## Molecular Engineering of Polymeric Membrane Materials for H<sub>2</sub> Purification and CO<sub>2</sub> Capture

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Growing evidence indicates that the  $CO_2$  emissions from the combustion of fossil fuels are contributing to global climate change. One approach to controlling  $CO_2$  emissions to the atmosphere is carbon capture, utilization and sequestration (CCUS) from large point sources, such as  $H_2$  and power plants. Pre-combustion capture from syngas has been considered as an economically viable route. In this scheme, gasification of fossil fuels produces syngas, which is further converted to  $CO_2$  and  $H_2$ . The  $CO_2$  must be removed and captured prior to  $H_2$  utilization for refinery hydrogenation reactions or combustion in the power plant turbines.

Membrane technology is an attractive approach to  $H_2$  purification and  $CO_2$  capture because of inherent advantages such as high energy efficiency. This presentation will systematically examine the need of membrane technology as a low-cost and energy-efficient separation technology enabling the CCUS via pre-combustion route. I will discuss how we rationally design polymeric membrane materials to achieve the combination of high  $CO_2$  permeability and high  $CO_2/H_2$  selectivity. More specifically, this talk will discuss the molecular engineering of poly(ethylene oxide) (PEO) containing polymers for improvement in mixed-gas  $CO_2/H_2$  separation performance. Interestingly, these materials exhibit unconventional increase in mixed-gas  $CO_2/H_2$  selectivity as  $CO_2$  feed partial pressure increases. The structure and property correlation in these PEO containing polymers will also be interpreted using a modified free volume model.