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1. Introduction

Biomimetics as an interdisciplinary concept involves learning from and imitating of nature using appropriate models. The mimicry of biological models does not only consist of the construction of engineering systems and modern technology, furthermore, it is able to be a guide for designing and architecturing in the complete range of natural sciences. In particular, the actual development of novel and uncommon chemical reactions is aimed to apply high optimized, biochemical processes to laboratory and industrially usable syntheses. Especially enzymes stand in the visual focus as highly efficient and almost extreme specialized catalysts, as nature has had a long time to develop and optimize a wide diversity of life. Nearly every organism owns different tissues, motion types, sensing functions and surviving instincts and the basis for all of these properties are highly efficient catalyzed chemical reactions. Organisms have not only to synthesize huge biopolymers for growth, but also to provide small molecules for signal transduction, energy transfers or regulation processes. Therefore, it is obvious that the study of those reactions not only gives insights in important biochemical processes, but also suggests new chemical pathways to use in synthesis or other related applications. The possibilities of biomimetics in chemistry are outlined on the basis of the enzyme carbonic anhydrase. As this enzyme is one of the most efficient catalysts originated by nature, the investigation of such a catalytical active protein enriches not only the knowledge of biological and biochemical mechanisms and their mode of action, but also gives detailed information about the active center inside of the enzyme. In that way, one can fathom out important factors for the growth and development as well as for the investigation of diseases and glitches.

In this chapter we describe the enzymatic metal mediated reaction of the cumulene system carbon dioxide with water and its extension to the bio-analogue activation of further isoelectronic cumulenes (carbonyl sulfide, isothiocyanate and allene).

\[
\text{CO}_2 + \text{H}_2\text{O} \rightleftharpoons \text{HCO}_3^- + \text{H}^+ \rightleftharpoons \text{H}_2\text{CO}_3
\]  

Most of the theoretical (DFT calculations) and experimental results were obtained by the authors of this chapter or in direct collaboration with former members of the Jena group (Dr. Stephan Schenk, Dr. Johannes Notni) and other researchers: Prof. Dr. Jürgen Kesselmeier (Mainz, Germany), Prof. Dr. Heinrich Vahrenkamp (Freiburg, Germany), Prof. Dr. Eckhard
Dinjus and Prof. Dr. Manfred Doering (Karlsruhe, Germany), Prof. Dr. Wolfram Koch (Frankfurt, Germany), Prof. Dr. James Weston (La Plata, Argentina), Prof. Dr. Detlef Schröder (Prag, Czech), Prof. Dr. Helmut Schwarz (Berlin, Germany), Prof. Dr. Andrea Bottoni (Bologna, Italy) and Dr. Sten Nilsson Lill (Göteborg, Sweden). We made significant efforts to cite the relevant contributions by others; nevertheless, the discussion of their results will be very brief in order to keep this chapter within a reasonable length.

2. The Model Enzyme Carbonic Anhydrase

Carbonic anhydrases (CA) are an ancient enzyme class most likely as old as life on earth. As every organism contains CA, they might be one of the most important enzyme classes. Naturally, these enzymes catalyze the reversible reaction of carbon dioxide with water to bicarbonate (see equation 1). They belong to the enzyme class of lyases, which generally catalyzes the cleavage of molecules.

At present all CAs are divided into six classes (α - ζ class). All classes developed independently and therefore are examples of convergent evolution (Hewett-Emmett & Tashian, 1996). The representatives of the α class are found in all mammals and can be subdivided into currently fifteen isoforms. Depending on the individual structure, each isoform fulfills other specialized tasks in biochemical processes. The isoform CA-I was identified in erythrocytes (Lowe et al., 1990) and is the mostly spread protein beside of globuline and responsible for the conversion of the badly water soluble gas carbon dioxide into bicarbonate, which is completely soluble and hence better transportable in water. For this reason CA-I is the base for efficient respiration. CA-II, which is one of the most active isoforms, owns turn over numbers near the diffusion controlled limit (k_{cat} = 620 000 sec^{-1})(Gibbons & Edsall, 1964) and an acceleration factor of 10^7 compared to the uncatalyzed reaction (Lindskog, 1997). Deficiency of CA-II leads to osteopetrosis with renal tubular acidosis and cerebral calcification (Sly et al., 1983).

While CA-IV is involved in the ionic transport mechanisms in the endothel of the cornea (Sun et al., 2008), CA-V is placed in the mitochondria and used as a supplier for HCO_3^- in the urea cycle (Dodgson & Forster, 1986). CA-VII and CA-VIII are involved in neuronal signal forwarding and the provision of neurotransmitters in the central nervous system (Ruusuvuori et al., 2004; Yan et al., 2007). Another point of interest is the role of CAs in the development of cancer, as the activity of CA-IX and CA-XII could be proven in several tumor cells and tissues, for example breast, kidney or cerebral cancer (Barnett et al., 2008; Brennan et al., 2006; Chrastina et al., 2003; Jarvela et al., 2008).

2.1 Insights into the Enzymatic Mechanism

All CAs contain one zinc(II) cation as a cofactor (Keilin & Mann, 1939) with the exception of ζ-CAs, which possess a diatomic metal center. The central metal is not constricted to zinc in those CAs due to low zinc concentrations in the habitat of organisms containing them (Lane et al., 2005). In the catalytic core of α-class CAs the zinc ion is complexed by three histidine amino acid residues (His94, His96, His119) and a water or rather hydroxide ligand depending on the intracellular pH value (see Figure 1).

After getting into the hydrophobic pocket, CO_2 becomes immobilized by a hydrogen bridging bond to Thr199, which plays the role of a gatekeeper to the active center. The zinc bound hydroxide ion then initiates the hydrolysis by a nucleophilic attack on CO_2. In the following step the produced bicarbonate is displaced by a water molecule, which is subsequently deprotonated by the help of a proton relais to His64 to give zinc bound hydroxide for the next...
2. The Model Enzyme Carbonic Anhydrase

2.1 Insights into the Enzymatic Mechanism

Fig. 1. An X-ray structure of CA-II depicted as a ribbon model (Eriksson et al., 1988). In the representation of the active center hydrogens are not displayed.

The step between the nucleophilic attack and the replacement of bicarbonate was the topic of a controversial discussion in the past. While Lipscomb et al. (Liang & Lipscomb, 1987; Lipscomb, 1983) favored a proton shift between the two oxygens, the direct coordination by an oxygen and a rotation step was postulated by Lindskog et al. (Lindskog, 1984; 1997). In fact, depending on the complexity of the used model and the used quantumchemical methods respectively, each mechanism can be the favored one. Additionally, the exchange of bicarbonate for water can occur without direct coordination of an oxygen of CO₂ (see Figure 2). For further mechanistic aspects see (Mauksch et al., 2001).

As the electronic configuration of the zinc(II) cation has a closed 3d shell, which is very stable, zinc compounds contain this ion nearly without any exception. Additional results of this stable state are a fairly low redox activity of the zinc(II) ion, no stabilization of the ligand field and a wide range of coordination numbers going from two to eight.

2.2 Design of Biomimetic Chemical Models

The properties of a capable model describing the biomimetics of CA depend significantly on the aim of the application. Firstly, it is important, whether the model is used in an experimental environment or for theoretical purposes, and secondly, it depends on the goal of examination.

At least a model for simulating the catalytic core of CA needs an appropriate ligand sphere and an additional nucleophilic ligand, so one of the most simplified models is the [Zn(NH₃)₃OH]⁺ complex (see Figure 3).

2.2.1 Model for Computational Purposes

This model is only suitable for theoretical aspects, as it is not stable in solution and consists beside the zinc ion and the hydroxide ligand only of three ammonia ligands, which are a simplification of the histidine amino acid ligand sphere. Nevertheless, the so called ammonia model fulfills the needs of a theoretical model very well. One major aspect regarding calculations in chemistry is the number of atoms in the molecule. As the computation time correlates to the number of atoms like N³ for density functional theory (DFT) and N⁵ or N⁶ for Post-Hartree-Fock (ab initio) methods, it is important to keep the molecule as small as possible for efficient calculations.

The second major aspect of choosing a model for computational chemistry is its ability to describe the geometric and electronic properties of a molecule as exact as possible in comparison.
to reality. Describing the reaction in the real enzyme, it is better to have a bigger system, which also pays attention to parts of the peptidic backbone and its role in the mechanistic pathway. A good choice for this aim is the model of Bottoni et al. (Bottoni et al., 2004), which includes the mechanistically important amino acid residues Glu106 und Thr199 (see Figure 3). If it is the aim to apply the biomimetic principle to a model in synthesis, it is important to have a stable and simple molecule, what provides the basic features of the catalytic core. Depending on the reaction, the nucleophilic ligand can be varied, e.g. to a thiolate ligand.

As this chapter describes the transfer of the biomimetic principles from the enzyme CA to a model used in synthesis, the Bottoni model is not appropriate. Beneath the ammonia model one can imagine a more complex one containing imidazole rings instead of ammonia ligands. Although it resembles the ligand sphere in the catalytic core of CA better, it is not worth the calculational effort. Firstly, the imidazole rings in the enzyme have a fixed position, whereas they can turn in the model, and secondly, they do not have any significant influence on the electronic situation at the nucleophilic part of the model compared to an ammonia ligand sphere (see Table 1).

The natural charges of the ammonia model are also very similar to those calculated for the model bearing a [12]aneN3 ligand (see Table 1). Contrary to the model using imidazole ligands, the ammonia and the [12]aneN3 model possess amino functions directly connected to the metal center, so the ammonia ligand sphere would resemble the situation at the [12]aneN3 complex better than the imidazole model. Keeping all these arguments in mind the ammonia complex is the appropriate model for our purposes. It is small and therefore can be calculated very quickly, and it is accurate enough for transferring the biomimetic principle from the catalytic core to a certain complex in synthesis. The ammonia model has already proved
Table 1. Natural charge distribution and Zn-O bond lengths in several CA models (calculated at the B3LYP/6-311+G(d,p) level) in comparison to the native CA.

<table>
<thead>
<tr>
<th>model</th>
<th>Zn</th>
<th>O</th>
<th>bond length [Å]</th>
</tr>
</thead>
<tbody>
<tr>
<td>human CA-II</td>
<td>+1.66</td>
<td>-0.99</td>
<td>1.878</td>
</tr>
<tr>
<td>ammonia</td>
<td>+1.62</td>
<td>-1.32</td>
<td>1.849</td>
</tr>
<tr>
<td>imidazole</td>
<td>+1.65</td>
<td>-1.31</td>
<td>1.879</td>
</tr>
<tr>
<td>[12]aneN₃</td>
<td>+1.61</td>
<td>-1.31</td>
<td>1.851</td>
</tr>
<tr>
<td>Bottini</td>
<td>+1.63</td>
<td>-1.32</td>
<td>1.910</td>
</tr>
</tbody>
</table>

In general, complexes containing a macrocyclic ligand possess a higher complex constant and stability and structure. But it also has a significant effect on the catalytic effectiveness of the model, as it defines the geometry in the close vicinity of the nucleophilic ligand. Most of the synthesized ligands have either a tripodal or macrocyclic geometry (see Figure 3). To the variety of ligands used for modelling CA belong tris(2-benzimidazolylmethyl)amine (BMA) (Brandsch et al., 1997; Echizen et al., 2004; Nakata et al., 1997), different azamacrocycles, such as 1,5,9-triazacyclododecane ([12]aneN₃) (Kimura et al., 1990; Richman & Atkins, 1974), further tris(imidazolyl)carbinole (TIC) (Tang et al., 1978), tris(imidazolyl)phosphane (TIP) (Kunz et al., 2003), 1,3,5-triaminocyclohexane (tach) (Bowen et al., 1996; Cronin et al., 1996; Cronin & Walton, 2003; Urbach et al., 1968) and the representatives of hydrotris(pyrazolyl)borates (Tp) (Trofimenko, 1966; 1967; Yang et al., 1995), tris(pyrazolyl)methanes (Tpm) (Sánchez-Méndez et al., 2004; Zvargulis et al., 1995) or tris(pyrazolyl)methanesulfonates (Tpmns) (Kláui et al., 2000). In aqueous solution, these complexes exist in a pH-dependent equilibrium with their protonated forms, [Zn([12]aneN₃)OH]⁺ shows a pKₐ value of 8.1 (Riccardi & Cui, 2007; Zhang & van Eldik, 1995). In aqueous solution, these complexes exist in a pH-dependent equilibrium with their protonated forms, [Zn([12]aneN₃)OH]⁺ shows a pKₐ value of 8.1 (Riccardi & Cui, 2007; Zhang & van Eldik, 1995).
what is important to prevent the metal and the ligand from dissociating, and therefore the presented models are appropriate, as they reach $K_D$ values in the nanomolar range.

Another thoroughly investigated ligand family consists of tripodal trispyrazolyl ligands bearing either a carbon or a boron atom and trisimidazolyl ligand respectively bearing a carbon or a phosphor atom. These complexes also have very good complex constants and they allow creating different reaction pockets by varying the residues of the pyrazole and imidazole rings. The creation of such hydrophobic pockets avoids the multiple complexation of the ligand by the zinc ion. Contrary to the macrocyclic complexes those compounds can occur as neutral or cationic complexes depending on the element used at the tripodal position. The formation of cationic complexes is of importance for correctly modelling the catalytic centre of CA. The application of a charged ligand would necessarily result in a model system that does not reflect the electronic properties of the active site.

Summarizing this, both model concepts - azamacrocyclic and tripodal ligands - are appropriate to be used to apply the reaction principle of CA as chemical biomimetics to develop new syntheses. Although they have a lot in common, the differences between them define the cases they can be used in. As the azamacrocyclic complexes do not possess a hydrophobic pocket around the nucleophilic ligand, they provide a better accessibility of the catalytic center. This might be interesting when reacting bulky substrates. But the absence of this pocket gains a disadvantage as not all complexes are stable. In particular the hydrosulfide and alkyl thiolate complexes are not available. As there exist not only three- but also four-dentate azamacrocyclic ligands, the corresponding complexes can also be used to simulate the biomimetics of human CA-II. Zinc complexes of cyclic azaethers possess very high complex constants because of the macrocyclic effect, so they can be used under very harsh conditions without getting destroyed.

In comparison, tripodal ligands can not only simulate a hydrophobic pocket, but provide the possibility to measure the reaction rates depending on the size and composition of it. Depending on the element used at the tripodal position, they are neutral or cationic systems. Thus some tripodal ligands provide complexes, which are independent of dealing with a counterion.

3. **Details about the Quantumchemical Calculations**

In analogy to preceding investigations on the mode of action of CA, we used density functional theory (DFT) calculations to propose a mechanism. All calculations were realized using the Gaussian 03 (Frisch et al., 2004) program package, whereas Natural Bond Orbital (NBO) analyzes were performed using NBO 5.0 (Glendening et al., 2001). Stable structure geometries were characterized by a frequency calculation, in which energetic minima must not have an imaginary mode, whereas the presence of exactly one imaginary frequency characterizes a transition state. The visualization of this mode normally illustrates the reaction path, but in several cases an intrinsic reaction coordinate (IRC) calculation is appropriate to verify the correct products and educts corresponding to a transition state.

4. **Carbonyl Sulfide (COS), an Important Atmospheric Trace Gas**

For a better understanding, in this part of our chapter we additionally will give a few explanations of the biological backgrounds. Due to the enormous catalytic effect of CA, almost all living organisms use these enzymes for important CO$_2$ exchange processes with the atmosphere. However, it is less well known that CA is also involved in the fixation of the atmo-
Fig. 3. Selection of possible CA models as well as several ligands used to construct CA simulating complexes.
spheric trace gas carbonyl sulfide (COS) (Sandoval-Soto et al., 2005). As more recently and thoroughly investigated, COS is the most stable reduced sulfur compound in the atmosphere with a lifetime of 1-7 years. CA plays a significant role in the regulation of COS in the atmosphere and in the global distribution of sulfur (Schenk et al., 2005). Interestingly, it contributes indirectly to the global climate due to the fact that degradation of COS accounts for sulfate particles in the stratospheric aerosol layer which reflects solar radiation. Its main sources are the oceans where it is thought to be generated by photooxidation of sulfur-containing amino acids. Main sink for COS is the vegetation. The consumption of COS by many organisms, such as higher plants, algae, and lichens (Kuhn & Kesselmeier, 2000), was found to be CA-dependent. Furthermore, CA plays an important role in the interpretation of toxicity of COS. During COS consumption, H₂S is formed, which has been recognized almost thirty years ago as being the actual toxin responsible for the noxious effect of this heterocumulene (Chengelis & Neal, 1980). This has been confirmed by the finding that inhibition of CA lowers the mortality of insects upon exposition to COS, by lowering the rate of hydrolysis. In addition, CA-dependence of H₂S formation upon COS consumption has been proven for the example of algae. It can therefore be considered as being accepted that COS is a natural substrate of CA, which catalyzes its irreversible hydration according to equation 2.

\[
\text{COS} + \text{H}_2\text{O} \rightarrow \text{CO}_2 + \text{H}_2\text{S}.
\] (2)

The mechanism of this CA-catalyzed COS fixation according to Figure 4 has been investigated by some of us using the established computational ammonia model (Schenk et al., 2004). The reaction follows the same principle as the CO₂ reaction. The attack of the zinc-bound hydroxide occurs exclusively at the C=S bond and results in a four-centre transition structure COS-2(ts), whereby a zinc-bound thiocarbonate COS-3 is formed. Although this structure could not be observed in an experimental investigation until now, there is more than circumstantial evidence for its intermediate existence based upon our detailed calculations and related structures observed in experiments. By a water-assisted proton transfer, CO₂ is formed which is expelled and the zinc hydrosulfide complex COS-4 is obtained. This species is stabilized by a strong ZnS bond. As a consequence, we initially assumed that the enzyme must be deactivated by this reaction. However, this contradicts the generally accepted finding that in the course of COS fixation H₂S is formed (Blezinger et al., 2000). Furthermore, we detected an interesting side path in this reaction's cascade: A water molecule helps to regenerate the original zinc hydroxide complex from the hydrosulfide complex COS-4 and the formation of H₂S. Nevertheless, at the beginning of our investigations the mode of reactivation of the catalytic
domain of the hydrosulfide analogue of CA was an open question, which is of special interest since some organisms apparently utilize this $\text{H}_2\text{S}$ production process and some additional reactions steps as their only source of sulfur (Kuhn & Kesselmeier, 2000).

In case the organism is growing in sulfate-rich environment, it necessarily will not require the sulfide that stems from COS fixation. The sulfur will then be expelled in form of $\text{H}_2\text{S}$, since this is the main process plants utilize to free themselves from excess sulfur of any origin (Saito, 2004). However, plants which grow on sulfate deficient soils actually depend on atmospheric sources of sulfur. $\text{H}_2\text{S}$ and particularly $\text{SO}_2$ are commonly considered as being the most important atmospheric sulfur compounds taken up by plants (Bertini et al., 1985). Nonetheless, in some areas where the atmospheric supply of these compounds is too low to meet the requirement of plants, $\text{H}_2\text{S}$ formed by the COS metabolism becomes an important factor. We thus elucidated a possible mechanism for the final step of this process in order to achieve its complete understanding also at a molecular scale.

As explained above and depicted in the scheme in Figure 4, the active catalyst $Z$ reacts with COS and is transformed into the zinc hydrosulfide $\text{COS-4}$ (stabilized by 68 kJ/mol) (Schenk et al., 2004). Protonation resulting in the hydrogen sulfide complex $\text{COS-5}$ requires only 31 kJ/mol. It can thus be assumed that $\text{COS-4}$ should be protonated at least to a small extent. The replacement of $\text{H}_2\text{S}$ by a water molecule is now enforced by two effects: First, the aquo complex $\text{COS-6}$ is stabilized by 21 kJ/mol which is hardly surprising in view of the exceptionally weak coordination of $\text{H}_2\text{S}$ ligands to zinc centres. Second, the dissociation of hydrogen sulfide is practically irreversible due to the fact that its concentration in the surrounding medium is negligible compared to water. The irreversibility of this ligand exchange ensures that all zinc-bound sulfur is eventually expelled, since the hydrosulfide protonation equilibrium (see equation 3) provides a small but constant amount of the neutral $\text{H}_2\text{S}$ ligand which can easily dissociate.
As a consequence, the protonation of COS-4 is the rate-determining step of the desulfuration reaction. Of course, this process is likely to proceed quite slowly. However, a minute overall reaction rate should be sufficient: due to the extraordinarily small atmospheric COS concentration, sulfur analog CA is present in only very small amounts. A fast regeneration is therefore not required. The final deprotonation of the intermediate COS-6 results in the catalyst COS-1, which, according to our calculations, is endergonic by 58 kJ/mol. This deprotonation has already been extensively discussed in the context of CA-catalyzed CO$_2$ fixation since it is the initial step in generation of the active site, i.e. a representative of this interesting class of nucleophilic cations for which CA is the natural paragon.

4.1 In vitro experimental investigations

Some years ago, a model reaction has been performed by Vahrenkamp and some of us which exemplifies the process of COS fixation (Bräuer et al., 2002). The trispyrazolylborato zinc hydroxide complex Tp$_{\text{Ph,Me}}$ZnOH has been applied as the CA model (Notni, Schenk, Protoschill-Krebs, Kesselmeier & Anders, 2007). It has been reacted with COS, whereby CO$_2$ and the corresponding hydrosulfide complex Tp$_{\text{Ph,Me}}$ZnSH comparable to COS-4 were formed (see Figure 5). In contrast to the [Zn([12]aneN$_3$)$_2$]$^{2+}$ system, the [TpZn]$^{+}$ moiety is stable enough to bear a hydrosulfide ligand. The Tp$_{\text{Ph,Me}}$ZnOH complex is thus excellently suited for modelling the calculated regeneration mechanism, particularly with regard to the similar deprotonation enthalpies encountered for the related trispyrazolylborato and azamacrocyclic complexes. At neutral pH, the reaction cascade stops at the stage of the hydrosulfide (COS-4). However, after repeating this reaction, we recently could demonstrate that at pH values below 4 the hydrosulfide ligand is apparently protonated. Upon treatment of trispyrazolylborato complex of COS-4 with small portions of HClO$_4$, H$_2$S is formed, which can be removed from the solution by an argon stream. Since the trispyrazolylborato ligand is hydrolyzed at very low pH, the reaction mixture has to be kept at pH = 3 or above, which requires constant measurement and dropwise addition of HClO$_4$.

Further contributions to this subject have been made by Carrao et al. (Hammes et al., 2002) and Parkin et al. (Bergquist et al., 2003), both of whom isolated and characterized different examples of the [TpZnOH$_2$]$^{+}$ structure type with various anions. The latter authors have also shown that addition of base deprotonates the aquo ligand. From all these works with the trispyrazolylborato ligand, we derive the assumption of intermediate existence of the cation COS-6. We subsequently recovered the hydroxide complex COS-1 from the alkalized solution of COS-6 and identified it by NMR spectroscopy.

4.2 In vivo experimental investigations

The release of H$_2$S from cleavage of COS was investigated with the green alga Chlamydomonas reinhardti as this organism contains extracellular CA localized on its outer cell membranes faced to environmental solution. Hence, the released H$_2$S can be expected to escape the biological metabolism quite easily. As shown in Figure 6, the release of H$_2$S is closely correlated to the consumption of COS but a substantial part of the expected H$_2$S is always missing. We assume either a metabolic consumption of H$_2$S and/or a loss of H$_2$S due to the technical needs using a headspace technique. The latter instance is most probable, as it is known that application of this technique may involve a loss of 30 to 40% of the H$_2$S formed. A similar release of H$_2$S upon COS consumption could be reported for marine algae, although

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4.1 In vitro experimental investigations

4.2 In vivo experimental investigations

Fig. 6. COS consumption and H$_2$S production in correlation to increasing COS concentrations by the green alga *Chlamydomonas reinhardtii* grown under 0.03 % CO$_2$. Amount of algae per measurement accounts for 1.26 µg Chl. ml$^{-1}$. Measurements were performed in the light with 5 W/m in 25 mm barbital buffer at pH = 7.5. COS and H$_2$S in aqueous solutions were determined by a dynamic headspace technique coupled to GC-FPD analysis.

on significant lower activity levels which can be discussed as being caused by missing extra-cellular CA.

4.3 Discussion of the DFT calculations: Experiment meets theory

A DFT-calculation which is related to those presented in this work has been reported earlier by Parkin and coworkers (Bergquist et al., 2003). They investigated the deprotonation of a complex of the [TpZnOH$_2$]$^+$ type according to equation 4.

$$[\text{TpZnOH}_2]^+ \rightleftharpoons [\text{TpZnOH}] + \text{H}^+$$

For this reaction, they calculated a free energy of 1172 kJ/mol. This is substantially different to the deprotonation energy value we obtained for the zinc-bound water of the azamacrocyclic compound COS-5 (58 kJ/mol). Although we used another model system, the energies should at least be of the same order of magnitude. Presumably, the difference results from the fact that in equation 4, the free energy of the proton has been set to zero by these authors. However, for calculations including solvent effects, the solvation free energy of the proton should be taken into account to obtain realistic deprotonation energies. For the total free energy of the proton in aquatic solution, we used a value of 1125 kJ/mol given in the literature. This value is the sum of the absolute free energy of the proton in the gas phase (26 kJ/mol) and its absolute solvation free energy (1099 kJ/mol). When this free energy value is added to the free energy of deprotonation reported by Parkin *et al.*, a corrected value of 47 kJ/mol is obtained. This is in very good agreement with our result. We therefore conclude that the selection of two
apparently completely different model systems, one with an azamacrocyclic and the other with a pyrazolyl type ligand, is not a problem. Apparently, the deprotonation energies of the zinc-bound water are very similar. Thus we assume that our experiment using the TpZnSH complex is suited to corroborate and illustrate the calculated mechanism. Although a true catalysis is not observed, the substrate COS can be transformed into CO$_2$ and H$_2$S in the presence of Tp$^{Ph,Me}$ZnOH just by altering the pH of the solution.

According to our calculations, the protonation free energies of the zinc-bound hydroxide and hydrosulfide differ by ca. 84 kJ/mol. This is in good agreement with our experimental observation that a fast desulfuration occurs only at pH values at which a zinc-bound water is not deprotonated. Nevertheless, a well-balanced pH at the active site of natural CA could allow both a predominantly deprotonated zinc-bound water ligand and small but sufficient protonation of the zinc-bound hydrosulfide. As we already pointed out above, we hold the view that a small amount of protonated hydrosulfide ligand at the zinc ion is sufficient for complete desulfuration of CA due to the fact that the dissociation of H$_2$S is practically irreversible.

In our opinion, the calculated mechanism is thus very likely to occur the way it is depicted in Figure 5.

The role of other amino acid residues in the catalytic mechanism has been addressed in studies by Bottoni (Bottoni et al., 2004) and Liedl (Tautermann et al., 2003). They have demonstrated that some of the residues, especially Glu106 and Thr199, are directly involved in some steps of the CO$_2$ fixation. It has also been commented upon that a histidine residue in the enzyme cavity near the active site (so-called proton shuttle) influences the $pK_a$ of the zinc-bound water. The residue which is located in a distance of approx. 7 Å from the zinc centre can be present in both protonated or deprotonated state. For both cases, different $pK_a$ values for the zinc-bound water have been measured (Bertini et al., 1985). It is very probable that its protonation state will also affect a zinc-bound hydrosulfide ligand. However, the conclusions drawn in all these studies did not introduce any change in the overall qualitative picture obtained with simpler models that neglect those amino acids. We hold the view that in order to exactly calculate such effects, an expanded model system taking into account the additional residues and a study of molecular dynamics would be required. This would currently exceed by far the computational resources available to us. In addition, we do not believe that such calculation would substantially alter the proposed mechanism. Our aim was to deliver the proof of principle for the hypothesis that hydrosulfide substitution of CA does not entail inhibition of the enzyme, and nothing but a water molecule is required for reactivation and formation of H$_2$S. We are sure that this proposal is sufficiently supported by our model calculations and experiment.

From the in vivo experiments, it is obvious that there is a correlation between COS consumption and H$_2$S release. As stated above, the missing amount of H$_2$S flow is not a problem since systematic errors in experiment and partial H$_2$S metabolism have been shown to be possible reasons for this finding. However, the most important observation is that there is apparently no deactivation of CA by COS: With increasing COS concentration, the plot of the H$_2$S release rates shows no signs of any saturation effects, i.e. non-proportionality to the COS consumption plot. This fact strongly corroborates the overall statement of this study.

5. Application of the Enzymatic Reaction Principle to further Examples of Isoelectronic Molecules

As seen in the sections above, the reaction principle of CA is not restricted to the molecule CO$_2$ but has been applied to COS by nature itself. So it is anticipated, that further isoelectronic molecules like allenes (R$_2$CCCR$_2$), isothiocyanates (R-NCS), carbodiimides (R-NCN-R), and
5. Application of the Enzymatic Reaction Principle to further Examples of Isolelectronic Molecules

Fig. 7. Catalytical hydration of CO$_2$ as well as the homologous biomimetic addition reaction to heterocumulenes. X = Cr$_2$, NR, O, S; Y = H, alkyl, aryl; M = Zn$^{2+}$, Co$^{2+}$; L = ligand

Fig. 8. Uncatalyzed reaction with water across a concerted four-membered cyclic transition state. X = CH$_2$, NH, O, S

other heterocumulenes should react resembling the mode of action of CA (see Figure 7). In principle, the structure of the adding compound is not restricted to water, as a lot of polar HX compounds, such as alcohols or H$_2$S and mercaptanes respectively, provide comparable properties. Hence these heterocumulenic structures are very similar, the addition of a HX compound to a heterocumulene catalyzed by a CA model can be written as depicted in Figure 7. In the next sections the reactions of two representatives will be presented.

5.1 Validation of the Catalytic Effect

A very important value for estimating the catalytic effect is the activation barrier of the rate determining step in the uncatalyzed reactions. Accordingly to the catalyzed reactions, the uncatalyzed reactions do not differ significantly between various heterocumulenes (see Figure 8). After formation of an encounter complex (EC) between water and the double bond system, a concerted transition state (TS), which normally is the rate determining step of the reaction, has to be surmounted to get to the first intermediates. In some cases, theses intermediates are the final products, in other cases further transition states with minor activation barriers follow.

Depending on the used heterocumulene, the Gibb’s free energies $\Delta G$ of the encounter complexes vary between 0 and 20 kJ/mol in comparison to the free non-interacting educts. However, these values might be slightly erroneous, as some DFT methods do not calculate weak intermolecular forces properly. Subsequently, the reaction coordinate leads to a four-membered
cyclic TS (see Figure 9), whose strained structure explains the high activation barrier of the reaction. Typical energy values for these structures can be found in Table 2. The resulting products are also shown in Figure 9. The addition reactions of allene to the products 6 and 7 are both exergonic (see Table 2) and propene-2-ol 6 tautomerizes under standard conditions to the more stable acetone. In case of isothiocyanates, the intermediates are still not exergonic, but after surmounting some minor transition states, several conformers of the exergonic carbamic thio acid can be reached (Eger et al., 2009).

To summarize this, the activation barriers of the uncatalyzed reactions of allenes and isothiocyanates are very high, as they are four-membered cyclic transition states and therefore possess Gibb’s free energies between 200 and 300 kJ/mol. Keeping the estimated activation barriers of carbon dioxide and carbonyl sulfide in mind, it should be possible to see a significant catalytic effect in the reactions of allenes and isothiocyanates.

5.2 The Selectivity Problem
Contrary to the case of carbon dioxide, allenes or isothiocyanates as educts for the nucleophilic attack of hydroxide or water provide a more complex scenario. As a heterocumulene, isothiocyanate possess nitrogen and oxygen on the outer positions of the cumulenic system and additionally has an imine group, which reduces the symmetry of the molecule and introduces more reaction possibilities (see Figure 9). Looking at allene, all known problems regarding alkenes and alkynes come to mind, thus chemo- (single or double addition), regio- (Markovnikov- or anti-Markovnikov products) and stereoselectivity (cis- or trans-addition products on stereotopic sides) play a role. For substituted allenes there exists a positional selectivity (Hashmi, 2000) as the attack can take place at two different positions of the allene molecule. Therefore, additions at one of the orthogonal double bonds will lead to constitutional isomers in the case of substituted allenes and as a consequence, this inclusion of regioselectivity doubles the number of isomers.

6. Isothiocyanates (R-NCS), the Link to Synthesis
As described previously, the reaction of isothiocyanates with water and other H-X compounds, i.e. alcohols and amines, is kinetically hindered. Water and alcohols do not react
under standard conditions, even when they are heated, it takes very long to see some product (Browne & Dyson, 1931; Hagemann, 1983; Rao & Venkataraghavan, 1962; Walter & Bode, 1967). This is only true as long as there is no acid or base present, which would open up other reaction possibilities. If the catalysis by a CA model is efficient, it would be the method of choice to hydrolyze or alcoholyze isothiocyanate systems under neutral conditions. This might be interesting for the synthesis of complex and acid or base sensitive molecules.

In comparison to carbon dioxide and carbonyl sulfide, isothiocyanates bear a residue on one of the outstanding hetero atoms. As this is an imine function, it increases the degree of freedom and therefore produces more possible pathways.

<table>
<thead>
<tr>
<th>educt</th>
<th>MeNCS 1</th>
<th>allene 2</th>
</tr>
</thead>
<tbody>
<tr>
<td>TS Superscript a</td>
<td>uN-2 Superscript h</td>
<td>uN-3 Superscript h</td>
</tr>
<tr>
<td>5.2 The Selectivity Problem</td>
<td></td>
<td></td>
</tr>
<tr>
<td>6. Isothiocyanates (R-NCS), the Link to Synthesis</td>
<td></td>
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</tr>
</tbody>
</table>

Table 2. Energies and geometries of the uncatalyzed reaction of methylisothiocyanate and allene with water.

Table 3. Natural Charges $\delta^{NC}$ for CO$_2$, COS, MeNCS, and allene.

<table>
<thead>
<tr>
<th>X, Y</th>
<th>X</th>
<th>C</th>
<th>Y</th>
</tr>
</thead>
<tbody>
<tr>
<td>carbon dioxide</td>
<td>X, Y = O</td>
<td>-0.56</td>
<td>1.13</td>
</tr>
<tr>
<td>carbon oxid sulfid</td>
<td>X = O, Y = S</td>
<td>-0.48</td>
<td>0.50</td>
</tr>
<tr>
<td>methylisothiocyanate</td>
<td>X = S, Y = N</td>
<td>-0.10</td>
<td>0.30</td>
</tr>
<tr>
<td>allene</td>
<td>X, Y = C</td>
<td>-0.51</td>
<td>0.07</td>
</tr>
</tbody>
</table>
6.1 Calculated Mechanistic Pathway

The calculations show only one encounter complex NCS-1, in which the isothiocyanate coordinates via the sulfur atom to two ammonia ligands using hydrogen bridging bonds. Coming from this encounter complex, three different transition states could take place. Whereas in NCS-1(ts) and NCS-2(ts) the C=S double bond adds to the Zn-O bond, the C=N double bond does this in the case of NCS-3(ts) (see Figure 10). These transition states resemble the rate determining steps in the reactions of carbon dioxide and carbonyl sulfide and also are the highest activation barriers in the pathway of isothiocyanate. Contrary to the situation in case of COS, which also possesses an unsymmetric cumulenic system, the energies of this transition states differ not significantly, so a prediction of selectivity depends not only on the energies of the rate determining steps, but also on the further reaction paths and thermodynamic control. Comparing the free enthalpies of the three transition states and the energies of the following reaction paths, it becomes obvious, that the attack on the C=S double bond is thermodynamically and kinetically slightly favored. Contrary to the fact, that the existence of the imine makes the situation at the rate determining step more complex, it simplifies it at the point, where the Lindskog and Lipscomb transition states enter the scenery right after the attack of the C=S double bond. As the disturbed symmetry of isothiocyanate opens up about eight possible pathways, the kinetically and thermodynamically most favorable will be discussed shortly here (see Figure 11).

Structure NCS-2(ts) is the rate determining step, as no other transition state builds up a higher activation barrier. $\Delta G = 82$ kJ/mol relative to the separated educts (ammonia model and methyl isothiocyanate), is not as good as the corresponding values estimated for carbon dioxide and carbonyl sulfide, but it is easily surmountable in a normal experimental environment. The catalytic effect becomes very clear, when comparing the activation barriers of the rate determining steps in the catalyzed and uncatalyzed reaction, as the gap between these values is about $\Delta \Delta G = 76$ kJ/mol. This is a significant decrease in energy. The reaction path proceeds further via a Lindskog reaction mechanism (NCS-4(ts)), which is rather lower than the corresponding Lipscomb proton shift. Nevertheless, the pathway surmounting NCS-4(ts) is the thermodynamically and kinetically favored one.

The found selectivity is only true for the reaction with methyl isothiocyanate, as calculation with several residues showed different results. In general, the inductive effect of the residue of the isothiocyanate changes the selectivity. The greater the ability of the residue to pull electrons out of the cumulenic system, the more an attack of the C=N double bond is preferred. This is mainly a result of the electronic structure in the cumulenic system. If the residue on the nitrogen atom pulls electron density out of the double bond system, it is mainly taken from...
Fig. 11. Pathway of the catalyzed reaction with methyl isothiocyanate. Level of theory is B3LYP/6-311+G(d,p), given values are Gibb’s free enthalpies in kJ/mol.

the C=S double bond. Thus NBO calculations show, that in such cases the C=N double bond has a strong triple, and the C=S double bond a strong single bond character (Eger et al., 2009). Further calculations with complexes not bearing a hydroxide ion but an hydrosulfide and an thiolate ion respectively, showed, that the biomimetics of CA are not only limited to hydroxide bearing complexes and thus the addition of water to cumulenic system. Furthermore a lot of different combinations of different nucleophiles and cumulenes are possible.

6.2 Experimental Results
As the reaction with a thiolate complex reduces the number of possible pathways significantly and those complexes recently proved their ability simulating CA biomimetic insertion reactions (e.g. with carbon disulfide) (Notni et al., 2006), this seems to be a good model complex to see, if isothiocyanate inserts even similar. Thiolate complexes bearing a four-dentate [12]aneN₄ ligand are known to work faster than the corresponding three-dentate complexed compounds (Notni, Günther & Anders, 2007).
Fig. 12. Insertion possibilities of isothiocyanate to a zinc thiolate complex.

The reaction shown in Figure 12 was carried out in dimethyl sulfoxide under standard conditions at room temperature. The insertion could be proved using GC/MS and Raman spectroscopy. For different isothiocyanates different reaction rates could be determined, as mostly isothiocyanates with an electron withdrawing residue as phenyl or p-nitro phenyl were able to insert easily at room temperature. Depending on the purpose of the reaction those activated cumulenes can react further with an HX compound, e.g. an alcohol or mercaptan.

7. Allene

Allene is the simplest hydrocarbon with cumulated double bonds. Since van’t Hoff has predicted the correct structures of allene and higher cumulenes, chemists are fascinated by the extraordinary properties like axial chirality of the elongated tetrahedron, if two different substituents at every terminal carbon exist. Allene with its isomer methyl acetylene accrues in large amounts in the C3-cut of the naphtha distillation. Currently both compounds are only hydrated to propene and propane respectively or flared off. Therefore the activation of allene has additionally to the biomimetic a strong economical aspect. Allene could be estimated as the parent compound for heterocumulenes with two cumulated double bonds. By the formal exchange of one or both terminal carbon atoms a vast number of heterocumulenes are available.

The first investigation of a possible biomimetic activation of allene with zinc catalysts was undertaken by Breuer et al. (1999). They found catalytical activity of zinc silicates with zinc acetate in methanol to give 2-methoxypropene and 2,2-dimethoxypropene in 85% yield.

7.1 Calculated Mechanistic Pathway

The presentation of the whole calculated reaction mechanism of the addition of water to allene goes beyond the scope of this chapter due to the immense number of reaction steps (see Jahn et al., 2008 for further reading). Therefore the description of mechanistical pathways is confined to the variants of the initial nucleophilic attack, which lead to mechanistical important intermediates. The results show that the initial attack is the rate determining step for the whole catalytic cycle.

The zinc catalyzed addition starts with an encounter complex A-1 between the zinc hydroxide complex and allene. This structure is the starting point for the different reaction variants, comparable to the uncatalyzed reaction described in section 5.1. Corresponding to the regio selectivity problem the attack to allene can take place at either the central or the terminal carbon atom (see Figure 13). The attack of the hydroxide on the terminal carbons leads to a concerted four-membered cyclic transition state A-2(ts) with an activation barrier of $\Delta G = 139$ kJ/mol.
Fig. 13. Calculated mechanism of the initial, rate determining steps of the activation of allene. \( \Delta G \) in kJ/mol. Level of theory is mPW1k/aug-CC-pVDZ.

This structure relaxes to the \( C_{2v} \)-symmetric, slightly exergonic intermediate A-3, in which the carbon backbone, the hydroxyl group, the metal ion and one nitrogen of the ligand span the symmetry plane. The hydroxyl group is placed between and in front of the ligands. There is only one possibility to close the catalytic cycle starting from intermediate A-3. This mechanism is an attack of a water molecule, which leads to a cleavage of the Zn-C bond. One water proton is shifted to the central carbon atom to give allylalcohol 7 and the remaining hydroxide regenerates the catalytic model.
Fig. 14. Mechanism of the racemization of A-9 via rotTS-I and rotTS-II. ∆G in kJ/mol. Level of theory is mPW1k/aug-CC-pVDZ.

Alternatively, the initial nucleophilic attack on the CA model complex can take place at the central carbon atom. Depending on the kind of the model complex, two different mechanisms of the initial reaction step can be found. This reaction path can either proceed via a stepwise or a concerted reaction mechanism, whereas the stepwise mechanism can only be found using the azamacrocyclic models. Contrary, the concerted one can be found in all cases. This shows the restrictions of the ammonia model.

Structure A-5(ts) is the first transition state of the stepwise variant. The activation barrier is ∆∆G = 18 kJ/mol) higher than for the concerted TS A-8(ts), which is interesting, as this TS has no sterical restrictions. As its carbon backbone stands approximately perpendicular to the Zn-O bond, structure A-5(ts) differs fundamentally in its geometry compared to the cyclic concerted TSs. The reaction coordinate is only defined by the difference of the distance between oxygen and the central carbon atom of allene. The TS relaxes to the intermediate A-6. With ∆G = 113 kJ/mol relative to the Gibb’s free energy of the separated reactants allene and zinc hydroxide complex, this intermediate is only poorly stable. Intermediate A-6 rearranges by a cascade of proton transfer steps between the substrate and the ligand to the intermediate A-7, which is one of the most stable structures in the calculated reaction path variants (∆G = -120 kJ/mol). Subsequently, the direct formation of acetone is facilitated by a proton shift from an attacking water molecule to the free methylidene group.

The third and most probable transition state between allene and the CA model complex is the concerted four-membered cyclic TS A-8(ts). Comparably to methylisothiocyanate, A-8(ts) resembles the rate determining step in the reactions of carbon dioxide and carbonyl sulfide. A-8(ts) possesses the lowest activation barrier of all three initial TSs (∆G = 124 kJ/mol). It finally relaxes into the intermediate A-9.

Contrary to all other intermediates of different heterocumulenes at comparable points of the reaction coordinate, structure A-9 has an outmost geometry. Whereas in all geometries of intermediates connected to the zinc ion by a heteroatom the former cumulated system and the metal ion are located in a plane, intermediate A-9 has a carbon atom connected to the zinc instead, which forces the plane spanned by the carbon backbone of the allene to stand perpendicular to the Zn-C bond and parallel to the plane spanned by the ligand respectively. A reason for that is the partial double bond character of the bonding between the central and the zinc-bound carbon atoms. As a consequence, A-9 is a chiral structure without an asymmetric center and therefore an example of planar chirality. However, the activation barrier of
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7.2 Experimental Results

The reaction of allene and [Zn([12]aneN₃)OH]ClO₄ as the CA model complex was investigated under heterogeneous conditions. Due to the gaseous aggregation state of the unsubstituted allene, a pressure cell was used. The analysis was done with Raman spectroscopic methods.

8. Conclusion

In summary, we have shown that the transformation of COS by carbonic anhydrase, which finally yields H₂S and CO₂, requires no further reactant than water in order to regenerate the most important zinc-bound hydroxide [L₂ZnOH]⁺ from the hydrosulfide complex. We conclude that CA is perfectly equipped by nature to perform the task of transformation of COS into H₂S. Furthermore, we regard this special function of CA to be perfectly linked to the plant sulfur metabolism. Therefore, this regeneration mechanism can be regarded as the missing link between CA-catalyzed COS fixation and plant sulfur metabolism, an aspect of fundamental significance for the understanding of some very important biological processes. Nature has chosen an elegant and efficient system for the hydration of CO₂ and COS, the [L₂Zn(OH)]⁺/CO₂ or COS/H₂O group of reactants. The catalyst is able to transform both cumulenes, though the relative energies of the corresponding reactions steps differ in some details significantly. Further we have shown that it is possible to apply biomimetic principles of high optimized, biochemical processes to the laboratory as well as industrially usable syntheses. The reaction principle of carbonic anhydrase is applicable to other isoelectronic molecules than CO₂, which are normally not processed by the enzyme. These biomimetic investigations about the enzyme carbonic anhydrase could serve as a paragon for the further research on biochemical model systems.
Acknowledgement

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9. References


URL: http://www.gaussian.com


URL: http://www.chem.wisc.edu/nbo5


Nature’s evolution has led to the introduction of highly efficient biological mechanisms. Imitating these mechanisms offers an enormous potential for the improvement of our day to day life. Ideally, by bio-inspiration we can get a better view of nature’s capability while studying its models and adapting it for our benefit. This book takes us into the interesting world of biomimetics and describes various arenas where the technology is applied. The 25 chapters covered in this book disclose recent advances and new ideas in promoting the mechanism and applications of biomimetics.