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One-pot synthesis of high-flux *b*-oriented MFI zeolite membranes for Xe recovery

Xuerui Wang,^{†,*} Pelin Karakiliç,[‡] Xinlei Liu,[†] Meixia Shan,[†] Arian Nijmeijer,[‡] Louis Winnubst,[‡] Jorge Gascon,^{†,§,*} and Freek Kapteijn^{†,*}

[†] Chemical Engineering Department, Delft University of Technology, Van der Maasweg 9, 2629 HZ Delft, The Netherlands

[‡] Department Inorganic Membranes, MESA+ Institute for Nanotechnology, University of Twente, P.O. Box 217, 7500 AE Enschede, The Netherlands

[§] KAUST Catalysis Center, Advanced Catalytic Materials, King Abdullah University of Science and Technology, Thuwal 23955, Saudi

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ABSTRACT: We demonstrate that *b*-oriented MFI zeolite membranes can be manufactured by *in-situ* crystallization using an intermediate amorphous SiO₂ layer. The improved in-plane growth by using a zeolite growth modifier leads to fusion of independent crystals and eliminates boundary gaps, giving good selectivity in the separation of CO₂/Xe mixtures. The fast diffusion of CO₂ dominates the overall membrane selectivity towards CO₂/Xe mixture. Because of the straight and short [010] channels, the obtained CO₂ permeation fluxes are several orders of magnitude higher than those of carbon molecular sieving membranes and polymeric membranes, opening opportunities for Xe recovery from waste anaesthetic gas.

1. INTRODUCTION

Xenon (Xe) is considered as an attractive and better anaesthetic than those standard used currently (such as N₂O and fluoroethers) in clinical medicine because of the low risk of hypoxia, not being a neurotoxin, and the absence of “memory” effects in nerve cells.¹ However, its broad application is significantly impeded by the extremely high price of Xe (5000 \$/kg)² associated with the low abundance in the Earth's atmosphere (0.087 ppmv) and energy-intensive cryogenic distillation used in its production.³ To make Xe less costly, equilibrium-based adsorption processes using metal-organic frameworks (MOFs),³ zeolites,⁴ or porous organic cages⁵ were explored to capture Xe directly from the air or discharged Xe-containing gas waste. In the case of waste anaesthetic gas, Xe (50-60% concentration) would be retrieved by selective removal of CO₂.⁶ Given the different molecular size and affinity, microporous zeolite membranes should be able to separate the smaller CO₂ molecules from the bigger Xe molecule (3.3 Å vs 4.1 Å) based on a kinetically controlled process.

Zeolites are crystalline microporous aluminosilicates with well-defined pore size and shape that have been widely used as catalysts, adsorbents, and membranes in many industrial processes, such as fluid catalytic cracking, natural gas upgrading, and bioethanol purification.⁷⁻⁹ Polycrystalline zeolite membranes are usually composed of anisotropic crystals with accessible apertures and channels randomly

aligned within the membrane layer, which limits access to the pores and increases diffusion path lengths.¹⁰ The development of synthetic protocols that allow the alignment of zeolite pores with the membrane surface is therefore of high interest.

Crystals of the archetype all-silica MFI zeolite exhibit hexagonal platelet morphologies with distinct surfaces, the (100) and (010) faces, and a third surface (*xOz*) with variable Miller index. The diffusion coefficient along the straight [010] channel is approximately three times higher than that on the channel perpendicular to this direction.¹¹ Thus, *b*-oriented MFI zeolite membranes with straight and short [010] channels are highly desired to maximize permeance (throughput) and separation factor (selectivity) simultaneously. Yoon *et al.*¹²⁻¹³ developed different methods for orientation-controlled monolayer assembly of zeolite crystals *via* covalent linkers or hand rubbing, which facilitate *b*-oriented MFI zeolite membrane formation by secondary growth.¹⁴⁻¹⁵ Subsequently, various approaches were developed to construct *b*-oriented MFI crystal/nanosheet layers, involving dynamic interfacial assembly,¹⁶⁻¹⁸ polymer-mediated assembly,¹⁹ Langmuir-Schaefer²⁰ or Langmuir-Blodgett²¹ deposition. An unparalleled MFI zeolite membrane with 200-nm-thickness was achieved *via* secondary growth of a zeolite nanosheet layer, obtained by filtration of high-aspect-ratio zeolite nanosheet suspension.²²⁻²³ Despite these elegant approaches, a cost-effective and easy to scale-up methodology for the production of *b*-oriented MFI zeolite membrane is still to be developed.

In addition to cumbersome seeding, another main challenge is to avoid orthogonal rotational intergrowth of MFI zeolite during the hydrothermal synthesis.^{15, 23} Yang *et al.*²⁴ pioneered microwave irradiation heating to prepare *b*-oriented MFI films in short synthesis times. Replacement of the traditional structure-directing agent (SDA) TPAOH by *bis*-*N,N*-propylammonium trihydroxide (*trimer*-TPAOH),¹⁴ triethanolamine (TEOA),²⁵ and TEAOH^{15, 21} has been investigated to enhance the epitaxial growth of *b*-oriented MFI seeds. Neutral²⁶ and fluoride-containing^{15, 27-28} solutions were explored as well to suppress the orthogonally rotational intergrowth of *b*-oriented MFI seeds during secondary growth. Recently, it was demonstrated that water vapour can trigger the epitaxial growth of *b*-oriented MFI seeds when using a sacrificial silica layer impregnated with SDAs as interphase and Si source, eliminating the gaps within the crystal/nanosheet layer and resulting in highly selective *b*-oriented MFI zeolite membranes.²⁹

Herein, we present a versatile *in-situ* crystallization approach to manufacture *b*-oriented MFI zeolite membranes on porous alumina supports. An amorphous silica layer was pre-coated by dip-coating, which would be further transformed to a *b*-oriented MFI zeolite membrane during the following, one step, hydrothermal treatment. We chose tributylphosphine oxide (TBPO) as zeolite growth modifier (ZGM) instead of the complex *trimer*-TPAOH¹⁴ or environmental hazardous HF^{15, 27-28} to improve the in-plane growth of MFI crystals within the membrane, leading to a bigger *b*-face dimension, which facilitates the fusion of independent crystals and eliminates boundary gaps. The aim of this study was not only to provide a preparation methodology of *b*-oriented MFI zeolite membranes, but also to demonstrate the medical applicability of these membranes in Xe recovery.

2. EXPERIMENTAL

2.1 Materials

The following chemicals from Sigma Aldrich were used as received without any purification: tetrapropylammonium hydroxide (TPAOH, 1.0 M solution in water), tetraethyl orthosilicate (TEOS, 98 %), tributylphosphine oxide (TBPO, 95 %). Ammonium fluoride (NH₄F, 96 %) was supported by VWR International BV. Macroporous α -Al₂O₃ discs with a diameter of 25 mm, a thickness of 2 mm, a porosity of 35 % and a pore size of 80 nm were supplied from Pervatech B. V. The Netherlands. Silica-coated α -Al₂O₃ plate supports (25 mm diameter, top silica layer of 1 nm pores, and a thicker bottom of 2.5 μ m pores) were purchased from Fraunhofer IKTS Hermsdorf.

2.2 Preparation of silica-coated alumina supports

A mesoporous γ -Al₂O₃ intermediate layer with a pore size of 5 nm and a thickness of 3 μ m was prepared by dip-coating the macroporous α -Al₂O₃ discs into a solution composed of boehmite and polyvinyl alcohol under cleanroom conditions (class 100); then, the discs were calcined at 650 °C for 3 h with a heating and cooling rate of 1 °C/min. The dip-coating and calcination were performed twice. The silica sol was prepared by an acid-catalyzed sol-gel reaction of TEOS in ethanol and nitric acid under continuous stirring for 3 h at 60 °C. After the

reaction, the silica sol was diluted with ethanol to have the final molar ratio of 1 TEOS: 3.8 EtOH: 6.2 H₂O: 0.085 HNO₃. Then, the γ -Al₂O₃ coated discs were dipped into the silica sol, followed by calcination at 600 °C for 3 h with a heating and cooling rate of 0.5 °C/min. The dip-coating and calcination processes were repeated once more to form a smooth microporous silica layer with a pore size of around 0.5 nm and a thickness of 70-100 nm (**Figure 1a**). Further details of the preparation of silica membranes can be found in our previous paper.³⁰

2.3 Preparation of *b*-oriented MFI zeolite membranes

MFI zeolite membranes were prepared by *in-situ* crystallization from a clear solution with a molar composition of 1 SiO₂: 0.2 TPAOH: 110 H₂O: 4 EtOH. TPAOH was added to deionized water followed by drop-wise addition of TEOS. The mixture was stirred at room temperature for at least 6 h. Then, 0-0.5 wt.% TBPO based on the total weight of the solution was added. The solution was transferred to a 45 mL Teflon-lined autoclave, wherein a silica-coated alumina disk was vertically placed. The autoclave was then placed in an oven pre-heated to 150 °C. After crystallization for a specific period, the membrane and powder was recovered and thoroughly washed with deionized water and ethanol. Different synthesis conditions for all the membranes were listed in Table S1. The silica gel attached on top surface was removed by washing with 0.2 M ammonium fluoride for 4 h. The structure-directing agent of TPA⁺ was removed by calcination: in the case of membranes, calcination was conducted at 400 °C for 2 h with a heating and cooling rate of 1 °C/min; however, 550 °C for 10 h was used for powders.

2.4 Crystallographic preferred orientation (CPO)

The preferred orientation of MFI crystals within the membrane layer was evaluated by $CPO_{(020)/(501)}$ based on powder X-ray diffraction (PXRD) patterns, which is defined in the following way:

$$CPO_{(020)/(501)} = \frac{I_{(020)}^{membrane} / I_{(501)}^{membrane} - I_{(020)}^{powder} / I_{(501)}^{powder}}{I_{(020)}^{membrane} / I_{(501)}^{membrane}} \quad (1)$$

Where $I_{(020)}^{membrane}$ and $I_{(501)}^{membrane}$ refer to the intensity of (020) and (501) peak of the membrane, whereas $I_{(020)}^{powder}$ and $I_{(501)}^{powder}$ refer to the peak intensity of MFI powder. If a peak was not detected, the intensity of that peak was set to 1 count/step according to the report of Hedlund *et al.*³¹

2.5 Gas separation performance

The as-synthesized *b*-oriented MFI zeolite membrane was sealed into a stainless steel cell using a Viton® O-ring. The effective membrane area for permeation was 3.14 cm². The temperature was controlled by an oven from room temperature to 200 °C. He, CO₂, N₂, Ar, Xe, and SF₆ single gas permeation through the membranes was measured in a steady state gas permeation setup. The pressure at the feed side (absolute pressure ranging from 1.2 bar to 3 bar) was controlled using a back-pressure controller and the permeate side was connected to a bubble flow meter at atmospheric pressure (absolute pressure of 1 bar) without sweep gas. The gas permeance (P , GPU) and ideal selectivity (S_{ij}) were defined as follows:

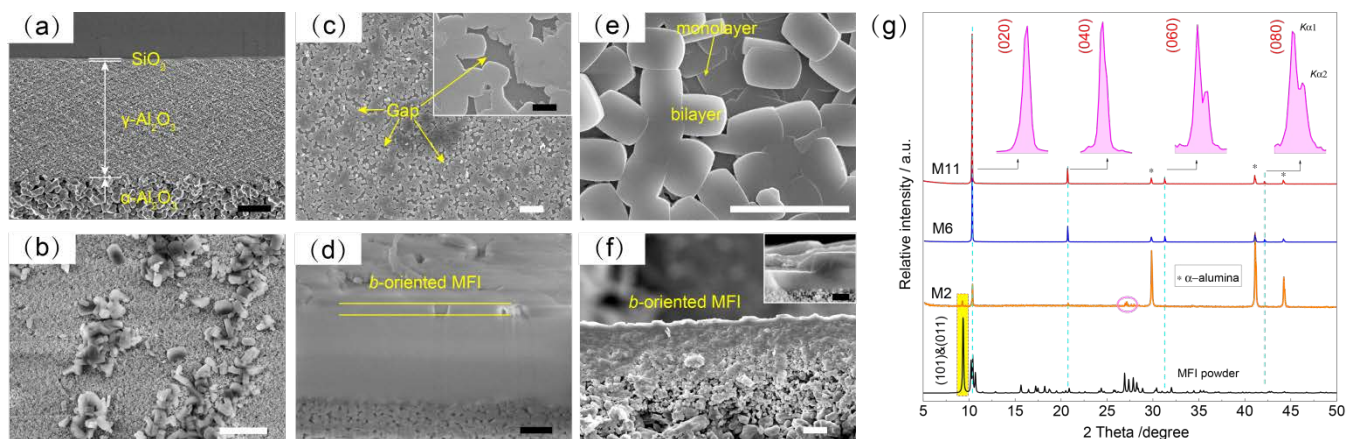


Figure 1. SEM and PXRD characterization of *b*-oriented MFI zeolite membranes. (a) silica-coated alumina support; (b) random-oriented MFI zeolite membrane on the bare alumina support synthesized for 8 h, M2; (c-d) *b*-oriented MFI zeolite membrane on silica-coated alumina support synthesized for 3 h without TBPO, M6; (e-f) *b*-oriented MFI zeolite membrane on silica-coated alumina support synthesized for 6 h with 0.1 wt.% TBPO, M11; scale bar in white and black color: 10 μm and 1 μm . (g) PXRD patterns of random- and *b*-oriented MFI zeolite membranes. Synthesis condition: 1 TEOS: 0.2 TPAOH: 110 H_2O at 150 $^\circ\text{C}$.

$$P_i = \frac{J_i}{\Delta p_i} \quad (2)$$

$$S_{ij} = \frac{P_i}{P_j} \quad (3)$$

where J_i is the permeation flux through the membrane, $\text{mol}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$; Δp_i is the transmembrane pressure difference of component i , Pa.

For CO_2/Xe mixtures, the permeation was investigated by the Wicke-Kallenbach technique. The total feed flow rate was 55 mL/min. Ar was used as sweep gas with a constant volumetric flow rate of 60 mL/min to eliminate concentration polarization and carry the permeate to a two-channel gas chromatograph (Interscience Compact GC) for composition determination. The channel equipped with a ShinCarbon ST 80/100 column (1 m \times 0.53 mm) and a thermal conductivity (TCD) detector was used to separate and analyze CO_2 and Xe. At each permeation condition, the system was stabilized for more than 2 h and the measurement was repeated at least ten injections. The permeance is defined using the same equation as single gas permeation. The separation factor (α_{ij}) is defined by the following equation:

$$\alpha_{ij} = \frac{y_i / y_j}{x_i / x_j} \quad (4)$$

Where x_i , x_j , y_i , and y_j are the molar fractions of the components at the feed side and permeate side, respectively.

2.6 Characterization

The morphology of MFI zeolite powder and membrane was observed by emission scanning electron microscopy (SEM, JSM-6010LA, JEOL). Prior to SEM analysis of the MFI zeolite membrane, a trench was milled in the membrane by accelerating concentrated gallium ions (30 kV, 0.75 nA) using a Dual Beam 3 Nova 200 FIB. Atomic force microscopy (AFM) images were collected in non-contact tapping mode using a Solver NEXT AFM instrument from NT-MDT. A silicon cantilever (HA_NC/50) with spring constants ranging from 0.4 to 2.7 N/m (resonant frequency of 140 kHz) was used. Nova Px 3.2.5 software was used for all of the data acquisition

and analysis. PXRD patterns were recorded in a Bruker-D8 Advance diffractometer using Co- $K\alpha$ radiation ($\lambda=1.78897 \text{ \AA}$). The 2θ range of 5° – 50° was scanned using a scan rate of 0.05 $\text{deg}\cdot\text{s}^{-1}$. Furthermore, Ar isotherm at 87 K was acquired with a 3 Flex (Micromeritics) apparatus using high purity Ar. The sample was first degassed under dry nitrogen flow at 350 $^\circ\text{C}$ for 10 h.

3. RESULTS AND DISCUSSION

3.1 *b*-oriented MFI zeolite formation from amorphous silica layer transition

MFI crystals were sparsely and randomly packed as hillocks on the bare alumina support after *in-situ* crystallization (Figure 1b and Figure S1a-b), in line with the previous report.³² Interestingly, a uniformly *b*-oriented MFI monolayer was obtained when an intermediate silica coating was applied to alumina support (Figure 1c, Figure S1c-d, Figure S2). The average crystal size along the a and c axes were 1.2 μm \times 0.9 μm , respectively, giving a lateral size of 1.08 μm^2 . The monolayer structure was well confirmed by FIB-SEM image of the cross-section, showing a thickness of 350 nm (Figure 1d and Figure S3). The nucleation of MFI zeolite crystals is believed to start at the interface of the silica gel layer and the TPAOH containing solution;³³ the crystallization proceeds until the monolayer structure anchors on the support.³⁴ It should be noted that the amorphous silica layer was partially dissolved and transformed into crystalline MFI crystals during the *in-situ* crystallization. The reactivity of such silica layer was further proved by the formation of *b*-oriented MFI crystals after hydrothermal treatment in a TPAOH solution in the absence of any additional Si source (Figure S4).

The visible gaps within the *b*-oriented MFI monolayer can be mitigated by extending the synthesis time (Figure S5) while still maintaining the *b*-orientation, as evidenced by a $\text{CPO}_{(020)/(501)}$ value higher than 99.6 % (Table S2).³¹ However, defect-free membranes could not be achieved, which is the result of the slow-down in-plane growth of MFI crystals anchoring on supports.³⁴ We believe that this is the reason why a *b*-oriented MFI zeolite film is merely achieved by *in-situ*

crystallization on an impermeable silicon wafer,³⁴ glass,³⁵ and alloy substrates³⁶ but not on porous support for separation.

3.2 Improved in-plane growth by ZGMs

Generally, twinning growth is an inherent growth behavior for MFI zeolites, which results from the orthogonally rotational intergrowth along the *c*-axis or addition of nanoparticles to the *b*-face.^{21, 23, 37} However, when 0.1 wt.% TBPO was added to the synthesis solution, *b*-oriented MFI zeolite membranes without visible gaps were achieved by *in-situ* crystallization at 150 °C for 6 h (Figure 1e). TBPO was chosen because of its highest efficiency over the other commercial ZGMs in reducing the [010] dimension of MFI zeolite.³⁸ The membrane structure in this case is slightly more complex, as additional *b*-oriented MFI crystals formed on top of the primary monolayer, as clearly observed by SEM imaging of the membrane cross-section (Figure 1f): 1.1±0.1 μm for the monolayer and 2.2±0.3 μm for the bilayer. The thickness increases further with synthesis time, for example, a membrane layer composed of bilayered and trilayered MFI zeolite crystals was obtained after *in-situ* crystallization of 10 h (Figure S6a-b).

Only the diffraction peaks from (020), (040), (060), and (080) faces could be observed from the PXRD patterns (Figure 1g, Figure S6c), further proving *b*-orientation of the MFI crystals. The splitting of the latter two diffraction peaks is due to the presence of two different wavelengths in the X-ray source (Co Kα1 and Kα2). To visualize the effect of TBPO on the in-plane growth of MFI zeolite, the lateral size vs synthesis time is plotted in Figure 2. The lateral size of MFI crystals linearly increases with synthesis time. Interestingly, the *b*-face dimension is always larger than that without zeolite growth modifiers. An increment of 210 % was achieved in the lateral size after addition of 0.1 wt.% TBPO, affording a well-intergrown *b*-oriented MFI monolayer with a *b*-face area of 11.9 μm². The improved in-plane growth is responsible for the elimination of visible gaps within the monolayer.

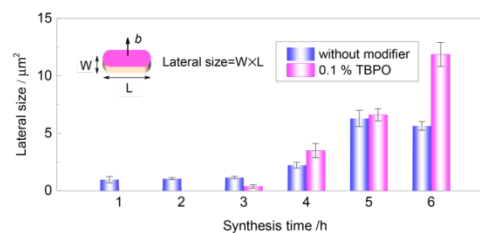


Figure 2. Lateral size of the crystals within *b*-oriented MFI zeolite membranes. Synthesis condition: 1 TEOS: 0.2 TPAOH: 110 H₂O at 150 °C.

3.3 Role of ZGMs

To understand the role of TBPO in the nucleation and growth, solid state ²⁹Si-NMR and Ar adsorption were conducted using MFI zeolite powder. Generally, a defective MFI zeolite would give a peak at δ = -102 ppm, which is ascribed to Q³ Si atoms bearing silanol groups.²⁷ However, we only observed the multiple peaks in the region between δ = -110 and -117 ppm, arising from the Q⁴ Si atoms (Figure 3a).³⁹ Previously, Tsapatsis *et al.*⁴⁰ reported that tetrabutylphosphonium hydroxide (TBPOH) triggers the rotational intergrowth of single-unit cell MFI lamellae, leading to a house-of-cards arrangement with a hierarchical structure containing micropore and mesopore simultaneously. The Ar isotherm at 87 K proves a uniform microporous structure and the absence of mesopores in our case (Figure 3b). AFM was further used to detect the surface morphology of the membrane. Terraces with a thickness of 1.5±0.1 nm (Figure 3c) are clearly observed from the *b*-face, indicating terrace spreading rather than surface nucleation dominates crystal growth.⁴¹ However, hillocks with a height of 15 nm were observed in the case of TBPO-free solution (Figure 3d), which can be well explained by the nonclassical growth mechanism of Sicalite-1 (all silica MFI zeolite) based on the aggregation of metastable silica nanoparticle precursors.³⁷

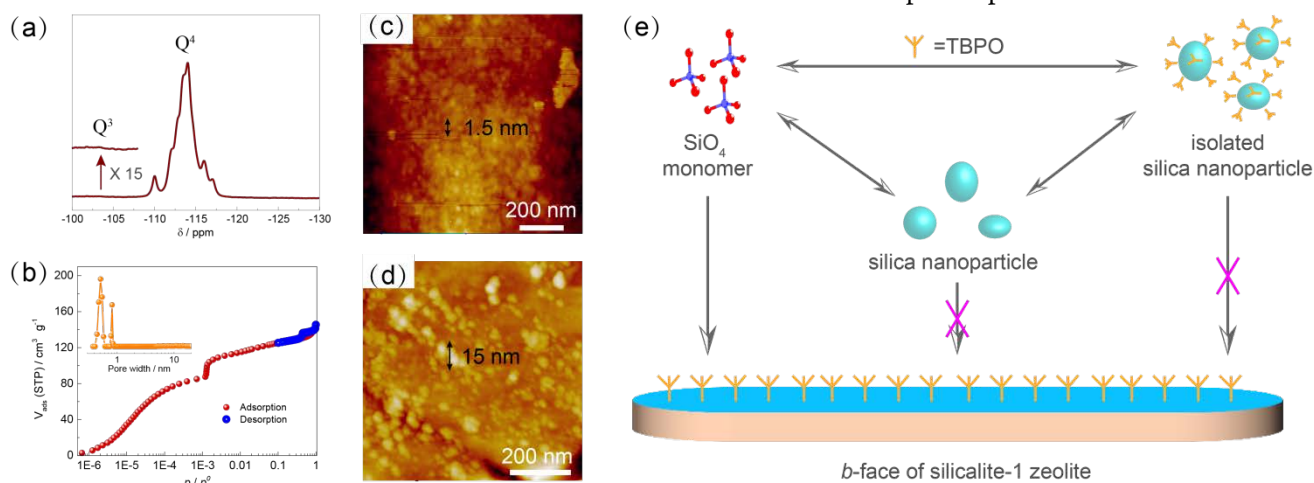


Figure 3. Mechanism of TBPO in the nucleation and growth of MFI crystals. (a-c) Magic angle spinning ²⁹Si NMR spectrum, Ar isotherm at 87 K, and AFM image of the *b*-face of MFI crystals synthesized from the solution containing 0.1 wt.% TBPO for 6 h; the steep step at $p/p^0=10^{-3}$ is arising from the fluid-to-crystalline-like phase transition of the adsorbed phase in the MFI micropores; (d) AFM image of the *b*-face of MFI crystals synthesized from TBPO-free solution for 3 h. Scale bars in white and black color indicate lateral and height dimension, respectively; (e) Illustration of the role of TBPO in MFI crystal nucleation and growth. Synthesis condition: 1 TEOS: 0.2 TPAOH: 110 H₂O at 150 °C.

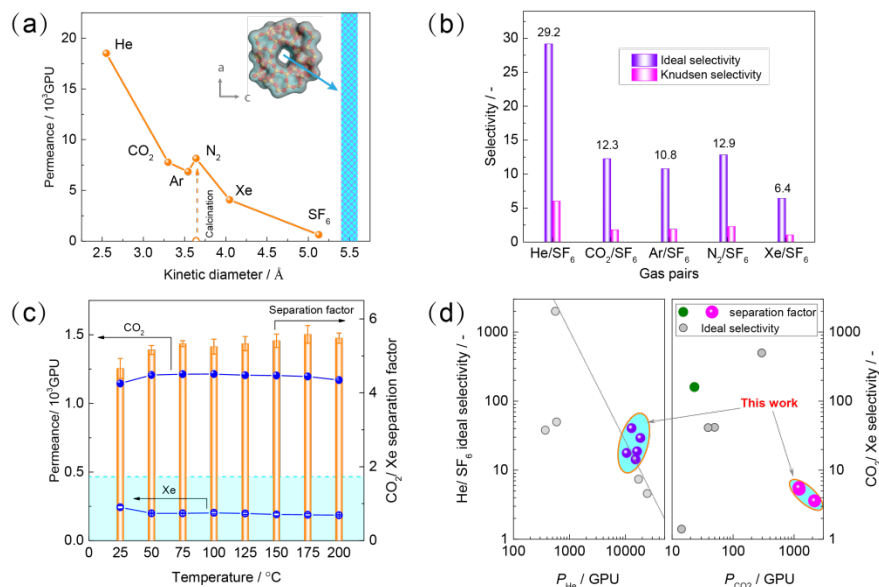


Figure 4. Separation performance of *b*-oriented MFI zeolite membranes. (a-b) single gas permeance and ideal selectivity of *b*-oriented MFI zeolite membrane M11 under an absolute feed pressure of 2 bar and room temperature; (c) temperature-dependent separation performance of *b*-oriented MFI zeolite membrane M11 for 50 / 50 CO₂/Xe mixture at an absolute feed pressure of 1.5 bar; the cyan colour indicates Knudsen selectivity of CO₂/Xe (1.7); (d) Comparison of five membrane performance in this work with other membranes, involving random-oriented MFI,⁴²⁻⁴⁴ SAPO-34,⁴⁵ carbon molecular sieving,⁴⁶ PIMs,⁴⁷ and PDMS⁴⁸ (raw data was shown in Table S3, S4, and S5).

Based on the above results, we propose the following role for TBPO (Figure 3e): TBPO molecules would cover the surface of wet silica nanoparticles by hydrogen bonding of the P=O groups to adsorbed water molecules and surface silanol groups. The maximal surface coverage of TBPO molecules on wet silica is as high as 43%.⁴⁹ The isolated silica nanoparticles would not participate in MFI zeolite nucleation and growth, thus, we speculate the isolation effect of TBPO leads to a lower concentration of active silica species and therefore delayed nucleation and mitigated twinning growth.²⁵ In this line, nucleation and intergrowth within the membrane layer were significantly slowed down in the presence of TPBO (Figure S7). This also explains why a longer synthesis time was used to achieve continuous *b*-oriented MFI membranes. Meanwhile, the *b*-face of MFI crystal is potentially favorable for the adsorption of ZGMs as proved by DFT calculation⁵⁰ and experimental results.³⁸ The adsorbed TBPO molecules on *b*-faces impede the addition of soluble silica units (classical mechanism) and agglomerated precursors (nonclassical mechanism) until they are replaced by SDAs, leading to a mitigated twinning growth and the enhancement of in-plane growth.^{38, 51} The absence of phosphorous in the membrane demonstrates that TBPO acts as a surface modifier rather than as SDA (XPS results, Figure S8).

3.4 Single gas permeation

N₂ permeance was as low as 1.4 GPU (1 GPU = 3.3928 × 10¹⁰ mol·m⁻²·s⁻¹·Pa⁻¹) for the SDAs-containing membrane, demonstrating absence of pin-holes (open symbol in Figure 4a). After complete removal of SDAs (Figure S9), single gas permeances decrease in the sequence of He, N₂, CO₂, Ar, Xe, and SF₆; they are independent on the *trans*-membrane pressure difference (Figure S10), indicating the absence of viscous flow. A clear cut-off was observed at a kinetic diameter of ~5 Å. The ideal selectivity of these light gas molecules over SF₆ is always

higher than Knudsen selectivity (Figure 4b, Table S3), further demonstrating a dense membrane layer. It was worth to note that the *in-situ* crystallization approach exhibited a good reproducibility to prepare *b*-oriented MFI zeolite membranes (Figure 4d), which is essential for the practical production.

3.5 Separation of CO₂/Xe mixture

Despite great achievements on separation of Xe from Kr or air by chabazite SAPO-34,^{45, 52-54} AIPO-18,⁵³ and ZIF-8 membranes,^{53, 55-56} CO₂/Xe mixture separation using microporous membranes has been overlooked in the past decades.⁴⁶ Herein, we propose a novel Xe recovery approach from exhaled anaesthetic gas based on *b*-oriented MFI zeolite membranes. This can be used to continuously remove the major impurity of CO₂ from the closed-circuit anesthesia system. As shown in Figure 4c, the separation factor of CO₂ to Xe was 5.6, higher than the ideal and Knudsen selectivity. The permeance of pure CO₂ exhibits a maximum as a function of temperature (Figure S11d), proving surface diffusion dominates CO₂ transport in MFI channels.⁵⁷ Like for other adsorptive gas mixtures in MFI,⁵⁸⁻⁵⁹ the competitive adsorption between CO₂ and Xe plays an important role in CO₂/Xe mixture separation. The preferential adsorption of Xe by MFI zeolite (Figure S12) would compromise CO₂ transport through the membranes, leading to much lower permeance than in case of single gas.^{45, 58} Since the mixed gas selectivity factor is higher than the ideal gas selectivity, the CO₂-selective separation is attributed to the high CO₂ diffusivity in MFI zeolite as demonstrated by breakthrough studies.^{4, 58} This is further supported by the slight increase of separation factor with temperature, which is the result of a stronger reduction in Xe adsorption because of its higher heat of adsorption than that of CO₂.⁴ The same situation applies for Kr/Xe separation in other nanoporous crystalline membranes, such as ZIF-8 and SAPO-34, as elucidated by molecular simulation.⁶⁰

The ideal selectivity of CO₂/Xe could be as high as ~500 for the benchmark 8-member ring zeolite membranes, e.g. SAPO-34;⁴⁵ however, the gas permeance would be significantly deteriorated by the mixed gas. For example, the permeance was mere 11.5 GPU⁴⁵ and 17.1 GPU⁵⁴ for Kr/Xe mixtures. The same situation occurred for carbon molecular sieving membranes⁴⁶ and polymeric membranes (e.g., PIM-1,⁴⁷ PIM-7,⁴⁷ and PDMS⁴⁸) for the separation of CO₂/Xe. The *b*-oriented MFI zeolite membrane in this work exhibited a CO₂ permeance of 1213 GPU, which is several orders of magnitude higher than the above membranes (Figure 4d). The higher permeation flux would significantly reduce the investment of membrane-based units and their footprint. The good reproducibility and long-term hydrothermal stability (>260 h, Figure S13) endow *b*-oriented MFI zeolite membranes a great potential for practical application of Xe recovery from exhaled anaesthetic gas.

4. CONCLUSIONS

In summary, we have reported a facile *in-situ* crystallization for the manufacture of *b*-oriented MFI zeolite membranes. The membrane exhibited an exceptional CO₂ permeation flux because of the short and oriented straight channels. The one-pot synthesis approach in this study paves a way to prepare *b*-oriented MFI zeolite membranes, showing great potential for application. The demonstrated Xe recovery from CO₂ allows its increased use as an anaesthetic in medical application.

ASSOCIATED CONTENT

Supporting Information. Detailed description of supporting characterizations (SEM, PXRD, XPS, FTIR, TGA, gas permeation property) used in the present study. This material is available free of charge via the Internet at <http://pubs.acs.org>.

AUTHOR INFORMATION

Corresponding Author

* Xuerui Wang, x.wang-12@tudelft.nl

* Jorge Gascon, jorge.gascon@kaust.edu.sa

* Freek Kapteijn, f.kapteijn@tudelft.nl

ORCID

Xuerui Wang: 0000-0003-2220-7531

Jorge Gascon: 0000-0001-7558-7123

Freek Kapteijn: 0000-0003-0575-7953

Author Contributions

The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

Notes

The authors declare no competing financial interest.

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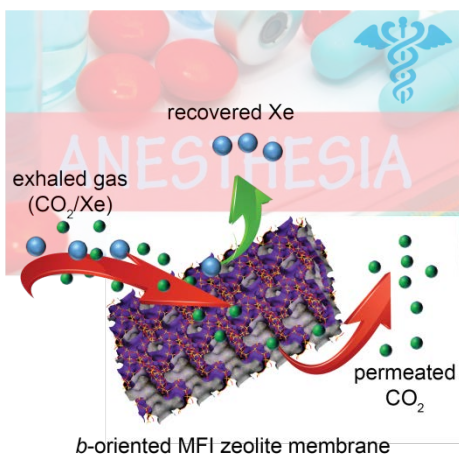
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