



Wolfson Department of Chemical Engineering Seminar

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<https://gtiit.zoom.us/j/95463226139>

Development of nanocomposite gas separation membranes based on polymers of intrinsic microporosity (PIMs)

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PhD Final Seminar

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Global environmental problems induced by huge amounts of CO₂ emissions have attracted increasing attention. Membrane separation has been investigated as a promising strategy for carbon capture and storage (CCS) due to its properties of small footprint, environmentally friendly, high efficiency, and low cost, which shows the potential for upscaling. Polymers of intrinsic microporosity (PIMs) are attractive in gas separation for their high fractional free volume, while the relatively unsatisfactory selectivity cannot be ignored. The molecular imprinting technique (MIT) is a versatile approach to introducing three-dimensional holes into polymer matrixes, which contain specific recognition sites for the templates. This work employed MIT to prepare a molecular imprinting polymer containing rich CO₂ adsorption sites for fabricating PIM-1-based CO₂ separation mixed matrix membranes (MMMs). The obtained MMM enhanced CO₂ permeability from 6000 to 10000 barrer. In the next step, the UiO-66-NH₂ was introduced as the core for surface molecular imprinting. With the imprinted UiO-66-NH₂ as filler, the obtained MMMs showed good CO₂/N₂ separation performance with CO₂ permeability of over 12,000 barrer and CO₂/N₂ selectivity of about 32, which is beyond the 2019 Robinson upper bound. To further enhance the membrane selectivity, thermal-treated amidoxime-modified PIM-1 (AOPIM-1) was used as the polymer matrix instead of PIM-1. Though a certain sacrifice in permeability was observed, the CO₂/N₂ selectivity improved significantly, demonstrating that thermal crosslinking effectively optimizes the membrane's pore structure.