefore, in this study, an alternative material is presented to potentially replace the current metal oxide adsorbents.

Microporous organic polymers (MOPs)^{18,19} are a relatively new class of porous materials that have a large surface area, small pore size, and low skeletal density.²⁰ This type of material

has already been used for various applications, such as gas storage, gas separation, catalysis, sensing, clean energy, CO₂ capture and so on.²¹⁻²⁵ However, to make MOPs eco-friendlier, the development of natural polymers is an alternative choice for CO₂ capture.

Natural rubber (NR) is a natural commodity that has tremendous economic and strategic importance due to its unique characteristics such as a high strength, flexibility, and elasticity.²⁶ In addition, the NR latex foam can be used as a good absorbent material, including as a chemical,²⁷ sound²⁸ and oil^{29,30} absorbent. However, in general, rubber products cannot be used at high temperatures due to their low thermal properties. Silica is a reinforcing filler that is widely used in rubber products,^{31,32} since it can improve not only the mechanical but also the thermal properties of rubber products,³³ and it has been reported to be an important physisorption material for CO₂ adsorption.^{34,35} However, the selectivity of CO₂ adsorption is low due to the absence of active sites for CO₂ capture. Therefore, amine-modified silica was investigated in this study to enhance the adsorption capacity and selectivity for CO₂.

Based on the literature studies, NR latex foam is not widely used as a gas adsorbent especially for CO₂. This is an importance of this work to present the unique properties of NR for CO₂ capture. Accordingly, the originality of this work was the preparation of an alternative material for CO₂ capture using NR, because NR can be easily formed into a highly porous foam for trapping gases or liquids. Moreover, NR is an eco-friendly material, compared to conventional solid sorbents, and can be used in various applications at ambient temperature and

^aDepartment of Chemical Technology, Faculty of Science, Chulalongkorn University, Bangkok 10330, Thailand. E-mail: sirilux.p@chula.ac.th

^bCenter of Excellence on Petrochemical and Materials Technology, Chulalongkorn University, Bangkok 10330, Thailand

^cGreen Materials for Industrial Application Research Unit, Faculty of Science, Chulalongkorn University, Bangkok 10330, Thailand



pressure. CO₂ is generally released from diverse sources, such as vehicles, industrial factories, or everyday life of humans. Thus, NR has potential as a solid sorbent in the preparation of CO₂ capture materials. In contrast, conventional CO₂ capture materials are typically developed as solid states that are suitable only for use at high temperatures. Until now, there have been only a few reports on the use of NR composite materials as potential CO₂ adsorbents,³⁶ and a mechanism for CO₂ adsorption in NR composite materials is still required. Accordingly, the aim of this study was to develop an eco-friendly material for CO₂ adsorption at ambient temperature and pressure. The NR latex foam was prepared using either an overhead stirrer or cake mixer, designated as OS or CM, respectively. To improve the CO₂ adsorption performance of the CM, it was filled with either unmodified silica (CM-USi) or (3-aminopropyl)triethoxysilane (APTES)-modified silica (CM-MSi). The MSi particles were characterized by Fourier-transform infrared spectroscopy (FT-IR) and transmission electron microscopy (TEM). The morphology of the OS, CM, CM-USi, and CM-MSi foams were also studied by scanning electron microscopy (SEM). The effects of the mixed gas (12.88 (v/v) CO₂/nitrogen (N₂)) flow rate on the CO₂ adsorption performance of the different NR latex foams with or without the USi or MSi filling were investigated using a stainless-steel reactor at ambient temperature and pressure. Finally, the mechanism for CO₂ adsorption by the different NR latex foams is proposed.

2. Experimental

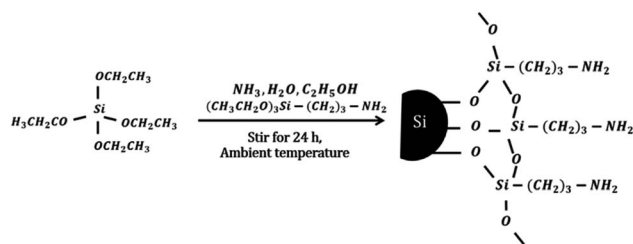
2.1. Materials

NR latex with a 60% dry rubber content was used as a raw material. The chemicals for rubber compounding, such as potassium oleate (K-oleate) as a surface active agent, sulfur as a crosslinking agent, zinc diethylthiocarbonate (ZDEC) as an accelerator, zinc 2-mercaptobenzothiazone (ZMBT) as an accelerator, Wingstay L (butylated reaction product of *para* cresol and dicyclopentadiene) as an antidegradant, diphenyl guanidine (DPG) as an accelerator, zinc oxide (ZnO) as an activator, and sodium silicofluoride (SSF) as an gelling agent, were purchased from the Rubber Research Institute, Bangkok, Thailand and dispersed in water. Tetraethoxysilane (TEOS), used as a silica precursor, and APTES, used as a modifying agent, were from Sigma-Aldrich (Saint Louis, USA). Ammonia solution (28 wt%), used as a basic catalyst, and absolute ethanol (EtOH), used as a solvent, were purchased from Qrec Chemical Ltd. (Chonburi, Thailand). All chemical reagents were used as received without purification.

2.2. Preparation of MSi particles

The MSi particles were prepared by mixing 0.013 mol TEOS, 0.702 mol EtOH, 0.052 mol ammonia solution, 0.389 mol deionized water and 0.007 mol APTES at ambient temperature and stirring for 24 h.³³ For preparation of the USi particles, APTES was not added to the solution. The obtained USi or MSi particles were then separated by centrifugation, washed in deionized water, and dried in an oven at 110 °C. The idealized

mechanism for silica modification by APTES is shown schematically below.



2.3. Rubber foam preparation

Table 1 shows the formulation of the NR compound. First, the K-oleate dispersion was added to the NR latex and beaten for 10 min by either CM or OS to make a foam as shown in Fig. 1. The rotation speeds of CM and OS were at 100 and 500 revolutions per minute (rpm). Then the sulfur, ZDEC, ZMBT, Wingstay L, DPG, ZnO and SSF dispersions were slowly added and continuously beaten for 2 min. The total NR compound volume after mixing by CM increased five-fold, while that by OS increased two-fold compared to the initial volume. The rubber compound was then poured into a glass mold and cured in a hot-air oven at 100 °C for 2 h to cure, washed with water to eliminate the unreacted chemicals and dried in a hot-air oven at 70 °C for 20 h. For preparation of the filled CM composite foam, USi or MSi particles were initially mixed with the K-oleate, and thereafter the same procedure was performed as above to prepare the CM-USi or CM-MSi composites, respectively.

2.4. Characterization

The functional groups of the MSi particles were characterized by FT-IR (PerkinElmer, Shelton, USA) analysis. Each sample was prepared in a KBr disk under high pressure, and the FT-IR

Table 1 Formulation for the NR latex foam compound^g

Ingredient	CM ^a (phr) ^b	OS ^c (phr)	CM-USi (phr)	CM-MSi (phr)
60% DRC NR latex ^d	100	100	100	100
10% K-oleate dispersion	15	15	15	15
50% sulfur dispersion	4	4	4	4
50% ZDEC dispersion	2	2	2	2
50% ZMBT dispersion	2	2	2	2
50% Wingstay L dispersion	2	2	2	2
33% DPG dispersion	2	2	2	2
50% ZnO dispersion	10	10	10	10
12.5% SSF dispersion	8	8	8	8
USi ^e	—	—	5	—
MSi ^f	—	—	—	5

^a NR latex foam prepared by cake mixer. ^b Parts by weight per hundred parts of rubber. ^c NR latex foam prepared by overhead stirrer. ^d All chemicals are calculated based on 60% dry rubber content (DRC) of the NR latex. ^e Unmodified silica particles. ^f Modified silica particles by (3-aminopropyl)triethoxysilane (APTES). ^g Conditions: the rubber compound was mixed at ambient temperature and cured for 2 h at 100 °C.



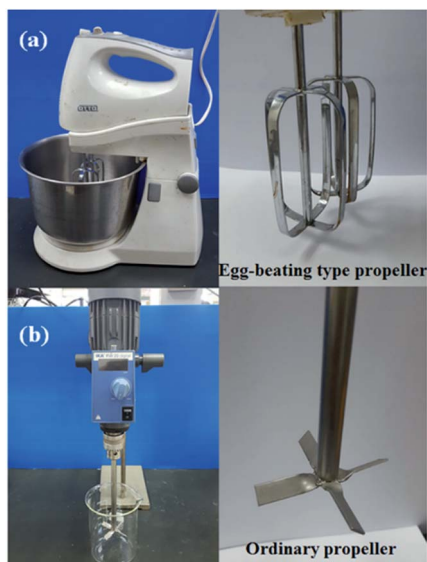


Fig. 1 The feature of the (a) cake mixer with egg-beating type propeller and (b) overhead stirrer with ordinary propeller.

spectra were recorded from 4000 to 400 cm^{-1} at a resolution of 4 cm^{-1} . The carbon, hydrogen, and nitrogen contents on the silica surface were examined by a PerkinElmer (PE-2400, Waltham, USA) CHNS/O elemental analyzer. The morphology of the MSi particles was examined by TEM (FEI TECNAI T20, Eindhoven, Netherland), operated at 120 kV. The samples were suspended in EtOH and then dropped onto a copper grid to dry at room temperature. The morphology of each NR latex foam was also studied by SEM (JEOL, JSM-6480LV, Tokyo, Japan) at an accelerating voltage of 15 kV. The samples were placed on a stub and coated with gold before measurement.

2.5. CO₂ adsorption-desorption performance

Fig. 2 shows a schematic representation of the experimental apparatus used to determine the CO₂ adsorption. In order to eliminate any physisorbed or chemisorbed CO₂ in the sorbent materials, the dried sorbent material (2.0 g) was treated under

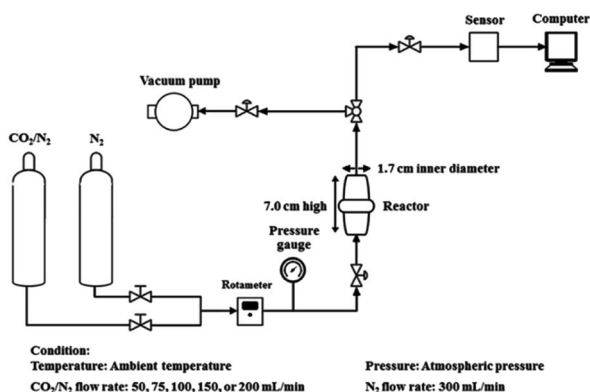


Fig. 2 Schematic representation of the CO₂ adsorption apparatus used in this study.

vacuum at 60 °C, packed into a stainless-steel reactor (1.7 cm inner-diameter and 7.0 cm-height), and treated under vacuum followed by a N₂ flow (300 mL min⁻¹). The gas flow was then switched to a mixed gas (12.88 : 87.12 (v/v) CO₂ : N₂) at various flow rates (75, 100, 150 and 200 mL min⁻¹). The CO₂ adsorption capacity was calculated based on the difference between the total inlet and outlet CO₂ amounts. The mean residence time was determined by flowing the mixed gas (12.88 : 87.12 (v/v) CO₂ : N₂) through the stainless steel reactor without any sorbent. Therefore, the CO₂ adsorption capacity was directly proportional to the area difference above the CO₂ breakthrough curves. The adsorption process was operated at atmospheric pressure and ambient temperature. The released CO₂ from the reactor was measured *via* an online sensor (K33 BLG sensor, Florida, USA).

The regeneration of the sorbent material was performed by a combination of vacuum and temperature swing adsorption processes in the stainless-steel reactor. The sorbent material was heated at 60 °C under a reduced pressure condition with a N₂ flow rate of 10 mL min⁻¹ and then cooled down to ambient temperature. After that, the sorbent material was subjected to a repeat CO₂ adsorption process. This CO₂ adsorption-desorption was continued for 12 cycles in this study.

3. Results and discussion

3.1. Characterization of the MSi particles

Representative FT-IR spectra of the USi and MSi particles are shown in Fig. 3. The O–OH stretching of moisture and SiO–H stretching of USi appeared at 3434 cm^{-1} .^{37–39} Furthermore, the absorption band corresponding to the deformation vibrations of the adsorbed water molecules appeared at 1630 cm^{-1} , while the Si–O in-plane stretching vibrations of the silanol Si–OH group appeared at 948 cm^{-1} .^{37–39} Additionally, the characteristic peaks of the Si–O–Si asymmetric, symmetric, and bending modes of silica were shown at 1107, 802, and 474 cm^{-1} , respectively.⁴⁰

After the modification of the silica surface with APTES to yield the MSi particles, additional absorption bands were observed at 2945 cm^{-1} and 1480 cm^{-1} , which represent the C–H

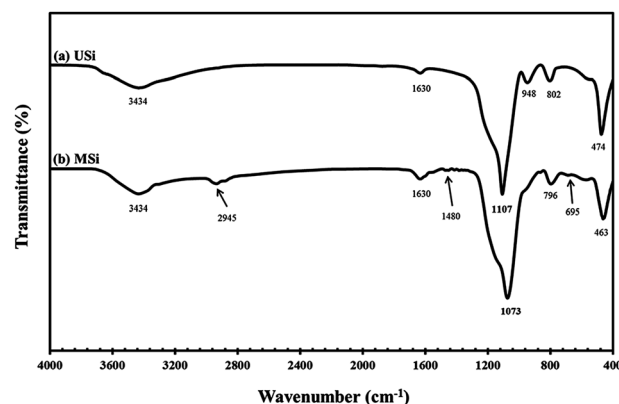


Fig. 3 Representative FT-IR spectra of the (a) USi and (b) MSi particles.



Table 2 The elemental analysis (by weight) and particle size of the unmodified (USi) and modified (MSi) silica particles

Sample	Elemental analysis ^a (wt%)			Particle size ^b (nm)
	C	H	N	
USi	0.00	1.09	0.00	176 ± 23
MSi	10.88	3.28	3.98	208 ± 36

^a By CHN analysis. ^b By TEM analysis.

stretching vibration⁴¹ and NH₂ deformation of hydrogen-bonded amine groups, respectively.⁴² The band at 695 cm⁻¹ was attributed to the Si–C stretching vibration of the methyl group bonded to a silicon atom.⁴³ However, the –NH₂ groups at 3300–3800 cm⁻¹ were not evident in the MSi particles due to the interference of moisture or water molecules.⁴⁴ Table 2 shows the elemental analysis of the USi and MSi particles, where the USi particles showed a hydrogen content of only 1.09 wt%, which came from the water adsorbed on the USi particles' surface. After modification, the C, H, N contents were increased,

supporting that the modification of the silica particles by APTES was completely achieved.

Fig. 4 shows the morphology of the USi and MSi particles as observed by TEM. The USi and MSi particles were of a uniform spherical shape, with average anhydrous particle sizes of about 176 ± 23 nm and 208 ± 36 nm, respectively. The fact that the size of the MSi particles was larger (*ca.* 1.18-fold) than that of the USi particles was due to the organic group of the APTES modifier covering the silica surface.

3.2. Characterization of the different NR (OS and CM) foams

The NR latex foams were prepared by OS and CM in order to evaluate the effect of the NR latex foam morphology on the CO₂ adsorption performance. The morphologies of the OS and CM rubber foams are shown in Fig. 5. Both types of the NR latex foam showed a spherical-shaped open cell structure, but the cell size of the OS foam (78 ± 29 μm) was about 4.3-fold smaller than that of the CM foam (338 ± 33 μm). It is interesting to note that the number of pores per cell in the CM foam was more than that in the OS foam. Moreover, the cells in the OS foam were generally interconnected with each other, which may have decreased the available surface area of the sorbent material for CO₂ adsorption. In contrast, the cells of the CM foam showed a well-defined cell wall⁴⁵ and a large capacity for potential CO₂ capture. The morphology of the CM foam after filling with USi or MSi particles was similar to that of the neat CM foam (Fig. 5(c) and (d)).

3.3. CO₂ adsorption performance

In order to test the CO₂ adsorption performance of the different prepared NR latex foams, they were evaluated in a mixed gas flow in a stainless-steel reactor at ambient temperature and pressure. The effects of the mixed gas flow rate on the CO₂ adsorption capacity of the different NR latex foams were evaluated. The CO₂ adsorption capacity of the OS and CM foams increased when the mixed gas flow rate was increased from 75 to 100 mL min⁻¹ (Fig. 6), which was due to the increased driving force of the mixed gas through the NR latex foam. However, the CO₂ adsorption capacity of both types of the NR latex foam gradually decreased as the mixed gas flow rate increased above 100 mL min⁻¹. This may be because at a higher mixed gas flow rate a shorter residence time was obtained.⁸ Interestingly, the CO₂ adsorption capacity of the CM foam was higher than that of the OS foam at all mixed gas flow rates, which was presumably due to the higher number of pores per cell able to capture more CO₂, whereas the OS foam had a smaller cell size with a lower number of pores per cell (Fig. 7). The only interaction between CO₂ and the different NR latex foams was physisorption. Accordingly, in this study, the CM foam was chosen to evaluate the effect of the addition of USi or MSi fillers in an attempt to improve the CO₂ adsorption capacity.

Fig. 8 shows the breakthrough curves of the CM-USi and CM-MSi foams. A breakthrough curve shows the relationship between the ratios of the CO₂ concentration at the outlet to the initial CO₂ concentration at the inlet of the reactor (*C/C*₀) over time. Initially, no CO₂ was detected in the outlet gas because the

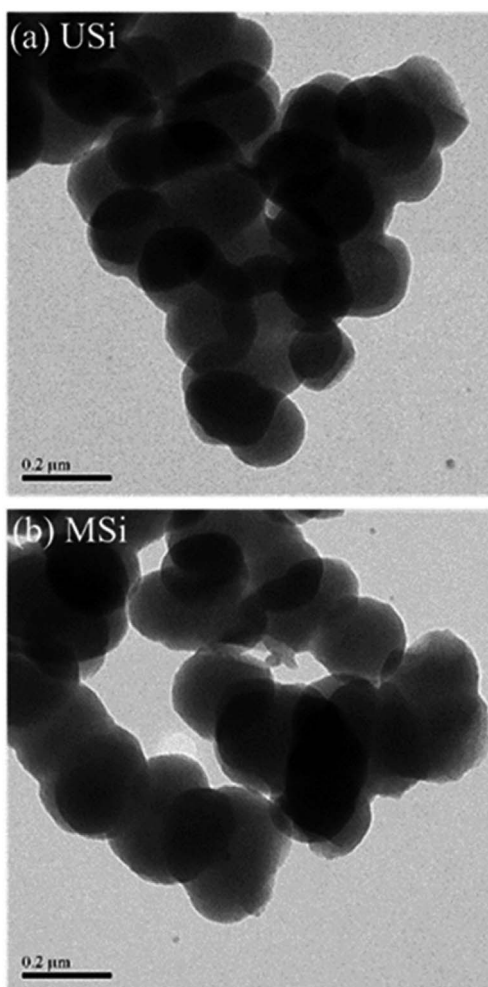


Fig. 4 Representative TEM micrographs (19 000× magnification; scale bar = 0.2 μm) of the (a) USi and (b) MSi particles.



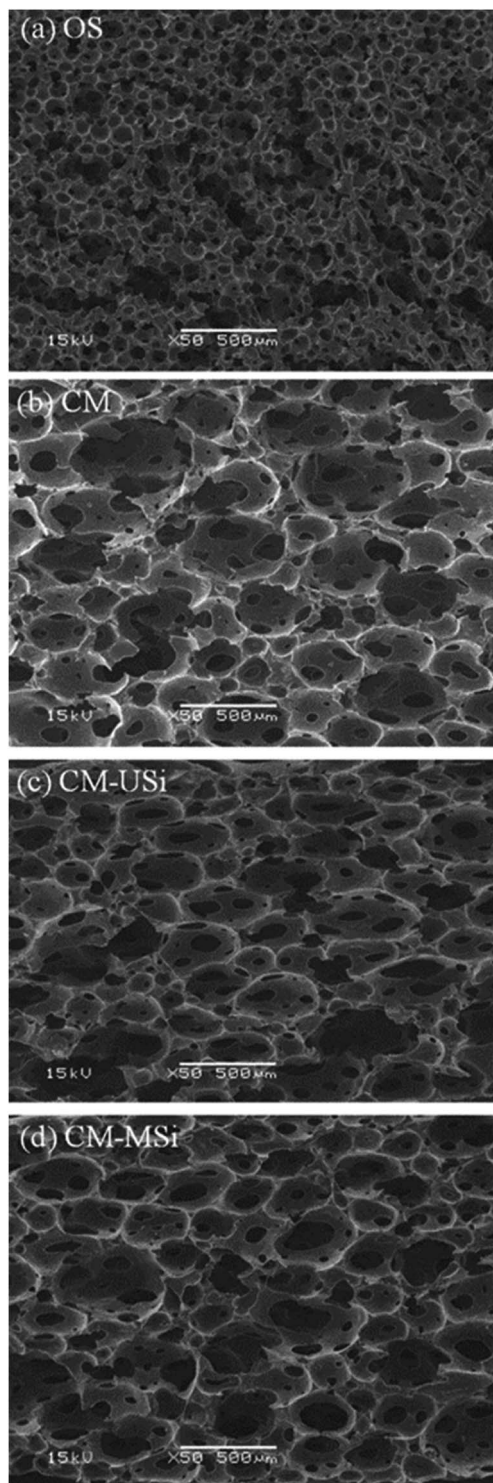


Fig. 5 Representative SEM micrographs (50 000 \times magnification; scale bar = 500 μ m) of the NR latex foam formed by (a) OS and (b) CM, and of the CM foam filled with either (c) USi or (d) MSi particles by different techniques.

inlet CO_2 was completely adsorbed on the respective NR latex foam. The breakthrough time (the time at which the effluent CO_2 concentration reaches a 10% allowable breakthrough concentration⁴⁶) increased when the CM foam was filled with

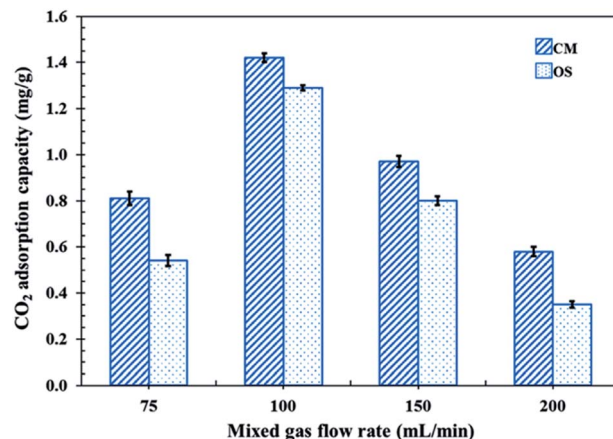


Fig. 6 CO_2 adsorption capacity of the CM and OS foams at different mixed gas flow rates (50, 75, 100, 150 and 200 mL min^{-1}) under ambient temperature and pressure.

either USi or MSi particles, and was ranked in the order CM-MSi > CM-USi > CM. Thereafter, the curves slowly and continuously increased, since the CO_2 adsorption performance slowly decreased, resulting in an increasing CO_2 concentration in the outlet gas. Finally, the curves became constant at 1.0 (outlet CO_2 concentration equals that of the inlet) because the adsorbents were saturated with CO_2 . The total CO_2 adsorption capacity was

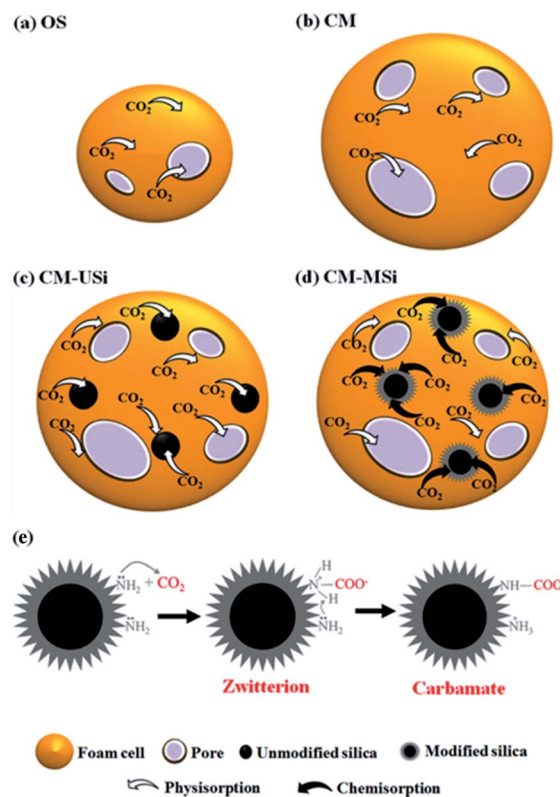


Fig. 7 Proposed mechanisms of CO_2 adsorption for the (a) OS, (b) CM, (c) CM-USi and (d) CM-MSi foams, and (e) chemisorption between CO_2 and modified silica in CM-MSi foam.



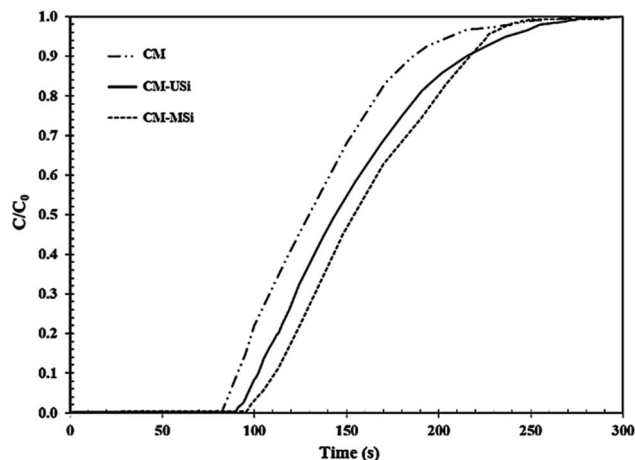


Fig. 8 Breakthrough curves of the NR latex foam, and the rubber composite of the CM foam filled with either USi (CM-USi) or MSi (CM-MSi) particles at a mixed gas flow rate of 100 mL min^{-1} under ambient temperature and pressure.

calculated from the area above the breakthrough curves and is shown in Table 3. The CO_2 adsorption capacity of the CM-MSi foam was clearly the highest followed by that of the CM-USi foam and these were 2.87 and 2.6-fold higher, respectively, than that of CM foam. These results indicated that the high adsorption capacity of the CM-MSi foam was due not only to physisorption between CO_2 and the CM foam, but also to chemisorption between CO_2 and the amine groups to form a carbamate on the surface of the MSi particles,^{47,48} whereas the only interaction between CO_2 and the CM-USi foam was physisorption (Fig. 7).

Fig. 9 shows the CO_2 adsorption–desorption of CM, CM-USi and CM-MSi foams over 12 successive cycles. The CO_2 adsorption capacity of the CM foam dropped 27.9% after five cycles of regeneration, while the CO_2 adsorption capacity of CM-USi and CM-MSi foams decreased by 11.9 and 12.4% after 10 and 9 cycles of regeneration, respectively. This suggested that CM-USi and CM-MSi foams exhibited a better regenerative ability than the CM foam. Accordingly, the presence of USi or MSi particles gave a higher stability to reuse the sorbent

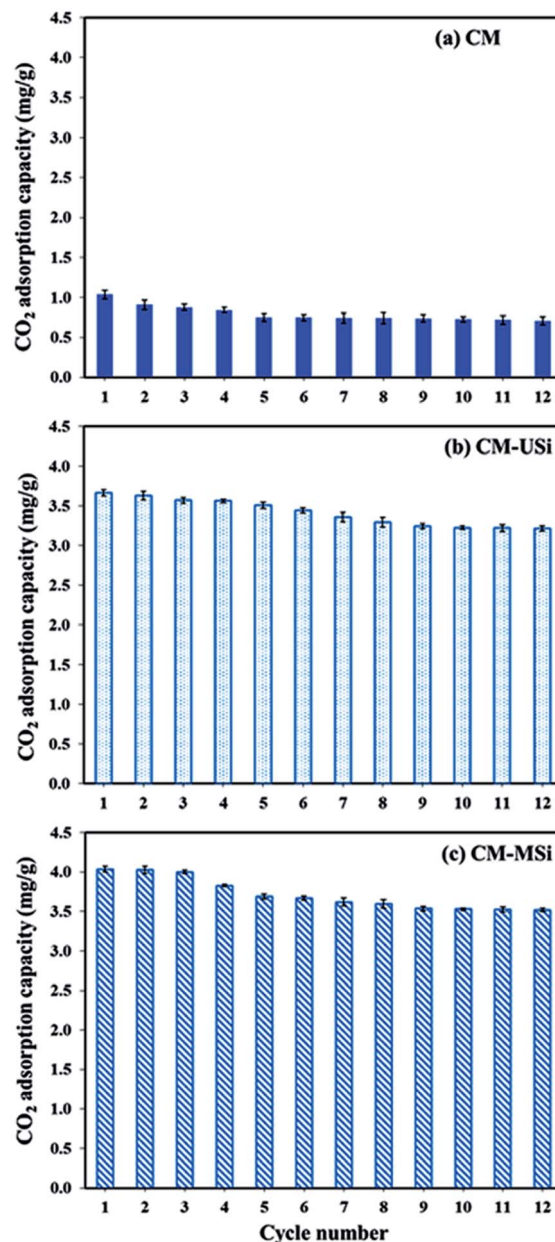


Fig. 9 The CO_2 uptake of (a) CM, (b) CM-USi and (c) CM-MSi foams over 12 adsorption–desorption cycles; adsorption: a mixed gas flow rate of 100 mL min^{-1} under ambient temperature and pressure; desorption: $60 \text{ }^\circ\text{C}$ under vacuum for 20 min.

Table 3 CO_2 adsorption capacity of the different NR latex foams (OS and CM) and the CM foam filled with either USi or MSi particles, at a mixed gas flow rate of 100 mL min^{-1} under ambient temperature and pressure

Sample	CO_2 adsorption capacity ^a (mg g^{-1})
OS ^b	1.29 ± 0.1
CM ^c	1.42 ± 0.3
CM-USi ^d	3.69 ± 0.2
CM-MSi ^e	4.08 ± 0.2

^a mg of CO_2 per g of adsorbent. ^b NR latex foam prepared by overhead stirrer. ^c NR latex foam prepared by cake mixer. ^d NR latex foam prepared by cake mixer filled with unmodified silica particles. ^e NR latex foam prepared by cake mixer filled with silica particles modified by (3-aminopropyl)triethoxysilane (APTES).

materials compared to the CM foam. However, when comparing the overall regeneration performance of all the sorbents, the CO_2 adsorption capacity dropped to constant values without a further decrease in the CO_2 adsorption capacity. This shows the advantage when using these sorbent materials in a real operation.

Table 4 shows the comparison of CO_2 adsorption capacities of various sorbent materials. Each sorbent material showed the different CO_2 adsorption capacities due to the different surface properties, nature of sorbent material (porous powder, film, or foam) or testing condition. When comparing the capture



Table 4 Comparison of CO₂ adsorption capacities of various sorbent materials

Sorbent materials	Temperature (°C)	Pressure (bar)	CO ₂ adsorption capacity (mg g ⁻¹)	Ref.
Polyamide-6	25	1.0	1.00	49
PA/CNT-PEI-100 ^a	25	1.0	10.0	49
MCM-41	30	0.1	5.28	50
S12 ^b	40	1.0	9.24	51
N-OMPs ^c	25	0.15	29.5	52
CM-MSi ^d	Ambient	1.0	4.08	This study

^a Polyamide-6/carbon nanotube composite impregnated with polyethyleneimine (100 wt%). ^b Nano silica particles with the particle size of 12 nm. ^c Nitrogen-doped ordered mesoporous polymers. ^d NR latex foam prepared by cake mixer filled with silica particles modified by (3-aminopropyl) triethoxysilane (APTES).

capacity of various kinds of sorbent material, CM-MSi can be a candidate for using as a solid sorbent because the adsorbent can be prepared as the green and non-toxic adsorbent material. In addition, this material still can modify to further increase the CO₂ capture capacity.

4. Conclusions

In this study, NR-based composite materials were designed and developed for CO₂ capture at ambient temperature and pressure. Based on a SEM analysis, the CM foam showed a larger cell size with many pores per cell, resulting in a higher CO₂ adsorption capacity compared to the OS foam. The optimum condition to obtain the highest CO₂ adsorption capacity of the CM foam (1.42 mg g⁻¹) was the use of the mixed gas (12.88 : 87.12 (v/v) CO₂ : N₂) at a flow rate of 100 mL min⁻¹ at ambient temperature and pressure. To further improve the CO₂ adsorption performance of the CM foam, USi or MSi particles were filled in the rubbery matrix, whereupon the CO₂ adsorption capacity increased by more than 2.6 and 2.87-fold, respectively, compared to the unfilled CM foam. The high adsorption capacity of CM-MSi was due to the presence of both physical and chemical adsorption, while the interaction of CO₂ and CM-USi represented only physical adsorption. Accordingly, the CM-MSi composite foam can be used as a solid sorbent or designed for CO₂ capture in high CO₂ release areas, such as gas stations, car parks, traffic jams or hospitals.

Conflicts of interest

The authors have declared no conflict of interest.

Acknowledgements

The authors gratefully acknowledge the funding support from the Thailand Research Fund (Grant No. RSA6180030) and Malaysia-Thailand Joint Authority (MTJA), Research Cess Fund.

References

1 D. Leasona, N. M. Dowell, N. Shaha, C. Petit and P. S. Fennell, *Int. J. Greenhouse Gas Control*, 2017, **61**, 71–84.

- C. H. Yu, C. H. Huang and C. S. Tan, *Aerosol Air Qual. Res.*, 2012, **12**, 745–769.
- D. M. D'Alessandro, B. Smit and J. R. Long, *Angew. Chem., Int. Ed.*, 2010, **49**, 6058–6082.
- J. De Rivaa, J. Suarez-Reyes, D. Morenoa, I. Díaz, V. Ferroa and J. Palomar, *Int. J. Greenhouse Gas Control*, 2017, **61**, 61–70.
- H. Kim, H. Shi and J. Y. Lee, *Int. J. Greenhouse Gas Control*, 2016, **45**, 181–188.
- L. Hauchhum and P. Mahanta, *Int. J. Energy Environ. Eng.*, 2014, **5**, 349–356.
- K. M. Lee, Y. H. Lim, C. J. Park and Y. M. Jo, *Ind. Eng. Chem. Res.*, 2012, **51**, 1355–1363.
- X. Wang, L. Chen and Q. Guo, *Chem. Eng. J.*, 2015, **260**, 573–581.
- C. F. Song, Y. Kitamura, S. H. Li and K. Ogasawara, *Int. J. Greenhouse Gas Control*, 2012, **7**, 107–114.
- B. Belaissaoui, Y. L. Moullec, D. Willson and E. Favre, *J. Membr. Sci.*, 2012, **415–416**, 424–434.
- X. Gao, X. Zou, H. Ma, S. Meng and G. Zhu, *Adv. Mater.*, 2014, **26**, 3644–3648.
- R. Wang, H. Y. Zhang, P. H. M. Feron and D. T. Liang, *Sep. Purif. Technol.*, 2005, **46**, 33–40.
- G. Zhao, X. Huang, X. Wang and X. Wang, *J. Mater. Chem. A*, 2017, **5**, 21625–21649.
- G. Zhao, W. Zhou, Y. Sun, X. Wang, H. Liu, X. Meng, K. Chang and J. Ye, *Appl. Catal., B*, 2018, **226**, 252–257.
- E. Abbasi, A. Hassanzadeh, S. Zarghami, H. Arastoopour and J. Abbasian, *Fuel*, 2014, **137**, 260–268.
- C. Wang, B. Dou, Y. Song, H. Chen, Y. Xu and B. Xie, *Ind. Eng. Chem. Res.*, 2014, **53**, 12744–12752.
- C. Gunathilake and M. Jaroniec, *J. Mater. Chem. A*, 2016, **4**, 10914–10924.
- T. A. Saleh, *Advanced Nanomaterials for Water Engineering, Treatment, and Hydraulics*, IGI Global, 2017, ISBN13: 9781522521365.
- T. A. Saleh and V. K. Gupta, *Nanomaterial and Polymer Membranes: Synthesis, Characterization, and Applications*, Elsevier, 2016, ISBN: 978-0128047033.
- C. F. Martin, E. Stöckel, R. Clowes, D. J. Adams, A. I. Cooper, J. J. Pis, F. Rubiera and C. Pevida, *J. Mater. Chem.*, 2011, **21**, 5475–5483.



- 21 Z. Xiang and D. Cao, *J. Mater. Chem. A*, 2013, **2**, 2691–2718.
- 22 X. Zou and G. Zhu, *Adv. Mater.*, 2018, **30**, 1700750.
- 23 N. B. McKeown and P. M. Budd, *Chem. Soc. Rev.*, 2006, **35**, 675–683.
- 24 T. A. Saleh, A. Sari and M. Tuzen, *J. Environ. Chem. Eng.*, 2017, **5**, 1079–1088.
- 25 T. A. Saleh and S. Ali, *J. Environ. Chem. Eng.*, 2018, **6**, 5361–5368.
- 26 K. C. Baranwal and H. L. Stephens, *Basic Elastomer Technology*, Rubber Division, Akron, 1st edn, 2001.
- 27 J. Johns and V. Rao, *Int. J. Polym. Mater. Polym. Biomater.*, 2011, **60**, 766–775.
- 28 X. Zhang, Z. Lu, D. Tian, H. Li and C. Lu, *Appl. Polym. Sci.*, 2013, **127**, 4006–4014.
- 29 S. Tomyangkul, P. Pongmuksuwan, W. Harnnarongchai and K. Chaochanchaikul, *J. Reinf. Plast. Compos.*, 2016, **35**, 672–681.
- 30 S. A. Riyajan and P. Keawittarit, *Polym. Int.*, 2016, **65**, 491–502.
- 31 N. Rattanasoma, T. Saowapark and C. Deeprasertkul, *Polym. Test.*, 2007, **26**, 369–377.
- 32 S. Prasertsri and N. Rattanasom, *Polym. Test.*, 2012, **31**, 593–605.
- 33 A. Tunlert, P. Prasassarakich and S. Poompradub, *Mater. Chem. Phys.*, 2016, **173**, 78–88.
- 34 Y. Belmabkhout, R. Serna-Guerrero and A. Sayari, *Chem. Eng. Sci.*, 2009, **64**, 3721–3728.
- 35 R. S. Franchi, P. J. E. Harlick and A. Sayari, *Ind. Eng. Chem. Res.*, 2005, **44**, 8007–8013.
- 36 D. G. P. M. Perera, T. O. Kumanayaka and S. Walpalage, *Int. J. Sci. Res.*, 2015, **5**, 1–5.
- 37 R. A. Oweini and H. E. Rassy, *J. Mol. Struct.*, 2009, **919**, 140–145.
- 38 T. A. Saleh, *Desalin. Water Treat.*, 2015, **57**, 10730–10744.
- 39 T. A. Saleh, *Environ. Sci. Pollut. Res.*, 2015, **22**, 16721–16731.
- 40 N. D. Hegde and A. V. Rao, *Appl. Surf. Sci.*, 2006, **253**, 1566–1572.
- 41 A. M. Donia, A. A. Atia, A. M. Daher, O. A. Desouky and E. A. Elshehy, *Int. J. Miner. Process.*, 2011, **101**, 81–88.
- 42 S. C. Hsu, C. Lu, F. Su, W. Zeng and W. Chen, *Chem. Eng. Sci.*, 2010, **65**, 1354–1361.
- 43 G. Hernández and R. Rodríguez, *J. Non-Cryst. Solids*, 1999, **246**, 209–215.
- 44 T. Theppradit, P. Prasassarakich and S. Poompradub, *Mater. Chem. Phys.*, 2014, **148**, 940–948.
- 45 H. M. Lim and M. Y. Amir-Hashim, *J. Rubber Res.*, 2011, **14**, 41–50.
- 46 F. Su, C. Lu, S. C. Kuo and W. Zeng, *Energy Fuels*, 2010, **24**, 1441–1448.
- 47 N. N. Linneen, R. Pfeffer and Y. S. Lin, *Chem. Eng. J.*, 2014, **254**, 190–197.
- 48 M. Yao, Y. Dong, X. Feng, X. Hu, A. Jia, G. Xie, G. Hu, J. Lu, M. Luo and M. Fan, *Fuel*, 2014, **123**, 66–72.
- 49 G. Zainab, N. Iqbal, A. A. Babar, C. Huang, X. Wang, J. Yu and B. Ding, *Compos. Commun.*, 2017, **6**, 41–47.
- 50 M. R. Mello, D. Phanon, G. Q. Silveira, P. L. Llewellyn and C. M. Ronconi, *Microporous Mesoporous Mater.*, 2011, **143**, 174–179.
- 51 I. H. Arellano, J. Huang and P. Pendleton, *RSC Adv.*, 2015, **5**, 65074–65083.
- 52 F. Liu, K. Huang, Q. Wu and S. Dai, *Adv. Mater.*, 2017, **29**(1–8), 1700445.

