## CO<sub>2</sub> capture by membranes and the perspectives

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One of the feasible methods to mitigate  $CO_2$  emission must be  $CO_2$  Capture & Storage (CCS), and effective  $CO_2$  capture technologies have been investigated, which account for a majority of the CCS cost. Membrane separation would be suitable for  $CO_2$  capture in terms of energy consumption, footprint and cost in comparison to current  $CO_2$  capture technology, liquid amine scrubbing.  $CO_2$ separation over  $H_2$  by polymeric membranes has been studied in this research group for pre-combustion  $CO_2$  capture at an integrated gasification combined cycle plant. The mechanism of preferential  $CO_2$  permeation was elucidated as shown in Fig. 1. Under humidified conditions,  $CO_2$ turns to bicarbonate ion, which is the major migrating species through the membrane. When the amines have hydroxyl groups, a seven-membered ring is formed to facilitate bicarbonate ion production upon hydrolysis. The resulting polymeric membranes show high  $CO_2$  permeability even under pressurized conditions.

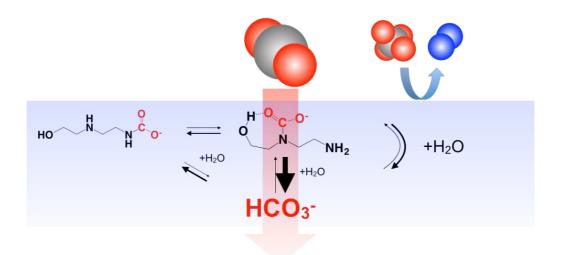


Fig. 1. Schematic drawing of CO<sub>2</sub> permeation mechanism through 2-(2-aminoethylamino)ethanol-containing polymeric membranes under humidified conditions.

For pilot-scale demonstration of the  $CO_2$  separation membranes developed, membrane modules should be prepared by the "*in-situ* modification (IM)" method. A thin  $CO_2$  separation (or active) layer is formed on inner surface of hollow fibers by passing through the membrane material solutions. The IM method is scalable, and the resulting hollow-fiber membrane modules also display excellent  $CO_2$  separation performance over  $H_2$ ,  $N_2$ , and  $CH_4$ .