Novel Porphyrinic Porous Organic Frameworks for High Performance Separation of Small Hydrocarbons

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A series of Porphyrinic-CTF materials are synthesized under ionothermal condition with high surface areas (>3200 m²/g) and tunable pore sizes. The ZnP-CTFs exhibit high adsorption capacities and selectivity towards C₃H₈ and C₂H₆ over CH₄ as IAST prediction. Furthermore, we explore the utility of ZnP-CTF for gas chromatographic separation of the small hydrocarbons mixture based on their different van der waals interactions and polarizability. More importantly, the fast breakthrough test further proves that the ZnP-CTF-400 and ZnP-CTF-500 can separate the small hydrocarbons under kinetic dynamic conditions.

Small hydrocarbons (C₁, C₂ and C₃) are important energy sources and chemical feedstock in petrochemical industry. The main sources of these small hydrocarbons are natural gas and pyrolysis gas. Natural gas is actually a naturally gas mixture consisting primarily of methane, but including varying amounts of C₂H₆, C₃H₈. On the other hand, besides CH₄, the pyrolysis gas also contains C₂ ~ C₃ hydrocarbons and other hydrocarbons. Before transportation and utilization of these resources, they must undergo processing for upgrading and purification from their raw gases mixtures. Ordinarily, a simple temperature swing distillation can recover most of heavier hydrocarbons. The small hydrocarbons with low boiling points are generally purified via cryogenic distillation to obtain high-grade single-component gas product, which requires high energy consumption. The pressure swing adsorption (PSA) process is a cost-effective and highly efficient technique for separating small hydrocarbons from their mixture. In the PSA process, the separation efficiency basically depend on solid adsorbents with specific adsorption properties. Thus, extensive efforts have been made to develop efficient solid adsorbents for PSA technology.

Typically, PSA is carried out under kinetic dynamic conditions and run a cycle in several minutes. To meet this challenge, the adsorbent must possess a fast separation ability. Chemisorption based adsorbents are not really suitable for this process. Porous solid materials, such as zeolites, activated carbons have been proved highly impressive adsorbents for physical adsorption and separation. However, even these traditional adsorbents exhibit a low separation capacity and selectivity in the process of separating small hydrocarbons mixtures. Metal organic frameworks (MOFs) or porous coordination polymers (PCPs) have been exploited as a new type of sorbents for hydrocarbon separation. Recently, porous organic frameworks (POFs), assembled by organic building blocks via covalent bonds, have been proposed as new carriers for gas capture and storage. Because of their controllable pore connectivity, chemical tailorability, high stability and large adsorption capacity, POFs have been emerged as appropriate candidates for the PSA processes.

Porphyrin is a kind of interesting macrocycle ligand with rich coordination chemistry in the square-planar coordination site. A large number of metal ions could be readily combined into the porphyrin center, which could serve as open metal site in POFs. Herein, we report a series of novel Zn porphyrinic-CTF materials (ZnP-CTF) with high BET surface areas and tunable pore sizes. The ZnP-CTF materials exhibit high adsorption capacities for C₂H₆ and C₃H₈ over CH₄. In addition, the selectivities of C₂H₆ and C₃H₈ towards CH₄ are evaluated by ideal adsorption solution theory (IAST). Furthermore, the excellently stable POF material exhibits high performance in Gas Chromatographic (GC) and kinetic fast breakthrough separation of small hydrocarbon mixture.

Results

Synthesis and characterization of ZnP-CTFs. The ZnP-CTF materials were synthesized by the dynamic trimerization of 5,10,15,20-Tetrakis(4-cyanophenyl)porphyrin under ionothermal conditions. Under high temperature, the catalytic ZnCl₂ melts and then induces the polymerization of aromatic nitrile groups.
Meanwhile, the Zn ions could combine into the porphyrin ring center, and the Zn-porphyrin skeletons are produced (Figure 1). As documented, the character of the product was influenced by the temperature of polymerization. At high temperature polymerization, the irreversible carbonization reactions ensued, coupled with the reversible trimerization of nitriles, allowing the reorganization of the dynamic triazine network to form mesopore. To tune the pore size and surface area of ZnP-CTFs, the polymerizations were carried out at 400 °C, 500 °C and 600 °C for 40 h, respectively. The resulting products were denoted as ZnP-CTF-400, ZnP-CTF-500 and ZnP-CTF-600. The formation of polytriazine network was confirmed by FT-IR analysis (Figure S1). The disappearance of carbonitrile stretching band at 2228 cm⁻¹ and the appearance of an absorption band at around 1531 cm⁻¹ pointing to the characteristic of the triazine units, demonstrate the completion of the trimerization reaction. A new peak at 1005 cm⁻¹ which is attributed to the absorbance of Zn–N bond in metalloporphyrin, proves the successful combination of Zn ion into the porphyrin center. For the better illustration of chemical structures of this polymer, ZnP-CTF-400 was characterized by solid-state ¹³C and ¹⁵N cross-polarization magic-angle spinning (CP/MAS) NMR. As shown in Figure S2, the NMR signal of carbon around 169.6 ppm is observed, which is assigned to carbon atom in triazine rings. The carbon signals assigned to porphyrin are also detected in the spectrum (Figure S2a). The intact structure is additionally confirmed by the ¹⁵N NMR spectrum (Figure. S2b). The signal at 136.3 ppm can be ascribed to triazine moiety, the signal at 98.1 ppm is characteristic chemical shift of N atom in the porphyrin unit. Due to the decomposition of ZnP-CTFs at high temperature, we can not obtain a well-resolution carbon NMR signals for ZnP-CTF-500 and ZnP-CTF-600. Elemental analysis and Zn inductively coupled plasma (ICP) spectroscopy were performed to confirm the composition of ZnP-CTF skeletons. The weight percentages of C, H, N and Zn contents for ZnP-CTF-400 are close to the theoretical values (Table S1). Zn 2p3/2 XPS data reveal that most of the Zn ions are coordinated in porphyrin (Figure S3). When the temperature increased to 500 °C and 600 °C, the Zn content of ZnP-CTF materials decreased due to pyrolysis of porphyrin ring. In addition, the microstructure is studied by transmission electron microscopy (Figure 2). As can be seen, a worm-like porous texture is observed in ZnP-CTFs materials. All the ZnP-CTF materials are amorphous as confirmed by powder X-ray diffraction (PXRD) (Figure S4). Thermogravimetric analysis (TGA) of ZnP-CTF materials under air condition indicated that they have excellent thermal stability above 400 °C (Figure S5).

The porosity and pore structures of ZnP-CTFs were investigated by nitrogen sorption measurement at 77 K. As shown in Figure 3, ZnP-CTF-400 exhibits Type-I isotherm, which is the typical feature of microporous materials. When temperature increases to 600 °C, the isotherms of ZnP-CTF-600 displays type IV isotherm, indicating the formation of mesopores. The existence of mesopore can also be demonstrated by the plots of the NL-DFT pore size distributions of ZnP-CTF-600 (Figure S6). For clear illustration, a series of pore parameters derived from the nitrogen isotherms are listed in Table 1, including Brunauer-Emmett-Teller (BET) and Langmuir surface areas, pore sizes, total pore volumes and micropore volumes. The Langmuir surface area of ZnP-CTF-500 reaches 3218 m²/g, which is higher than some other porphyrin-based porous organic frameworks (Table S2).

Gas adsorption. Given ZnP-CTF materials have large surface area, abundant nitrogen atoms and micro- and mesoporous bi-modal pore system in the frameworks, we examined their CH₄, C₂H₆ and...
C$_3$H$_8$ adsorption isotherms at 273 K and 298 K, respectively. As shown in Figure 4a–4c and Figure S8, all gas adsorption isotherms are reversible, indicating the materials can be easily regenerated. At 298 K and 101 kPa, the C$_3$H$_8$ uptake for ZnP-CTF-400, ZnP-CTF-500 and ZnP-CTF-600 is 112 cm$^3$/g, 161 cm$^3$/g and 103 cm$^3$/g, respectively. The C$_2$H$_6$ uptake is 70 cm$^3$/g for ZnP-CTF-400, 90 cm$^3$/g for ZnP-CTF-500 and 54 cm$^3$/g for ZnP-CTF-600 under the same conditions. It is found that the ZnP-CTF materials show low adsorption capacity for CH$_4$ under this condition. The CH$_4$ uptake is 11 cm$^3$/g for ZnP-CTF-400, 15 cm$^3$/g for ZnP-CTF-500 and 9 cm$^3$/g for ZnP-CTF-600, respectively. Because ZnP-CTFs possess larger surface areas, they could take more C$_2$ and C$_3$ hydrocarbons than NaX zeolite and some previously reported MOFs but smaller than MOF-74 materials$^{11–15}$. Importantly, the uptake amounts at P = 101 kPa vary so much indicating that there is an odds for the purification of the small hydrocarbon mixtures according to the number of carbon atoms.

Based on the adsorption isotherms at 273 K and 298 K, the isosteric heats of adsorption (Q$_st$) of ZnP-CTFs are calculated to estimate their affinity with probe molecules. As shown in Figure 4d–4f, a
notably high Qs values for ZnP-CTF-400 (C3H8: 38–45 kJ/mol; C2H6: 25–30 kJ/mol and CH4: 19–25 kJ/mol) are obtained. The strong interaction between ZnP-CTF-400 and gas molecules could be attributed to uniform micropores in the framework. Comparing with ZnP-CTF-500 and ZnP-CTF-600, the smaller pore size of ZnP-CTF-400 would lead to an overlap of the van der Waals forces of neighboring gas molecules, thus giving rise to more stronger interaction.

Ideal adsorbed solution theory simulation. The adsorption selectivities of ZnP-CTF materials for C2H6 and C3H8 over CH4 in an equimolar gas mixture are evaluated by ideal adsorption solution theory (IAST). As shown in Figure 4g–4i, the selectivities of C3H8 towards CH4 are in 290 for ZnP-CTF-400, 513 for ZnP-CTF-500 and 733 for ZnP-CTF-600, at 298 K and 101 kPa, respectively. The high selectivities of ZnP-CTFs make the separation of C3H8 from natural gas highly feasible. The C2H6/CH4 selectivity of ZnP-CTF-400, ZnP-CTF-500 and ZnP-CTF-600 are 33, 26 and 25 under the same conditions (Table S3). We also investigate the C3H8/C2H6 separation ability at 298 K of ZnP-CTF materials. ZnP-CTF-600, which has more mesopores, exhibits a higher C3H8/C2H6 selectivity (15) than ZnP-CTF-500 (11) and ZnP-CTF-400 (9) under 101 kPa and 298 K.

Gas chromatographic separation. Encouraged by the adsorption interactions of ZnP-CTF materials for C3H8, C2H6 and CH4, together with their high selectivities, we also explored the ability of ZnP-CTF-400 for kinetic separation of the small hydrocarbon molecules in Gas Chromatographic. The GC separation of C1 ~ C3 are performed by a chromatographic column (180 × 2 mm) packed with 190 mg ZnP-CTF-400. As shown in Figure 5, the ZnP-CTF-400 column can clearly separate small hydrocarbons mixture due to their different carbon atoms. Both C2H6 and C3H8 achieve baseline separation from CH4. The ZnP-CTF-400 column gives high-resolution separation of three mixtures: C2H6/CH4 (Rs = 2.8), C3H8/CH4 (Rs = 7.6), C2H6/C3H8 (Rs = 4.3). The high-resolution of such columns can be explained as follows: the retention of analyte on the column mainly depends on their different van der Waals interactions with the porous ZnP-CTF skeletons. The long alkane C3H8 has strong van der Waals interactions with porous walls than CH4 and C2H638–33. On the other hand, the Zn ions in ZnP-CTF offer active sites to polarize gas molecule34,35, the polarizability of C3H8 is larger than CH4 and C2H6 (polarizability α in units of 10^-24 cm^3: α(C3H8) = 6.3 > α(C2H6) = 4.5 > α(CH4) = 2.6). The more polarizable molecules have strong interaction with ZnP-CTF material through dispersion forces. This behavior is similar to the open metal sites effect in MOFs coated GC stationary phase36–39.

Breakthrough experiments. Considering the potential use of ZnP-CTF material in practical separation processes, we should also evaluate it under kinetic dynamic conditions at operating temperatures and pressures. The breakthrough curve is a typical way of estimating the gas separation ability for adsorbents under flowing gas conditions, which are related to the PSA process40. To further evaluate the performances of ZnP-CTF materials in an actual adsorption-based separation process, we studied dynamic breakthrough separation experiments of gas mixture at ambient conditions. For a fast separation in the PSA process, the breakthrough tests of different hydrocarbon mixtures were performed using a fixed-bed column packed with ZnP-CTF-400 or ZnP-CTF-500. ZnP-CTF-400 possesses strong hydrocarbons adsorption enthalpy and ZnP-CTF-500 has large uptake capacities. The pressure in the column was maintained at 3.1 MPa which is relevant to real purification of natural gas and LNG (3–4 MPa). The breakthrough curves of ZnP-CTF-400 and ZnP-CTF-500 are shown in Figure 6. As expected from result of GC separation, in each case, CH4 penetrated through the bed firstly, whereas the higher hydrocarbons retained in adsorbent. The results of CH4/C2H6 mixture (CH4/C2H6 = 80:20 (vol)) test in the two columns with the space velocity of 5 min^-1 are shown in Figure 6a and 6d. After injection of the gas mixture, the appeared to be 100% pure CH4 was detected both in ZnP-CTF-400 and ZnP-CTF-500 columns. Upon saturation of gas adsorbed in ZnP-CTF-400 column, C2H6 "brokethrough" around 17 min while the ZnP-CTF-500 column can retain C2H6 until 21 min, which means ZnP-CTF-500 have a large adsorption capacity.
In the case of CH$_4$/C$_3$H$_8$ mixture (CH$_4$/C$_3$H$_8 = 90:10$) with space velocity of 6 min$^{-1}$, during 4–21 min, the ZnP-CTF-400 retained C$_3$H$_8$ and the outlet stream contained pure CH$_4$ (Figure 6b). After 23 min, the concentration of the outlet gas reached the feed gas composition, indicating the saturation of column. The ZnP-CTF-500 column can retain C$_3$H$_8$ for 25 min and pure CH$_4$ was collected during 5–25 min (Figure 6e), indicating both of packing beds can also separate CH$_4$ from CH$_4$/C$_3$H$_8$ mixture. To verify the two materials can separate of these small hydrocarbons in ternary mixed system, we measured breakthrough curves for the mixture of CH$_4$/C$_2$H$_6$/C$_3$H$_8$ (25:50:25) with space velocity of 6 min$^{-1}$.

As shown in Figure 6c and 6f, CH$_4$ with the weakest interaction with adsorbents breakthrough earliest and C$_3$H$_8$ with the strongest interaction with adsorbents retain for 18 min. Due to completely separation of CH$_4$ is achieved, it is possible to produce pure methane from this three-component mixture during the PSA process. The impressive performances of ZnP-CTF-400 and ZnP-CTF-500 in separation of the small hydrocarbons in ternary mixed system, we measured breakthrough curves for the mixture of CH$_4$/C$_2$H$_6$/C$_3$H$_8$ (25:50:25) with space velocity of 6 min$^{-1}$.

**Discussion**

In conclusion, we successfully synthesize novel ZnP-CTF materials through ionothermal reaction. By controlling synthesis temperature, the tunable surface areas and pore sizes of ZnP-CTF materials are achieved. More valuably, ZnP-CTF materials show different adsorption capacities for the small hydrocarbons according to the number of carbon atoms. The selective hydrocarbons adsorption of ZnP-CTF materials are demonstrated by IAST prediction. Furthermore, we explore the utility of ZnP-CTF-400 for high performance GC separation of small hydrocarbons mixture owing to their different van der Waals interactions and polarizability. The fast dynamic breakthrough tests further prove the ZnP-CTF-400 and ZnP-CTF-500 can separate the small hydrocarbons based on their hierarchical isosteric heats in an actual adsorption-based separation process. For the first time, we systematically investigated POFs for the hydrocarbons separation under both equilibrium and kinetic conditions. The high separation capability and selectivity offers ZnP-CTF-500 as a promising candidate for energy efficient separation of small hydrocarbons in PSA process.

**Methods**

**Materials.** All starting materials were purchased from commercial suppliers and used without further purification unless otherwise noted. Anhydrous ZnCl$_2$ was purchased from Alfa Aesar. 5,10,15,20-Tetrakis(4-cyanophenyl)porphyrin$^{41}$ was synthesized according to the previously reported method.

**Synthesis of ZnP-CTFs.** 5,10,15,20-Tetrakis(4-cyanophenyl)porphyrin (1 mmol) and ZnCl$_2$ (8 mmol) were transferred into a quartz ampule (3 $\times$ 12 cm) under an inert atmosphere. The ampule was evacuated, sealed, and heated to the desired temperature (400, 500, 600 °C) for 40 h. The ampule was then cooled to room temperature and opened. The reaction mixture was subsequently ground and then stirred with water for 24 h. Further stirring in 0.1 M HCl for 15 h was carried out to remove the residual salt. After this purification step, the resulting black powder was filtered, washed successively with water and CH$_3$OH, and dried in vacuum at 150 °C. ZnP-CTFs were isolated in 85–90% yield.
Ideal adsorption solution theory calculations. The ideal adsorption solution theory (IAST) was used to predict the binary mixture adsorption from the experimental pure-gas isotherms. Previous studies have shown that the IAST can predict mixture gas adsorption in many nanoporous materials, including porous-organic frameworks. The single-component isotherms were fitted using a dual-site Langmuir-Freundlich equation (1):

\[
q = q_{\text{m}1} \frac{b_1 p^{1/n_1}}{1 + b_1 p^{1/n_1}} + q_{\text{m}2} \frac{b_2 p^{1/n_2}}{1 + b_2 p^{1/n_2}}
\]

Here, \(p\) is the pressure of the bulk gas at equilibrium with the adsorbed phase (kPa), \(q\) is the adsorbed amount per mass of adsorbent (mmol/g), \(q_{\text{m}1}\) and \(q_{\text{m}2}\) are the saturation capacities of sites 1 and 2 (mmol/g), \(b_1\) and \(b_2\) are the affinity coefficients of sites 1 and 2 (1/kPa), and \(n_1\) and \(n_2\) represent the deviations from an ideal homogeneous surface.

The IAST adsorption selectivity, \(S\), for binary mixtures of \(CO_2(1)/N_2(2)\), is defined as follows equation (2):

\[
S = \frac{q_1}{q_1 + q_2} \left( \frac{p_1}{p_2} \right)^{n_1} \left( \frac{p_2}{p_1} \right)^{n_2}
\]

where \(q_1\) and \(p_1\) (i = 1, 2) are the mole fractions of component 1 and 2 in the adsorbed and bulk phases, respectively. The IAST calculations were carried out for binary mixture containing equimolar gas.

Details of the breakthrough curve measurement. Breakthrough curve measurements were performed using a home-made gas-flow system. In practice, the experiment is conducted as follows: The adsorbent ZnP-CTF-400 was filled into the column with a length of 150 mm and an internal diameter of 5 mm. Before test, the column was activated in flow of Ar for 24 h. The pressure in the column was maintained at 3.1 MPa. The experiments were performed at 298 K. For the CH4/C2H6 column was activated in flow of Ar for 24 h. The pressure in the column was experiment is conducted as follows: The adsorbent ZnP-CTF-400 was filled into the details of the breakthrough curve measurement.

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Author contributions
H.M. designed and performed most of experiments. S.M., G.Z. and F.S. analyzed date. H.R. and H.M. co-wrote the paper. All authors contributed to results analysis and discussion. All authors reviewed the manuscript.

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