Impact of Post-Synthesis Modification of Nanoporous Organic Frameworks on Selective Carbon Dioxide Capture

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IMPACT OF POST-SYNTHESIS MODIFICATION OF NANOPOROUS ORGANIC FRAMEWORKS ON SELECTIVE CARBON DIOXIDE CAPTURE

A thesis submitted in partial fulfillment of the requirements for the degree of Master of Science at Virginia Commonwealth University.

by

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List of Abbreviations

BET ................................................................................. Brunauer–Emmett–Teller
BILP .............................................................................. Benzimidazole-Linked Polymer
BLP ................................................................................. Borazine-Linked Polymer
CCS .............................................................................. Carbon Capture and Sequestration
CMP .............................................................................. Conjugated Microporous Polymer
COF .............................................................................. Covalent Organic Framework
DCM ............................................................................... Dichloromethane
DFT .............................................................................. Density Functional Theory
FT-IR .............................................................................. Fourier Transform-Infrared Spectroscopy
IAST .............................................................................. Ideal Adsorbed Solution Theory
MEA ............................................................................... Monoethanolamine
MOF ............................................................................... Metal-Organic Framework
NLDFT ........................................................................... Non-Local Density Functional Theory
NMR ............................................................................... Nuclear Magnetic Resonance Spectroscopy
NPOF .............................................................................. Nanoporous Organic Framework
PAF ............................................................................... Porous Aromatic Framework
PECONF ................................................................. Porous Electron-Rich Covalent Organo Nitridic Framework
PIM ............................................................................... Polymers of Intrinsic Microporosity
POP ............................................................................... Porous Organic Polymer
PPN.................................................................Porous Polymer Network
PSM .....................................................................Post-Synthesis Modification
SEM .................................................................Scanning Electron Microscopy (Microscope)
TGA ....................................................................Thermogravimetric Analysis
XRD......................................................................X-Ray Diffraction
Abstract

IMPACT OF POST-SYNTHESIS MODIFICATION OF NANOPOROUS ORGANIC FRAMEWORKS ON SELECTIVE CARBON DIOXIDE CAPTURE

By Timur İslamoğlu, M.Sc.

A thesis submitted in partial fulfillment of the requirements for the degree of Master of Science at Virginia Commonwealth University

Virginia Commonwealth University, 2013

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Porous organic polymers containing nitrogen-rich building units are among the most promising materials for selective CO₂ capture and separation applications that impact the environment and the quality of methane and hydrogen fuels. The work described herein describes post-synthesis modification of Nanoporous Organic Frameworks (NPOFs) and its impact on gas storage and selective CO₂ capture. The synthesis of NPOF-4 was accomplished via a catalysed cyclotrimerization reaction of 1,3,5,7-tetrakis(4-acetylphenyl)adamantane in Ethanol/Xylenes mixture using SiCl₄ as a catalyst. NPOF-4 is microporous and has high surface area (SBET = 1249 m² g⁻¹). Post-synthesis modification of NPOF-4 by nitration afforded (NPOF-4-NO₂) and subsequent reduction resulted in an amine-functionalized framework (NPOF-4-NH₂) that exhibits improved gas storage capacities and high CO₂/N₂ (139) and CO₂/CH₄ (15) selectivities compared to NPOF-4 under ambient conditions. These results demonstrate the impact
of nitro- and amine- pore decoration on the function of porous organic materials in gas storage and separation application.
Chapter 1

Introduction

1.1 Global CO₂ Concerns

The increase in CO₂ concentration in the atmosphere is an urgent global problem that needs to be addressed immediately. Over the last fifty years, CO₂ concentration has increased from about 310 ppm to over 390 ppm and CO₂ emission from power plants has been identified as one of major contributors.¹ Currently, ~85% of the world's energy comes from fossil fuel sources and realistically, this situation will not change until sustainable and cleaner energy sources and technologies become widely available at a reasonable cost which will take time.² An alternative way to reduce CO₂ emission is switching from coal-fired power plants to natural gas-fired power plants since natural gas has a lower carbon footprint. For the same reason, the use of natural gas in automotive applications is highly desired; however, natural gas found in reservoirs has tangible amounts of CO₂, N₂, and H₂S that need to be minimized before transport and use. Natural gas purification processes involve CO₂ and H₂S removal to prevent pipeline corrosion and to increase the energy density of this methane-rich gas making its onboard storage and use more effective. As a result of the economical and environmental impacts of gaseous impurities, selective CO₂ capture and sequestration (CCS) have attained great attention over the past decades. Particularly, the selective separation of CO₂ from N₂, CH₄, and H₂ has gained interest since carbon dioxide is the
main impurity in flue gases (gases that exit to the atmosphere via an exhaust port such as smoke stacks from power plants) as well as natural gas. Advancements in this area by the scientific and industrial communities can limit the release of this greenhouse gas to the atmosphere and can lead to the production of high purity natural gas in a more economically and friendly manner.\textsuperscript{3,5}

1.2 CO\textsubscript{2} Capture Technologies

Different technologies and associated materials exist for CO\textsubscript{2} separation and captures as schematically illustrated in Figure 1.1. With the current technology, the most widely employed method to capture CO\textsubscript{2} from flue gas is by chemisorption where CO\textsubscript{2} is chemically absorbed by solutions that contain amine organic solvent (typically 30 wt\% in water) such as monoethanolamine (MEA) which is the most common amine compound used. In such systems, the flue gas is scrubbed with an amine solution, where up to 98\% of CO\textsubscript{2} can be captured.\textsuperscript{6} The CO\textsubscript{2}-laden solvent is then pumped to a second vessel where heat is applied to regenerate the amine solution by releasing absorbed CO\textsubscript{2}. Even though this process is the current state-of-the-art technology, there are several concerns because of the enormous energy required for solvent regeneration. This concern stems from the fact that the regeneration of amine solutions consumes great amount of energy due to the high heat capacity of water. Moreover, the volatile nature of amines in addition to their decomposition during regeneration processes result in the loss of absorbent over time. In addition, amine sorbents themselves are corrosive, not stable in the presence of sulfur containing compounds, and during regeneration when the sorbent is heated, they can be oxidized by oxygen present it the gas stream.\textsuperscript{7} Recent studies have indicated that electricity cost would increase by \textasciitilde80\% if CO\textsubscript{2} were
to be captured by aqueous MEA absorption processes that eliminate CO$_2$ from the flue gas of coal-fired power plants and such processes can consume about 40% of the energy output of the plant.$^8$ The expected properties of an ideal liquid absorbent can be summarized as high capture capacity and selectivity towards the targeted gas, regeneration with a minimal amount of input energy, thermal and chemical stability, and a straightforward synthetic route from accessible precursors. Today's inventories of adsorbents for carbon dioxide capture application, unfortunately, are far from combining two or more of these properties together.$^9$
Membrane-based CO$_2$ separation has also been studied extensively as an efficient way to capture CO$_2$. Membranes can be prepared from a wide range of materials such as organic polymers, zeolites, ceramics and they can be porous or non-porous. The driving force for CO$_2$ in the system can be achieved by having higher partial pressure on one side of the membrane relative to the other side which can be obtained by pressurizing the one side and/or applying vacuum on the other side. Membranes cannot be optimized for large volume of gas separation since they cannot usually achieve high degree of separation. Therefore, multiple treatments are
necessary and also they suffer from sulfur and other trace contaminants. In addition, their feasibility decreases when CO₂ concentration in the feed stream are below 20%.

Another approach to capture CO₂ from flue gas is the use of porous solids. With this approach, the regeneration energy penalty is greatly reduced compared to MEA solution process because water is no longer part of the capturing process. The development of efficient systems with solid adsorbents depends on the design of new materials with high stability, reversible and high adsorption capacity, and high selectivity under industrial conditions. Zeolites and activated carbon have also been extensively studied for CO₂ capture. Even though zeolites, porous alumina silicate materials, perform better than MEA absorbents, their low CO₂ capacity and instability in the presence of water limit their usage. Activated carbon shows better CO₂ adsorption capacity compared to zeolites, but it generally suffers from low selectivity of CO₂ over N₂ and CH₄. Among the emerging alternatives that can mitigate the aforementioned problems are Metal-Organic Frameworks (MOFs). MOFs are crystalline hybrid materials that combine organic ligands (linkers) and metal ions (nodes) and they have attracted intense research interest due to their tunable chemical and physical properties. Their extremely high surface area, permanent porosity, thermal stability, well-defined structures, and tunable pore functionality have made them plausible adsorbents for CO₂ and other gases such as methane and hydrogen. However, the limited chemical stability of MOFs under practical conditions due to non-covalent nature of framework significantly limits their application to CO₂ capture and separation processes. To address this limitation, porous organic polymers (POPs), have been identified as alternatives for carbon capture from flue gas as well as natural gas sweetening. POPs
are highly cross-linked, amorphous polymers possessing nanopores with high surface area. Unlike MOFs, POPs are metal-free purely organic frameworks linked by covalent bonds and display remarkable physicochemical stability. Due to their stable covalent nature, POPs are in general not subject to hydrolysis in the presence of water and remain intact under acidic or basic conditions. Under thermodynamic and kinetic controls, POPs can be designed to form crystalline networks known as covalent-organic frameworks (COFs) such as those linked by B-O, C-N, and B-N bonds.\textsuperscript{15-17} It should be mentioned that these COFs have limited stability under aqueous conditions and lose their porosity. In contrast, POPs’ physical and chemical stability allow for their chemical modification, a process that imparts desirable properties such as pore hydrophobicity which is central for gas separation and storage, sensing, and catalysis.

1.3 Porous Solid Adsorbents for CO\textsubscript{2} Capture

Porous materials, including POPs, are classified depending on the size of the pores. According to IUPAC classification, microporous solids have pore width not exceeding 2 nm; mesoporous materials are materials that exhibit pore width ranging from 2 - 50 nm, and macroporous materials having pores larger than 50 nm.\textsuperscript{18}

Synthetic porous materials (MOFs, COFs, POPs, etc) have attracted considerable attention due to their tunable chemical and physical nature that is not accessible for other porous materials such as zeolites, porous silica, and activated carbon. They can be processed easily; for instance, they can be produced in a molded monolithic form that makes them very attractive for use in a wide range of applications. In addition, some soluble porous polymers can be processed by solvent-based
techniques without losing their porosity, in contrast to those of naturally occurring porous materials listed above.\textsuperscript{19}

The list for synthetic porous organic materials includes those developed by El-Kaderi such as Benzaimidazole-Linked Polymers (BILP),\textsuperscript{20} Borazine-Linked Polymers (BLPs),\textsuperscript{21,22} and Nanoporous Organic Frameworks (NPOF),\textsuperscript{23} and many other systems: Porous Aromatic Framework (PAF),\textsuperscript{24} Porous Polymer Networks (PPN),\textsuperscript{25} Polymer Intrinsic Microporosity (PIM),\textsuperscript{26} Covalent Triazine Frameworks (CTZ),\textsuperscript{27} and Conjugated Microporous Polymers (CMP).\textsuperscript{28} Almost all synthetic strategies to obtain porous materials employ either ditopic or polytopic functional organic linkers in order to attain extended frameworks. Desired textural properties such as extremely high surface area, permanent porosity, and narrow pore size distribution can be achieved by varying the molecular building units during synthesis. Combined together, these parameters dictate the structure-function relationship in POPs and determine their effectiveness in CO\textsubscript{2} capture applications.

In order for a porous organic polymer to be useful for carbon capture from flue gas as well as natural gas stock, the following criteria must be met.\textsuperscript{6}

- The adsorbent needs to have high CO\textsubscript{2} adsorption capacity in order to make adsorbent column smaller.
- The adsorbent needs to have high selectivity toward CO\textsubscript{2}. Since CO\textsubscript{2} is not the major component, highly selective adsorbent will increase the quality of CO\textsubscript{2} capture.
- The adsorbent needs to be chemically stable since reactive chemicals such as oxygen and sulfur are also present as contaminants.
• The adsorbent needs to have mechanical stability in order to be densely packed for maximum volumetric capacity.
• The adsorbent itself and building blocks should be readily synthesized from abundant and accessible chemicals to make the process cost-effective.
• The regeneration of adsorbent should be achieved with minimal additional energy input to decrease the energy penalty of the total system.
• The mass and heat transfer of adsorbent should be good. Densely packed adsorbent should allow the adsorption and desorption of the guest molecules. In addition, heat conductivity is needed since temperature can be used to regenerate the adsorbents.

1.4 Synthesis of Porous Organic Materials

Several synthetic methods have been reported for the synthesis of porous organic materials and they mainly entail metal catalysts. Other methods are metal-free processes and represent a more “economical” way for the mass production of porous adsorbents. Probably, having many different routes to synthesize POPs is one of the most attractive features of this field. These synthetic routes include Yamamoto coupling,24 Suzuki coupling,29 imidazole ring formation,30 Friedel-Crafts alkylation,31 dibenzo-dioxane formation,32 imidization,33 amidation,33 Sonogashira-Hagihara coupling,28 imine formation,34 nitrile cyclotrimerization27 and acetyl cyclotrimerization.35

In the following section, representative synthetic methodologies of porous organic materials are presented to highlight some of their advantages and shortcomings. One of the most extensively used methods for the synthesis of POPs is metal catalyzed cross-coupling reactions. In 2009, Qiu et al. reported a new class of POPs referred as PAF-1
(Porous Aromatic Framework).\textsuperscript{24} PAF-1 was synthesized by Yamamoto coupling (Scheme 1.1) using tetra(bromo phenyl)methane as building block and it has an exceptionally high BET surface area of 5600 m\textsuperscript{2} g\textsuperscript{-1} and remarkable gas uptake capacity (H\textsubscript{2}, CH\textsubscript{4}, CO\textsubscript{2}) under high pressure. Due to its robust covalent nature, PAF-1 showed high thermal and chemical stability. However, carbon-based PAFs do not perform well in gas storage applications at low pressure (1.0 bar) due to their uniform and non-functionalized pore walls that lead to low isosteric heats of adsorption for small gases.

**Scheme 1.1:** Yamamoto coupling formation of PAF-1 and PPN-4.

Zhou and coworkers have synthesized several PPNs by employing the same method. They have extended the library of PAFs and modified their synthetic procedures that resulted in one of the highest surface area (SA\textsubscript{BET} = 6461 m\textsuperscript{2} g\textsuperscript{-1}) known to date as reported for PPN-4 (Scheme 1.1).\textsuperscript{36} The very high surface area of PAF-1 and PPN-4 was attributed to the diamond-like framework topology which provides accessible pores with highly cross-linked structures. Another key point in the synthesis of PPN-4 was that the C-C coupling was carried out at room temperature to slow down the kinetic of the reaction and hence the overall network growth.
Very recently, El-Kaderi and coworkers introduced a new class of porous organic material called Benzaimidazole-Linked Polymers (BILPs),\textsuperscript{20} prepared by metal-free catalyzed condensation of aryl-o-diamine and arylaldehyde that resulted imidazole-ring formation (Scheme 1.2). Both experimental and theoretical studies have proven that having polar functional groups on pore surfaces increases the CO\textsubscript{2} uptake dramatically, especially at low pressures, and makes the material more selective for CO\textsubscript{2} over N\textsubscript{2} and CH\textsubscript{4}. As such, the imidazole-functionalized pore walls in BILPs make this type of material one of the most attractive candidates for CO\textsubscript{2} capture application due to their selective nature and high CO\textsubscript{2} capacity. BILPs showed CO\textsubscript{2} uptake as high as 5.4 mmol g\textsuperscript{-1} at 273 K and 1 bar and very good selectivity as high as 113 for CO\textsubscript{2} over N\textsubscript{2} and 17 for CO\textsubscript{2} over CH\textsubscript{4}.\textsuperscript{37}

**Scheme 1.2:** A representative route for the synthesis of BILPs.
Kaskel and coworkers published a report where cyclotrimerization of bifunctional acetyl compounds were employed to synthesize porous organic polymers as illustrated in Scheme 1.3.\textsuperscript{35}

Scheme 1.3: Cyclotrimerization of acetyl groups to construct porous framework.

The polymers reported by Kaskel are hydrophobic in nature and exhibit only moderate surface area (up to 865 m\textsuperscript{2} g\textsuperscript{-1}). Unlike PAFs, PPNs, and CMPs where cross-coupling reactions were used for the construction of the frameworks, Kaskel's polymerization strategy does not employ expensive or precious catalysts and therefore, is more viable for large-scale production processes. However, non-functional carbon based frameworks are less desirable for CO\textsubscript{2} capture due to their low affinity and low selectivity at ambient settings.

As mentioned above, selective CO\textsubscript{2} capture and capacity are the most important features for CO\textsubscript{2} adsorbents. High surface area materials do not necessarily lead to high CO\textsubscript{2} uptake since gas uptake capacity depends on the favorable interactions between pore walls and guest gas molecules. Among the new directions for enhancing these factors has been pore functionalization with polar groups such as -OH, -NH\textsubscript{2}, -
SO$_3$, etc. Incorporation of these functionalities into the framework can be achieved by pre-synthesis modification (Figure 1.2) of the building blocks wherein CO$_2$-philic moieties are attached to building blocks before framework formation, automated functionalization as in the case of imidazole ring formation, or through post-synthesis modification (PSM) (Figure 1.3) processes in which functional groups are tethered to the pore walls after framework assembly.$^{38}$

**Figure 1.2:** Schematic representation of pre-synthesis modification. Adopted from reference 38.

**Figure 1.3:** Schematic representation of post-synthesis modification. Adopted from reference 38.
1.5 Post-Synthesis Modification of NPOFs

Significant efforts have been devoted to the development of functional porous materials for use in gas storage and separation applications. Post-synthesis modification (PSM) is one way of synthesizing these functional NPOFs. The chemical and thermal stability of NPOFs have allowed researchers to introduce heterogeneity to the surface of the pores by PSM even under harsh reaction conditions in order to enhance the CO$_2$ uptake capacity as well as selectivity. Even though PSM of NPOFs have been studied more recently it is still in its early stage compared to MOFs. As stated above, targeted framework functionalities within the pores can be engineered through the functionalization of building block before the network formation which is known as pre-synthesis functionalization. However, there are some problems associated with this method. First, functionalization of building blocks often necessitates alternate reaction conditions (i.e. protection of some functional groups), and these reactions are often time consuming and non-trivial. In addition, the synthesis of NPOFs typically requires harsh conditions such as high temperature, acidic or basic media wherein the functional groups on the building blocks may not survive or lead to side reactions. Therefore, developing new methods for PSM of NPOFs is highly desirable.
Zhou and coworkers have shown the successful incorporation of sulfonic acid and its lithium salt within the pores of PPN-6 (Scheme 1.4).\(^{39,40}\) PPN-6-SO_3H and PPN-6-SO_3Li exhibited significant increase in isosteric heat of adsorption and showed a substantial increase in the uptakes of H_2 and CO_2.\(^{39}\) This dramatic increase in the heat of adsorption and CO_2 uptake capacity is attributed to strong electrostatic interactions between sulfonic acid/CO_2 and lithium/CO_2. The same group reported another study on PSM of PPNs. The latter work described PPN-6 functionalization with alkyl amine chains and the modified networks resulted in an excellent CO_2 uptake (4.3 mmol g\(^{-1}\)) and selectivity of CO_2/N_2 (442, calculated by Ideal Adsorbed Solution Theory) at 1 bar and 295 K (Scheme 1.4).\(^{40}\)
1.6 Thesis Problem

A significant need exists for improving CO₂ capture from flue gas as CO₂ emission from power plants is the main cause for increasing CO₂ concentration in the atmosphere. Employing porous solid adsorbents to capture CO₂ is one of the most promising techniques; however, selective adsorption of CO₂ may not be possible if the framework does not have functional groups within the pores to attract CO₂. In addition, integrating these functionalities into the pores to engineer their chemical and physical environment is not always possible by using pre-functionalized building blocks. With these concerns in mind, the specific aims of the work described herein are summarized below:

1) To prepare highly porous networks that are chemically and physically robust, using viable and cost effective methods. The chemical nature of these materials should enable PSM processes to improve gas binding and selectivity.

2) Post-synthesis modification of the frameworks described above should offer moderate binding affinities that lead to high CO₂ capacity while permitting adsorbent regeneration without energy input in the form of heating.

3) Establish experimental methods to investigate the physical and textural properties of the frameworks before and after PSM and evaluate their performance in small gas uptake and selective binding.
Chapter 2

Synthesis and Characterization of a Novel Nanoporous Organic Framework and its Post-Synthesis Modification

2.1 Introduction

For a porous solid to be employed in flue gas CO$_2$ capture application, chemical and physical stability is also as important as CO$_2$ capture capacity because a typical flue gas sample is composed of 73-77% N$_2$, 7.4-15% CO$_2$, 6.2-14.6% H$_2$O, ~420 ppm SO$_2$, 50-300 ppm CO, 60-70 ppm NO$_x$, and ~4.45 ppm O$_2$ which depends on whether it is from gas-fired or coal-fired flue gas.$^{41}$ Therefore, adsorbents are expected to maintain their porosity in the presence of these gases. In this regard, purely organic porous polymers are superior to MOFs because of their chemical stability under harsh conditions due to their covalent nature. By taking these concerns into consideration we chose tetraphenyl adamantane (TPA) as the starting building unit to synthesize NPOF-4 because of the rigid nature of the TPA core. TPA then was acetylated to afford 1,3,5,7-tetrakis(4-acetylphenyl)adamantane (TAPA). The latter was used to construct NPOF-4 through acetyl groups’ cyclotrimerization. The tetrahedral structure of TPA can lead to the formation of 3-D frameworks that possess high accessible free volume for gas storage and separation. Post-synthesis modification of NPOF-4 was needed in order to improve selective CO$_2$ capture and storage. NPOF-4 functionalization was successfully
achieved by nitration of the benzene rings directly attached to the adamantane core, and by subsequent reduction of the nitro groups to amines.

2.2 Synthesis

Starting materials and solvents, unless otherwise noted, were obtained from Acros Organics and used without further purification. 1,3,5,7-Tetraphenyladamantane was synthesized according to published method.\textsuperscript{42} SiCl\textsubscript{4} was purchased from Sigma-Aldrich. SnCl\textsubscript{2} was purchased from MP Biomedicals, LLC. Absolute ethanol was purchased from Pharmaco Products. Solvents were dried by distillation from Na(Xylenes) or Na/benzophenone (THF). Air-sensitive samples and reactions were handled under an inert atmosphere of nitrogen using either glovebox or Schlenk line techniques.

2.2.1 Synthesis of Building Block

The synthetic strategy for the preparation of TAPA is summarized in Scheme 2.1. A 250 mL reaction flask was charged with AlCl\textsubscript{3} (16.4 g, 123 mmol), placed in an ice bath and 90 mL of acetyl chloride (AcCl) was added to afford a colorless solution. A mixture of tetraphenyl adamantane (2.50 g, 5.67 mmol) and freshly distilled DCM (100 mL) was added to this solution at 0 °C and stirred for 10 minutes at 0 °C, and then it was allowed to warm to room temperature and stirred for 16 h. The reaction mixture was slowly poured into ice (200 mL) while stirring and then was stirred further for 30 minutes at room temperature. The organic layer was extracted with DCM, and wash with 10 % NaHCO\textsubscript{3} solution. The resulting yellow solution was dried over MgSO\textsubscript{4}, and filtered over glass frit. Excess DCM was removed by evaporation under reduced pressure. The resulting product was crystallized from Chloroform/Ethanol mixture to
afford off-white crystals in 68% yield (2.35 g, 3.86 mmol) upon concentration. $^1$H NMR (CDCl$_3$, 300 MHz) $\delta$ (ppm) 7.98 (d, $J_{HH} = 6.0$, 8H), 7.595 (d, $J_{HH} = 9.0$, 8H), 2.61 (s, 12H), 2.24 (s, 12H). $^{13}$C NMR (CDCl$_3$, 75 MHz) $\delta$ (ppm) 197.9, 154.0, 135.7, 128.9, 125.5, 46.8, 39.8, 26.9. Anal. Calcd. for C$_{42}$H$_{40}$O$_4$0.5CHCl$_3$: C, 76.36; H, 6.11. Found: C, 76.12; H, 6.07. TOF MS ES+ m/z for C$_{42}$H$_{40}$O$_4$ Calcd 609.30, [M+H]$^+$ 609.35. 43

**Scheme 2.1:** Reaction scheme for synthesis of TAPA, NPOF-4, NPOF-4-NO$_2$, and NPOF-4-NH$_2$. 
2.2.2 Synthesis of Nanoporous Organic Framework (NPOF-4)

The synthetic strategy for the preparation of NPOF-4 is summarized in Scheme 2.1. A 100 mL reaction flask was charged with TAPA (300 mg, 0.493 mmol), 60 mL absolute Ethanol and 60 mL Xylenes (dried over Na). The colorless solution was cooled to 0 °C, and SiCl₄ (18.3 mL, 157.8 mmol) was added dropwise. The solution was allowed to stir for 30 minutes at 0 °C, and then 1 hour at room temperature giving a black colored solution which was further refluxed at 80 °C for 16 h. The solution was allowed to cool down to room temperature and then was filtered over a glass frit. The bright yellow powder was washed with Ethanol, 2.0 M HCl, 2.0 M NaOH, water, and DCM, and then dried at 70 °C in the oven to afford NPOF-4 (240 mg, 90% yield) as a yellow powder. Anal. Calcd. for C₁₂₆H₉₆: C, 93.99; H, 6.01. Found: C, 83.09; H, 6.11.³⁵

2.2.3 Post-Synthesis Nitration of NPOF-4

The synthetic strategy for the post-synthesis nitration of NPOF-4 is summarized in Scheme 2.1. A 50 mL round bottom flask was charged with 12.5 mL of concentrated H₂SO₄, and then cooled to 0 °C. To this solution, 180 mg NPOF-4 was added in small portions followed by a dropwise addition of 9 mL 70% HNO₃, and was stirred for 6 h at 0 °C. The mixture was then poured into 100 mL ice. The resulting mixture was stirred for 30 minutes at room temperature, and then washed with water and EtOH. The resulting brown powder was soaked in ethanol/water mixture for 16 h and then was filtered and dried in air at 80 °C for 12 h to give NPOF-4-NO₂ (225 mg, 93% yield) as a brown powder. Anal. Calcd. for C₁₂₆H₈₄N₁₂O₂₄: C, 70.39; H, 3.94; N, 7.82. Found: C, 67.52; H, 4.61; N, 7.84. ⁴⁴
2.2.4 Post-Synthesis Reduction of NPOF-4-NO₂

The synthetic strategy for the post-synthesis reduction of NPOF-4-NO₂ is summarized in Scheme 2.1. A 100 mL reaction flask was charged with 210 mg NPOF-4-NO₂, 50 mL Ethanol, and 3.0 g SnCl₂. The mixture was refluxed at 70 °C for 18 h. The resulting green suspension was filtered, then suspended in 20 mL concentrated HCl and stirred for 16 h at room temperature. Then HCl was replenished and stirring continued for 5 h at room temperature. The mixture was filtered and washed with 2 M NaOH solution, then sufficient amount of water. The resulting polymer was washed with EtOH, dried in air at 80 °C for 6 h to give NPOF-4-NH₂ (180 mg, 100% yield) as a dark brown powder. Anal. Calcd. for C₁₂₆H₁₀₈N₁₂: C, 84.53; H, 6.08; N, 9.39. Found: C, 65.55; H, 4.96; N, 8.39. Note: the low carbon content may be due to several issues that still need to be addressed; among these issues are partial reduction of the nitro groups, residual tin chloride, or trapped HCl.

2.3 Characterization of Building Block and NPOFs

The chemical composition and structural aspects of these nanoporous frameworks were investigated by spectral and analytical methods. Elemental microanalyses were performed at the Midwest Microlab, LLC. Liquid ¹H and ¹³C NMR spectra were obtained on a Varian Mercury-300 MHz NMR spectrometer (75 MHz carbon frequency). Solid-state ¹³C cross-polarization magic angle spinning (CP-MAS) NMR spectra for solid samples were taken at Spectral Data Services, Inc. Spectra were obtained using a Tecmag-based NMR spectrometer operating at a H-1 frequency of 363
MHz, using a contact time of 1 ms and a delay of three seconds for the CPMAS experiment. All samples were spun at 7.0 kHz. Thermogravimetric analysis (TGA) were carried out using a TA Instruments Q-5000IR series thermal gravimetric analyzer with samples held in 50 μL platinum pans under an atmosphere of air (heating rate 5 °C /min). For Scanning Electron Microscopy imaging (SEM), samples were prepared by dispersing the material onto a sticky carbon surface attached to a flat aluminum sample holder. The samples were then coated with platinum at 1×10⁻⁵ mbar of pressure in a nitrogen atmosphere for 90 seconds before imaging. Images were taken on a Hitachi SU-70 Scanning Electron Microscope. Powder X-ray diffraction data were collected on a Panalytical X'pert pro multipurpose diffractometer (MPD). Samples were mounted on a sample holder and measured using Cu Kα radiation with a 2θ range of 1.5-35. FT-IR spectra were obtained on a Nicolet-Nexus 670 spectrometer furnished with an attenuated total reflectance accessory. Porosity and gas sorption experiments were collected using a Quantachrome Autosorb 1-C volumetric analyzer using adsorbates of UHP grade. In a typical experiment, a sample of polymer (~ 60 mg) was loaded into a 9 mm large bulb cell (from Quantachrome) of known weight and then hooked up to the Autosorb 1-C and degassed at 120 °C for 12 hours. The degassed sample was refilled with helium, weighed precisely, and then transferred back to the analyzer. The temperatures for adsorption measurements were controlled by using a refrigerated bath of liquid nitrogen (77 K), liquid Argon (87 K), or temperature controlled water bath (273 and 298 K).
Figure 2.1: $^1$H NMR for 1,3,5,7-tetrakis(4-acetylphenyl)adamantane in CDCl$_3$. 
Figure 2.2: $^{13}$C NMR for 1,3,5,7-tetrakis(4-acetylphenyl)adamantane in CDCl$_3$. 
Figure 2.3: FT-IR Spectra of TAPA, NPOF-4, NPOF-4-NO$_2$ and NPOF-4-NH$_2$.
Figure 2.4: SEM images of NPOF-4, NPOF-4-NO$_2$ and NPOF-4-NH$_2$. 

SEM images of NPOF-4

SEM images of NPOF-4-NO$_2$

SEM images of NPOF-4-NH$_2$
Figure 2.5: TGA Traces of NPOF-4, NPOF-4-NO\textsubscript{2} and NPOF-4-NH\textsubscript{2}.
Figure 2.6: XRD Patterns for NPOF-4, NPOF-4-NO$_2$ and NPOF-4-NH$_2$. 
Figure 2.7: Solid state $^{13}$C CP-MAS NMR spectra of NPOF-4.
Figure 2.8: Solid state $^{13}\text{C}$ CP-MAS NMR spectra of NPOF-NO$_2$. 
2.4 Results and Discussion

Cyclotrimerization of acetyl functional groups which is frequently used to construct macromolecules through benzene ring formation has been employed here for synthesizing the adamantane-based nanoporous organic polymers (Scheme 2.1). Acid catalyzed cyclotrimerization of three acetyl groups results in the formation of a 1,3,5-substituted benzene units and the elimination of three water molecules.

NPOF-4 was synthesized in good yields following a modified procedure reported recently by Kaskel as depicted in Scheme 2.1. The slow addition of SiCl$_4$ at 0 °C

Figure 2.9: Solid state $^{13}$C CP-MAS NMR spectra of NPOF-NH$_2$. 
prevents rapid premature oligomers formation and leads to a more controlled pore formation and enhanced porosity. Although there are reports describing room temperature cyclotrimerization of acetylated aromatic molecules to generate isolated molecules, high temperatures are frequently used to either accelerate the rate of reactions or to overcome the steric and electronic effects of substituents. NPOP-4 is insoluble in common organic solvents such as DMF, THF, toluene, ethanol, etc, and remains intact upon washing with aqueous HCl and NaOH as evidenced by spectral and physical studies. Its high chemical stability enables post-synthesis modification particularly to introduce new functional groups that are known to enhance gas storage and separation properties. For example, pore surface chemical modification of organic frameworks with amino groups has significantly enhanced CO₂ binding in PPNs and POPs. There has been considerable interest recently in the synthesis of amino functionalized porous materials because of their interactions with small gas molecules consisting of polar bonds, particularly to CO₂ capture and selective separation of CO₂ from gas mixtures. Consequently, we have first modified NPOF-4 by nitration (NPOF-4-NO₂) then reduced the resulting network to the corresponding amino-functionalized framework (NPOF-4-NH₂).

Nitration was performed through the use of concentrated HNO₃/H₂SO₄ mixture to yield brown powder in a quantitative yield. The nitrated polymer NPOF-4-NO₂ was then reduced to NPOF-4-NH₂ using SnCl₂ in ethanol under refluxing at 70 °C for 18 h which resulted in a dark brown powder in 100% yield after drying at 80 °C. While the synthesis of NPOF-4-NO₂ and NPOF-4-NH₂ under harsh acidic conditions attests the chemical robustness of the resultant polymers, the thermal stabilities were examined by thermo
gravimetric analysis (TGA). The parent NPOP-4 is stable up to 400 °C, while the nitrated and reduced forms NPOF-4-NO₂ and NPOF-4-NH₂ showed a lower thermal stability (~200 °C, Figure 2.5). Scanning electron microscopy (SEM) of the polymers revealed aggregated particle morphology in the range of 0.4-1.1 µm (Figure 2.4) which seem to remain intact under nitration and reduction steps. As expected, NPOFs are amorphous as evidenced by powder X-ray diffraction studies (Figure 2.6).

The chemical functionalization of the polymers were investigated by FT-IR and solid state ¹³C NMR spectroscopic methods. Figure 2.3 shows the FT-IR spectra of the monomer and the synthesized polymers. Upon polymerization, the intensity of the band at 1680 cm⁻¹ (C=O) of TAPA is substantially decreased in NPOF-4 as a result of acetyl groups cyclotrimerization. This was further supported by the disappearance of 26 ppm ¹³C NMR peak of TAPA in NPOF-4 (Figure 2.7). Successful incorporation of nitro groups were confirmed by the appearance of new FT-IR bands at 1532 and 1350 cm⁻¹ (Figure 2.3) which have been assigned to asymmetrical and symmetrical stretching of NO₂, respectively, in NPOF-4-NO₂. Upon reduction, these bands disappear and new bands at 3350 and 3230 cm⁻¹ appear which confirm the conversion of the nitro groups to the amine functionality (Figure 2.3). Solid-state ¹³C NMR spectra showed the broadening of the peaks at around 150 ppm upon nitration and reduction probably due to the overlap of peaks of substituted carbons (Figure 2.8 and Figure 2.9). The chemical composition of the functionalized networks was investigated by elemental analysis that indicated the content of nitrogen by mass to be 7.84% for NPOF-4-NO₂ (~0.54 N per phenyl directly attached to adamantane) as illustrated in Scheme 2.1. The nitrogen content, as expected, increased to 8.39% upon reduction. Notably, this method seems
to be more effective than the previous methods reported for PAFs which resulted in only 0.25-0.3 N per phenyl rings).
Chapter 3
Gas Storage Capabilities for Nanoporous Organic Frameworks

3.1 Introduction

Great interest has been expressed in the gas storage capabilities of many porous media. Recently, the storage of carbon dioxide has received great attention because of its greenhouse nature and impact on the environment. Additionally, the use of argon gas to examine porosity can provide useful information about the textural properties of gas adsorbents and very reliable data on surface area and pore size distribution.

Nitrogen is the probe gas most commonly used for surface area and pore size distribution (PSD) analysis because it is inexpensive, readily available, inert, and well-studied in the adsorption literature. The nitrogen isotherm at 77 K (liquid nitrogen temperature), which is again convenient since liquid nitrogen is inexpensive, is widely used as a standard for micro and mesopore size analysis, however, it is also generally accepted that nitrogen adsorption is not satisfactory for quantitative assessment of the ultramicropores (pore widths <0.7 nm). Although the kinetic diameter of nitrogen and argon are similar (0.36, 0.34 nm, respectively) the adsorption behavior of these probe gases is different because argon is a monoatomic and spherical molecule without a quadrupole moment. On the other hand, the nitrogen adsorption and the orientation of the adsorbate molecules on the surface is affected by interactions between functional groups on the surface and the quadrupole moment of the nitrogen molecule (4.7 x 10^{-40} Cm^2). Therefore, argon has been extensively used for characterizing the pore...
structure of heterogeneous adsorbents to avoid the abovementioned problems. Thus, we have decided to use Argon isotherms at 87 K to characterize the porosity of our materials because of their heterogeneous surfaces especially after the incorporation of functional groups within the pores.47

To assess gas storage capabilities, sorption equipment is employed. The pressure range that dedicated low pressure sorption equipment can reach is typically around $10^{-6}$ to 1 bar. Gas storage by high pressure gas sorption measurements is beyond the scope of the research described herein.

For the present work, all sorption isotherms were determined in an automated low pressure volumetric gas adsorption instrument (Quantachrome Autosorb 1-C). An image of the instrument with essential components labeled is shown in Figure 3.1. Sorption isotherms were collected at constant temperatures which typically range from 77 K to 298 K. The temperature choice within this range is determined by the achievability and maintainability of the desired temperature. For example, submerging a sample in liquid nitrogen bath will keep the temperature at 77 K until all of the nitrogen boils off. Similarly, ice-water bath is used to maintain the temperature at 273 K. In order to test the applicability of materials to real life conditions isotherms are also collected at higher temperatures i.e. 25 °C which can be achieved and maintain by a thermal circulator bath compatible with the sorbent analyzer instrument. The result of an isothermal experiment is a graph with pressure on the x-axis and volumetric gas uptake (cc g$^{-1}$) on the y-axis then the uptake can be converted to desired units. The instrument measures the amount of gas adsorbed onto (or desorbed from) a solid surface at pre-requested equilibrium vapor pressure. The instrument admits or removes a known
volume of gas at a constant temperature and measure the corresponding pressure. The pressure in the sample cell changes as the instrument admits or removes gas in repetitive small quantities until the threshold around the requested pressure is reached. Then the amount of gas adsorbed or desorbed calculated by the difference between the amount of gas admitted or removed and the amount required to fill the void space.

**Figure 3.1:** Image of Autosorb-1C with essential components labeled.
3.2 Surface Area and Pore Size Characterization

To investigate the permanent porosity of NPOFs, Argon sorption data was acquired using a Quantachrome Autosorb 1-C at 87 K. Pore size distributions were calculated using the Non-Local Density Functional Theory (NLDFT)\(^{48}\) on the adsorption branch with a cylindrical/spherical pore model for all NPOFs. This model was used in conjunction with a similar model on the adsorption branch of the carbon dioxide isotherm at 273 K as has been reported previously in literature to obtain lower level pore size distributions.\(^{49,50}\) The choice of one model over another is generally determined by the shapes of the theoretical computer-generated models. Argon sorption studies were performed on activated samples. The activation process for all samples involved removing guest molecules by heating at 120 °C for 18 hours under vacuum (10\(^{-5}\) torr). Porous textural properties of NPOFs are summarized in Table 3.1. The rapid argon uptakes at very low pressure (below 0.1 x 10\(^{-2}\) bar) are indicative of microporosity and applying the Brunauer-Emmett-Teller (BET) model within the pressure range of \(P/P_0=0.01\)\(-\)0.15 resulted in apparent surface areas \(S_{\text{BET}}\) of 1249 m\(^2\) g\(^{-1}\) (NPOF-4), 337 m\(^2\) g\(^{-1}\) (NPOF-4-NO\(_2\)), and 554 m\(^2\) g\(^{-1}\) (NPOF-4-NH\(_2\)). The surface area decreased significantly upon pore functionalization with nitro groups which can restrict pore accessibility by Argon. Further modification upon nitro group reduction resulted in a surface area value that is intermediate between those of NPOF-4 and NPOF-4-NO\(_2\). Pore size distribution was estimated by fitting the argon uptake branch of the isotherms with NLDFT (cylindrical/spherical pore geometry with zeolites/silica model) and found to be in a broad range of \(~8.0\)-16.8 Å for NPOF-4, \(~7.7\)-10.8 Å for NPOF-4-NO\(_2\) and \(~7.7\)-15.2 Å for NPOF-4-NH\(_2\) (Figure 3.3). Pore volumes were calculated from single-point
measurements \((P/P_0 = 0.95)\) and found to be 0.78, 0.21, and 0.28 cc g\(^{-1}\) for NPOF-4, NPOF-4-NO\(_2\), and NPOF-NH\(_2\), respectively. The porous properties of NPOF-4 and its functionalized derivatives are comparable to those of porous organic networks and the chemical nature of the latters can be advantageous for small gas storage and separation applications as described below.

**Table 3.1:** Surface areas, pore size distributions and pore volumes for NPOF networks.

- \(^a\)Calculated by BET method.
- \(^b\)Calculated by Langmuir method.
- \(^c\)Calculated from Argon adsorption at \(P/P_0 = 0.95\).
- \(^d\)Calculated from NLDFT.

<table>
<thead>
<tr>
<th>Network</th>
<th>(S_{\text{BET}}) (m(^2) g(^{-1}))(^a)</th>
<th>(S_{\text{Lang}}) (m(^2) g(^{-1}))(^b)</th>
<th>PSD (Å)(^c)</th>
<th>(V_{\text{total}}) (cm(^3) g(^{-1}))(^d)</th>
</tr>
</thead>
<tbody>
<tr>
<td>NPOF-4</td>
<td>1249</td>
<td>1685</td>
<td>8.0-16.8</td>
<td>0.78</td>
</tr>
<tr>
<td>NPOF-4-NO(_2)</td>
<td>337</td>
<td>461</td>
<td>7.7-10.8</td>
<td>0.21</td>
</tr>
<tr>
<td>NPOF-4-NH(_2)</td>
<td>554</td>
<td>719</td>
<td>7.7-15.2</td>
<td>0.28</td>
</tr>
</tbody>
</table>

Figure 3.2 shows the argon adsorption/desorption isotherms for NPOF-4, NPOF-4-NO\(_2\), and NPOF-NH\(_2\) at 87 K. All isotherms are fully reversible and exhibit hysteresis that is well pronounced for NPOF-4 most likely due to the flexible nature of the framework as documented for numerous porous organic materials.\(^{35}\)
Figure 3.2: Argon uptake isotherms for NPOFs; adsorption (filled) and desorption (empty) data points were taken at 87 K and 0-1 bar.
In spite of their amorphous nature, the surface area of the networks under investigation, particularly for NPOF-4, are considerably high and comparable with a wide range of porous organic networks such as PIMs (618-1760 m² g⁻¹),⁵¹ cage compounds (1375 m² g⁻¹),⁵² imine-linked microporous polymer organic frameworks (POFs, 466-1521 m² g⁻¹),⁵³ covalent organic frameworks (COF-300, 1360 m² g⁻¹),⁵⁴ diimide-based polymers (750-1407 m² g⁻¹),⁵⁵ functionalized conjugated microporous polymer (CMP) networks (522-1043 m² g⁻¹)⁵⁶ and porous electron-rich covalent organo nitridic frameworks (PECONFs, 499-851 m² g⁻¹).⁵⁷

**Figure 3.3:** Pore size distribution for NPOF-4, NPOF-4-NO₂ and NPOF-4-NH₂.
Pore size distribution by NLDFT are based on fitting the sorption experimental data to mathematical functions. As such, the graphical comparison for the fitting method of the NLDFT calculations as well as the curves for the BET calculations are shown in the following figures.

**Figure 3.4:** NLDFT calculated isotherm for NPOF-4 overlaid with the experimental Argon isotherm. A fitting error less than 1% indicates validity of the model.
Figure 3.5: Multipoint BET plot for NPOF-4 calculated from the Argon adsorption in the range 0.04-0.16 P/P₀.
Figure 3.6: Langmuir plot for NPOF-4 calculated from the Argon adsorption in the range 0.05-0.25 P/P_0.

NPOF-4
Ar at 87 K
S_A_{Langmuir} = 1685 m^2/g
R^2 = 0.999499
Figure 3.7: NLDFT calculated isotherm for NPOF-4-NO$_2$ overlaid with the experimental Argon isotherm. A fitting error less than 1% indicates validity of the model.
Figure 3.8: Multipoint BET plot for NPOF-4-NO$_2$ calculated from the Argon adsorption in the range 0.04-0.16 P/P$_o$. 

NPOF-4-NO$_2$
Ar at 87 K
S$_{\text{BET}}$ = 337 m$^2$/g
R$^2$ = 0.99959
Figure 3.9: Langmuir plot for NPOF-4-NO$_2$ calculated from the Argon adsorption in the range 0.05-0.25 P/P$_o$. 

NPOF-4-NO$_2$
Ar at 87 K
SA$_{Langmuir}$ = 461 m$^2$/g
$R^2$ = 0.99903
Figure 3.10: NLDFT calculated isotherm for NPOF-4-NH$_2$ overlaid with the experimental Argon isotherm. A fitting error less than 1% indicates validity of the model.

NPOF-4-NH$_2$
Ar uptake at 87 K

Original
Fitted

Actual Fitting Error = 0.104 %
Figure 3.11: Multipoint BET plot for NPOF-4-NH$_2$ calculated from the Argon adsorption in the range 0.04-0.16 P/P$_o$. 

NPOF-4-NH$_2$

Ar at 87 K

SA$_{BET} = 554$ m$^2$/g

$R^2 = 0.999669$
3.3 Hydrogen, Carbon Dioxide, and Methane Sorption of Nanoporous Organic Frameworks

To investigate the performance of NPOFs in the capture of different gases, sorption experiments on activated samples for carbon dioxide, methane, and hydrogen were performed. Hydrogen isotherms were performed at 77 and 87 K whereas carbon dioxide and methane isotherms were performed at 273 and 298 K. The combined isotherms for CO$_2$ (Figure 3.13 and Figure 3.14), CH$_4$ (Figure 3.15 and Figure 3.16), and H$_2$ (Figure 3.17 and Figure 3.18) are all fully reversible illustrating the facile gas
release typical for organic polymers which makes them energetically attractive for gas storage and separation applications.

**Figure 3.13:** Carbon dioxide isotherms for NPOFs at 273 K: adsorption (filled symbols) and desorption (empty symbols).
**Figure 3.14:** Carbon dioxide isotherms for NPOFs at 298 K: adsorption (filled symbols) and desorption (empty symbols).
Figure 3.15: Methane isotherms for NPOFs at 273 K: adsorption (filled symbols) and desorption (empty symbols).
Figure 3.16: Methane isotherms for NPOFs at 298 K: adsorption (filled symbols) and desorption (empty symbols).
Figure 3.17: Hydrogen isotherms for NPOFs at 77 K: adsorption (filled symbols) and desorption (empty symbols).
Figure 3.18: Hydrogen isotherms for NPOFs at 87 K: adsorption (filled symbols) and desorption (empty symbols).

Low-pressure gas sorption measurements for CO$_2$, H$_2$, CH$_4$, and N$_2$ were collected in order to investigate the impact of pore functionalization on gas storage and the preferential binding of CO$_2$ over CH$_4$ and N$_2$. The impact of pore functionalization on CO$_2$ uptake is depicted in Figure 3.13 which indicates that NPOF-4-NH$_2$ has the highest uptake of 12.78 wt% (2.9 mmol g$^{-1}$) at 1 bar and 273 K and is about 16% higher than the uptakes of NPOF-4-NO$_2$ and NPOF-4 despite of the latter’s much higher surface area. These observations originate from the polarizable nature of the CO$_2$ molecule and its large quadrupole moment. While pore functionalization with -NO$_2$ and -NH$_2$ has
drastic impact on the binding affinity of CO₂ (explained in Chapter 4), its impact on the CO₂ final gravimetric uptake at 1.0 bar was less significant. The microporous nature of NPOF-4 and its high surface area facilitate CO₂ uptake at ambient pressure. High CO₂ uptake at low pressure (0.15 bar) is desirable for low-pressure post-combustion application where the flue gas usually consists of ~15% CO₂, ~75% N₂ and ~10% of other gases.

To commensurate the pore size of functionalized NPOF-4 for CO₂ separation from N₂ and CH₄, introduction of chemical heterogeneity to the pore walls which polarizes the CO₂ molecule have significant effect on CO₂ uptake at low pressures. At 0.2 bar and 273 K NPOF-4-NO₂ adsorbs 1.15 mmol g⁻¹ (5.08 wt%) of CO₂ and NPOF-4-NH₂ adsorbs 1.32 mmol g⁻¹ (5.80 wt%) which corresponds to an increase of gravimetric capacity of 54% and 76%, respectively, with respect to that of the non-functionalized NPOF-4 which takes 0.75 mmol g⁻¹ (3.30 wt%) of CO₂. The relatively large dipole moment of NO₂ and NH₂ results in dipole-quadrupole interactions with CO₂, and remarkably increase the initial CO₂ uptake at low pressure range. At 1.0 bar and 273 K NPOF-4-NO₂ adsorbs 2.41 mmol g⁻¹ (10.63 wt%) and NPOF-4-NH₂ adsorbs 2.9 mmol g⁻¹ (12.78 wt%) of CO₂ which exceed functionalized CMPs (1.6-1.8 mmol g⁻¹), triptycene-based cage compounds (2.1 mmol g⁻¹), and functionalized amidoxime-PIM-1 (2.74 mmol g⁻¹). At 1.0 bar and 298 K NPOF-4-NH₂ adsorbs 1.88 mmol g⁻¹ (12.78 wt%) of CO₂ which is comparable to BPL carbon (SÅBET= 1150 m² g⁻¹; a common reference material for CO₂ uptake) that shows an uptake of 1.90 mmol g⁻¹ under the same conditions.
We have also studied the H\textsubscript{2} and CH\textsubscript{4} uptake in order to evaluate the impact of -NO\textsubscript{2} and -NH\textsubscript{2} functionalities on the binding affinity of H\textsubscript{2} and CH\textsubscript{4}. Both gases are being considered as an alternative fuel for automotive applications because of their abundance and clean nature. The H\textsubscript{2} uptakes by NPOF-4 (1.33 wt\%) and its functionalized frameworks were modest (NPOF-NO\textsubscript{2}: 0.93 wt\% and NPOF-NH\textsubscript{2}: 1.15 wt\%) at 273 K/1 bar although the functionalized materials have higher binding affinities as depicted in Figure 4.3.

In a similar fashion, the CH\textsubscript{4} uptakes at 273 K and 298 K and 1 bar were collected for all materials and the results are presented in Figure 3.15 and Figure 3.16. A comparison of the methane uptake capacities for NPOF-4 and its functionalized derivatives suggests that surface area is the predominant factor in attaining high H\textsubscript{2} and CH\textsubscript{4} storage capacities at low pressure (1.0 bar), while surface functionality plays a more important role for CO\textsubscript{2} capture.
Chapter 4
Isosteric Heat of Adsorption of Carbon Dioxide, Methane, and Hydrogen with Nanoporous Organic Frameworks

4.1 Introduction

The process of adsorption is exothermic, i.e. heat is usually released when a gas molecule is adsorbed on a surface due to the loss of molecular motion of the gas molecules when they get adsorbed on a surface. As a consequence of that, the stronger adsorbent-adsorbate interactions are, the larger the amounts of heat released will be. The amount of the heat released also depends on the surface which is already covered with adsorbate. Presumably, as the surface is covered by adsorbate molecules, the heat of adsorption reduces. Therefore, it is more convenient to represent the heat as an isosteric heat of adsorption ($Q_{st}$), that is, at constant surface coverage for different temperatures. As a result, at least two isotherms are needed at different temperature in order to evaluate $Q_{st}$.

Calculation of isosteric heat of adsorption is based on experimental isotherms. Often, these isotherms are fitted to mathematical models that are intended to describe the data. In some cases, more than one isotherm is required to calculate $Q_{st}$. In the rest of the cases using more than one isotherm is encouraged to ensure that isotherm fitting parameters are reasonably accurate.
The virial method has been extensively employed to determine isosteric heat of adsorption of porous materials.\textsuperscript{61-64} A virial expansion is a summation equation for which, in this case, the number of summation iterations is unknown:

$$\ln P = \ln N + \frac{1}{T} \sum_{i=0}^{m} a_i N_i + \sum_{i=0}^{n} b_i N_i$$

In this equation, $P$ is pressure in torr, $T$ is temperature in Kelvin, and $N$ is the mmol of gas adsorbed per gram of sample. The values for $m$ and $n$ are undefined (hence, the unknown number of summation iterations) and are varied such that $m \geq n$ and result in the best fit as determined by the sum of the squares of the errors. The values for $a_0$, $a_1,...,a_m$ and $b_0$, $b_1,...,b_n$ are fitting parameters for the virial expansion. Following the appropriate fitting, the values for $a_0$, $a_1,...,a_m$ are used in the calculation for the isosteric heat of adsorption, $Q_{st}$:

$$Q_{st} = -R \sum_{i=0}^{m} a_i N_i$$

The value of $m$ in this case matches the value found for the preceding equation such that all $a_i$ values are used. The calculated values are then plotted as they relate to the surface coverage, $N$, while the isosteric heat of adsorption at the point of zero-coverage is typically reported as a means of comparison.

In the case of the virial-type calculation for the isosteric heat of adsorption, only one temperature is necessary to perform the full calculation. However, the large combination of $a_i$ and $b_i$ values for varying $m$ and $n$ values results in a very subjective approach from the researcher’s perspective. In order to mitigate, the subjective human element in these calculations, a minimum of two isotherms taken at different
temperatures should be performed when utilizing the virial method. This practice of multiple temperature-isotherms would drop the subjectivity of the calculations for the virial method, however, it should be noted that a base level of subjectivity still exists.

4.2 Experimental Calculations for the Isosteric Heat of Adsorption of Carbon Dioxide, Methane, and Hydrogen

Isosteric heats of adsorption of CO$_2$, CH$_4$ and H$_2$ were calculated for the networks by using virial method. $Q_{st}$ values showed that both functionalized NPOFs networks gave higher heat of adsorption for CO$_2$ over non-functionalized NPOF-4 for the entire coverage range. Viral equation is fitted to isotherms which were taken at two different temperatures. The $Q_{st}$ values and gas uptakes for networks are summarized in Table 4.1.

<table>
<thead>
<tr>
<th>Network</th>
<th>SA$_{BET}$</th>
<th>H$_2$ at 1 bar</th>
<th>CO$_2$ at 1 bar</th>
<th>CH$_4$ at 1 bar</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>77 K 87 K</td>
<td>273 K 298 K</td>
<td>273 K 298 K</td>
<td>273 K 298 K</td>
</tr>
<tr>
<td>NPOF-4</td>
<td>1249 13.3</td>
<td>109.9 61.6</td>
<td>20.2 8.2</td>
<td>24.6</td>
</tr>
<tr>
<td>NPOF-4-NO$_2$</td>
<td>337 9.3</td>
<td>106.3 68.8</td>
<td>9.0 5.4</td>
<td>20.8</td>
</tr>
<tr>
<td>NPOF-4-NH$_2$</td>
<td>554 11.5</td>
<td>127.8 82.8</td>
<td>12.5 7.4</td>
<td>20.7</td>
</tr>
</tbody>
</table>

The pore modification also increases the CO$_2$ bonding affinity from 23.2 kJ mol$^{-1}$ to 32.2 and 30.1 kJ mol$^{-1}$. These observations are again due to the polarizable nature of the CO$_2$ molecule and its large quadrupole moment. The enhanced $Q_{st}$ values are expected due to both chemical and physical pore modifications. In addition to the electronic nature of –NH$_2$ and -NO$_2$, upon framework functionalization, the pore size of
NPOF-4 is reduced and this reduction provides higher adsorption potentials for CO$_2$ and thus higher uptake and bonding affinity were observed. The $Q_{st}$ drops with higher CO$_2$ loading for functionalized NPOFs and clearly indicates the significant interactions between the CO$_2$ molecules and the functionalized pore surface. In contrast to this observation, the chemically homogeneous and non-functionalized pore surface of NPOF-4 leads to a non-notable change in $Q_{st}$ upon increased CO$_2$ loading. The $Q_{st}$ values for functionalized NPOFs exceeds most of the organic polymers such as BILPs, functionalized CPMs but are much lower than polyamine-tethered PPNs and PAFs, alkylamine appended MOF which feature short-chain aliphatic amines having binding affinities similar to those of amine solutions being employed in CO$_2$ scrubbing (50-100 kJ mol$^{-1}$). It should be noted that PSM of PAF-1 with alkylmethylamine (PAF-1–CH$_2$NH$_2$) showed CO$_2$ uptake of 98 cc g$^{-1}$ which is almost double that of parent PAF-1 (55 cc g$^{-1}$) at 273 K / 1 bar even though the latter has a surface area three times higher. This dramatic increase in CO$_2$ uptake can be attributed to the much higher binding affinity of PAF-1–CH$_2$NH$_2$ compared to parent PAF-1 (15.6 kJ mol$^{-1}$). The notable difference in the binding affinity for CO$_2$ is most likely caused by the more reactive nature of the alkylamine employed for the functionalization of PAF-1 and PPN-6 as evidenced by the large initial $Q_{st}$ values at lower coverage. Tailoring the binding affinity for selective CO$_2$ separation is essential because high heats of adsorption can lead to a major drawback wherein regeneration processes of adsorbents require heating as in the case of amine solutions (ca. 50-100 kJ mol$^{-1}$) that are associated with considerable energy penalty. Recent findings by Wilmer, et al. have suggested that these high binding affinities are less desirable when hypothetical MOFs
were screened for CO\textsubscript{2} capture from flue gas or natural gas, and that only moderate surface areas would be needed (1000 to 1500 m\textsuperscript{2} g\textsuperscript{-1}).\textsuperscript{67} Ironically, the Q\textsubscript{st} of NPOF-4-NO\textsubscript{2} (32.5 kJ mol\textsuperscript{-1}) and NPOF-4-NH\textsubscript{2} (30.1 kJ mol\textsuperscript{-1}) are within the desirable range and suggest that both materials can have novel properties for use in selective CO\textsubscript{2} separation from natural gas or flue gas. Additionally, the fully reversible nature of the CO\textsubscript{2} isotherms of both samples at ambient conditions (Figure 3.13 and Figure 3.14) also indicates that CO\textsubscript{2}–adsorbent interactions are weak enough to allow for adsorbents regeneration without applying heat.

The Q\textsubscript{st} values for H\textsubscript{2} were calculated from adsorption data collected at 77 and 87 K by the virial method. At zero-coverage, the Q\textsubscript{st} values are 7.17, 8.3 and 8.09 kJ mol\textsuperscript{-1} for NPOF-4, NPOF-4-NO\textsubscript{2} and NPOF-NH\textsubscript{2}, respectively (Figure 4.3). These values are higher than the values reported for organic polymers in general and similar to those having functionalized pores. In a similar fashion, the CH\textsubscript{4} uptakes at 273 K and 298 K and 1 bar were collected for all materials and the corresponding heats of adsorption was again calculated by using the virial method and the results are presented in Figure 4.2. The methane Q\textsubscript{st} values suggest that pore functionalization with polar functionalities reduces the interaction with methane as a result of the non-polar nature of the CH\textsubscript{4} molecule which is consistent with recent reports on methane storage by non-functionalized PAFs.
Figure 4.1: Carbon dioxide isosteric heat of adsorption curves of NPOFs.
Figure 4.2: Methane isosteric heat of adsorption curves of NPOFs.
Figure 4.3: Hydrogen isosteric heat of adsorption curves of NPOFs.
Figure 4.4: Virial analysis curve fitting of CO$_2$ adsorption isotherms for NPOF-4, NPOF-4-NO$_2$ and NPOF-4-NH$_2$ (blue circles: 273 K, olive squares: 298 K).
**Figure 4.5:** Virial analysis curve fitting of CH$_4$ adsorption isotherms for NPOF-4, NPOF-4-NO$_2$ and NPOF-4-NH$_2$ (blue circles: 273 K, olive squares: 298 K).
Figure 4.6: Virial analysis curve fitting of H₂ adsorption isotherms for NPOF-4, NPOF-4-NO₂ and NPOF-4-NH₂ (blue circles: 77 K, olive squares: 87 K).
Chapter 5

Gas Separation and Selectivity Capabilities for Nanoporous Organic Frameworks

5.1 Introduction

As a result of the economical and environmental impacts of gaseous impurities, selective gas separation continues to gain great attention by the scientific community.\textsuperscript{62,68-96} In particular, the separation of CO\textsubscript{2} from N\textsubscript{2}, CH\textsubscript{4} has become a global concern since carbon dioxide exists as an impurity in flue gases as well as in natural gas. New technologies and material design in this area can result in a reduction in CO\textsubscript{2} concentration in the atmosphere and can provide a facile approach to the production of high purity natural gas. Owing to their tunable properties to capture targeted gases, porous organic polymers are an attractive method for this purification process. Calculations for the separation and selective capture capabilities of a porous polymer media are performed based on the polymer's pure gas isotherms. Typically, there are three different methods for calculating selectivity: (1) initial slopes method,\textsuperscript{97} (2) Henry's Law constants where experimental isotherms are fitted to the virial-type expansion that was mentioned in Chapter 4 for the calculation of isosteric heat of adsorption,\textsuperscript{63,98} and (3) ideal adsorbed solution theory (IAST) where single-component isotherms is used to predict the mixture adsorption equilibria.\textsuperscript{98,99} The latter two methods will not be
discussed here and we have used the initial slopes method to carry out selectivity studies.

The initial slopes method for calculating selectivity employs the varying levels of gas uptake in the low pressure range (typically from 0 to about 0.1 bar). In this pressure range, the isotherms are assumed to be linear. As such, data points in this range are fit according to the linear equation: \( y = mx + b \), where \( y \) is the gas uptake in mmol g\(^{-1}\), and \( x \) is the pressure in bar, \( m \) is the slope of the curve, and \( b \) is the y-intercept of the curve. Selectivity is then calculated based on the ratio of the slopes of the curves for the respective gases. Selectivities based on this method are pursued based on the fact that it represents the simplest and fastest method of the three methods listed above. Nevertheless, the assumption that the isotherms are linear in the low pressure range can affect selectivity levels. Additionally, the applicable pressure range is very subjective and may hinder the calculation’s reliability. As a result, utilizing the initial slopes method is typically performed as a quick assessment to pursue selectivity for the polymer or as a confirmation of one of the other two calculation methods.

In addition to this method, recent reports have reported selectivity studies using the equation:

\[
S = \frac{q_1 / q_2}{p_1 / p_2}
\]

Equation 1

where \( S \) is the selectivity factor, \( q_i \) represents the quantity adsorbed of component \( i \), and \( p_i \) represents the partial pressure of component \( i \).\(^8\) For post-combustion CO\(_2\) capture, the partial pressure of CO\(_2\) and N\(_2\) are 0.15 bar and 0.75 bar, respectively. We have used both methods to study the selectivity of NPOFs.
5.2 Results and Discussion

Effective CO\(_2\) adsorbents are expected to have high selectivity for CO\(_2\) over other gases such as N\(_2\) (~75 % in flue gas) and CH\(_4\) (~95% in natural gas) along with high CO\(_2\) capacity for gas separation application. Therefore, we carried out CO\(_2\)/N\(_2\) and CO\(_2\)/CH\(_4\) selectivity studies for NPOF-4 before and after framework functionalization to evaluate the preferential CO\(_2\) adsorption at 273 and 298 K (Figure 5.1 - Figure 5.6). The general observation was that all materials show very low N\(_2\) uptake and tangible CO\(_2\) amounts at 0.15 bar (partial CO\(_2\) pressure in flue gas) especially for the functionalized NPOFs. The same trend was also noticed for the CH\(_4\) isotherms that show markedly lower CH\(_4\) uptakes than CO\(_2\) by all NPOFs. Selectivity studies were first calculated according to Equation 1.

For post-combustion CO\(_2\) capture, the partial pressure of CO\(_2\) and N\(_2\) are 0.15 bar and 0.75 bar, respectively. The CO\(_2\)/N\(_2\) selectivity studies resulted in very high values for NPOF-4-NO\(_2\) (133) and NPOF-NH\(_2\) (81), while NPOF-4 exhibited much lower selectivity (49) at 273 K (Table 5.1). Upon increasing the temperature to 298 K, the selectivity drops significantly to 14 (NPOF-4), 62 (NPOF-4-NO\(_2\)), and 29 (NPOF-4-NH\(_2\)). Furthermore, CO\(_2\)/N\(_2\) selectivity studies were determined by using the ratios of Henry law constants for which the constants can be calculated from the initial slopes of pure gas isotherms. This simple and convenient method has been widely employed to investigate the gas selective nature of a wide range of porous adsorbents including porous organic materials as we have reported recently for BILPs. According to initial slope calculations presented in Table 5.1, the CO\(_2\)/N\(_2\) selectivity levels for functionalized NPOFs are again much higher than the value calculated for NPOF-4 and are somewhat
higher than the values obtained from Equation 1. Nevertheless, both methods revealed the highly selective nature of NPOF-4-NO₂ and NPOF-4-NH₂ towards CO₂ over N₂. Noteworthy, these selectivity levels are among the highest by porous materials. For comparison, the most selective materials, NPOF-4-NO₂, outperforms other selective adsorbents at 273 K including BILPs (59-113),³⁷ PECOF-2 (109),⁵⁷ BPL carbon (17.8)⁶⁰ and ZIFs (17-50),⁵⁹ Bio-MOF-11 (81)⁷¹ and noncovalent porous materials (NPMs) (74).¹⁰⁰

By following the same methods described above, we calculated the CO₂/CH₄ selectivity of NPOFs at 273 K and 298 K (Table 5.1). In a typical natural gas purification process, the mole fractions of CO₂ and CH₄ are 0.05 and 0.95, respectively, resulting in CO₂ having a partial pressure of only 0.1 bar. The CO₂/CH₄ selectivity values from both methods; initial slope calculations and Equation 1, provided consistent data for each framework. The selectivity trend is in line with those observed for CO₂/N₂ studies and reveal that NPOF-4-NO₂ is the most selective material with selectivity levels in the range of 16-15 at 273 K that again outperforms the parent framework (NPOF-4 : 3-3) and its amine-functionalized derivative (NPOF-4-NH₂ : 13-11) (Table 5.1). The latter values fall within the range of recently reported porous materials such as diimide-based organic polymers, BPL carbon, ZIFs, and most MOFs. The outcome of selectivity studies surprisingly indicates that introducing the –NO₂ functionality into the pore walls of microporous organic frameworks is very advantageous for CO₂ separation applications. Most likely the higher selectivity of NPOF-4-NO₂ originates from the polar nature of the nitro functionality and due to its larger size when compared to –NH₂ which reduces pore size and induces dipole-dipole interactions involving CO₂. Although NPOF-4-NO₂ and
NPOF-4-NH₂ have remarkable selectivities, their capacity for CO₂ at the practical partial pressure of this gas in flue gas (0.15 bar) remains below the desired level (3.0 mmol g⁻¹) which can limit their applications as adsorbents. Our results again highlight the considerable challenge of attaining high selectivity levels without compromising adsorption capacity for CO₂ adsorbents. Selectivity results are summarized in the following tables.

**Table 5.1:** Selectivity results for NPOFs.  

aSelectivity was calculated from the Equation 1 at 273 K and 298 K. bSelectivity (mol mol⁻¹) was calculated from initial slope calculations at 273 and 298 K.

<table>
<thead>
<tr>
<th>Network</th>
<th>Selectivity⁹</th>
<th>Selectivityᵇ</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>CO₂/N₂</td>
<td>CO₂/CH₄</td>
</tr>
<tr>
<td></td>
<td>273 K</td>
<td>298 K</td>
</tr>
<tr>
<td>NPOF-4</td>
<td>49</td>
<td>14</td>
</tr>
<tr>
<td>NPOF-4-NO₂</td>
<td>133</td>
<td>62</td>
</tr>
<tr>
<td>NPOF-4-NH₂</td>
<td>81</td>
<td>29</td>
</tr>
</tbody>
</table>
Figure 5.1: Gas uptake selectivity studies for NPOF-4 at 273 K.
Figure 5.2: Gas uptake selectivity studies for NPOF-4 at 298 K.
Figure 5.3: Gas uptake selectivity studies for NPOF-4-NO$_2$ at 273 K.
Figure 5.4: Gas uptake selectivity studies for NPOF-4-NO$_2$ at 298 K.
Figure 5.5: Gas uptake selectivity studies for NPOF-4-NH$_2$ at 273 K.
Figure 5.6: Gas uptake selectivity studies for NPOF-4-NH$_2$ at 298 K.

\[
\text{CO}_2; \quad y = 4.57228x - 0.00252
\]
\[
\text{CH}_4; \quad y = 0.55692x - 0.00576
\]
\[
\text{N}_2; \quad y = 0.11553x + 3.27791 \times 10^{-4}
\]
Table 5.2: CO$_2$ and N$_2$ uptakes at 273 and 298 K for CO$_2$/N$_2$ selectivity studies by using Equation 1.

<table>
<thead>
<tr>
<th>Network</th>
<th>273 K</th>
<th>298 K</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>CO$_2$</td>
<td>N$_2$</td>
</tr>
<tr>
<td></td>
<td>Pressure (bar)</td>
<td>Uptake (mmol/g)</td>
</tr>
<tr>
<td>NPOF-4</td>
<td>0.15</td>
<td>0.59</td>
</tr>
<tr>
<td>NPOF-4-NO$_2$</td>
<td>0.15</td>
<td>0.97</td>
</tr>
<tr>
<td>NPOF-4-NH$_2$</td>
<td>0.15</td>
<td>1.09</td>
</tr>
</tbody>
</table>

Table 5.3: CO$_2$ and CH$_4$ uptakes at 273 and 298 K for CO$_2$/CH$_4$ selectivity studies by using Equation 1.

<table>
<thead>
<tr>
<th>Network</th>
<th>273 K</th>
<th>298 K</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>CO$_2$</td>
<td>CH$_4$</td>
</tr>
<tr>
<td></td>
<td>Pressure (bar)</td>
<td>Uptake (mmol/g)</td>
</tr>
<tr>
<td>NPOF-4</td>
<td>0.05</td>
<td>0.21</td>
</tr>
<tr>
<td>NPOF-4-NO$_2$</td>
<td>0.05</td>
<td>0.47</td>
</tr>
<tr>
<td>NPOF-4-NH$_2$</td>
<td>0.05</td>
<td>0.51</td>
</tr>
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</table>
Chapter 6

Concluding Remarks

The research presented in this thesis comprises the design, synthesis and post-synthetic modifications of functional Nanoporous Organic Frameworks (NPOFs). We have successfully synthesized a new NPOF from a novel building block, 1,3,5,7-tetrakis(4-acetylphenyl)adamantane (TAPA) by employing a metal-free synthesis approach. Subsequently, we have successfully used post-synthesis modification processes to functionalize the framework with –NO₂ and –NH₂ functionalities to investigate their effect on CO₂ uptake and the potential use of functionalized NPOFs in small gas storage and separation applications. The metal-free synthesis of NPOF-4 and its convenient pore surface modification resulted in a significant enhancement in CO₂ binding affinity: 32.5 and 30.1 kJ mol⁻¹ and selective binding over nitrogen and methane: CO₂/N₂ (139) and CO₂/CH₄ (15) at 273 K. These results confirmed that surface functionality overweighs the high surface area with regards to high CO₂ uptakes and binding affinity. Therefore, we conclude that having polar groups (-NO₂ and -NH₂) on pore walls notably enhances the initial CO₂ uptake and overall selectivity. These results highlight the potential use of functionalized NPOFs in post-combustion CO₂ separation and in natural gas purification processes. The functionalized NPOFs are also capable of storing up to 1.15 wt% H₂ at 77 K and 1 bar with high isosteric hydrogen heats of adsorption (8.1 and 8.3 kJ mol⁻¹). The relatively high initial uptake of H₂ by
functionalized NPOFs also indicates the significance of the physical and electronic nature of the pores in the design of future hydrogen storage sorbents.
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**Poster Presentations**


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