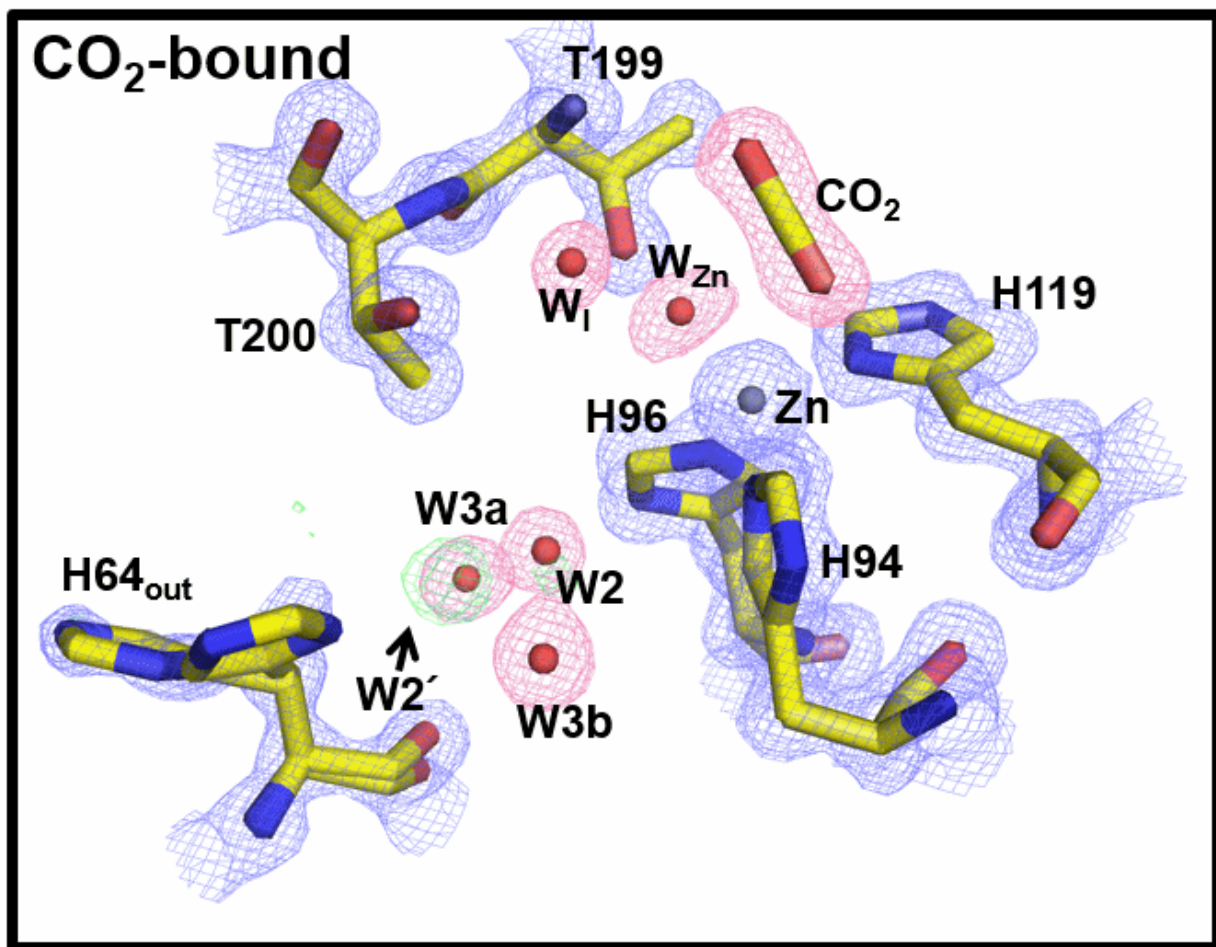


Study captures ultrafast motion of proteins

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Schematic image, showing protein movement during CO₂ release in carbonic anhydrase ii crystals. Credit: Ulsan National Institute of Science and Technology

A new study by an international team of researchers, affiliated with

Ulsan National Institute of Science and Technology (UNIST) has announced that they have succeeded for the first time in observing the structural changes in carbonic anhydrase.

The breakthrough comes from a research, conducted by Professor Chae Un Kim (School of Natural Science) of UNIST in collaboration with researchers from Soongsil University, Cornell University, and University of Florida.

Carbonic anhydrase, which is found within [red blood cells](#), is a crucial [enzyme](#) that stabilizes carbon dioxide (CO₂) concentrations. This enzyme catalyzes a reaction converting CO₂ and water into [carbonic acid](#), which dissociates into protons and [bicarbonate ions](#).

Moreover, it is also known that carbonic anhydrase is able to catalyze at a rate of 10⁶ reactions per second. In the absence of this catalyst, the conversion from CO₂ to bicarbonate, and vice versa, would be extremely slow and difficult.

One of the important functions of the enzyme in humans is to adjust the acidity of the chemical environment to prevent damage to the body, as well as to help transport carbon dioxide out from tissue cells to the lungs. Although [carbonic anhydrase](#) performs a lot of beneficial functions, defects in the enzyme are responsible for developing diseases, such as glaucoma, acidemia, or osteopetrosis.

Prof. Kim, the lead researcher of the study states, "The reaction rate of carbonic anhydrase is one of the fastest of all enzymes." He continues, "Due to the rapid movement of proteins, direct observation for such movement has been extremely difficult to obtain, protein scientists say."

In this study, Prof. Kim's team used their own method of "High-pressure Cryocooling" and "X-ray Crystallography" to capture the gaseous carbon

dioxide in crystals of carbonic anhydrase and follow the sequential structure changes as the [carbon dioxide](#) is released.

The results of the study will not only greatly contribute to the future biomedical research and new drug development, but will also help make carbon capture more economic.

According to Prof. Kim of UNIST, "This study also shows technical methods that may be applicable to other enzymes that bind and react to low-molecular weight substrates, such as CO₂ and NO₂ ."

More information: Chae Un Kim et al, Tracking solvent and protein movement during CO₂ release in carbonic anhydrase II crystals, *Proceedings of the National Academy of Sciences* (2016). [DOI: 10.1073/pnas.1520786113](#)

Provided by Ulsan National Institute of Science and Technology

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