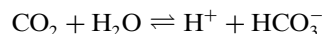


Carbonic anhydrase under pressure

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Carbonic anhydrase is among the most thoroughly investigated enzymes. It catalyzes the conversion between carbon dioxide and bicarbonate



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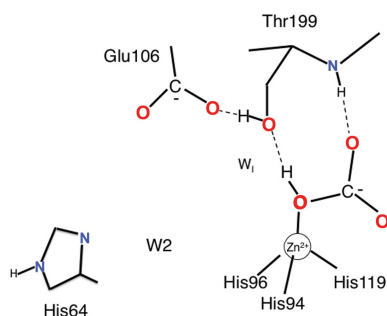
Early on it was realised that the rate of the spontaneous reaction was far too low to function physiologically. Thus, there must be an enzyme and it was identified by Meldrum & Roughton (1932). Now six distinct families of the enzyme are known (α , β , γ , δ , ζ and η). Furthermore, the human α -form is divided into 14 different isoenzymes with the same basic fold but with different cellular locations and properties. Human CA II (hCAII) is expressed in many cell types and is involved in many physiological processes. Its catalytic rate is limited by the rate of diffusion of substrates and products (Frost & McKenna, 2013). Evidently, the substrates or the products cannot bind strongly and the conformational changes need to be minimal.

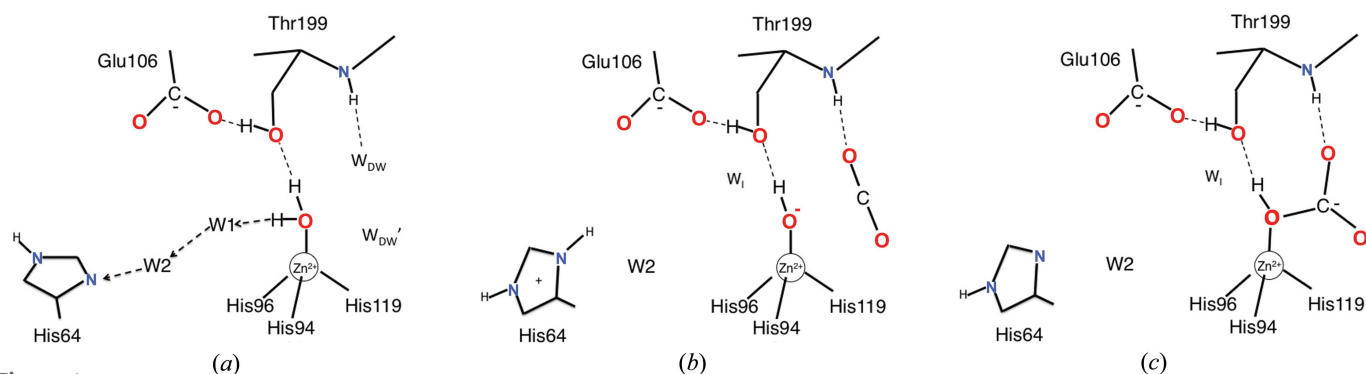
The first structure of hCAII provided the protein fold and the active-site zinc ion in a 15 Å deep cleft. A zinc-bound water (W_{Zn}) and a number of additional active-site water molecules were identified (Liljas *et al.*, 1972). Numerous studies of mutants and bound inhibitors have provided further insights (Liljas *et al.*, 1994). In the enzyme catalysis the W_{Zn} is the hydration water. It is activated by releasing a proton through the water structure to a histidine in the active site, His64, from where it is released into bulk solvent (Nair & Christianson, 1991). The $\text{p}K_a$ of W_{Zn} is around 7, as it is for His64 (Domsic *et al.*, 2010). W_{Zn} is hydrogen bonded to O γ 1 of Thr199, which is further hydrogen bonded to the negatively charged Glu106. This forces the zinc ligand to provide a proton for the hydrogen bond to Thr199. The CO_2 is bound in the deep hydrophobic part of the active site next to W_{Zn} (Kim *et al.*, 2005; Domsic *et al.*, 2008). The hydration of CO_2 or dehydration of HCO_3^- is efficiently done without significant structural changes to the protein. The entry or dissipation of CO_2 probably occurs along the hydrophobic side of the active site while HCO_3^- enters or leaves along the hydrophilic side. However, the activities in the active site during catalysis primarily need the dynamics of water molecules.

An article in the current issue of IUCrJ describes crystallographic analyses of hCAII at ultrahigh resolution where the crystals were exposed to CO_2 under 7 and 2.5 atm pressure (Kim *et al.*, 2018). These studies extend previous studies where the crystals were exposed to 15 atm pressure of CO_2 and frozen to liquid nitrogen temperature after various lengths of time (Kim *et al.*, 2016). The previous investigations provided the location of a fully bound CO_2 molecule at the expected position at the bottom of the active site. The present article illuminates the previous courageous high-pressure experiments with studies at intermediate pressure. The observations lead to a range of intermediate structures with lower occupancy of substrate and identify what rearrangements take place at partial occupancy of substrate. A unique feature of the article is the focus on rearrangements of the water structure during catalysis. This is something of importance for most enzymes.

An obvious rearrangement is the replacement of CO_2 by two water molecules, called 'deep water' (W_{DW} and $W_{\text{DW}'}$). They substitute for the two O atoms of CO_2 . The flexibility of the network of water molecules in the active site is described in detail. The waters adjust in several steps to the changes that occur in the active site. Several water molecules also appear to have neighbouring alternate positions.

The most dramatic conformational change is the flipping of His64 from an inward-facing orientation to an orientation towards bulk solvent to release a proton from the zinc water. Having received the proton, the positively charged imidazole is probably repelled




Figure 1

A likely catalytic mechanism for human carbonic anhydrase II. In the first step (a) the zinc water (W_{Zn}) releases a proton to His64 via W1 and W2. When carbon dioxide binds (b) it releases the two deep-water molecules. W1 disappears and is replaced by W_1 . His64 adopts the outward orientation to release the abstracted proton to bulk solvent. Then the W_{Zn} makes a nucleophilic attack on the carbon of the carbon dioxide to form bicarbonate (c). Three water molecules rapidly replace the bicarbonate and His64 returns to its inward orientation (a). Several additional water molecules are identified and participate in the dynamic water structure of the active site.

by the positive charge of the zinc ion to adopt the outward conformation to release the proton to bulk water. Subsequently it flips back to the original position.

The distance from W_{Zn} to His64 is too long for a direct proton transfer. Rather, the proton is shuttled from W_{Zn} to His64 via two bridging water molecules (W1 and W2; Fig. 1a). The series of intermediate structures studied show that the water molecule next to W_{Zn} (W1) is absent when His64 is in the outward position (Fig. 1b). In this state there can be no proton transfer, since His 64 is already protonated. Instead of W1 an intermediate water molecule, W_1 , appears. It also has an alternate site W_1' .

A new cycle of catalysis begins with a new W_{Zn} that replaces the bicarbonate. The site can be filled from any of several neighbour water molecules. W_{Zn} then gets deprotonated and CO₂ binds by displacing the two deep-water molecules (W_{DW} and W_{DW}'). W1 and W2 then occupy their normal positions where W2, closest to His64, has two different positions, W2 and W2'. The second position, W2', is related to the flipping of the histidine to the outward conformation, which allows more space.

His64 then accepts the proton from W_{Zn} through W1 and W2. At this stage His64 adopts its outward facing conformation to release its proton, which leads to changes in the water structure. W_{Zn} is now able to make a nucleophilic attack on the carbon of the carbon dioxide to yield bicarbonate (Fig. 1c). The binding of the bicarbonate is not optimal since it is liganded to the zinc ion by its protonated oxygen but not a negatively charged one. The protonated oxygen is forced to remain at the zinc due to its hydrogen bond to the obligate

hydrogen-bond acceptor O γ 1 of Thr199. This makes the binding weak, and three water molecules then readily displace the bicarbonate. This completes the catalytic process.

The high rate of the enzyme is evidently due to the weak binding of the substrates, as well as the firm structure of its active site, the only moving part being His64. Easily moving water molecules during catalysis are also a requirement for the high rate.

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