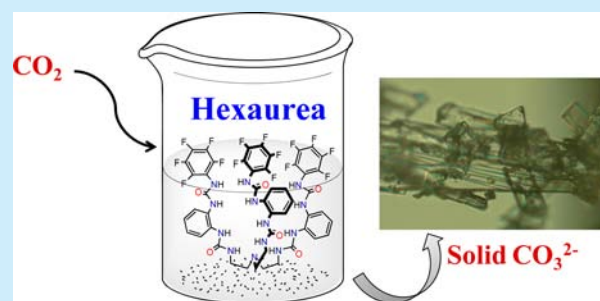


Absorption of Atmospheric CO₂ as Carbonate Inside the Molecular Cavity of a New Tripodal Hexaurea ReceptorAvijit Pramanik,[†] Maryam Emami Khansari,[†] Douglas R. Powell,[‡] Frank R. Fronczek,[§] and Md. Alamgir Hossain^{*,†}[†]Department of Chemistry and Biochemistry, Jackson State University, Jackson, Mississippi 39217, United States[‡]Department of Chemistry and Biochemistry, University of Oklahoma, Norman, Oklahoma 73019, United States[§]Department of Chemistry, Louisiana State University, Baton Rouge, Louisiana 70803, United States

S Supporting Information

ABSTRACT: A new hexaurea receptor has been synthesized, which absorbs atmospheric CO₂ to produce an air-stable solid carbonate complex under normal conditions. Structural analysis of the carbonate complex with this receptor suggests that the carbonate is fully encapsulated within its highly organized intramolecular cavity via 12 strong NH···O bonds in the range of 2.703(3)–2.989(3) Å from six urea units, with each anionic oxygen coordinated via four NH···O bonds with two urea groups.

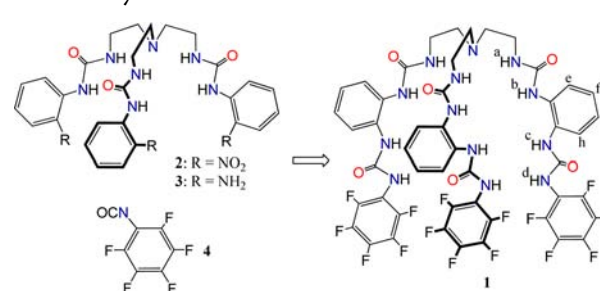


Continuous increase of carbon dioxide (CO₂) concentration in the atmosphere resulting from various human activities poses a threat to global warming.¹ CO₂ is also a major source of water acidification by dissolving in water to form carbonic acid (H₂CO₃).² Therefore, there is a current need to develop new chemical devices for the recycling of carbon dioxide to produce chemical commodities.³ Recently epoxide-based molecules have been used to react with CO₂ in the presence of a catalyst to produce new polycarbonates and cyclic carbonates.⁴ Gale et al. reported that simple monofunctional urea-based neutral compounds are capable of absorbing CO₂ in the presence of primary aliphatic amines to form carbamates [N(CO₂)–].⁵ Increasing the functional groups, tren-based tris-ureas/thioureas,⁶ such as pentafluorophenyl-substituted tris-urea,^{6a} *m*-nitrophenyl-substituted tris-urea,^{6b} phenyl-substituted tris-thiourea,^{6c} and *p*-cyanophenyl-substituted tris-thiourea^{6d} have recently been shown to bind carbonate anions, forming 2:1 (host/guest) complexes. Although the carbonate is coordinated via 12 hydrogen bonds (NH···O < 3.2 Å) in most cases, two receptors are required to provide complementary binding sites for the anion. Therefore, the complete coordination for a carbonate anion can ideally be achieved by a receptor possessing 12 complementary binding sites around a single cavity.

It is well-documented that increasing the effective binding sites in a host leads to the enhancement of its binding ability for a guest due to the chelate effect.⁷ From this viewpoint, we have been interested in synthesizing polyurea-functionalized receptors based on the commercially available “tren” as a core. Herein, we report a highly organized hexaurea receptor possessing 12 H-bond donors, which absorbs atmospheric CO₂ in the form of carbonate encapsulated in a single cavity surrounded by perfectly arranged six urea units. Within this self-generated intramolecular

cavity, the unique orientation of 12 binding sites provides an ideal complementarity for the trigonal planar carbonate anion.

The new hexaurea receptor **1** was synthesized by a three-step strategy (Scheme 1). The tris(2-aminoethyl)amine (tren) and *o*-

Scheme 1. Synthesis of the Hexaurea **1**

nitrophenyl isocyanate were reacted to give a nitro-functionalized tris-urea **2**, which was reduced with hydrazine hydrate and Pd/C (10%) to produce an amino-functionalized tris-urea **3**.⁸ The final coupling was achieved by reacting **3** with pentafluorophenyl isocyanate **4** to form an *o*-phenylene-bridged hexaurea **1**.

Single crystals of carbonate complex [1(CO₃)](n-Bu₄N)₂ were obtained from slow evaporation of a DMSO solution of **1** with an excess amount of *n*-tetrabutyl ammonium fluoride (TBAF) in a vial at room temperature in 2 days. The needle-like crystals were isolated by simple decantation technique and were found to be air-stable. Although, the primary goal was to obtain a

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fluoride complex, crystallographic analysis revealed that, instead of fluoride, one carbonate was encapsulated within the cavity of **1**. The bound carbonate was originated from atmospheric CO₂ absorbed by the basic TBAF–receptor solution which was left open to the atmosphere. The formation of carbonate in the presence of TBAF is well-documented in the literature.⁶

Structural analysis from single-crystal X-ray diffraction of the carbonate complex reveals that the complex crystallizes in an orthorhombic *Pna*2₁ space group to give a molecular formula, [1(CO₃)](*n*-Bu₄N)₂, in which one divalent carbonate is fully encapsulated.⁹ Two tetrabutyl ammonium ions remain outside the cavity to balance the anionic charges. In the receptor unit, all protons on NH groups are pointed toward the cavity and serve as H-bond donors for the bound anion.

As shown in Figure 1a, the encapsulated carbonate is held tightly inside the cavity by strong 12 NH...O bonds ($d_{\text{NH}\cdots\text{O}} =$

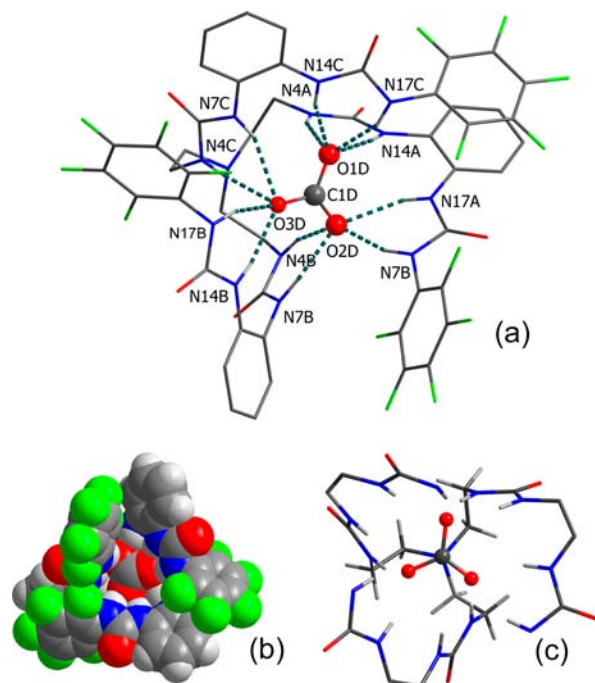


Figure 1. (a) Single-crystal X-ray structure of the carbonate complex of **1** (counteranions and hydrogen atoms on carbons are omitted for clarity), (b) space-filling model of encapsulated carbonate complex, (c) partial view of pseudo-C₃-symmetric arrangement of tripodal cavity with the encapsulated carbonate.

2.703(3)–2.989(3) Å from 12 NH groups of urea units (Table 1). Each oxygen of the carbonate is coordinated to four NH's from two different arms of the hexaurea receptor, with the two inner NH's from one arm and two outer NH's from other arm. Within a single arm of the receptor, the pentafluorophenyl and *o*-phenylene rings linked with an outer urea group are almost perpendicular to each other, while these two aromatic rings from two different arms are stacked together via $\pi\cdots\pi$ or C–H $\cdots\pi$ interactions ($\pi_{\text{C}}\cdots\pi_{\text{C}} = 3.602$ Å, $\text{C}_{8\text{E}}\text{--H}\cdots\pi = 3.841$ Å, and $\text{C}_{5\text{E}}\text{--H}\cdots\pi = 3.855$ Å). Such arrangements of the aromatic rings make the receptor preorganized for the complete participation of all six ureas in coordinating the internal anion. The space-filling view of the complex (Figure 1b) illustrates the encapsulated carbonate inside the cavity of **1**, showing the stacking of the aromatic groups. The trigonal planar carbonate is almost perpendicular to the axis of the tertiary nitrogen (N4C) of **1** and the carbon

Table 1. H-Bonding Parameters (Å, deg) for [1(CO₃)]^{2–}

D–H \cdots O	H \cdots O	D \cdots O	$\angle\text{DHO}$
N4A–H4A \cdots O1D	2.20(3)	2.966(3)	152(3)
N7A–H7A \cdots O1D	2.01(3)	2.852(2)	165(3)
N14A–H14A \cdots O2D	2.16(3)	2.935(3)	155(2)
N17A–H17A \cdots O2D	1.84(3)	2.703(3)	161(3)
N7B–H7B \cdots O2D	1.99(3)	2.869(3)	162(2)
N4B–H4B \cdots O2D	2.19(3)	2.949(3)	154(3)
N14B–H14B \cdots O3D	1.93(3)	2.801(3)	157(2)
N17B–H17B \cdots O3D	2.00(3)	2.788(3)	154(3)
N4C–H4C \cdots O3D	2.03(3)	2.822(3)	162(3)
N7C–H7C \cdots O3D	2.21(3)	2.989(3)	153(3)
N14C–H14C \cdots O1D	1.97(3)	2.783(2)	160(3)
N17C–H17C \cdots O1D	1.92(3)	2.746(3)	156(3)

(C1D) of CO₃^{2–}, forming a pseudo-C₃-symmetric complex (Figure 1c).¹⁰

Proton NMR titrations were performed to examine the stability and stoichiometry of the receptor–carbonate complex in DMSO-*d*₆ using Et₄NHCO₃. Upon the addition of Et₄NHCO₃ (20 mM) to the receptor (2 mM), a new set of NMR spectra appeared due to slow exchange on the NMR time scale (Figure 2).¹¹ All four NH signals were shifted significantly to downfield

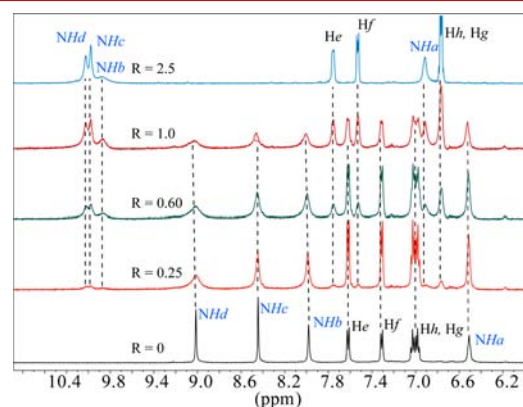


Figure 2. Partial ¹H NMR spectra of **1** with an increasing amount of Et₄NHCO₃ ($R = [\text{Et}_4\text{NHCO}_3]_0/[\textbf{1}]_0$) in DMSO-*d*₆.

($\Delta\delta\text{NH}_a = 0.41$ ppm, $\Delta\delta\text{NH}_b = 1.88$ ppm, $\Delta\delta\text{NH}_c = 1.53$ ppm, and $\Delta\delta\text{NH}_d = 1.08$ ppm), indicating the interactions of all NH groups with the anion. Presumably, the chelation of the bound anion, as also observed in the solid-state structure of the complex of **1**, results into the formation of a kinetically stable complex on the NMR time scale.⁸ The relative change in the integration intensity of NH resonances of the **1**–bicarbonate complex and the free **1** allowed us to determine the binding constant (Figure 3).¹¹ The experimental data provided the best fit to a 1:1 (host/guest) binding model,¹² yielding a binding constant $K = 1780$ M^{–1}. The 1:1 binding in solution was further supported by a Job's plot analysis (Figure S17). It is noted that, because of the unavailability of a suitable DMSO soluble CO₃^{2–} salt, the HCO₃[–] as a tetraethyl ammonium (Et₄N⁺) salt was used in the NMR titration studies, as previously used by other groups⁶ for tren-based ligands. Thus, the determined binding constant ($K = 1780$ M^{–1}) is the result of the interactions of **1** with singly charged HCO₃[–] as opposed to CO₃^{2–} observed in the crystal. The time-dependent NMR spectra of **1** and Et₄NHCO₃ in DMSO-*d*₆ showed no change in the NMR signals, suggesting that HCO₃[–]

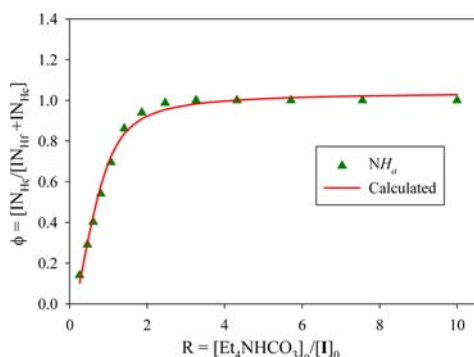


Figure 3. ^1H NMR titration curve of **1** (2 mM) with an increasing amount of Et_4NHCO_3 ($R = [\text{Et}_4\text{NHCO}_3]_0/[\mathbf{1}]_0$) in $\text{DMSO}-d_6$. The titration plot was obtained from the relative change in the integration intensity $\Delta\phi$ ($\Delta\phi = I_{\text{NHc}}/[I_{\text{NHf}} + I_{\text{NHc}}]$, where I_{NHc} is the intensity of NHc signal in the carbonate–**1** complex and I_{NHf} is the intensity of the corresponding NH signal in the free **1**).

was not deprotonated to form CO_3^{2-} during the titration process (Figure S18).

The binding affinity of **1** for Et_4NHCO_3 was compared with its precursor tris-urea receptor **2** (Scheme 1) under similar conditions in $\text{DMSO}-d_6$. The results show that this receptor binds HCO_3^- with $K = 226 \text{ M}^{-1}$ (Figure S24 in Supporting Information), which is much weaker than 1780 M^{-1} observed for **1**. An *n*-butyl-functionalized tris-thiourea reported previously was shown to bind bicarbonate with $K = 564 \text{ M}^{-1}$ in $\text{DMSO}-d_6$.^{6c} This observation further demonstrates that the unique orientation of the binding sites in the hexaurea **1** makes it an excellent binder for the carbonate anions due to the possible chelate effect observed in traditional transition metal complexes.^{7b} We also performed ^1H NMR titrations of **1** for other anions, showing strong affinity for HSO_4^- (2250 M^{-1}). The association constants of **1** with Cl^- and CH_3COO^- were 256 and 200 M^{-1} , respectively. However, the receptor showed no significant binding for I^- , ClO_4^- , and NO_3^- . Determination of binding constants for F^- and H_2PO_4^- was hampered due to the broadening of NMR peaks.

The formation of the complex in $\text{DMSO}-d_6$ was further evaluated by a series of ^{13}C NMR spectra (Figure 4). Partial ^{13}C NMR of Et_4NHCO_3 and free **1** are shown in Figure 4a,b, respectively. The sharp signal at 157.19 ppm in the free Et_4NHCO_3 shifted to 168.38 ppm ($\Delta\delta = 11.19 \text{ ppm}$) after the addition of **1** equiv of the ligand (Figure 4c), indicating the

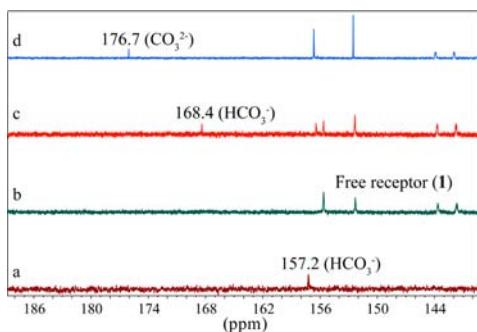


Figure 4. Partial ^{13}C NMR spectra (125 MHz, $\text{DMSO}-d_6$) of (a) Et_4NHCO_3 , (b) free receptor **1**, (c) **1** in the presence of 1 equiv of Et_4NHCO_3 , and (d) solid carbonate complex of the receptor ($[\mathbf{1}(\text{CO}_3)](n\text{-Bu}_4\text{N})_2$).

encapsulation of HCO_3^- in the receptor's cavity.^{6a} Figure 4d displays the ^{13}C NMR of $[\mathbf{1}(\text{CO}_3)](n\text{-Bu}_4\text{N})_2$ used for single-crystal diffraction, showing a peak more downfield at 176.7 ppm for CO_3^{2-} . This difference could be due to the charges (HCO_3^- vs CO_3^{2-}) and the counteranions (Et_4N^+ vs $n\text{-Bu}_4\text{N}^+$) present in the two different samples.

The solid-state FT-IR analysis was also performed to examine the interactions of the receptor with carbonate. The free receptor shows two peaks at 1650 and 1600 cm^{-1} for two carbonyl groups ($-\text{C}=\text{O}$),¹³ which shift to higher wavenumbers at 1696 cm^{-1} ($\Delta\nu_{\text{CO}} = 46 \text{ cm}^{-1}$) and 1630 cm^{-1} ($\Delta\nu_{\text{CO}} = 30 \text{ cm}^{-1}$), respectively (Figure S15). Further, the significant upward shift ($\Delta\nu_{\text{N-H}} = 74 \text{ cm}^{-1}$) of broad NH's stretching frequency from 3330 cm^{-1} (free **1**) to 3404 cm^{-1} (complex) was observed, suggesting the strong $\text{N-H}\cdots\text{O}$ interactions between NH groups and the carbonate.

The ability of **1** to uptake of the atmospheric CO_2 was examined by dissolving the receptor with TBAF in DMSO and keeping the mixture in open air (Figure 5). The solid products

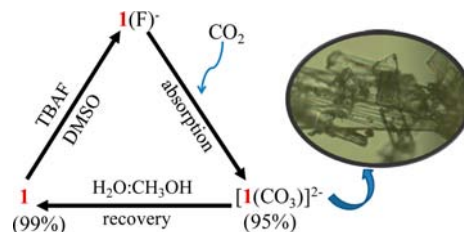


Figure 5. Uptake of CO_2 and the recovery of **1**. Magnified **1** carbonate crystals are shown at the right side.

deposited at the interface of the liquid and air were collected to give almost 99% carbonate complex. The complex was characterized by ^1H NMR and also verified from X-ray diffraction analysis (CCDC 969272), giving an identical structure described in the preceding section. Treating the complex with water and methanol (1:1, v/v) gave over 95% recovery of **1**. In previous reports, a pentafluorophenyl-substituted tris-urea^{6a} and a *m*-nitrophenyl-substituted tris-urea^{6b} were shown to fix atmospheric CO_2 , forming 98 and 90% complexes, respectively.

In conclusion, we have developed a novel receptor with six urea groups possessing an intramolecular cavity that is suitable to absorb atmospheric CO_2 as CO_3^{2-} under mild basic conditions. The highly functionalized receptor molecule utilizes all 12 NH-binding sites to interact with the internal carbonate anion, forming an air-stable complex. The solid-state structure described herein represents an exceptional example of the encapsulated carbonate in a single molecule, providing an ideal complementarity of binding sites for the anion. Given the significant role of CO_2 in environmental, chemical, and biological systems,^{3,14} the present findings hold promise for developing novel chemical devices for the recycling of atmospheric carbon dioxide.

■ ASSOCIATED CONTENT

§ Supporting Information

Crystallographic file in CIF format, and synthetic procedures and NMR titration spectra in pdf format. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Notes

The authors declare no competing financial interest.

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