An ATR-FTIR Study on the Effect of Molecular Structural Variations on the CO₂ Absorption Characteristics of Heterocyclic Amines, Part II

Kelly Robinson,[a, b] Adam McCluskey,[b] and Moetaz I. Attalla*[a]

This paper reports on an ATR-FTIR spectroscopic investigation of the CO₂ absorption characteristics of a series of heterocyclic diamines: hexahydropropyrimidine (HHPY), 2-methyl and 2,2-dimethylhexahdropyrimidine (MHHPY) and DMHHPY, hexahdropyridazine (HHPZ), piperazine (PZ) and 2,5- and 2,6-dimethylpiperazine (2,6-DMPZ and 2,5-DMPZ). By using in situ ATR-FTIR the structure-activity relationship of the reaction between heterocyclic diamines and CO₂ is probed. PZ forms a hydrolysis-resistant carbamate derivative, while HHPY forms a more labile carbamate species with increased susceptibility to hydrolysis, particularly at higher CO₂ loadings (>0.5 mol CO₂/mol amine). HHPY exhibits similar reactivity toward CO₂ to PZ, but with improved aqueous solubility. The α-methyl-substituted MHHPY favours HCO₃⁻ formation, but MHHPY exhibits comparable CO₂ absorption capacity to conventional amines MEA and DEA. MHHPY show improved reactivity compared to the conventional α-methyl-substituted primary amine 2-amino-2-methyl-1-propanol. DMHHPY is representative of blended amine systems, and its reactivity highlights the advantages of such systems. HHPZ is relatively unreactive towards CO₂. The CO₂ absorption capacity Cₐ (mol CO₂/mol amine) and initial rates of absorption R₀ (mol CO₂/mol amine min⁻¹) for each reactive diamine are determined: PZ: Cₐ = 0.92, R₀ = 0.045; 2,6-DMPZ: Cₐ = 0.86, R₀ = 0.025; 2,5-DMPZ: Cₐ = 0.88, R₀ = 0.018; HHPY: Cₐ = 0.85, R₀ = 0.032; MHHPY: Cₐ = 0.86, R₀ = 0.018; DMHHPY: Cₐ = 1.1, R₀ = 0.032; and HHPZ: no reaction. Calculations at the B3LYP/6-31+G** and MP2/6-31+G** calculations show that the substitution patterns of the heterocyclic diamines affect carbamate stability, which influences hydrolysis rates.

1. Introduction

The dominant sources of anthropogenic CO₂ emission are fossil-fuel combustion and industrial processes.[1, 2] Sequestration of this CO₂ is now a major target for reduction of atmospheric CO₂ levels. While it is clear that any significant long-term reduction in greenhouse gas emissions must involve changing our approach to energy production and consumption, technologies are required to reduce levels in the short term. Coal-fired power stations are the largest point-source emitters of CO₂ in Australia and worldwide.[3] The prospect of integrating post-combustion CO₂ capture (PCC) technology in both existing and new coal-fired power stations offers the potential to lower CO₂ emissions in the face of existing and predicted growth in the number of coal-fired power stations.[4]

Currently, aqueous amine-based PCC is viewed as the most promising and near-ready technology for the reduction of CO₂ emissions from coal-fired power stations. PCC involves separating CO₂ from a flue gas stream by chemical absorption and regenerating CO₂ from the absorbent by heating in a two-step process for subsequent storage or industrial use. PCC is industrially proven with absorbents such as aqueous monoethanolamine (MEA), used for decades for CO₂ removal from gas streams in small-scale commercial processes such as ammonia production and natural-gas processing.[5, 6]

Despite being an established technology, deployment of current industry-standard technology (30 wt% aqueous MEA) on a large scale applies a considerable efficiency penalty to the power generation process. Regeneration of PCC absorbent is energy-intensive,[7] and will result in up to 25% reduction in the net efficiency of a coal-fired power plant.[8, 9] Clearly, the absorption/regeneration characteristics of the amine-based PCC absorbent will influence the economic feasibility of this technology. One approach to reducing the energy requirements and cost of the PCC process is the development of more cost effective and better performing amines. There is considerable scope to develop absorbents that show higher CO₂ absorption, lower regeneration costs and greater chemical stability, particularly in the face of an increasing move towards demonstration-scale PCC plants.

The CO₂ absorption/desorption by aqueous amine-based absorbents has been, and continues to be, extensively studied in the search for improvements in PCC efficiency. The CO₂ absorption/desorption process is shown schematically in Figure 1. Typically, CO₂ reacts with aqueous amines to generate carba-
CO₂ absorption by aqueous amines is a reversible process, and the degree of reversibility is amine-dependent. Amines that form stable carbamates exhibit faster reaction rates, but a larger input of energy is required for absorbent regeneration. Conversely, amines that form more bicarbonate than carbamate exhibit slower reaction rates and require less energy for regeneration. Recent technological advances have allowed for the convenient and rapid analysis of these chemical species to be carried out in situ during the PCC absorption/desorption cycle by attenuated total reflectance Fourier transform infrared (ATR-FTIR) spectroscopy. In particular, the ability of ATR-FTIR to distinguish carbamate from bicarbonate formation accelerates the screening of potential PCC amines.\[10,17\]

We recently reported the application of ATR-FTIR in a model PCC absorbent system with substituted piperidines.\[10\] Herein we report on the in situ CO₂ absorption characteristics of a series of heterocyclic diamines (Figure 2): piperazine (PZ), 2,6-dimethyl- and 2,5-dimethylpiperazone (2,6-DMPZ and 2,5-DMPZ), hexahydropyrimidine (HHPY), 2-methylhexahydropyrinidine (MHHPY), 2,2-dimethylhexahydropyrimidine (DMHHPY) and hexahydropyridazine (HHPZ).

**2. Results and Discussion**

**2.1. Infrared Spectral Analysis**

The effect of structural variations on the CO₂ absorption characteristics of the heterocyclic diamines shown in Figure 2 were assessed in relation to the IR-identifiable products, that is, carbamate versus bicarbonate absorbance; CO₂ absorption capacity, defined as moles of CO₂ absorbed per mole of amine in solution (mol CO₂/mol amine); and the initial rate of CO₂ absorption (mol CO₂/mol amine min\(^{-1}\)). Full details of our experimental approach is given in the Experimental Section and also in our previous work.\[10\] Each experiment was conducted until equilibrium was established and a maximum CO₂ loading achieved. This was amine-dependent but typically required 45–

\[\text{Piperazine and Methyl Substituted Derivatives (Commercially available)}\]

\[\text{Hexahydropyrimidine and Methyl Substituted Derivatives (Synthesised)}\]

\[\text{Hexahydropyridazine (Synthesised)}\]
90 mins. Calculations were also performed to investigate the electronic/steric effects of the structural variations on the amine-carbamate derivatives.

2.1.1. Piperazine (PZ)

Our investigations commenced with parent heterocyclic diamine PZ (Figure 2). As can be seen from the partial (1750–950 cm\(^{-1}\)) FTIR spectrum collected during a typical CO\(_2\) absorption experiment with an aqueous PZ solution (1.5 mol L\(^{-1}\)), five major FTIR peaks evolve during CO\(_2\) absorption (Figure 3). The carbamate (NCOO\(^-\)) derivatives of heterocyclic monoamines have been identified as giving rise to several strong absorption bands in the 1600–1260 cm\(^{-1}\) region, including the asymmetric (\(\nu_{\text{as}}\text{COO}^-\), 1600–1500 cm\(^{-1}\)) and symmetric (\(\nu_{\text{sym}}\text{COO}^-\), 1450–1350 cm\(^{-1}\)) vibrations of the COO\(^-\) moiety and the N–COO\(^-\) stretching vibration (\(\nu_{\text{N-COO}^-}\), 1300–1260 cm\(^{-1}\)) of the NCOO\(^-\) derivative.\(^{[8]}\) The protonated amine (NH\(_3^+\)) generated on absorption of CO\(_2\) was found to give rise to an absorbance band in the 1479–1474 cm\(^{-1}\) region (NH\(_3^+\) bending mode). In an amine/CO\(_2\)/H\(_2\)O system, the bicarbonate (HCO\(_3^-\)) species was identified as giving rise to a broad peak in the 1360–1354 cm\(^{-1}\) region (\(\nu_{\text{as}}\)HCO\(_3^-\)).\(^{[16,17]}\) Assignment was based on the spectral data acquired for 1-methylpiperidine (tertiary amine)/CO\(_2\)/H\(_2\)O and 2-amino-2-methyl-1-propanol (AMP)/CO\(_2\)/H\(_2\)O systems. It is known that the absorption of CO\(_2\) by aqueous AMP (\(\alpha\)-dimethyl-substituted MEA derivative) leads to the formation of mostly HCO\(_3^-\) with no significant NCOO\(^-\) formation.\(^{[14,18,19]}\)

Herein these peaks can be related to the vibrational modes of the potential ionic reaction products, including PZ-carbamate (PZ-COO\(^-\)), protonated PZ (PZ-H\(^+\)) and bicarbonate (HCO\(_3^-\)). PZ, being a secondary diamine, should react with CO\(_2\) in solution to form NCOO\(^-\), predominately in the form of a protonated PZ-COO\(^-\) derivative (\(\text{H}_2\text{NR}_1\text{R}_2\text{NCOO}^-\)).\(^{[20,21]}\) One amine moiety acts as the absorption site for CO\(_2\), and the other as a proton acceptor. PZ has also been reported to form the dicarbamate (\(\text{OOC-PZ-COO}^-\)), which was detected by \(^1\)H and \(^13\)C NMR spectroscopy, at CO\(_2\) loadings of 0.2–0.8 mol CO\(_2\)/mol amine.\(^{[20,21]}\)

The FTIR spectra for the PZ/CO\(_2\)/H\(_2\)O system closely resemble those that we previously reported for the piperidine/CO\(_2\)/H\(_2\)O system, differing only in slight shifts in key IR stretching frequencies.\(^{[8]}\) At low levels of absorbed CO\(_2\) the PZ/CO\(_2\)/H\(_2\)O system exhibits the \(\nu_{\text{as}}\text{COO}^-\) (1524 cm\(^{-1}\)), \(\nu_{\text{sym}}\text{COO}^-\) (1432 cm\(^{-1}\)) and \(\nu_{\text{N-COO}^-}\) (1276 cm\(^{-1}\) and 1294 cm\(^{-1}\)) of the PZ-COO\(^-\) derivative and the NH\(_3^+\) vibration of PZ-H\(^+\) (1470 cm\(^{-1}\)). These peaks shift to 1546, 1425 and 1289 cm\(^{-1}\), respectively, with increasing CO\(_2\) absorption levels (Figure 3). As anticipated, a near-linear relationship between cumulative CO\(_2\) absorption and IR peak intensity is observed for the spectral peaks assigned to \(\nu_{\text{as}}\text{COO}^-\) and \(\nu_{\text{N-COO}^-}\). Increased peak absorbance is concomitant with the rate of NCOO\(^-\) formation at the reaction onset plateauing as a maximum CO\(_2\) loading of 0.92 mol CO\(_2\)/mol amine is approached (Figure 4). This near-linear relationship differs from that observed for the \(\nu_{\text{N-COO}^-}\) bands at 1276 and 1294 cm\(^{-1}\). From the data presented in Figures 3 and 4 the primary \(\nu_{\text{N-COO}^-}\) absorbance emerged at 1276 cm\(^{-1}\) and was the dominant peak, but only at CO\(_2\)/mol amine loadings of 0.4–0.5 mol CO\(_2\)/mol amine. At amine loadings greater than 0.5 CO\(_2\)/mol amine the \(\nu_{\text{N-COO}^-}\) absorbance decreases correspondingly with a sharp increase in intensity of the absorbance band at 1294 cm\(^{-1}\) and a frequency shift to 1289 cm\(^{-1}\). This trend in \(\nu_{\text{N-COO}^-}\) peak absorbance in the 1294–1276 cm\(^{-1}\) region is attributed to formation of \(\text{OOC-PZ-COO}^-\). The IR absorbance of PZ/CO\(_2\)/H\(_2\)O in this region differs from those of all other heterocyclic amine and diamine systems thus far reported, and the remaining subset of secondary heterocyclic amines analysed in this study (see below) displays only a single \(\nu_{\text{N-COO}^-}\) absorbance band in the 1283–1272 cm\(^{-1}\) region. As the IR stretching frequencies of PZ-dicarbamate had not been previously reported we turned to computational approaches to facilitate the assignment of key vibrational modes of PZ-COO\(^-\), in particular \(\nu_{\text{N-COO}^-}\). Calculations were performed at the B3LYP/6-31+G** and MP2/6-31+G** levels (gas phase, Spartan ’08).\(^{[22]}\)

The B3LYP/6-31+G** calculations assigned PZ-\(\nu_{\text{N-COO}^-}\) to a single band at 1282 cm\(^{-1}\) (no scaling), while MP2/6-31+G**
positioned this band at 1284 cm\(^{-1}\) (no scaling), similar in shape, but not as broad, as that which initially emerges at 1276 cm\(^{-1}\) in Figure 3. For the ‘OOC-PZ-COO’ species B3LYP/6-31+G** gave two sharp \(\nu_{N-COO}\) absorbances at 1297–1266 and 1348–1345 cm\(^{-1}\), which correlated well with MP2/6-31+G** calculated positions of 1302–1274 cm\(^{-1}\) and 1364–1355 cm\(^{-1}\). These values correlate well with the experimentally observed peaks at values of 1266, 1276 and 1294 cm\(^{-1}\), with the latter two shifting to 1289 cm\(^{-1}\) with \(\text{CO}_2\) absorption. The B3LYP/6-31+G** and MP2/6-31+G** calculations confirm our peak assignments for the PZ-CO\(_2\) carbamate absorption species above.

The evolution of a weak broad absorbance band in the 1360–1350 cm\(^{-1}\) region of the PZ/\(\text{CO}_2\)/\(\text{H}_2\text{O}\) IR spectral profile (Figure 3) was assigned to \(\nu_{C-O}\) of HCO\(_3^-\). This absorbance band was far less prominent than that we observed for the piperidine/\(\text{CO}_2\)/\(\text{H}_2\text{O}\) system. Additionally, this absorbance in the PZ system does not follow the trend observed with the corresponding piperidine system; that is, the depletion of \(\nu_{asCOO^-}\), \(\nu_{sCOO^-}\) and \(\nu_{asCOCO^-}\) absorbance bands with concomitant increase in the HCO\(_3^-\) absorbance band for \(\text{CO}_2\) loadings greater than 0.5 mol \(\text{CO}_2\)/mol amine.\(^{[10]}\) The increase in HCO\(_3^-\) absorbance in the piperidine system is attributable to hydrolysis of the initially formed carbamate, which strongly suggests, consistent with the IR data presented herein, that the PZ system forms a hydrolysis-resistant carbamate.

2.1.2. 2,6- and 2,5- Dimethyl-Substituted Piperazine Derivatives (2,6-DMPZ and 2,5-DMPZ)

The effect of alkyl substituents on the \(\text{CO}_2\) absorption characteristics of PZ was examined with 2,6-dimethylpiperazine (2,6-DMPZ) and 2,5-dimethylpiperazine (2,5-DMPZ) (Figure 2). The IR spectral profiles obtained for 2,6-DMPZ/\(\text{CO}_2\)/\(\text{H}_2\text{O}\) and 2,5-DMPZ/\(\text{CO}_2\)/\(\text{H}_2\text{O}\) are shown in Figures 5 and 6, respectively.

The IR spectral data collected for the 2,6-DMPZ/\(\text{CO}_2\)/\(\text{H}_2\text{O}\) system (Figure 5a and b) are almost identical to those recorded for the equivalent PZ system. While the \(\nu_{asCOO^-}\) and \(\nu_{sCOO^-}\)bands were less intense and only a single \(\nu_{asCOCO^-}\) absorbance band evolves in the 1289–1276 cm\(^{-1}\) region, all major absorbances of the 2,6-DMPZ system are within 6 cm\(^{-1}\) of those of the PZ system: \(\nu_{asCOO^-}\) at 1526 cm\(^{-1}\); \(\nu_{sCOO^-}\) at 1425 cm\(^{-1}\); \(\nu_{asCOCO^-}\) at 1279 cm\(^{-1}\); \(\nu_{sCOCO^-}\) at 1425 cm\(^{-1}\); \(\nu_{asCOCO^-}\) absorbance at 1354 cm\(^{-1}\). The \(\nu_{asCOCO^-}\) and \(\nu_{sCOCO^-}\) absorbances shift to 1550 and 1289 cm\(^{-1}\), respectively, with increasing \(\text{CO}_2\) absorption. The lack of a second \(\nu_{asCOCO^-}\) absorbance is reflected in the relationship between absorbance and cumulative \(\text{CO}_2\) absorption (Figure 5).

Due to steric congestion arising from the two \(\alpha\)-CH\(_3\) moieties, initial \(\text{CO}_2\) absorption most likely occurred at the less hindered and more nucleophilic amine moiety, resulting in NCOO\(^-\) formation. This reduced nucleophilicity and hence reactivity towards \(\text{CO}_2\) hindered dicarbamate formation, correlating with the observation of a single \(\nu_{asCOCO^-}\) peak in the IR spectrum. The reduced prevalence of dicarbamate formation resulted in increased hydrolysis and HCO\(_3^-\), as evidenced by rapid growth of the \(\nu_{C-O}\) band at 1354 cm\(^{-1}\) (Figure 5a, b). The \(\alpha\)-dimethylamine moiety acted catalytically, in a manner analogous to that reported for sterically hindered amines, to accelerate formation of HCO\(_3^-\).\(^{[10]}\)

The subtle structural variations between 2,6-DMPZ and 2,5-DMPZ resulted in a significant change in the IR profile. In the case of the 2,5-DMPZ/\(\text{CO}_2\)/\(\text{H}_2\text{O}\) system the most dominant feature is HCO\(_3^-\) absorbance, as evidenced by the intense peak in
the 1400–1300 cm\(^{-1}\) region. There is little evidence to support formation of a stable carbamate (Figure 6).\[^{10, 17}\]

### 2.1.3. Hexahydropyrimidine (HHPY)

The HHPY/CO\(_2\)/H\(_2\)O system displayed some similarity with the PZ/CO\(_2\)/H\(_2\)O system in terms of signal positioning but with evidently weaker signals, due in part to the lower concentration of amine (HHPY) available for this study. Notwithstanding this, Figure 7 shows evolution of \(v_{\text{asCOO}}\) at 1570–1520 cm\(^{-1}\), \(v_{\text{asCOO}}\) at 1427 cm\(^{-1}\) and \(v_{\text{NH-bend}}\) at 1293 cm\(^{-1}\) of HHPY-COO\(^-\); the NH\(^+\) bending mode of HHPY-H\(^+\) at 1479 cm\(^{-1}\); and HCO\(_3^-\) absorbance at 1354 cm\(^{-1}\). The HCO\(_3^-\) absorbance band was more prominent than that observed for the PZ/CO\(_2\)/H\(_2\)O system, that is, HHPY forms a more labile NCOO\(^-\) derivative that is more susceptible to hydrolysis.

### 2.1.4. 2-Methylhexahydropyrimidine (MHHPY)

The IR spectrum of the MHHPY/CO\(_2\)/H\(_2\)O system is dominated by the broad HCO\(_3^-\) absorbance band in the 1400–1300 cm\(^{-1}\) region (Figure 8), which is characteristic of \(\alpha\)-substituted amines such as AMP.\[^{10, 17}\] MHHPY is the methyl-substituted analogue of HHPY. Given the intensity of the HCO\(_3^-\) band in Figure 8, MHHPY was readily hydrolysed under the study conditions, with HCO\(_3^-\) formation dominating on absorption of CO\(_2\).

### 2.1.5. 2,2-Dimethylhexahydropyrimidine (DMHHPY)

Given the structural similarity between DMHHPY and MHHPY, we anticipated predominant HCO\(_3^-\) formation on CO\(_2\) absorption by DMHHPY. However, the IR profile obtained for the DMHHPY/CO\(_2\)/H\(_2\)O system (Figure 9a) was significantly different to that obtained with MHHPY (Figure 8) and the HHPY and PZ systems (Figures 7 and 3, respectively). Here DMHHPY is acting more in keeping with a blended-amine PPC absorbent system. Close examination of the in-house synthesized DMHHPY revealed the presence of unconverted 1,3-diaminopropane (DAP) which had been unavoidably carried forward to the final product. Hence, the contamination of DMHHPY with DAP explains the observed blended-system-like IR profile (Figure 9a).

Re-examination of the IR spectrum of the DMHHPY/CO\(_2\)/H\(_2\)O system identified the NH bending mode of 1,3-diaminopropane at 1602 cm\(^{-1}\). While 1,3-diaminopropane [approximately 37%, 1.91 g (\(\text{H}^1\) NMR)] was the minor component within the
DMHHPY (63%, 3.24 g)/CO₂/H₂O system, it formed predominantly the corresponding carbamate (DAP) on CO₂ absorption.

To allow potential deconvolution of the DAP and DMHHPY signals in the original DMHHPY/CO₂/H₂O IR profile, data were collected separately for a DAP/CO₂/H₂O system at a DAP concentration of 0.6 mol L⁻¹ (Figure 9b). It is apparent that the original DMHHPY system is dominated by the reactivity of DAP (cf. Figure 9a, b). Both systems show evolution of νasCOO⁻ (1565 and 1568 cm⁻¹, respectively), νsCOO⁻ (1440 cm⁻¹) and νNH₃⁺ bending mode of protonated DAP (1492 cm⁻¹).

For the blended DMHHPY/CO₂/H₂O system, weaker absorbance bands were also observed to emerge at 1385 and 1370–1350 cm⁻¹ at CO₂ loadings above 1.0 mol CO₂/mol amine. These new peaks are consistent with NCOO⁻ hydrolysis and HCO₃⁻ formation. Carbamate hydrolysis was not observed for the pure DAP/CO₂/H₂O system.

In the initial DMHHPY/CO₂/H₂O system (Figure 9a) the DAP-COO⁻ absorbance bands dominate the IR spectrum. However in the DAP/CO₂/H₂O systems the carbamate absorbances are considerably weaker, the initial effect of which was thought to be that the DAP concentration on the DMHHPY system appears to be significantly higher than the 0.6 mol L⁻¹ evident in Figure 9b. However, a similar difference in intensity between the carbamate absorbance bands of a blended AMP (2.4 mol L⁻¹)/PZ (0.6 mol L⁻¹) system (Figure 10, further described below) versus an unblended PZ (0.6 mol L⁻¹) system was also observed (Figure 11). Carbamate absorbance in the unblended PZ system was found to be considerably weaker than that observed for the AMP/PZ blended system, despite equivalent PZ concentrations (6 mol L⁻¹). Based on the percentage concentrations determined by ¹H NMR spectroscopy the ratio of DMHHPY to DAP in the blended system was 0.95/0.85 mol L⁻¹ (total concentration 1.8 mol L⁻¹).

For comparative purposes an AMP/PZ blended amine system (2.4 mol L⁻¹/0.6 mol L⁻¹, respectively) was also investigated. Similarly to the blended DMHHPY absorbent, the AMP/PZ blend consists of an amine that forms predominately HCO₃⁻ on absorption of CO₂ (AMP, major constituent) and an amine that forms predominately NCOO⁻ (PZ, minor constituent). A similar trend in IR absorbance was observed in the spectral data collected for the AMP/PZ/CO₂/H₂O system (Figure 10) to that described above for the blended DMHHPY/CO₂/H₂O system. Despite PZ being the minor constituent of the amine blend, the PZ-carbamate absorbance bands dominated the IR spectral profile. Figure 10 shows the evolution of νasCOO⁻ (1533 cm⁻¹), νsCOO⁻ (1421 cm⁻¹) and νNH₃⁺ (1276 cm⁻¹ and shifts to 1263 cm⁻¹) of the PZ-COO⁻ derivative and the NH₂⁺ vibration of PZ-H⁺ (1471 cm⁻¹). HCO₃⁻ absorbance was seen to emerge in the 1386–1330 cm⁻¹ region after a CO₂ loading of about 0.5 mol CO₂/mol amine. For comparison Figure 11a and b present the IR spectral profile for an unblended AMP/CO₂/H₂O system (3 mol L⁻¹) and PZ/CO₂/H₂O system (0.6 mol L⁻¹), respectively. The HCO₃⁻ absorbance band was more prominent in the IR spectra of the AMP/PZ/CO₂/H₂O system compared to that of the DMHHPY/CO₂/H₂O system. This was most likely due to the difference in amine concentrations, with the AMP/PZ blend having a total concentration of 3 mol L⁻¹ and the DMHHPY/1,3-diaminopropane blend a total concentration of 1.5–1.8 mol L⁻¹.
2.1.6. Hexahydropyridazine

HHPZ absorbed no CO₂ during a typical CO₂ absorption/FTIR experiment. HHPZ is a hydrazine derivative that is reported to have a \( pK_a \) value of 7.9,\(^{[23]} \) which is much lower than that of PZ (9.73),\(^{[24]} \) HHPY (9.75)\(^{[25]} \) or 2,5-DMPZ (9.66).\(^{[26]} \) The low basicity of HHPZ compared to the other amines (\( pK_a > 9.5 \)) would significantly reduce the reactivity of the amine towards CO₂.

2.2. Absorption Capacity and Absorption Rate

Having established the ability of our diamines to absorb CO₂, the initial absorption rate \( R_a \) and absorption capacity \( C_A \) were determined. The \( R_a \) value was measured by a thermal gravimetric analysis (TGA) method, and \( C_A \) was measured simultaneously with the IR spectral data (see Experimental Section). These data are presented in Table 1. For comparison, the reactivity of conventional absorbents MEA, DEA and AMP (\( \alpha \)-dihydroxy-substituted MEA) are also included.

| Amine   | \( C_A \) [mol CO₂/mol amine] | \( R_a \) [mol CO₂/mol amine min⁻¹] |
|---------|-----------------------------|------------------------------------|
| PZ      | 0.92                        | 0.045                              |
| 2,6-DMPZ | 0.86\(^{[1]} \)            | 0.025\(^{[4]} \)                   |
| 2,5-DMPZ | 0.88                        | 0.018\(^{[1]} \)                   |
| HHPY    | 0.85                        | 0.032                              |
| MHHPY   | 0.86                        | 0.018                              |
| DMHHPY  | 1.33                        | 0.032                              |
| HHPZ    | 0                          | 0                                  |
| MEA     | 0.56\(^{[12],[1]} \)       | 0.027                              |
| DEA     | 0.60\(^{[12],[1]} \)       | 0.015                              |
| AMP     | 0.84\(^{[12],[1]} \)       | 0.006                              |

[a] Mol CO₂/mol amine; data used to calculate \( C_A \) were measured in the absorption reactor/FTIR system. [b] Mol CO₂/mol amine, min⁻¹; data used to calculate \( R_a \) were measured by microscale TGA. Initial absorption rates were calculated by using linear regression to determine the slope of the absorption capacity curve. \( R^2 > 0.995. [c] A precipitate formed during CO₂ absorption/FTIR. [d] A precipitate formed during the CO₂ and N₂ runs of the TGA experiment. [e] A precipitate formed during the CO₂ run of the TGA experiment. This could be due to the evaporation of water. [f] 3 mol L⁻¹ concentration analysed.

The current industry-standard PCC amine MEA returned \( C_A = 0.56 \) mol CO₂/mol amine and \( R_a = 0.027 \) mol CO₂/mol amine min⁻¹. From the data amassed for PZ, 2,6-DMPZ, HHPY and DMHHPY in Table 1, superior \( C_A \) and \( R_a \) values were observed for all these diamines relative to MEA. Superior \( C_A \) values were also observed for 2,5-DMPZ and MHHPY, but with lower \( R_a \) values. HHPZ did not react with CO₂, and DMHHPY was blended with DAP. Diamine \( C_A \) values ranged from 0.85 (HHPY) to 0.92 (PZ) mol CO₂/mol amine and \( R_a \) values from 0.018 (2,5-DMPZ) to 0.045 (PZ) mol CO₂/mol amine min⁻¹. While in absolute terms PZ was the standout pure diamine with the highest \( C_A \) (0.92 mol CO₂/mol amine) and \( R_a \) (0.045 mol CO₂/mol amine min⁻¹), \( C_A \) and \( R_a \) are not the sole factors to be considered in determining the most efficient PCC diamine absorbent; aqueous solubility and stability of the carbamate also play a role. HHPY displays higher water solubility than PZ, on the basis of observations when preparing 1.5 mol L⁻¹ amine solutions. HHPY was readily soluble at this concentration, as opposed to PZ, which required heating and stirring for dissolution. Additionally, HHPY displays high \( C_A \) (0.85 mol CO₂/mol amine) and \( R_a \) (0.032 mol CO₂/mol amine min⁻¹; Table 1) and showed clear evidence of formation of a hydrolysis-susceptible carbamate (see above and Figure 7).

The PZ analogues 2,6-DMPZ and 2,5-DMPZ displayed lower \( R_a \) values of 0.025 and 0.018 mol CO₂/mol amine min⁻¹, respectively. As these analogues differ only in the number of methyl substituents (PZ has none) and their positioning, these data suggested that introduction of methyl moieties had an adverse effect on the initial rate of CO₂ absorption. 2,6-DMPZ has two methyl groups \( \alpha \) to a single NH group, while 2,5-DMPZ has one methyl group \( \alpha \) to each NH group. The measured \( R_a \) values indicate that the effect of addition of \( \alpha \)-methyl groups is cumulative, with \( R_a \) dropping from 0.032 (PZ) to 0.025 (2,6-DMPZ) to 0.018 mol CO₂/mol amine min⁻¹ (2,5-DMPZ). Concurrent with the reduction in \( R_a \) was an increased prevalence towards HCO₃⁻ formation for 2,5-DMPZ (see above and Figure 6). The reactivity of 2,6-DMPZ towards CO₂ was found to be similar to that of MEA, with the exception of a higher \( C_A \) value. The propensity of 2,5-DMPZ for HCO₃⁻ formation was similar to that observed with MHHPY (see above and Figure 8), which was reflected in the almost identical \( C_A \) (0.88 and 0.86 mol CO₂/mol amine respectively) and \( R_a \) (0.018 and 0.018 mol CO₂/mol amine min⁻¹, respectively) values obtained for these amines. The reactivity of 2,6-DMPZ towards CO₂ was found to be similar to that of MEA, with the exception of a higher \( C_A \) value. 2,6-DMPZ was found to form predominantly carbamate on absorption of CO₂ (Figure 5), similar to HHPY (Figure 7). While the propensity for carbamate hydrolysis and subsequent HCO₃⁻ formation is also similar to that observed with HHPY (cf. Figures 5 and 7), which was reflected in the almost identical \( C_A \) (0.88 and 0.85 mol CO₂/mol amine, respectively) values, 2,6-DMPZ returned a lower \( R_a \) value (0.025 and 0.032 mol CO₂/mol amine min⁻¹, respectively). Despite forming predominately HCO₃⁻, the initial absorption rates obtained for both MHHPY and 2,5-DMPZ were much higher than that obtained for AMP and comparable to those of MEA and DEA (Table 1).

Of the diamines examined, DMHHPY exhibited the highest \( C_A \) (1.1 mol CO₂/mol amine value) and an \( R_a \) value higher than that of MHHPY and comparable to that of HHPY. This is an artefact of the serendipitous blending with DAP, which contributes significantly to the observed CO₂ absorption capacity. Aqueous DAP has \( C_a = 0.95 \) mol CO₂/mol amine. The bicarbonate-forming DMHHPY further promotes CO₂ absorption, resulting in \( C_A > 1.0 \) mol CO₂/mol amine. The 1′ amino groups of DAP contribute towards DMHHPY’s increased initial reaction rate compared to HHPY.
2.3. Effect of Structural Variations on Carbamate Structures

The effect of diamine structural variation on the ability to form stable carbamates was examined at the B3LYP/6-31+G* and MP2/6-31+G** levels of theory. Geometry optimisations were initially performed on PZ, 2,6-DMPZ, 2,5-DMPZ, HHPPY, MHHPY, DMHPY and HHPZ. Table 3 lists selected atomic properties including electrostatic potential (ESP) partial charges on the amino nitrogen atoms and the exposed area on the nitrogen atoms for these diamines. The trends in results obtained at the two levels of theory were found to be in good agreement with one another.

Diamines can react with CO2 in aqueous solution to form three possible forms of carbamate species: amine carbamate (HNR,RNCOO−), protonated amine carbamate (H−HNR,RNCOO−), and dicarbamate. Of these three forms, "HHR,R,NCOO− was expected to be the main reaction product, with one amino group acting as the binding site for CO2 and the other as a proton acceptor. Geometry optimisations were next performed for the protonated amine-carbamate ("H-HNR,R,NCOO−) and amine-carbamate (HNR,R,NCOO−) species. For the lowest-energy conformer of each "HNR,R,NCOO− and HNR,R,NCOO− derivative, Table 3 provides the calculated N−COO− bondlength (r_{N−C−A}), t_{C1−O−C1−O2} [Å], and ESP partial negative charges on both oxygen atoms as a measure of charge delocalisation.

As anticipated, methyl substitution significantly increased the partial negative charges on the amino groups of 2,6-DMPZ (N1 = −0.718; N2 = −0.752), 2,5-DMPZ (N1 = −0.851; N2 −0.728) and DMHPY (N1 = −0.851; N2 = −0.881) and reduced the exposed area.

Table 2. ESP charges and exposed areas on the nitrogen atoms [Å2] for optimised forms of the diamines analysed herein.

| Amine          | ESP charge on N | Exposed area on N [Å2] |
|----------------|----------------|-----------------------|
|                | B3LYP | MP2 | B3LYP | MP2 |
| PZ N1          | −0.584 | −0.640 | 4.71 | 4.80 |
| N2             | −0.584 | −0.640 | 4.71 | 4.80 |
| 2,6-DMPZ N1    | −0.718 | −0.752 | 4.21 | 4.25 |
| N2             | −0.752 | −0.778 | 4.70 | 4.82 |
| 2,5-DMPZ N1    | −0.746 | −0.785 | 4.45 | 4.54 |
| N2             | −0.746 | −0.785 | 4.45 | 4.54 |
| HHPPY N1       | −0.874 | −0.907 | 4.91 | 4.96 |
| N2             | −0.874 | −0.907 | 4.91 | 4.96 |
| MHHPY N1       | −0.851 | −0.881 | 4.77 | 4.83 |
| N2             | −0.728 | −0.759 | 4.57 | 4.65 |
| DMHPY N1       | −0.959 | −0.979 | 4.50 | 4.57 |
| N2             | −0.957 | −0.982 | 4.51 | 4.57 |
| HHPZ N1        | −0.525 | −0.551 | 6.82 | 6.89 |
| N2             | −0.386 | −0.402 | 5.71 | 5.77 |

Table 3. Calculated N−COO− bond lengths r_{N−C−A} [Å], r_{C1−O−C1−O2} [Å] and ESP partial charge on both oxygen atoms for optimised geometries of the "HNR,R,NCOO− and HNR,R,NCOO− derivatives of the subset of diamines analysed.

| Carbamate derivative | r_{N−C−A} [Å] | MP2 | r_{C1−O−C1−O2} [Å] | MP2 | ESP charge on O1/O2 | MP2 |
|---------------------|---------------|-----|-------------------|-----|---------------------|-----|
| H−−PZ-carbamate     | 1.513         | 1.507 | 1.247/1.247       | 1.256/1.256 | −0.690/−0.690       | −0.681/−0.681 |
| PZ-carbamate        | 1.471         | 1.477 | 1.257/1.257       | 1.264/1.264 | −0.743/−0.743       | −0.739/−0.739 |
| H−−2,6-DMPZ-carbamate | 1.508       | 1.505 | 1.248/1.249       | 1.256/1.256 | −0.688/−0.689       | −0.680/−0.680 |
| 2,6-DMPZ-carbamate  | 1.469         | 1.477 | 1.258/1.258       | 1.264/1.264 | −0.772/−0.772       | −0.767/−0.767 |
| H−−2,5-DMPZ-carbamate | 1.523      | 1.527 | 1.247/1.247       | 1.255/1.255 | −0.691/−0.692       | −0.699/−0.698 |
| 2,6-DMPZ-carbamate  | 1.461         | 1.463 | 1.250/1.260       | 1.266/1.266 | −0.753/−0.753       | −0.749/−0.749 |
| H−−2,5-DMPZ-carbamate | 1.505       | 1.503 | 1.249/1.247       | 1.259/1.253 | −0.699/−0.665       | −0.702/−0.656 |
| 2,5-DMPZ-carbamate  | 1.465         | 1.470 | 1.259/1.258       | 1.266/1.264 | −0.750/−0.735       | −0.746/−0.731 |
| H−−HHPPY-carbamate  | 1.507         | 1.503 | 1.227/1.277       | 1.235/1.284 | −0.641/−0.735       | −0.625/−0.736 |
| HHPPY-carbamate     | 1.475         | 1.478 | 1.257/1.259       | 1.263/1.267 | −0.762/−0.767       | −0.744/−0.751 |
| H−−MHHPY-carbamate  | 1.417         | 1.417 | 1.211/1.354       | 1.211/1.354 | −0.650/−0.772       | −0.640/−0.782 |
| MHPPY-carbamate     | 1.463         | 1.469 | 1.258/1.260       | 1.264/1.266 | −0.753/−0.770       | −0.746/−0.759 |
| H−−DMHHPY-carbamate | 1.535         | 1.550 | 1.235/1.244       | 1.243/1.253 | −0.653/−0.699       | −0.648/−0.702 |
| DMHPY-carbamate     | 1.501         | 1.500 | 1.252/1.261       | 1.258/1.268 | −0.758/−0.819       | −0.743/−0.816 |
| H−−HHPZ-carbamate   | 1.390         | 1.396 | 1.215/1.350       | 1.221/1.353 | −0.575/−0.660       | −0.566/−0.670 |
| HHPZ-carbamate      | 1.460         | 1.470 | 1.254/1.263       | 1.260/1.267 | −0.741/−0.759       | −0.736/−0.758 |

(a) Isomer 1: R1 = R2 = CH3 and R3 = R4 = H (Figure 12). (b) Isomer 2: R1 = R2 = CH3 and R3 = R4 = H (Figure 12). (c) R1 = R3 = CH3 and R2 = R4 = H (Figure 12). (d) Hydrogen bonding between O2 of the carbamate moiety and a proton of the NH + group (Figure 12).
on the amino nitrogen atom of 2,6-DMPZ (N1 4.21 Å; N2 4.70 Å), 2,5-DMPZ (N1, N2 4.45 Å), MHHPY (N1 4.77 Å; N2 4.47 Å) and DMHHPY (N1 4.50 Å; N2 4.51 Å) relative to PZ (N1, N2 4.71 Å) at the B3LYP level. These changes in electronic effects both have impact on the resonance structure of the carboxylate moiety and hence overall stability of the carbamate derivative (see below).

Studies herein (Figure 3 and 4) demonstrated that PZ forms a hydrolysis-resistant NCOO− derivative, and thus the optimised geometries of H−-PZ-COO− and PZ-COO− were used as the baseline against which the remaining amine NCOO− derivatives were compared. For both H−-PZ-COO− and PZ-COO− resonance stabilisation of the carboxylate moiety is evident, with calculations showing identical charges on O1 and O2 as well as displaying identical r_C=O values. The r_C=O of 1.247 and 1.257 Å reveal partial double-bond character for H−-PZ-COO− and PZ-COO−, respectively (Table 3). The N−COO− bond length of both PZ-COO− species was of single-bond character (standard single-bond r_N=C in PZ is 1.466 Å (B3LYP) and 1.465 Å (MP2)). H−-2,6-DMPZ-COO− and 2,6-DMPZ-COO− also exhibit this stable resonance structure, albeit with slightly shorter N−COO− bond lengths. These findings are in keeping with our IR studies on 2,6-DMPZ and PZ (Figure 5), which gave very similar outcomes, with the exception of the emergence of a small HCO3− absorbance band. The protonated and unprotonated 2,6-DMPZ-COO− forms of isomer 2 (Figure 12), the minor reaction component, contributed to HCO3− formation.

H−-HHPY-COO− displays lower levels of charge delocalisation across the two oxygen atoms. Changes in charge distribution and bond length were noted with r_C=O = 1.227 Å and r_O2 = 1.277 Å, which were mirrored in the change in electron density at O1 (−0.641) and O2 (−0.735) (B3LYP). The shift in electron distribution was much less pronounced in the HHPY-COO− species with an r_C=O of 1.257 Å and a r_O2 of 1.259 Å which was mirrored in the change in electron density at O1 (−0.762) and O2 (−0.767) (B3LYP). The N−COO− bond lengths of H−-HHPY-COO− (1.507 Å) and HHPY-COO− (1.475 Å) are similar to that of the PZ-COO− species (1.471 Å). These findings are in keeping with our IR studies (Figure 7), in which HHPY was identified as forming a more labile NCOO− derivative that is more susceptible to hydrolysis than that of PZ. The lowest energy conformer obtained for H−-HHPY-COO−, as opposed to HHPY-COO− species, exhibited intramolecular hydrogen bonding between the COO− group and the NH3+ moiety. The low-energy conformers of H−-MHHPY-COO− and H−-MHHPZ-COO− were also found to exhibit the same intramolecular hydrogen bonding. The resonance structure of the H−-MHHPY-COO− was less delocalised with r_C=O = 1.211 Å and r_O2 = 1.354 Å, which was mirrored in the changes in electron density at O1 (−0.650) and O2 (−0.772) (B3LYP). The N−COO− bond length of H−-MHHPY-COO− was found to be much shorter (1.417 Å). The shift in electron distribution of the carboxylate resonance structure was again much less pronounced in the MHHPY-COO− species compared to the H−-MHHPY-COO− species. MHHPY-COO− has a shorter N−COO− bond length (1.463 Å) than PZ-COO− (1.471 Å) and MHHPY-COO− (1.475 Å). MHHPY forms predominantly HCO3− on absorption of CO2 as does 2,5-DMPZ. Both species display shorter N−COO− bonds and lower levels of resonance stabilisation.

In our IR studies DMHHPY was found to be representative of a blended amine system. Nonetheless, optimised geometries of the H−-DMHHPY-COO− and DMHHPY-COO− were still analysed, and both exhibited reduced resonance in the carboxylate moiety with r_C=O = 1.235 Å and r_O2 = 1.244 Å, which was mirrored in the changes in electron density at O1 (−0.653) and O2 (−0.699), and with r_C=O = 1.252 Å and r_O2 = 1.261 Å, which was mirrored in the changes in electron density at O1 (−0.758) and O2 (−0.819) (B3LYP), respectively. Given the structural similarity of DMHHPY and MHPHY, DMHHPY was expected to form predominantly HCO3− on absorption of CO2.

Experimentally HHPZ was unreactive towards CO2. Calculations revealed H−-HHPZ-COO− to have a significantly shorter N−COO− bond length of 1.390 Å (significant double-bond character), as well as the largest displacement in electron distribution of the carboxylate resonance structure with r_C=O = 1.215 Å and r_O2 = 1.350 Å, which was mirrored in the changes in electron density at O1 (−0.575) and O2 (−0.660) (B3LYP). This was much less pronounced in the HHPZ-COO− species; nonetheless, in a diamine system it is typical for one amino group to act as binding site for CO2 while the other is protonated. These data support our experimental observations.

3. Conclusions

A series of heterocyclic diamines (PZ, 2,6-DMPZ, 2,5-DMPZ, HHPY, MHHPY, DMHHPY and HHPZ) have been evaluated as potential PCC absorbents by in situ ATR-FTIR spectroscopy. Of these diamines, PZ displayed both the highest CO2 absorption capacity (Cw = 0.92 mol CO2/mol amine) and highest initial absorption rate (R0 = 0.045 mol CO2/mol amine min−1). These values represent a significant enhancement over currently used amines such as MEA. PZ forms a hydrolysis-resistant carbamate, as well as a dicarbamate. This behaviour is unique to PZ. Hydrolysis of the carbamate derivative of HHPZ was observable in the IR spectra collected during CO2 absorption. HHPZ displayed similar CO2 absorption characteristics to PZ, but with a higher propensity for HCO3− formation. The introduction of α-methyl substituents increased the propensity towards carbamate hydrolysis and HCO3− formation. Additionally α-methyl substitution decreased R0 with PZ analogues 2,6-DMPZ and 2,5-DMPZ displaying lower R0 values of 0.025 and 0.018 mol CO2/mol amine min−1, respectively. Increasing the number of methyl groups α to the NH group also increases the rate of HCO3− formation. Despite forming predominately HCO3−, the R0 of MHHPY (0.018 mol CO2/mol amine min−1) was much higher than that of the corresponding α-dimethyl substituted 1° amine AMP (R0 = 0.006 mol CO2/mol amine min−1) and comparable with that of the industrially relevant MEA and DEA. The serendipitously blended DAP/DMHHPY exhibited the highest Cw (1.1 mol CO2/mol amine) and excellent R0 (0.032 mol CO2/mol amine min−1). HHPZ was found to be relatively unreactive towards CO2. In all instances our calculations at the B3LYP/6-31+G* and MP2/6-31+G** levels of theory supported our experimental observation. Finally, we
Experimental Section

General: All starting materials were purchased from Sigma Aldrich and used without further purification. Solvents were bulk and distilled prior to use. 1H and 13C NMR spectra were recorded on a Bruker Avance AMX 300 MHz spectrometer at 300.1315 and 75.4762 MHz, respectively. Chemical shifts δ are reported relative to internal standards. Mass spectra were recorded on a Shimadzu LCMS-2010 EV spectrometer and obtained by the ESI method. IR spectra were recorded by ATR-FTIR on a Mettler-Toledo FTIR spectrometer.

Preparation of Hexahydropyrimidine (HHYP): Formaldehyde (37 wt% solution, 1.5 mol, 126 g) was added dropwise to an ice-cooled, stirred solution of anhydrous 1,3-diaminopropane (1 mol, 74.1 g) over 30 mins. The reaction mixture was then stirred for 24 h at room temperature and cooled in an ice bath, and NaOH was added (1 mol, 40 g). The organic layer was transferred to a Dean–Stark apparatus and azeotropically distilled with cyclohexane (100 mL). The cyclohexane was removed in vacuo and the residue fractionally distilled to afford 2-methylhexahydropyrimidine (16.17 g, 71%) as a clear oil (b.p. 56–59°C/30 mm Hg). 1H NMR (300 MHz, CDCl3): δ = 1.36 (2H, m), 1.96 (2H, brs), 2.82 (4H, t, J = 5.5 Hz), 3.63 ppm (2H, s); 13C NMR (75 MHz, CDCl3): δ = 26.81, 45.65, 62.79 ppm; MS (ESI+): m/z 87 (M+1); FTIR: ν = 3266 cm⁻¹.

Preparation of 2-Methylhexahydropyrimidine (MHHPY): Acetaldehyde (0.215 mol, 9.5 g) in diethyl ether (100 mL) was added dropwise to ice-cooled 1, 3-diaminopropane (0.2 mol, 14.8 g). The reaction mixture was then stirred over K₂CO₃ (0.4 mol, 55.4 g) for 24 h at room temperature. The solvent was removed in vacuo and the residue fractionally distilled to afford 2-methylhexahydropyrimidine (17.5 g, 87%) as a clear oil (b.p. 56–59°C/30 mm Hg, lit. [28]: b.p. 60°C/30 mm Hg). 1H NMR (300 MHz, CDCl3): δ = 0.80 (3H, d, J = 6.0 Hz), 1.09 (2H, m), 2.48 (2H, m), 2.76 (4H, m), 3.25 ppm (1H, q, J = 6.0 Hz); 13C NMR (75 MHz, CDCl3): δ = 22.75, 26.80, 45.41, 67.09 ppm; MS (ESI+): m/z 87 (M+1); FTIR: ν = 3266 cm⁻¹.

Preparation of 2-Dimethylhexahydropyrimidine (DMHHYP): A solution of acetone (0.215 mol, 12.5 g) in diethyl ether (100 mL) was added dropwise to ice-cooled 1, 3-diaminopropane (0.2 mol, 14.83 g). The reaction mixture was then stirred over K₂CO₃ (0.4 mol, 55.4 g) for 24 h at room temperature. The solvent was removed in vacuo and the residue fractionally distilled to afford 2, 2-dimethylhexahydropyrimidine (16.17 g, 71%) as a clear oil (b.p. 50–54°C/25 mbar, lit. [28]: b.p. 65°C/25 mm Hg). 1H NMR (300 MHz, CDCl3): δ = 0.90 (6H, s), 1.05 (2H, m), 1.45 (2H, s), 2.60 ppm (4H, t, J = 5.7 Hz); 13C NMR (75 MHz, CDCl3): δ = 27.26, 36.87, 39.99, 64.31 ppm; MS (ESI+): m/z 115 [M+1]; FTIR: ν = 3271, 1602 cm⁻¹.

Preparation of Hexahydropyridazine (HHPZ): Dibromobutane (0.1 mol, 21.6 g) was added dropwise to a stirred solution of diethylhydrazine dicarboclylate (0.1 mol, 17.6 g), K₂CO₃ (0.2 mol, 27.7 g) and acetonitrile (100 mL) at room temperature. The reaction mixture was then heated to reflux for 24 h, cooled and filtered. The solvent was removed in vacuo and the residue fractionally distilled to afford 1, 2-dicarbethoxyhexahydropyridazine (18.5 g, 80%) as a clear oil (b.p. 110–114°C/2–3 mbar, lit. [29]: 106–114°C/3 mm Hg). 1H NMR (300 MHz, CDCl₃): δ = 1.07 (6H, m), 1.49 (4H, s), 2.74 (4H, brs), 3.97 ppm (4H, m); 13C NMR (75 MHz, CDCl₃): δ = 14.36, 23.19, 44.76, 61.93, 155.15 ppm; MS (ESI+): m/z 231 [M+1]; FTIR: ν = 1706 cm⁻¹.

1,2-Dicarbethoxyhexahydropyridazine (0.13 mol, 30.1 g) was added to a solution of NaOH (0.72 mol, 20.8 g), water (20 mL) and methanol (150 mL). The reaction mixture was then heated to reflux for 20 h, cooled and the precipitated inorganic salt removed by filtration. The filtrate was heated to reflux for a further 20 h, cooled, filtered and the solvent removed in vacuo. The residue was extracted with dichloromethane (40 mL×3), concentrated in vacuo and fractionally distilled to afford hexahydropyridazine (6.75 g, 60%) as a yellow oil (b.p. 38–40°C/11 mbar, lit. [30]: b.p. 38°C/8 mm Hg).

1H NMR (300 MHz, MeOD): δ = 2.39 (4H, m), 3.73 (4H, m), 5.72 ppm (2H, s); 13C NMR (75 MHz, MeOD): δ = 23.06, 47.00 ppm; MS (ESI+): m/z 87 [M+1]; FTIR: ν = 3269 cm⁻¹.

Microscale Thermogravimetric Analysis (TGA): A Setaram Labsys TG-DTA/DSC thermogravimetric analyser (TGA) was used in isothermal mode at 40°C to analyse aqueous CO₂/amine reactivity on a microscale (100 µL). 1.5 mol L⁻¹ aqueous amine solutions were exposed to a gas stream of 15 vol% CO₂ (>99.9% purity, BOC Australiia) in N₂ at atmospheric pressure. A gas flow rate of 30 mL min⁻¹ was used for all experiments. To determine total CO₂ uptake two separate TGA experiments were performed for each amine. The first experimental run determined the mass loss due to evaporation when the test solution was exposed to a 100% N₂ gas stream. The second experimental run determined the mass increase of the test solution when exposed to CO₂ (15 vol% CO₂ in N₂ gas stream) over the same length of time. Each experiment was performed on a fresh 100 µL aliquot of the test solution in a 100 µL alumina crucible (Setaram). From the data collected an absorption curve was then calculated for each amine by subtracting the mass at time t of the evaporation run from the mass at time t of the absorption run. Initial absorption rates could then be calculated by using linear regression to determine the slope of the absorption curve.

Absorption/FTIR Experiments: The absorption reactor apparatus used to analyse aqueous CO₂/amine reactivity has been described previously,[10] Briefly, a gas stream of 13 vol% CO₂ (>99.9% purity, BOC Australia) in N₂, with a flow rate of 1.8 L.min⁻¹ was bubbled through a 1.5 mol L⁻¹ aqueous amine solution in a glass reactor vessel, maintained at 40°C by a temperature-controlled water bath (Techno). The CO₂ content of both the gas inflow and outflow was measured by using a Horiba VA 3000 CO₂ analyser. The difference between the CO₂ concentration of the reactor gas inflow and gas outflow was used determine the amount of CO₂ absorbed by the amine solution (mol CO₂/mol amine). Each experiment was run until the measured CO₂ concentration in the outflow returned to the original percentage value, that is, equilibrium was reached. A typical run lasted between 45 and 90 mins. Solution volumes of 30 mL were used for all experiments.

For the duration of each absorption experiment an ATR diamond-tipped IR probe, coupled via a mirrored K6 conduit to an i10 FTIR spectrometer (Mettler-Toledo), was immersed in the aqueous amine solution. In situ IR measurements were obtained simultaneously with the CO₂ absorption measurements, with the FTIR spectrometer set to continuously collect spectra for the duration of the absorption experiment over the spectral range of 4000–650 cm⁻¹. Each spectrum was recorded as the average of 256 scans over a sampling interval of fifteen seconds with a resolution of 4 cm⁻¹. The amines investigated were synthesised according to the procedures detailed above, with the exception of commercially available PZ, 2,6-DMPZ and 2,5-DMPZ (Sigma-Aldrich).
CO₂ Absorption Characteristics of Heterocyclic Amines

CO₂ Absorption Characteristics of Heterocyclic Amines

Acknowledgements

This project is part of the CSIRO Coal Technology Portfolio and received funding from the Australian Government as part of the Asia-Pacific Partnership on Clean Development and Climate. The views expressed herein are not necessarily the views of the Commonwealth, and the Commonwealth does not accept responsibility for any information or advice contained herein.

Keywords: absorption · amines · carbon dioxide fixation · IR spectroscopy · nitrogen heterocycles

[1] J. G. Canadell, C. Le Quere, M. R. Raupach, C. B. Field, E. T. Buitenhuis, P. Ciais, T. J. Conway, N. P. Gillett, R. A. Houghton, G. Marland, Proc. Natl. Acad. Sci. USA 2007, 104, 18866 – 18870.
[2] M. R. Raupach, G. Marland, P. Ciais, C. Le Quere, J. G. Canadell, G. Klepper, C. B. Field, Proc. Natl. Acad. Sci. USA 2007, 104, 10288 – 10293.
[3] CO₂CRC, CSIRO, 2006, Sate of the Art Review: National Post Combustion Capture and Storage Demonstration Program, Revision 2-1, PCC Ltd, Australia.
[4] International Energy Agency, 2007, IEA Greenhouse Gas R&D Programme: CO₂ Capture Ready Plants. http://www.iea.org/publications/free_all_papers.asp.
[5] D. Gielen, J. Podkanski, F. Unander, Energy Technology Analysis: Prospects for CO₂ Capture and Storage, International Energy Agency (IEA), Paris, 2004.
[6] E. Blomen, C. Hendriks, F. Neele, Energy Procedia 2009, 1, 1505 – 1512.
[7] M. Wang, A. Lawal, P. Stephenson, J. Sidders, C. Ramshaw, Chem. Eng. Res. Des. 2011, 89, 1609 – 1624.
[8] J. Davison, Energy 2007, 32, 1163 – 1176.
[9] M. Finkenrath, Cost and Performance of Carbon Dioxide Capture from Power Generation, IEA Energy Papers, No. 2011/05, 2011.
[10] K. Robinson, A. McCluskey, M. Attalla, ChemPhysChem 2011, 12, 1088 – 1099.
[11] D. J. Fauth, T. P. Filburn, M. L. Gray, S. W. Hedges, J. Hoffman, H. W. Pennline, T. Filburn, Abstr. Pap. Air and Waste Management Association 100th Annual Conference and Exhibition, Pittsburgh, 2007.
[12] D. Bonenfant, M. Mireault, R. Hausler, Ind. Eng. Chem. Res. 2003, 42, 3179 – 3184.
[13] E. F. da Silva, H. F. Svendsen, Ind. Eng. Chem. Res. 2006, 45, 2497 – 2504.
[14] P. D. Vaidya, E. Y. Kenig, Chem. Eng. Technol. 2007, 30, 1467 – 1474.
[15] T. L. Donaldson, Y. N. Nguyen, Ind. Eng. Chem. Fund. 1980, 19, 260 – 266.
[16] G. F. Versteeg, L. A. J. Van Dijck, W. P. M. Van Swaaij, Chem. Eng. Commun. 1996, 144, 113 – 158.
[17] K. Robinson, P. Jackson, M. Attalla, Abstr. Pap. CHEMeca 2009, Perth, Australia, 2009.
[18] A. K. Chakraborty, K. B. Bischoff, G. Astarita, J. R. Darnewood, J. Am. Chem. Soc. 1988, 110, 6947 – 6954.
[19] G. Sartori, D. W. Savage, Ind. Eng. Chem. Fund. 1995, 34, 1277 – 1289.
[20] S. Bishnoi, G. T. Rochelle, Chem. Eng. Sci. 1997, 52, 5531 – 5543.
[21] V. Ermatchkov, /C141. P/C216rez-Salado Kamps, G. Maurer, Ind. Eng. Chem. Res. 2000, 39, 1643 – 1661.
[22] J. M. Locke, R. J. Crumbie, R. Griffith, T. D. Bailey, S. Boyd, J. D. Roberts, J. Org. Chem. 2007, 72, 4156 – 4162.
[23] K. N. Zelenin, V. V. Alekseyev, I. V. Ukraintsev, I. V. Tselinsky, J. Chem. Thermodyn. 2003, 35, 1277 – 1289.
[24] Wavefunction Inc, Q-chem Inc, Spartan ’08 for Windows, Macintosh and Linux, Irvine, CA, 2008.
[25] R. F. Evans, Aust. J. Chem. 1967, 20, 1643 – 1661.
[26] J. Edwards, D. Ormsby, 2008, Organic Bases, http://ifs.massey.ac.nz/outreach/resources/chem/orgbases.php.
[27] J. M. Locke, R. J. Crumbie, R. Griffith, T. D. Bailey, S. Boyd, J. D. Roberts, J. Org. Chem. 2007, 72, 4156 – 4162.
[28] K. Robinson, A. McCluskey, M. Attalla, Abstr. Pap. CHEMeca 2009, Perth, Australia, 2009.
[29] F. Khalili, A. Henni, A. L. L. East, J. Chem. Eng. Data 2009, 54, 2914 – 2917.
[30] R. F. Evans, Aust. J. Chem. 1967, 20, 1643 – 1661.
[31] J. Edwards, D. Ormsby, 2008, Organic Bases, http://ifs.massey.ac.nz/outreach/resources/chem/orgbases.php.
[32] J. M. Locke, R. J. Crumbie, R. Griffith, T. D. Bailey, S. Boyd, J. D. Roberts, J. Org. Chem. 2007, 72, 4156 – 4162.
[33] K. Robinson, A. McCluskey, M. Attalla, Abstr. Pap. CHEMeca 2009, Perth, Australia, 2009.
[34] F. Khalili, A. Henni, A. L. L. East, J. Chem. Eng. Data 2009, 54, 2914 – 2917.
[35] R. F. Evans, Aust. J. Chem. 1967, 20, 1643 – 1661.
[36] J. Edwards, D. Ormsby, 2008, Organic Bases, http://ifs.massey.ac.nz/outreach/resources/chem/orgbases.php.
[37] J. M. Locke, R. J. Crumbie, R. Griffith, T. D. Bailey, S. Boyd, J. D. Roberts, J. Org. Chem. 2007, 72, 4156 – 4162.
[38] K. Robinson, A. McCluskey, M. Attalla, Abstr. Pap. CHEMeca 2009, Perth, Australia, 2009.
[39] F. Khalili, A. Henni, A. L. L. East, J. Chem. Eng. Data 2009, 54, 2914 – 2917.
[40] R. F. Evans, Aust. J. Chem. 1967, 20, 1643 – 1661.
[41] J. Edwards, D. Ormsby, 2008, Organic Bases, http://ifs.massey.ac.nz/outreach/resources/chem/orgbases.php.
[42] K. Robinson, A. McCluskey, M. Attalla, Abstr. Pap. CHEMeca 2009, Perth, Australia, 2009.
[43] F. Khalili, A. Henni, A. L. L. East, J. Chem. Eng. Data 2009, 54, 2914 – 2917.
[44] R. F. Evans, Aust. J. Chem. 1967, 20, 1643 – 1661.