



**HAL**  
open science

## Room-Temperature Synthesis of BPH Zeolite Nanosheets Free of Organic Template with Enhanced Stability for Gas Separations

Edwin Clatworthy, Maxime Debost, Nicolas Barrier, Stéphanie Gascoin, Philippe Boullay, Aurélie Vicente, Jean-Pierre Gilson, Jean-Pierre Dath, Nikolai Nesterenko, Svetlana Mintova

► **To cite this version:**

Edwin Clatworthy, Maxime Debost, Nicolas Barrier, Stéphanie Gascoin, Philippe Boullay, et al.. Room-Temperature Synthesis of BPH Zeolite Nanosheets Free of Organic Template with Enhanced Stability for Gas Separations. ACS Applied Nano Materials, 2021, 4 (1), pp.24-28. 10.1021/ac-sanm.0c02925 . hal-03143339

**HAL Id: hal-03143339**

**<https://hal.science/hal-03143339>**

Submitted on 16 Feb 2021

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

# Room-Temperature Synthesis of BPH Zeolite Nanosheets Free of Organic Template with Enhanced Stability for Gas Separations

Edwin B. Clatworthy,<sup>†</sup> Maxime Debost,<sup>†</sup> Nicolas Barrier,<sup>‡</sup> Stéphanie Gascoin,<sup>‡</sup> Philippe Boullay,<sup>‡</sup> Aurélie Vicente,<sup>†</sup> Jean-Pierre Gilson,<sup>†</sup> Jean-Pierre Dath,<sup>‡</sup> Nikolai Nesterenko,<sup>‡</sup> Svetlana Mintova<sup>†\*</sup>

<sup>†</sup> Normandie Université, ENSICAEN, UNICAEN, CNRS, Laboratoire Catalyse et Spectrochimie (LCS), 14050 Caen, France

<sup>‡</sup> Normandie Université, ENSICAEN, UNICAEN, CNRS, Laboratoire de Cristallographie et Science des Matériaux (CRISMAT), 14050 Caen, France

<sup>‡</sup> Total Research and Technology, Feluy, B-7181 Seneffe, Belgium

Supporting Information Placeholder

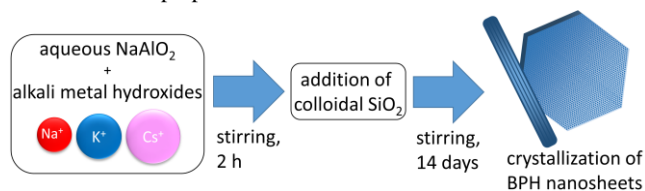
**ABSTRACT:** BPH-type synthetic aluminosilicate zeolite (Linde Q) nanosheets are synthesized at room temperature without organic structure-directing agents (OSDAs). The BPH nanosheets (7 nm thick, 90 nm across, Si/Al = 1.3) are formed in a colloidal suspension using a mixture of structure-directing cations (Cs<sup>+</sup>, K<sup>+</sup> and Na<sup>+</sup>). *In situ* XRD reveals the high stability of the BPH nanosheets up to 500 °C under vacuum (0.021 mbar) due to the presence of Cs<sup>+</sup>. The high CO<sub>2</sub> adsorption capacity (3.78 mmol/g) and stability of the BPH nanosheets open the door for the preparation of oriented thin films and membranes for gas separation applications.

Zeolites have been the subject of significant research for the last 70 years because of their applications in key petrochemical processes and consumer products.<sup>1</sup> Such research includes the synthesis of zeolites with nanoscale dimensions to achieve higher surface area and improved diffusion properties, a high silica content to improve their thermal stability, and new structures.<sup>2-3</sup> To achieve these outcomes organic structure-directing agents (OSDAs) are frequently employed, however, their use is undesirable because of their high cost and necessary post-synthetic removal resulting in increased energy-use, emissions (NO<sub>x</sub>, CO<sub>x</sub>), and can reduce the zeolite crystallinity and provoke leaching of Al. While high-silica zeolites are necessary for high temperature processes, low-silica zeolites, which can offer a greater number of acid sites and compositional diversity, are typically used in separation and purification processes.<sup>4-5</sup> In comparison to many high-silica zeolites, the synthesis of low-silica zeolites is typically mediated by alkali metal cations that act as structure directing agents (SDAs) in highly alkaline media.<sup>6</sup> An example of such a low-silica zeolite is Linde Q with beryllophosphate-H (BPH) framework topology; a synthetic aluminosilicate with extra-framework potassium (K<sup>+</sup>) cations, and a ratio of Si/Al ≈ 1.0. The BPH framework consists of 12 membered-ring (12MR) channels along the z-axis perpendicularly intersected by three 8MR channels.<sup>7-9</sup> Reported by Breck *et al.* in 1961, Linde Q is prepared from a K<sup>+</sup>-containing alkaline aluminosilicate precursor mixture hydrothermally treated at 50 °C for 162 h. The synthesis and structural characterization of Linde Q, (also referred to as K-I and zeolite Q) was investigated further in subsequent reports.<sup>10-15</sup> It was revealed that Linde Q has low thermal stability in air; undergoing structural collapse above 150 °C.<sup>12, 15</sup> Efforts to improve the

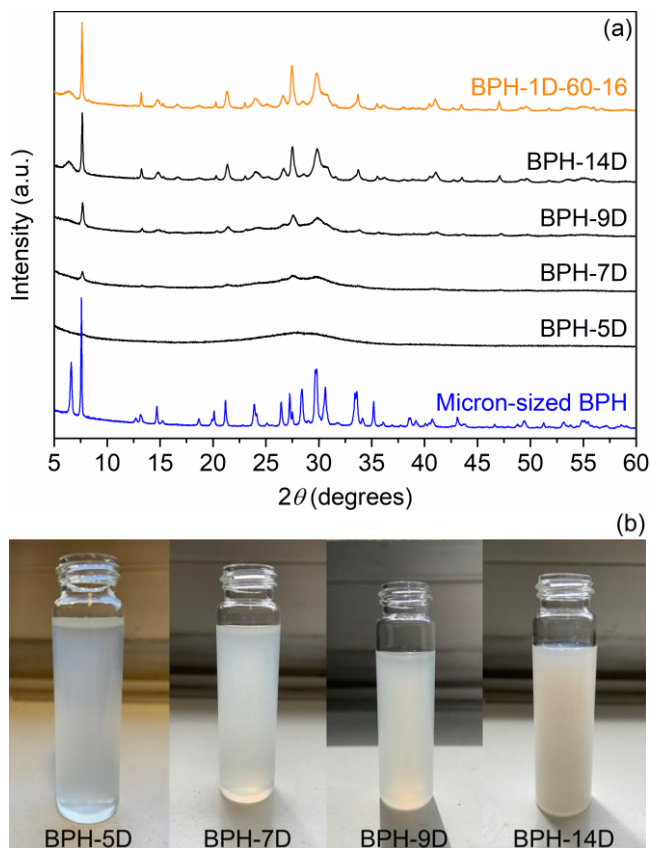
thermal stability of Linde Q have involved post-synthetic ion-exchange with rare-earth metal salts, and calcination in the presence of a salt.<sup>16, 17</sup> Alternatively, synthesis procedures using OSDAs in combination with post-synthetic treatments has afforded higher silica variants (UZM-4 and PST-12, Si/Al up to 10).<sup>4, 18-21</sup> Regarding the particle size, procedures using OSDAs and seed crystals have afforded hexagonal crystals of UZM-4 50–200 nm wide.<sup>22</sup>

Here we report the room-temperature and OSDA-free synthesis of BPH (Linde Q) containing a combination of three cations (Cs<sup>+</sup>, K<sup>+</sup> and Na<sup>+</sup>), yielding nanosheets approximately 7 nm thick and 90 nm across. The as synthesized BPH nanosheets possess high thermal (up to 500 °C) stability due to the presence of Cs<sup>+</sup> extra-framework cations and are prospective candidates for the preparation of oriented thin films and membranes for gas separation applications.

The synthesis of the BPH nanosheets involved the following steps (Scheme 1): (i) preparation of clear aqueous colloidal suspensions by rapid stirring of alkaline aluminate mixtures by combining sodium aluminate (NaAlO<sub>2</sub>), sodium hydroxide (NaOH), potassium hydroxide (KOH) and cesium hydroxide (CsOH), followed by the addition of colloidal silica (SiO<sub>2</sub>); (ii) crystallization of BPH nanosheets from the clear colloidal suspensions under continuous stirring at room temperature between 5 to 21 days (samples labelled as BPH-5D, BPH-7D, BPH-9D, BPH-14D). To explore the limits of the synthesis a sample was prepared for 21 days demonstrating negligible differences in the obtained material. For comparison, a sample was prepared hydrothermally by taking the clear precursor suspension stirred for 1 day at room temperature followed by treatment at 60 °C for 16 hours (BPH-1D-60-16). In addition, the conventional micron-sized BPH was prepared as a reference.<sup>7</sup>



**Scheme 1.** Synthesis of BPH nanosheets at room temperature.



**Figure 1.** (a) XRD patterns of the BPH nanosheets (top to bottom) BPH-1D-60-16, BPH-14D, BPH-9D, BPH-7D, BPH-5D, and micron-sized BPH; (b) photographs of BPH-5D, 7D, 9D, and 14D samples showing the change in colloidal appearance related to the degree of crystallinity at room temperature.

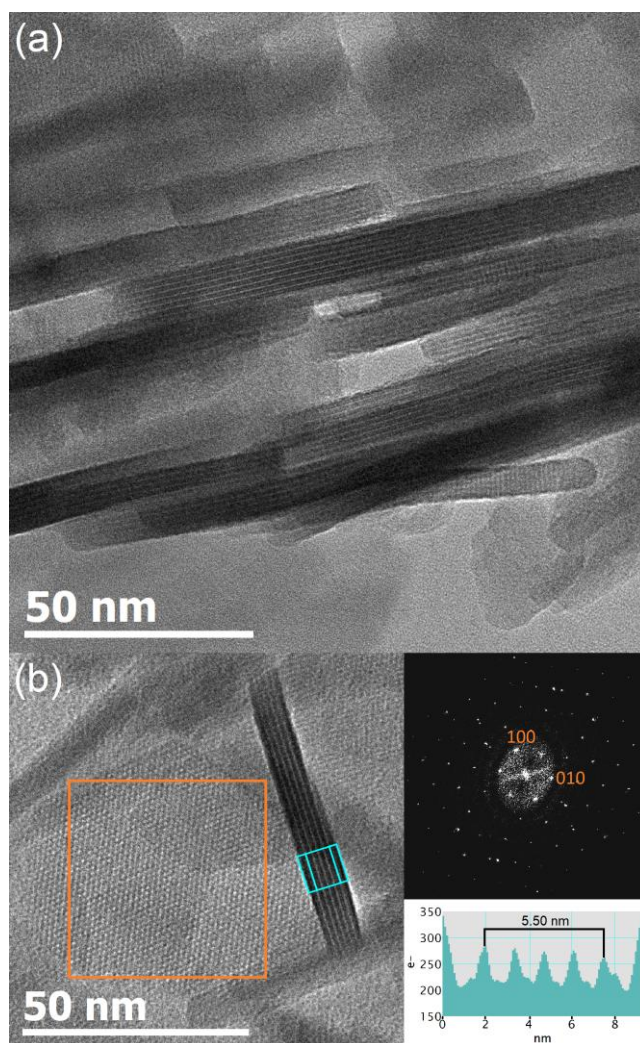
XRD analysis reveals the formation of crystalline BPH after approximately 5 days at room temperature with the emergence of the (100) reflection at  $\approx 7.6^\circ 2\theta$  (Figures 1a, S1). Coincident with the increasing intensity of the reflections is a change in the appearance of the reaction mixture from water clear to opaque (Figure 1b), related to the formation of colloiddally stable zeolite particles in the suspension. Plotting the crystallinity against time reveals an approximately linear trend over the initial 14 days (Figure S2). The zeolite yield *vs* crystallization time at room temperature shows two distinct regions of behavior from 5 to 9 days and 14 to 21 days, showing a significant increase in the yield of solid material between 9 and 14 days attributed to the assembly of the zeolite framework from structural building units (Figure S2).

The XRD pattern of the hydrothermally treated sample is identical to the samples synthesized at room temperature for 14 and 21 days (Figure 1A). Analysis of the XRD data of samples BPH-14D, BPH-21D and BPH-1D-60-16 (Figures S3–5) was performed using the Le Bail method; the structure was fitted with the space group *P*321 (trigonal). Comparison of the lattice parameters (*a*, *c*) of the nanosheet samples with the micron-sized BPH reveals significant differences: *a* is reduced while *c* is increased *e.g.* *a* = 13.3688, *c* = 13.471 Å for BPH-14D *vs* *a* = 13.4909, *c* = 13.3788 for the micron-sized BPH (Table S1).<sup>14</sup> Additionally, the unit cell volume of the nanosheet samples is smaller; 2085.1 for BPH-14D *vs*  $\approx 2108.78 \text{ \AA}^3$  for the micron-sized BPH (Table S1).<sup>15</sup> Similar differences between the nanosheet samples (BPH-21D and BPH-1D-60-16) and the micron-sized BPH were also observed. The smaller lattice parameter *a* and the unit cell volume of the

nanosheet samples is consistent with the substitution of  $\text{Al}^{3+}$  ions with  $\text{Si}^{4+}$  ions in the framework, noted by Lewis *et al.* for UZM-4.<sup>4</sup> This is attributed to the higher Si/Al ratio, determined by ICP analysis, of the nanosheet samples compared to micron-sized BPH (1.3 *vs* 1.0, Table S2).

<sup>29</sup>Si MAS NMR analysis of the nanosheet sample BPH-14D reveals a more complex spectrum compared to the micron-sized BPH which displays three sharp resonances at  $-84.8$ ,  $-86.0$  and  $-89.9$  ppm consistent with three crystallographically distinct  $\text{Si}^{4+}$  sites fourfold coordinated by  $\text{Al}^{3+}$ .<sup>14-15</sup> In comparison, the BPH-14D nanosheets display three asymmetric resonances at  $-85.1$ ,  $-89.8$ , and  $-94.1$  ppm and several other weaker signals (Figure S6a). The shift of the three most intense resonances to lower field is attributed to the higher Si/Al ratio ( $> 1.0$ ). In comparison to the ICP results, deconvolution of the <sup>29</sup>Si spectrum of BPH-14D gives a Si/Al ratio of  $\approx 1.3$ . <sup>27</sup>Al MAS NMR analysis of the BPH-14D nanosheets revealed a single resonance at 59.4 ppm (Figure S6b). Compared to the micron-sized BPH (61.2 ppm), the resonance in the <sup>27</sup>Al spectrum of BPH-14D is shifted to slightly lower field, again consistent with a higher Si/Al ratio of 1.3.<sup>14-15</sup> No signal corresponding to extra-framework Al was observed.

TEM analysis of BPH-14D revealed the presence of well-defined and hexagonally shaped nanosheets (90–130 nm wide, 7–



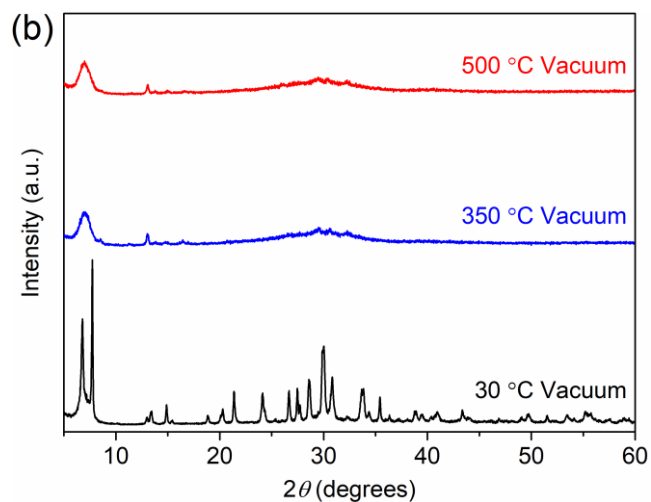
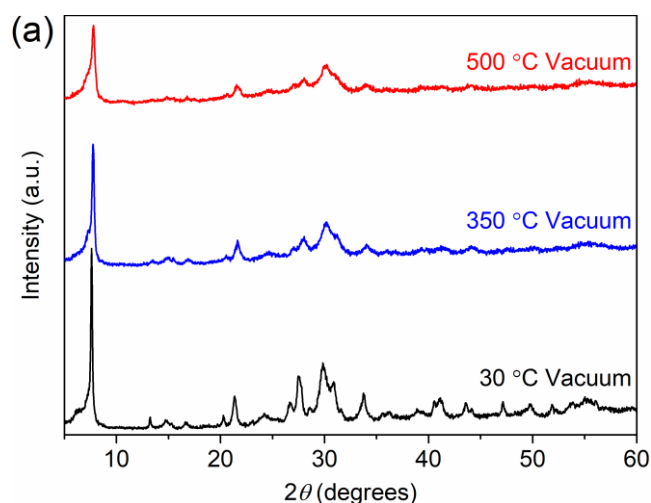
**Figure 2.** (a) TEM image highlighting the morphology and thickness of nanosheets in sample BPH-14D. (b) TEM analysis of the

nanosheet sample BPH-14D including FFT (orange square) and cross-section analysis (blue rectangle).

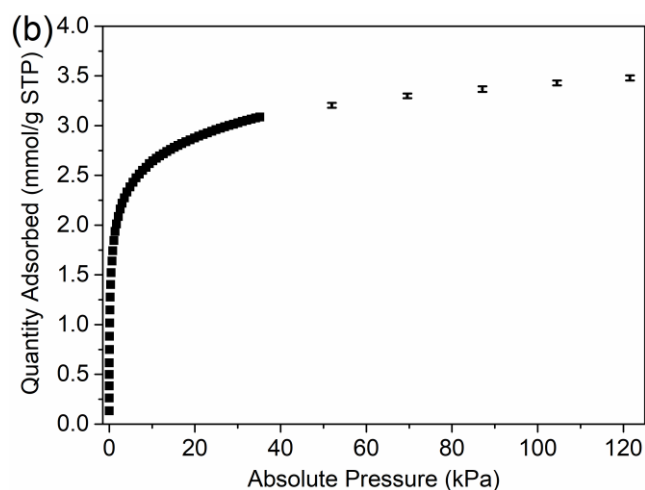
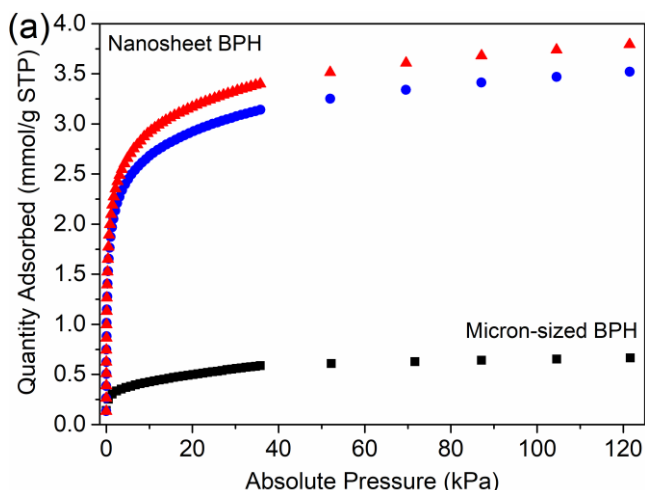
11 nm thick, Figures 2a, b). Smaller particles (< 90 nm at maximum width) with irregular shapes were also observed. For the hydrothermally treated sample BPH-1D-16-60 the morphology was similar to the sample BPH-14D, however, the sheets were larger (50–220 nm wide, 5–19 nm thick, Figure S7). In comparison, the micron-sized BPH was composed of plate-like crystals (Figure S8).

As mentioned in the literature,<sup>16</sup> by exchanging the extra-framework  $K^+$  for  $La^{3+}$  it was shown that the micron-sized BPH could be stabilized above 150 °C. The stability of the BPH nanosheets was investigated by *in situ* variable temperature XRD (30–600 °C). Under medium vacuum (0.021 mbar) the nanosheets are significantly more stable compared to the micron-sized BPH, retaining  $\approx 60\%$  of their initial crystallinity at 350 °C and  $\approx 44\%$  of their initial crystallinity at 500 °C (Figures 3a, S9). In comparison, the micron-sized BPH exhibits structural collapse beginning at 150 °C and almost complete amorphization at 350 °C under medium vacuum (Figure 3b, S10).

With the improved thermal stability of the nanosheets under medium vacuum their adsorption properties were evaluated. TGA of BPH-14D under low vacuum (1.4 mbar) reveals weight loss below 100 °C attributed to the release of surface water (Figure S11). Further weight loss is observed at 150 °C with almost



**Figure 3.** XRD patterns recorded at 30 °C (black), 350 °C (blue) and 500 °C (red) under medium vacuum (0.021 mbar) of (a) BPH-14D nanosheets, and (b) micron-sized BPH.



**Figure 4.** (a)  $CO_2$  adsorption isotherms (0 °C) of the nanosheet samples BPH-1D-60-16 (red triangles), BPH-14D (blue circles), and micron-sized BPH (black squares). (b) Average  $CO_2$  adsorption isotherm of three consecutive measurements of the nanosheet sample BPH-14D, the last five data points present the standard deviation.

complete dehydration occurring by 200 °C. Similarly, the micron-sized BPH shows almost complete dehydration by 200 °C but with weight loss occurring at  $\approx 100$  and 150 °C. The different behavior of the two samples is attributed to the different surface area allowing for more water molecules to be bound to the external surface of the nanosheets and a more rapid diffusion of water through the zeolite.

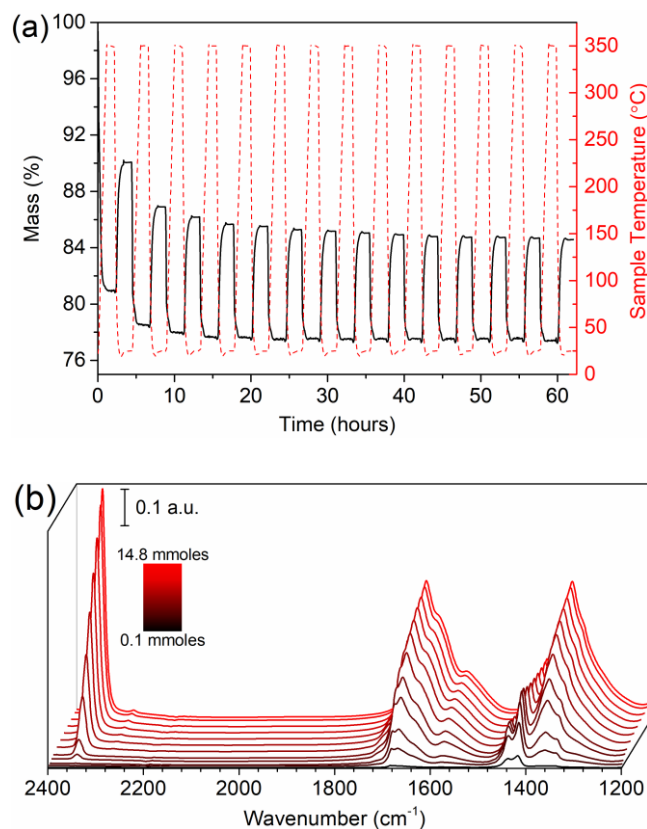
Prior to adsorption analysis the samples were pre-treated at 350 °C under high vacuum (2.67  $\mu$ bar).  $N_2$  adsorption analysis shows that the BPH nanosheets exhibit a mixture of type I and IV isotherms with a type H1 hysteresis attributed to interparticle mesopores (Figure S12, Table S3). In comparison, the micron-sized BPH possesses a negligible BET specific surface area and low micropore volume. Similarly, the  $CO_2$  adsorption capacity of the BPH nanosheets is significantly greater, up to 3.78 mmol/g, as compared to the micron-sized BPH (Figure 4a) which undergoes structural collapse (Figure S13); these results clearly demonstrate the superior thermal stability of the nanosheets under vacuum. The  $CO_2$  adsorption capacity of the BPH nanosheets is compara-



ble to commercial Na<sup>+</sup>/K<sup>+</sup>-LTA and Cs-CHA at 0 °C,<sup>23-25</sup> but less than Na<sup>+</sup>-LTA (4A) at 0 °C, and FAU (NaY and NaX) zeolites at 25 °C.<sup>26</sup> The nanosheets demonstrate consistent adsorption behavior over three consecutive measurements with less than 1% standard deviation (Figure 4b). The adsorption behavior of the nanosheets was investigated further by performing multiple TG cycles of heating (350 °C) under low vacuum (1.4 mbar) and flowing CO<sub>2</sub> at 25 °C (Figure 5a). The nanosheets display an initial decrease in CO<sub>2</sub> capacity attributed to the formation of carbonate species (*vide infra*) followed by stable behavior.

To determine the cause of the greater stability of the BPH nanosheets, Cs ion-exchange of the micron-sized BPH was performed and the product analyzed by CO<sub>2</sub> adsorption and XRD (Supporting Information). The Cs-micron-sized BPH exhibited a similar CO<sub>2</sub> adsorption capacity compared to the BPH nanosheets prepared at room temperature (Figure S14). XRD revealed the Cs-micron-sized BPH after CO<sub>2</sub> adsorption had lost a greater degree of crystallinity compared to the BPH nanosheets (Figure S15). These results demonstrate that the greater stability of the BPH nanosheets compared to the conventional micron-sized BPH is due to the presence of the extra-framework Cs<sup>+</sup>.

*In situ* FTIR spectroscopy of the BPH nanosheets with CO<sub>2</sub> revealed the formation of chemisorbed species between 1800 and 1200 cm<sup>-1</sup> (Figure 5b).<sup>27</sup> The bands at 1416 and 1441 cm<sup>-1</sup> are attributed to carboxylate and free carbonate species; rapidly reaching maximum absorbance before equilibrium. The most prominent bands have maxima at 1363 and 1668 cm<sup>-1</sup>, attributed to bridged bidentate carbonates. The band at 1363 cm<sup>-1</sup> was accompanied by shoulders at 1339 and 1374 cm<sup>-1</sup> while the band at 1668 cm<sup>-1</sup> exhibited shoulders at 1639, 1682 cm<sup>-1</sup>, and a weaker



**Figure 5.** (a) TG cycles of BPH-14D treated at 350 °C under low vacuum (1.4 mbar) followed by flowing CO<sub>2</sub> (40 mL/min) at 25 °C, mass: black solid line, sample temperature: red dashed line.

(b) *In situ* FTIR spectra of BPH-14D under increasing CO<sub>2</sub> atmosphere, pre-treated at 350 °C under high vacuum (10<sup>-3</sup> μbar).

shoulder at 1716 cm<sup>-1</sup>. A broad and less intense band was observed at 1585 cm<sup>-1</sup> indicative of carboxylate species. Physisorbed CO<sub>2</sub>, including <sup>13</sup>CO<sub>2</sub>, was identified by the ν<sub>3</sub> asymmetric stretches at 2346 and 2280 cm<sup>-1</sup>. Removal of the CO<sub>2</sub> under high vacuum followed by a second activation treatment revealed the retention of free (1435, 1421 cm<sup>-1</sup>) and bridged bidentate (1663 cm<sup>-1</sup>) carbonates which may have implications for selective separation of CO<sub>2</sub> due to carbonate-promoted cation displacement (Figure S14).<sup>28</sup>

In conclusion, we have achieved the room-temperature OSDA-free synthesis of BPH nanosheets possessing high thermal stability under vacuum and CO<sub>2</sub> adsorption capacity. Nanosheets with dimensions approximately 7 nm thick and 90 nm wide possess a Si/Al ratio of 1.3 and superior thermal stability up to 500 °C under vacuum compared to conventional micron-sized BPH (Linde Q) due to the presence of Cs<sup>+</sup> extra-framework cations. The nanosheets demonstrate considerable CO<sub>2</sub> adsorption capacity, comparable to commercial Na<sup>+</sup>/K<sup>+</sup>-LTA and Cs-CHA zeolites,<sup>23-25</sup> while investigation of the thermal behavior under different degrees of vacuum (low, medium, high) indicates that the application of a higher vacuum results in a greater retention of the initial crystallinity and sorption capacity of the nanosheets. The nanosheets are colloiddally stable and are ideal candidates for the preparation of oriented thin films and membranes for gas separation applications.

## ASSOCIATED CONTENT

### Supporting Information

Contains information pertaining to materials, equipment and experimental procedures, Le Bail refinement data, ICP results, XRD and *in situ* variable temperature XRD under vacuum, TEM, SEM, <sup>29</sup>Si and <sup>27</sup>Al MAS NMR, TGA, N<sub>2</sub> and Ar adsorption data. The Supporting Information is available free of charge on the ACS Publications website at DOI:

## AUTHOR INFORMATION

### Corresponding Author

\*E-mail: mintova@ensicaen.fr

### Notes

The authors declare no competing financial interests.

## ACKNOWLEDGMENT

Financial support from TOTAL and Industrial Chair ANR-TOTAL "Nanoclean Energy" is acknowledged as well as from the Normandy Region through the RIN Recherche Program.

## REFERENCES

- (1) Vermeiren, W.; Gilson, J.-P., Impact of Zeolites on the Petroleum and Petrochemical Industry. *Top. Catal.* **2009**, *52* (9), 1131-1161.
- (2) Mintova, S.; Gilson, J.-P.; Valtchev, V., Advances in nanosized zeolites. *Nanoscale* **2013**, *5* (15), 6693-6703.
- (3) Li, J.; Corma, A.; Yu, J., Synthesis of new zeolite structures. *Chem. Soc. Rev.* **2015**, *44* (20), 7112-7127.
- (4) Blackwell, C. S.; Broach, R. W.; Gatter, M. G.; Holmgren, J. S.; Jan, D. Y.; Lewis, G. J.; Mezza, B. J.; Mezza, T. M.; Miller, M. A.; Moscoso, J. G., Open-Framework Materials Synthesized in the TMA<sup>+</sup>/TEA<sup>+</sup> Mixed-Template System: The New Low Si/Al Ratio Zeolites UZM-4 and UZM-5. *Angew. Chem. Int. Ed.* **2003**, *42* (15), 1737-1740.

- (5) Barrett, P.; Huo, Q.; Stephenson, N., Recent advances in low silica zeolite synthesis. In *Stud. Surf. Sci. Catal.*, Elsevier: 2007; Vol. 170, pp 250-257.
- (6) Flanigen, E. M., A Review and New Perspectives in Zeolite Crystallization. In *Molecular Sieves*, Meier, M. W.; Uytterhoeven, J. B., Eds. ACS Publications: 1973; Vol. 121, pp 119-139.
- (7) Breck, D. W.; Acara, N. A. Crystalline Zeolite Q. US2991151A, 1961.
- (8) Harvey, G., The synthesis and structure of beryllphosphate-H - A new open-framework zeolite. *Z. Kristallogr.* **1988**, *182* (1-4), 123-124.
- (9) Harvey, G.; Baerlocher, C.; Wroblewski, T., Structure solution and Rietveld refinement of beryllphosphate-H zeolite. *Z. Kristallogr. - Cryst. Mater.* **1992**, *201* (1-4), 113-124.
- (10) Zhdanov, S. P.; Ovsepyan, M., Some synthetic potash zeolites and their properties. *Dokl. Akad. Nauk* **1964**, *157* (4), 913-916.
- (11) Zhdanov, S. P.; Buntar-Samulevich, N.; Ovsepyan, M., Synthetic chabasites and their adsorptive properties. *Dokl. Akad. Nauk* **1965**, *161* (2), 384-387.
- (12) Barrer, R.; Cole, J.; Sticher, H., Chemistry of soil minerals. Part V. Low temperature hydrothermal transformations of kaolinite. *J. Soc. Chem. A* **1968**, 2475-2485.
- (13) Bosmans, H. J.; Tambuyze, E.; Paenhuys, J.; Ylen, L.; Vancluys, J., Zeolite Formation in the System  $K_2O-Na_2O-Al_2O_3-SiO_2-H_2O$ . In *Molecular Sieves*, Meier, M. W.; Uytterhoeven, J. B., Eds. ACS Publications: 1973; Vol. 121, pp 179-188.
- (14) Andries, K.; Bosmans, H.; Grobet, P., The crystal structure of zeolite Linde Q: A proposal based on powder X-ray diffraction and  $^{27}Al$  and  $^{29}Si$  MAS nmr spectroscopy. *Zeolites* **1991**, *11* (2), 124-131.
- (15) Andries, K.; De Wit, B.; Grobet, P.; Bosmans, H., The synthesis and properties of synthetic zeolite Linde Q. *Zeolites* **1991**, *11* (2), 116-123.
- (16) Vaughan, D. E. W. ECR-33: a stabilized rare-earth exchanged Q type zeolite. US5382420A, 1995.
- (17) Huo, Q.; Stephenson, N. A. Stabilization of molecular sieves by salt addition. US6632767B2, 2003.
- (18) Lewis, G. J.; Moscoso, J. G.; Miller, M. A.; Wilson, B. A. Crystalline aluminosilicate zeolitic composition: UZM-4. US6419895B1, 2002.
- (19) Wilson, B. A.; Lewis, G. J.; Jan, D.-Y.; Wilson, S. T.; Patton, R. L. Crystalline aluminosilicate zeolitic composition: UZM-4M. US6776975B2, 2004.
- (20) Miller, M. A.; Moscoso, J. G.; Koster, S. C.; Gatter, M. G.; Lewis, G. J., Synthesis and characterization of the 12-ring zeolites UZM-4 (BPH) and UZM-22 (MEI) via the charge density mismatch approach in the Choline- $Li_2O-SrO-Al_2O_3-SiO_2$  system. In *Stud. Surf. Sci. Catal.*, Elsevier: 2007; Vol. 170, pp 347-354.
- (21) Park, M. B.; Ahn, S. H.; Ahn, N. H.; Hong, S. B., Charge density mismatch synthesis of MEI-and BPH-type zeolites in the  $TEA^+-TMA^+-Li^+-Sr^{2+}$  mixed-structure-directing agent system. *Chem. Commun.* **2015**, *51* (17), 3671-3673.
- (22) Moteki, T.; Okubo, T., From Charge Density Mismatch to a Simplified, More Efficient Seed-Assisted Synthesis of UZM-4. *Chem. Mater.* **2013**, *25* (13), 2603-2609.
- (23) Rzepka, P.; Wardecki, D.; Smeets, S.; Müller, M.; Gies, H.; Zou, X.; Hedin, N.,  $CO_2$ -Induced Displacement of  $Na^+$  and  $K^+$  in Zeolite [NaK]-A. *J. Phys. Chem. C* **2018**, *122* (30), 17211-17220.
- (24) Shang, J.; Li, G.; Singh, R.; Xiao, P.; Liu, J. Z.; Webley, P. A., Determination of composition range for "molecular trapdoor" effect in chabazite zeolite. *J. Phys. Chem. C* **2013**, *117* (24), 12841-12847.
- (25) Debost, M.; Klar, P. B.; Barrier, N.; Clatworthy, E. B.; Grand, J.; Lainé, F.; Brazda, P.; Palatinus, L.; Nesterenko, N.; Boullay, P., Synthesis of discrete CHA zeolite nanocrystals without organic templates for selective  $CO_2$  capture. *Angew. Chem. Int. Ed.* **2020**, *132*.
- (26) Walton, K. S.; Abney, M. B.; LeVan, M. D.,  $CO_2$  adsorption in Y and X zeolites modified by alkali metal cation exchange. *Microporous Mesoporous Mater.* **2006**, *91* (1-3), 78-84.
- (27) Stevens Jr, R. W.; Siriwardane, R. V.; Logan, J., In situ Fourier transform infrared (FTIR) investigation of  $CO_2$  adsorption onto zeolite materials. *Energy Fuels* **2008**, *22* (5), 3070-3079.
- (28) Bryukhanov, I. A.; Rybakov, A. A.; Larin, A. V., Carbonate-Promoted Drift of Alkali Cations in Small Pore Zeolites: Ab Initio Molecular Dynamics Study of  $CO_2$  in NaKA Zeolite. *The Journal of Physical Chemistry Letters* **2019**, *10* (9), 2191-2195.

---

Insert Table of Contents artwork here

