

Introduction of π -Complexation into Porous Aromatic Framework for Highly Selective Adsorption of Ethylene over Ethane

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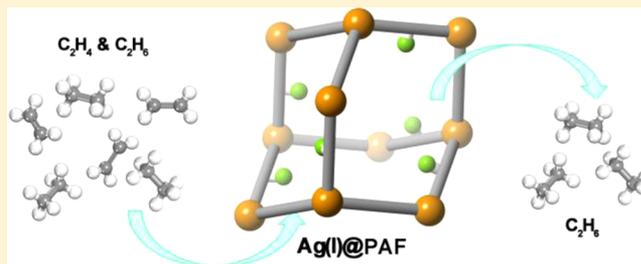
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ABSTRACT: In this work, we demonstrate for the first time the introduction of π -complexation into a porous aromatic framework (PAF), affording significant increase in ethylene uptake capacity, as illustrated in the context of Ag(I) ion functionalized PAF-1, PAF-1-SO₃Ag. IAST calculations using single-component-isotherm data and an equimolar ethylene/ethane ratio at 296 K reveal that PAF-1-SO₃Ag shows exceptionally high ethylene/ethane adsorption selectivity (S_{ads} : 27 to 125), far surpassing benchmark zeolite and any other MOF reported in literature. The formation of π -complexation between ethylene molecules and Ag(I) ions in PAF-1-SO₃Ag has been evidenced by the high isosteric heats of adsorption of C₂H₄ and also proved by in situ IR spectroscopy studies. Transient breakthrough experiments, supported by simulations, indicate the feasibility of PAF-1-SO₃Ag for producing 99.95%+ pure C₂H₄ in a Pressure Swing Adsorption operation. Our work herein thus suggests a new perspective to functionalizing PAFs and other types of advanced porous materials for highly selective adsorption of ethylene over ethane.



INTRODUCTION

Ethylene, one of the most widely used feedstock molecules in the petrochemical industry, is usually obtained via steam cracking and thermal decomposition of ethane.¹ The similar molecular sizes and volatilities make the separation of ethylene/ethane mixtures one of the most challenging chemical separations at large scale.² Current technology uses cryogenic distillation performed under the conditions of high pressure (23 bar) and low temperature (−25 °C), resulting in an extremely cost and energy intensive process.³ Extensive efforts to develop low energy approaches for efficient ethylene/ethane separation at higher temperature and normal atmospheric pressure have focused on membrane separation,⁴ organic solvent-based sorbents,⁵ and porous solid adsorbents.⁶ Among these approaches, porous solid adsorbents attract particular interest because of their great potential to afford much lower cost and energy consumption.

Over the past decade, advanced porous materials such as metal–organic frameworks (MOFs)⁷ and porous organic polymers (POPs)⁸ [e.g., porous aromatic frameworks (PAFs),⁹ conjugated microporous polymers (CMPs),¹⁰ porous polymer networks (PPNs),¹¹ and porous organic frameworks (POFs)¹²] have been explored as new classes of solid adsorbents for applications in gas storage,¹³ gas separation,¹⁴ carbon capture,¹⁵ catalysis,¹⁶ and so forth. Compared with conventional solid adsorbents of zeolites and mesoporous silica materials,¹⁷ MOFs⁷ and POPs⁸ feature the amenability of design and modular nature, adjustable pore sizes, functionalizable pore surfaces, and high surface areas. These features also make them hold great promise for hydrocarbon separation,¹⁸ including the separation of ethylene/ethane mixtures.¹⁹

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Adsorption-based separation of ethylene/ethane using MOFs focuses on the preferential interactions between open metal sites and ethylene molecules, and high ethylene uptake capacities and ethylene/ethane selectivities have been demonstrated in MOFs with open metal sites.^{19a–f}

In comparison with MOFs, POPs, despite the amorphous nature for most of them, feature robust covalent framework structures showing high water, moisture, and chemical stability;⁸ they could also be readily scaled up using one-pot reactions. However, the lack of preferential binding sites for ethylene molecules leads to poor ethylene/ethane adsorption selectivity.^{8c,19g} A recently reported copper(catecholate) decorated POP demonstrates enhanced ethylene/ethane selectivity;²⁰ nevertheless, the absolute selectivity remains low, presumably due to the moderate interactions between open Cu(II) sites and ethylene molecules. Therefore, in order to achieve high ethylene/ethane selectivity in POPs, stronger binding sites for ethylene molecules are desired.

It has been well-documented that Cu(I) and Ag(I) ions can form π -complexation with the carbon–carbon double bonds of olefin molecules in solutions,^{5,6,21} and these systems have been employed for absorptive separations of olefins from paraffins, which are however inefficient because of the poor contact between the hydrocarbons and the liquid absorbents.⁵ We postulate that if such kinds of π -complexation can be introduced into POP, the π -complexation will afford strong interactions between the ethylene molecules and the framework, whereas the porous structure of POP can maximize the contact of between the ethylene molecules and the framework, thereby resulting in high ethylene/ethane selectivity. In this contribution, we demonstrate for the first time the introduction of π -complexation into POPs, as illustrated in the context of functionalizing the highly porous PAF, PAF-1²² with Ag(I) ions. The resultant PAF-1-SO₃Ag not only exhibits significant enhancement of ethylene uptake capacity compared to the parent PAF-1, but also demonstrates exceptional ethylene/ethane adsorption selectivity, far surpassing benchmark zeolite and any other MOF and POP reported thus far. The formation of π -complexation between ethylene molecules and Ag(I) ions in PAF-1-SO₃Ag has been proven by heat of adsorption analysis and in situ IR spectroscopic studies.

RESULTS AND DISCUSSION

Materials Preparation and Physicochemical Characterization. PAF-1²² [(cross-linked poly tetraphenylmethane) also known as (a.k.a.) PPN-6²³] is an amorphous POP possessing a hypothetical diamondoid-topology structure with very high surface area and exceptional stability in water/moisture and acidic/basic media. PAF-1-SO₃Ag can be readily achieved by Ag(I) ion exchange of sulfonate-grafted PAF-1 (hereafter denoted PAF-1-SO₃H) following the procedures reported previously (Supporting Information (SI) Scheme S1).^{23a,24a}

N₂ gas sorption isotherms at 77 K (Figure 1) reveal Brunauer–Emmett–Teller (BET) surface areas of 4714, 1087, and 783 m²·g⁻¹ for PAF-1, PAF-1-SO₃H, and PAF-1-SO₃Ag, respectively. Pore size distribution analysis (Horvath–Kawazoe model) indicates that the pore size is reduced from ~15 Å for PAF-1 to ~8 Å for PAF-1-SO₃H, whereas the pore size of PAF-1-SO₃Ag is predominantly distributed around ~8 Å, suggesting negligible pore size change after the Ag(I) ion exchange process (SI Figure S1).

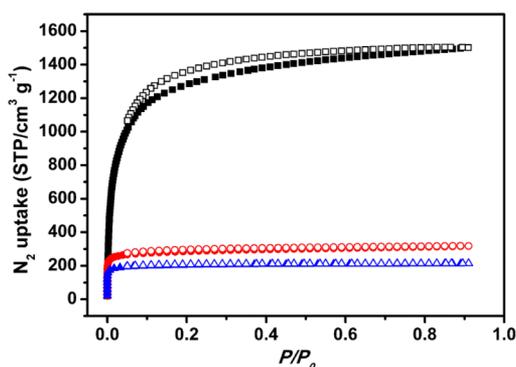


Figure 1. N₂ sorption isotherms at 77 K for PAF-1 (black), PAF-1-SO₃H (red), and PAF-1-SO₃Ag (blue).

The presence of Ag(I) in PAF-1-SO₃Ag was confirmed by X-ray photoelectron spectroscopy (XPS) analysis, which shows a silver signal at binding energies of 368.8 and 374.8 eV (SI Figure S2) corresponding to the peaks of Ag 3d_{5/2} and Ag 3d_{3/2}, respectively. Fourier transform infrared spectroscopy (FTIR) of PAF-1-SO₃Ag shows the obvious characteristic peak of SO₃⁻ group at 1086 and 1186 cm⁻¹, respectively (SI Figure S3). Solid ¹³C NMR spectra of PAF-1-SO₃Ag and PAF-1-SO₃H show similar central carbon atom signals at δ = 65 ppm and the signals of aromatic carbon (δ = 121 ppm to 147 ppm), indicating the preservation of framework structure after Ag(I) ion exchange (SI Figure S4). Inductively coupled plasma mass spectrometry (ICP-MS) and elemental analysis (EA) indicate that ~50% SO₃H were exchanged into SO₃Ag.

Ethylene and Ethane Adsorption. The low-pressure ethylene sorption isotherms were collected at 296 K. The incorporation of Ag(I) ion into PAF-1 results in a significant enhancement of ethylene adsorption capacity despite the remarkable decrease in surface area. At 296 K and 1 atm, the ethylene uptake amounts of PAF-1 and PAF-1-SO₃H are 57 and 66 cm³·g⁻¹, respectively (SI Figure S5). In contrast, PAF-1-SO₃Ag exhibits a significantly higher ethylene uptake capacity of 91 cm³·g⁻¹ (4.1 mmol·g⁻¹) under the same conditions (Figure 2). PAF-1-SO₃Ag surpasses the ethylene uptake capacity of zeolite 5A²⁵ (~2.3 mmol·g⁻¹ at 303 K and 1 atm) and compares to that of zeolite NaX²⁶ (~4.2 mmol·g⁻¹ at 305 K and 1 atm), two benchmark zeolites widely studied for ethylene/ethane separation. In addition, PAF-1-SO₃Ag outperforms the copper(catecholate) decorated POP, CuA₁₀B₁,²⁰ in ethylene uptake, which exhibits an ethylene adsorption amount

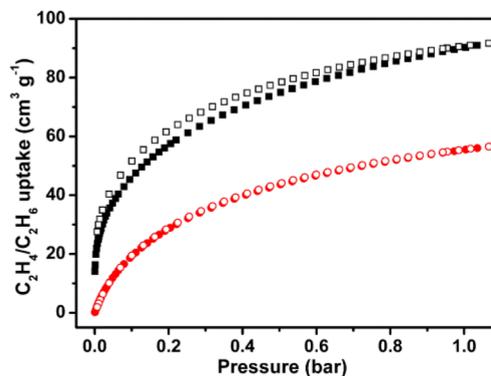


Figure 2. C₂H₄ (black) and C₂H₆ (red) sorption isotherms of PAF-1-SO₃Ag at 296 K. Filled: adsorption; unfilled: desorption.

of $\sim 1.8 \text{ mmol}\cdot\text{g}^{-1}$ at 0.79 atm and 298 K. The ethylene uptake capacity of PAF-1-SO₃Ag at 296 K and 1 atm is relatively lower compared to that of some high surface area MOFs possessing open metal sites (e.g., $7.2 \text{ mmol}\cdot\text{g}^{-1}$ of MgMOF-74, $7.2 \text{ mmol}\cdot\text{g}^{-1}$ of Cu-BTC, and $5.8 \text{ mmol}\cdot\text{g}^{-1}$ of NOTT-102),^{19a} but the MOFs containing open metal sites usually experience partial framework degradation after exposed to moisture, inevitably leading to drastic decreases in ethylene uptake capacity upon reuse. In contrast, the fact that PAF-1-SO₃Ag was prepared via ion exchange in aqueous solution suggests its water stability. This, together with its moisture stability, is further confirmed by the reproducibility of the ethylene sorption isotherms for PAF-1-SO₃Ag even after exposure to an air environment with 80% humidity for 2 days (SI Figure S6).

To test the recyclability of PAF-1-SO₃Ag, we simulated temperature and vacuum swings with an ASAP2020 analyzer, by saturating with ethylene up to 1.1 bar at 296 K followed by a high vacuum for 3 h at 105 °C. After 5 cycles, there was no apparent loss in capacity (SI Figure S7), indicating the complete desorption during each regeneration cycle. Upon the basis of the differential scanning calorimetry (DSC) analysis, energies of 2.67 MJ/kg are needed to release ethylene and regenerate PAF-1-SO₃Ag (SI Figure S8).^{19b}

Interestingly, different from the ethylene adsorption, the trend of ethane uptake by the three samples follows the order of PAF-1 > PAF-1-SO₃H > PAF-1-SO₃Ag at 296 K and 1 atm (SI Figure S9). The smallest ethane uptake amount observed for PAF-1-SO₃Ag is primarily attributed to its lower surface area when compared with PAF-1 and PAF-1-SO₃H. This result also suggests that the incorporation of Ag(I) ions would not increase the ethane uptake capacity.

Ethylene/ethane adsorption selectivities were calculated using ideal adsorbed solution theory (IAST)²⁷ for PAF-1-SO₃Ag, PAF-1, and PAF-1-SO₃H (Figure 3). For an equimolar

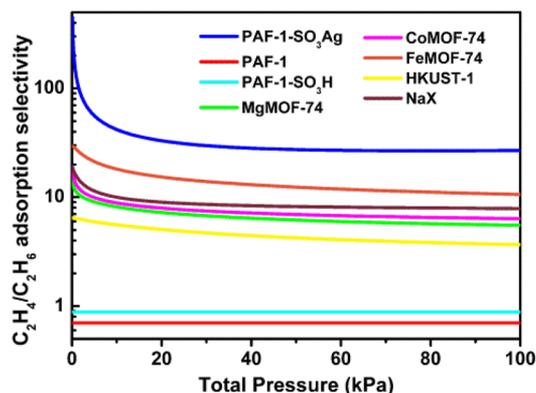


Figure 3. Comparison of the IAST calculations for C₂H₄/C₂H₆ adsorption selectivities for PAF-1-SO₃Ag with PAF-1, PAF-1-SO₃H and other porous materials^{19a} at 296 K.

mixture of ethylene and ethane at 296 K, the adsorption selectivity (S_{ads}) obtained for PAF-1-SO₃Ag is 27 at 100 kPa, far exceeding those calculated for both PAF-1 ($S_{\text{ads}} = 0.7$) and PAF-1-SO₃H ($S_{\text{ads}} = 0.88$). The ethylene/ethane adsorption selectivity of PAF-1-SO₃Ag at 296 K and 100 kPa is also significantly higher than those of zeolite NaX,^{19a} the MOFs^{19a} FeMOF-74 [a.k.a. Fe₂(dobdc)], CoMOF-74 [a.k.a. Co₂(dobdc)], MgMOF-74 [a.k.a. Mg₂(dobdc)], CuBTC (a.k.a. HKUST-1) (Figure 3), and the POP CuA₁₀B₁,²⁰ exhibiting ethylene/ethane selectivities of 8, 11, 6.4, 5.6, 3.6,

and 3.8, respectively. It is worth noting that the ethylene/ethane adsorption selectivities of PAF-1-SO₃Ag are considerably higher than those of zeolite NaX and other MOFs over the entire pressure range with the adsorption selectivity value at 1 kPa ($S_{\text{ads}} = 125$) even about an order of magnitude higher (Figure 3).

In practice, the combination of adsorption selectivity and uptake capacity of gas mixtures contribute to the characteristics of ethylene/ethane separation.^{18c,19a} Figure 4 shows the IAST

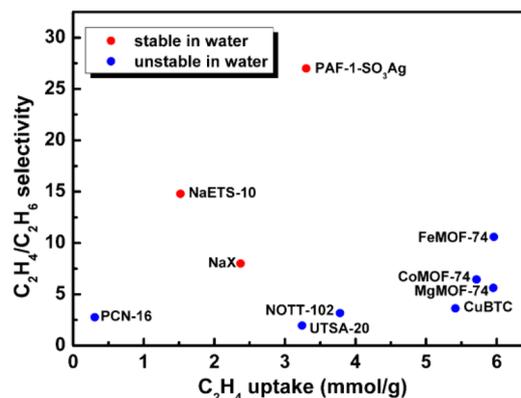


Figure 4. IAST calculations of the C₂H₄/C₂H₆ adsorption selectivity versus the gravimetric uptake capacity of ethylene for adsorption from an equimolar C₂H₄/C₂H₆ mixture at the total bulk gas phase at 296 K and 100 kPa^{19a} (Note: the uptake capacity of ethylene for FeMOF-74 is at 318 K^{19b}).

calculations of the ethylene/ethane adsorption selectivity versus the gravimetric uptake capacity of ethylene for adsorption from an equimolar ethylene/ethane mixture at the total bulk gas phase at 296 K and 100 kPa for PAF-1-SO₃Ag and several benchmark microporous adsorbent materials.^{19a} Both adsorption selectivity and gravimetric uptake capacity of PAF-1-SO₃Ag are significantly higher than two important zeolites of NaETS-10²⁸ and NaX.^{19a,29} The volumetric ethylene uptake capacity of PAF-1-SO₃Ag (SI Figure S11), which is estimated based on the density of the compressed PAF-1-SO₃Ag pellet, also surpasses that of NaETS-10²⁸ and NaX.^{19a,29} Albeit the ethylene uptake capacity of PAF-1-SO₃Ag is lower than that of some MOF materials, much higher ethylene adsorption selectivity alongside excellent water stability represent advantages in practice over most MOFs³⁰ investigated so far.

Ethylene–Framework Interactions. We reasoned that the exceptional ethylene adsorption properties of PAF-1-SO₃Ag should stem from the strong interactions between ethylene molecules and the framework of PAF-1-SO₃Ag as a result of the formation of π -complexation between the d orbitals of Ag(I) and the π orbitals of carbon–carbon double bonds in ethylene.^{5,6} We estimated the isosteric heats of adsorption (Q_{st}) based upon Clausius–Clapeyron equation by differentiation of the dual-Langmuir–Freundlich fits of the isotherms at two different temperatures,^{18c,19a} 296 and 318 K (SI Figure S12) with T -dependent parameters. As shown in Figure 5, at close to zero loading, the Q_{st} for ethylene in PAF-1-SO₃Ag is 106 kJ·mol⁻¹, remarkably higher than that of PAF-1 (14 kJ·mol⁻¹) and PAF-1-SO₃H (23 kJ·mol⁻¹). The Q_{st} for ethylene in PAF-1-SO₃Ag is consistent with that observed in other Ag(I)-based π -complexation systems,^{21,31} suggesting the formation of π -complexation between the ethylene molecules and Ag(I) ions in PAF-1-SO₃Ag. The Q_{st} exceeds that in MOFs with open

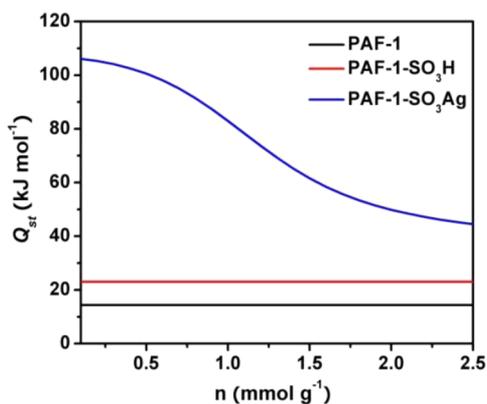


Figure 5. Isothermic heats adsorption, Q_{st} of C_2H_4 for PAF-1, PAF-1-SO₃H, and PAF-1-SO₃Ag.

metal sites, e.g., FeMOF-74 (45 $\text{kJ}\cdot\text{mol}^{-1}$)^{19b} or (47 $\text{kJ}\cdot\text{mol}^{-1}$),^{19a} MgMOF-74 (42 $\text{kJ}\cdot\text{mol}^{-1}$),^{19a} CoMOF-74 (41 $\text{kJ}\cdot\text{mol}^{-1}$),^{19a} CuBTC (39 $\text{kJ}\cdot\text{mol}^{-1}$).^{19a} These results highlight that, compared with open metal sites, Ag(I) ions can boost the interactions with ethylene molecules more in a porous framework via the formation of π -complexation. In contrast with the high Q_{st} for ethylene, PAF-1-SO₃Ag shows a significantly lower Q_{st} for ethane with a value of 27 $\text{kJ}\cdot\text{mol}^{-1}$ (SI Figure S14); thus validating that the Ag(I) ions serve as a preferential binding sites, selectively adsorbing ethylene over ethane thereby resulting in high ethylene/ethane adsorption selectivities.

To further prove the formation of π -complexation between the ethylene molecules and Ag(I) ions in PAF-1-SO₃Ag, in situ IR measurements of ethylene adsorption at room temperature were conducted. The $-\text{CH}_2$ out-of plane wagging mode at 949 cm^{-1} was found as the most sensitive mode, responding to the interaction between ethylene and the substrate surface.³² As shown in Figure 6, ethylene adsorption on PAF-1 and PAF-1-

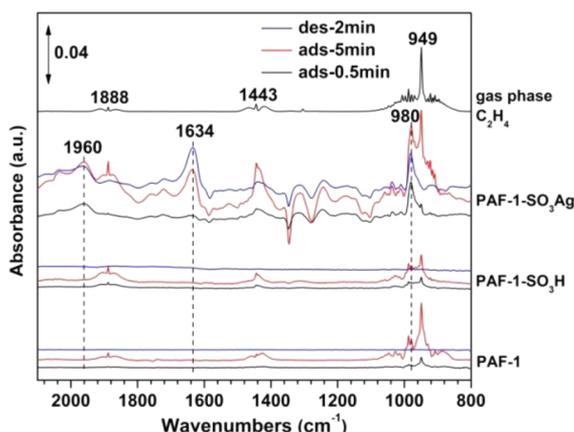


Figure 6. IR spectra from ethylene adsorption and desorption on PAF-1, PAF-1-SO₃H, and PAF-1-SO₃Ag at room temperature. IR spectrum from gas phase ethylene is also shown for reference.

SO₃H exhibits IR features similar to that of gas phase C_2H_4 , indicating a weak interaction, which is further evidenced by the complete removal of ethylene IR features after room temperature desorption in helium purge. In contrast, upon initial adsorption, PAF-1-SO₃Ag shows strongly perturbed CH_2 mode at 980 cm^{-1} . The intensity is even comparable with the gas-phase mode at 949 cm^{-1} at saturation. Two extra IR features at

1960 cm^{-1} (combination mode of $-\text{CH}_2$ wagging) and 1634 cm^{-1} ($\text{C}=\text{C}$ stretching), not observed on the PAF-1 and PAF-1-SO₃H, further confirm ethylene adsorption on PAF-1-SO₃Ag. These new IR bands due to adsorbed ethylene persist well after room temperature desorption, indicating a strong interaction between ethylene and PAF-1-SO₃Ag. The blue-shift of the $-\text{CH}_2$ wagging mode can be attributed to the combinative $d-\pi$ and $d-\pi^*$ interaction between Ag and ethylene,^{33–36} thus confirming the formation of π -complexation between the ethylene and Ag(I) ions in PAF-1-SO₃Ag.

Ethylene/Ethane Breakthrough Experiments and Simulations. To evaluate the performance of PAF-1-SO₃Ag in an actual adsorption-based separation process, breakthrough experiments were performed in which an equimolar ethylene/ethane mixture was flowed over a packed bed of the solid with a total flow of 2 mL/min at 296 K. As shown in Figure 7, PAF-1-SO₃Ag can effectively separate an equimolar mixture of ethylene and ethane into the pure component gases of greater than 99% purity.

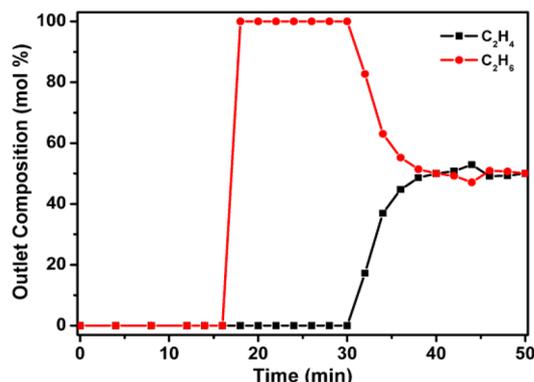


Figure 7. Experimental data on transient breakthrough of an equimolar C_2H_4/C_2H_6 mixture in an adsorber bed packed with PAF-1-SO₃Ag in the adsorption phase of a PSA operation.

We also carried out breakthrough simulations for C_2H_4/C_2H_6 mixtures in a fixed bed (SI Figure S17) to further demonstrate the feasibility of producing 99.95%+ pure C_2H_4 in a Pressure Swing Adsorption (PSA) operation. The simulated breakthrough curves are in reasonably good agreement with the experimental data (SI Figure S18a). During the adsorption cycle, C_2H_6 at purities >99% can be recovered for a certain duration of the adsorption cycle, as indicated by the arrow in SI Figure S18b. In addition, ethylene of 99.95%+ purity, required as feedstock to the polymerization reactor, can also be recovered during the time interval indicated by the arrow in SI Figure S19 in the desorption cycle. Video animations of the breakthrough simulations can be viewed in the HTML version of this work.

CONCLUSIONS

In summary, we have demonstrated for the first time the introduction of π -complexation into POPs for highly selective adsorption of ethylene over ethane, as illustrated in the context of Ag(I) ion functionalized porous aromatic framework, PAF-1-SO₃Ag. PAF-1-SO₃Ag exhibits significantly higher ethylene/ethane adsorption selectivity at 296 K than benchmark zeolite and any other MOF and POP reported in literature. The high ethylene/ethane adsorption selectivity of PAF-1-SO₃Ag is traceable to the formation of π -complexation between Ag(I)

ions and the double bonds of ethylene molecules, which is reflected in the high isosteric heats of adsorption of C_2H_4 and evidenced by in situ IR spectroscopy studies. The feasibility of PAF-1- SO_3Ag for producing 99.95%+ pure C_2H_4 in a PSA operation has been demonstrated by breakthrough experiments that are supported by simulations. Albeit the light-sensitivity, utilization of costly Ni-COD catalyst, and high isosteric heats of adsorption could represent some potential challenges for the application of PAF-1- SO_3Ag in practice, these kinds of issues could be tackled via some engineering processes. Notwithstanding, our work presented herein provides a new perspective to functionalizing POPs for energy-saving ethylene/ethane and other olefin/paraffin separations. Ongoing work in our laboratory includes investigating PAF-1- SO_3Ag for separations of other olefin/paraffin mixtures and applying the approach of π -complexation to functionalizing other types of advanced porous materials for hydrocarbon separations.

EXPERIMENTAL SECTION

Materials and Syntheses. All starting materials, reagents, and solvents were purchased from commercial sources (Aldrich, Alfa, Fisher, and Acros) and used without further purification.

Synthesis of Tetrakis(4-bromophenyl)methane. Tetrakis(4-bromophenyl)methane was synthesized according to the procedures reported in the literature²² with some minor modification. To a three-necked round-bottom flask containing bromine (6.4 mL, 19.9 g), tetraphenylmethane (2.0 g, 6.24 mmol) was added stepwise with small portions under vigorous stirring at room temperature (25 °C). After the addition was completed, the resulting solution was stirred for 60 min and then cooled to 0 °C. At 0 °C temperature, ethanol (25 mL) was added slowly, and the reaction mixture was allowed to warm to room temperature overnight. Then, the precipitate was filtered off and washed subsequently with saturated aqueous sodium hydrogensulfite solution (25 mL) and water (100 mL). After drying at 80 °C for 24 h under vacuum (80 mbar), tetrakis(4-bromophenyl) methane was recrystallized in EtOH/ CH_2Cl_2 to afford a yellow solid, yield: 88%.

Synthesis of PAF-1. PAF-1 was synthesized according to the procedures reported in the literature^{23a} with some minor modification. Tetrakis(4-bromophenyl)methane (509 mg, 0.8 mmol) was added to a solution of 2,2'-bipyridyl (565 mg, 3.65 mmol), bis(1,5-cyclooctadiene)nickel(0) (1.0 g, 3.65 mmol), and 1,5-cyclooctadiene (0.45 mL, 3.65 mmol) in anhydrous DMF/THF (60 mL/90 mL), and the mixture was stirred overnight at room temperature under nitrogen atmosphere. After the reaction, 6 M HCl (60 mL) was added slowly, and the resulting mixture was stirred for 12 h. The precipitate was collected by filtration, then washed with methanol and water, and dried at 150 °C for 24 h under vacuum (80 mbar) to produce PAF-1 as a white powder, yield: 80%.

Synthesis of PAF-1- SO_3H . PAF-1- SO_3H was synthesized according to the procedures reported in the literature^{23a,24} with some minor modification. To an ice-cooled mixture of PAF-1 (100 mg) in dichloromethane (15 mL), chlorosulfonic acid (1.0 mL) was added dropwise. The resulting mixture was stirred at room temperature for 3 days. Then, the mixture was poured over ice, and the solid was collected, washed with water thoroughly, and dried 150 °C for 24 h under vacuum (80 mbar) to produce PAF-1- SO_3H as blue powder, yield: 96%.

Synthesis of PAF-1- SO_3Ag . To the 15 mL CH_3CN/H_2O (1:1) solution, 100 mg PAF-1- SO_3H and 800 mg $AgBF_4$ were added. The mixture was stirred under room temperature for 48 h, and then the solid was collected by filtration followed by washing with CH_3CN and water. The whole process was performed carefully under dark environment. This exchange process was repeated three times, and then dried at 110 °C under vacuum (80 mbar) for further test, yield: 94%. EA: C: 47.25%; H: 3.19%; N: 0.53%; S: 17.11%; ICP-MS: Ag: 29.20%.

Gas Adsorption. Gas sorption measurements were performed using an ASAP 2020 volumetric adsorption analyzer. High-purity

grade gases of N_2 (99.999%), C_2H_4 (99.5%), and C_2H_6 (99.5%) were used for the collection of respective sorption isotherms.

In Situ IR Experiments. IR spectra of ethylene adsorption were collected using a Thermo Nicolet Nexus 670 spectrometer in diffuse reflectance mode (DRIFTS). The PAF-1- SO_3Ag sample, ca. 5 mg, was treated in a DRIFTS cell (HC-900, Pike Technologies) at 423 K in helium (30 mL/min) for 1 h to removal water and other adsorbates. The sample was then cooled down to room temperature for ethylene adsorption. The adsorption was conducted by flowing 10% ethylene/He (30 mL/min) over the sample for 5 min and then desorption was done in flowing helium. IR spectra were recorded continuously to follow the surface changes during the adsorption and desorption process. All reported IR spectra are difference spectra referenced to a background spectrum collected at room temperature after pretreatment but prior to ethylene adsorption.

Fitting of Pure Component Isotherms. The measured experimental isotherm data for C_2H_4 , and C_2H_6 on PAF-1- SO_3Ag were fitted with the dual-Langmuir–Freundlich isotherm model:

$$q = q_{A,sat} \frac{b_A p^{1/\nu_A}}{1 + b_A p^{1/\nu_A}} + q_{B,sat} \frac{b_B p^{1/\nu_B}}{1 + b_B p^{1/\nu_B}} \quad (1)$$

The fit parameters for C_2H_4 and C_2H_6 are specified in SI Table S1. SI Figure S13 presents a comparison of the experimentally determined component loadings for C_2H_4 and C_2H_6 on PAF-1- SO_3Ag at 296 K with the isotherm fits using parameters specified in SI Table S1. The fits are excellent over the entire range of pressures.

The pure component isotherm data for PAF-1, and PAF-1- SO_3H could be fitted with single site Langmuir model; the fit parameters are provided in SI Tables S2 and S3, respectively.

Calculations of Adsorption Selectivity. The selectivity of preferential adsorption of C_2H_4 (component 1) over C_2H_6 (component 2) in a mixture containing 1 and 2, can be formally defined as follows:

$$S_{ads} = \frac{q_1/q_2}{p_1/p_2} \quad (2)$$

In eq 2, q_1 and q_2 are the component loadings of the adsorbed phase in the mixture. The calculations of S_{ads} are based on the use of the Ideal Adsorbed Solution Theory (IAST) of Myers and Prausnitz.²⁷

Estimation of Isothermic Heats of Adsorption, Q_{st} . The isosteric heat of adsorption, Q_{st} , were calculated using the Clausius–Clapeyron equation by differentiation of the dual-Langmuir–Freundlich fits of the isotherms at two different temperatures, 296 and 318 K with T -dependent parameters.

$$Q_{st} = RT^2 \left(\frac{\partial \ln p}{\partial T} \right)_q \quad (3)$$

Breakthrough Experiments. In a typical experiment, 400 mg of PAF-1- SO_3Ag was swiftly ground and packed into a quartz column (6 mm I.D. \times 220 mm) with silica wool filling the void space. The sample was in situ activated under vacuum (6.5×10^{-4} Pa) at 110 °C for 2 h. Then, Helium flow (2 mL/min) was introduced the system to purge the adsorbent until the temperature of the column was decreased to 23 °C. The breakthrough test was started by introducing a 1:1 C_2H_4/C_2H_6 mixture gas at a total flow rate of 2.0 mL/min and switching off the He gas. Effluent from the column was monitored using a GC with a flame ionization detector. The dead volume of this setup was determined to be 18.6 cm^3 .

ASSOCIATED CONTENT

Supporting Information

Characterization details, additional gas sorption isotherms, simulated breakthrough curves, TGA plots, XPS and IR spectra plots, and supporting figures. This material is available free of charge via the Internet at <http://pubs.acs.org>.

W Web-Enhanced Feature

Video animations of transient breakthrough for the adsorption/desorption cycle are available in the HTML version of this paper.

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Notes

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