Molecularly engineered switchable photo-responsive membrane in gas separation for environmental protection

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Abstract

In recent years, stimuli-responsive materials have garnered interest due to their ability to change properties when exposed to external stimuli, making it useful for various applications including gas separation. Light is a very attractive trigger for responsive materials due to its speedy and non-invasive nature as well as the potential to reduce energy costs significantly. Even though light is deemed as an appealing stimulus for the development of stimuli-responsive materials, this avenue has yet to be extensively researched, as evidenced by the fewer works done on the photo-responsive membranes. Of these, there are even less research done on photo-responsive materials for the purpose of gas separation, thus, we have collected the examples that answer both these criteria in this review. This review covers the utilisation of photo-responsive materials specifically for gas separation purposes. Photochromic units, their integration into gas separation systems, mechanism and research that have been done on the topic so far are discussed.

Keywords: Gas separation, Membrane technology, Photo-responsive, Stimuli-responsive

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1. Introduction

Even though various conventional processes have been utilised for the purpose of gas separation [1-4], membrane processes have emerged and remained as an attractive alternative [5-11]. In fact, membranes have managed to make its way into the industry in larger-scale applications [12-16], proving its capability to match the performances of the conventional processes while overcoming their myriad disadvantages. For example, the efforts from Du Pont Co. [12] and Ube Industries [17] justify the potential of membrane in gas separation.

This sparks the interest to further improve the membrane processes to yield higher productivity by introducing reversibly switch functional materials in the membrane for a more convenient and energy-saving process [18-21]. The aim is to go beyond the host-guest interaction related stimuli, whereby researchers are intent to expand the concept of the responsive framework to the structural transformation controlled by the external stimuli [22, 23]. Indeed, it opens possibilities to implant a unique functionality in a membrane.

Recent research is geared towards exploiting stimuli-responsive materials whose interfacial properties and subsequent separation performances can be controlled by external stimuli such as the changes of temperature, light, pH, solution ionic strength, chemical cues or electric and magnetic fields [18, 24-26]. By integrating the stimuli-responsive functional groups in the pores or on the surface of a membrane, the resulting stimuli-responsive membrane can take advantage of the interplay among the membrane pores. Thus, it results in changes of the structural conformation of a membrane or the reactivity of the stimuli-responsive groups to control the mass-transfer and separation processes of a membrane [27].

A trait of a stimuli-responsive membrane that is very appealing is the high rate and selectivity with which the reversible changes occur [15, 28]. For example, Bassanetti and co-workers [29] has successfully produced a flexible crystalline prototypal structure for reversible CO$_2$, CH$_4$
and Xe uptake. In the study, the expandable porous adamantoid framework manifested its response to selected gases to promote reversible conformational changes, including enlargement of the channel cross-section, which can thus be considered for eco-friendly gas capture and separation.

A unique thermo-responsive gating adsorption behaviour that depends on the thermal motion of the extra-framework counterions of metal–organic framework (MOF) was presented by Gao and co-workers [30] to observe selectivity uptake of CO$_2$ over N$_2$, H$_2$, and Ar. The NO$_3^-$ ions in thermos-responsive MOF, [Cu$_2$(BTR)$_2$]·2NO$_3^-$·xG (1), located around the adsorption pores exhibited intense motion of the nitrate from closed pore to open pore structure in response to the thermal stimulus. Subsequently, this regulated structural transition from closed to open pore state allowed the passage of specific gas penetrant. At 195 K, results showed no obvious sorption of CO$_2$, implying the closed pore state of MOF. In contrast, an abrupt increase in CO$_2$ uptake was observed when temperature was approaching 231 K, referring to the “opened pores” structure which was dynamically accessible to CO$_2$ molecules. Without causing deformation of the nested structure, authors have demonstrated a reversible gating adsorption by thermally triggering the displacement of the NO$_3^-$ around the MOF pores, thus enabling a strategic separation and encapsulation of CO$_2$.

However, designing a functional stimulus-responsive material still poses a challenge as they require complicated coordination between the responsive material and the stimulus to get an output that can be measured [31-33]. The preservation of responsiveness function after incorporation on surfaces, at interfaces or in the material itself is key [34]. Thus, having in-depth knowledge of the interacting mechanisms between the stimulus and the corresponding responsive materials can be useful in outlining the inherent physicochemical properties of the responsive materials. This widens the opportunity to further manipulate the performance of
the material to explore the many possibilities for designing smart responsive systems.

However, as the subsequent sections of this review will reveal, works on photo-responsive systems for gas separation are limited. Other than the lack of understanding the molecular-level mechanism of photo-responsivity that causes the changes in photo-chromic units upon exposure to light, researchers have encountered obstacles such as the degradation of the photo-chromic unit [21] and reduction in the system’s performance due to the incorporation of photo-chromic unit [35].

In this review, the synthesis and utilisation of stimuli-responsive materials that undergo physicochemical changes when exposed to external stimuli for gas separation are highlighted. Several reviews have already broached the topic of stimuli-responsive materials in membrane system, however, those mainly focused on liquid-phase separation. Very few have focused exclusively on stimuli-responsive materials for the purpose of gas separation, especially for light- or photo-responsive ones. From this review, some of the limitations in photo-responsive systems for gas separation are presented and addressed, such as the possible mechanisms behind photo-reaction, the less common photo-chromic units as well as the common ones used for gas separation and incorporation of these photo-chromic units into a system that is suited for gas separation. A better in-depth look for a more specific light- and photo-responsive-based membrane system can be obtained in this work, which could possibly spark innovation for further improvement in smart membrane for gas separation processes.

2. Stimuli-Responsive Materials in Gas Separation

In past decades, significant progress has been reported in the fundamental understanding of organic molecules which are capable of reversibly transforming into two or more stable states in response to external signals (temperature, pH, light, etc.) [36]. This opens the possibility of
designing sophisticated responsive membrane tools by modifying the chemical composition, 
design and morphology of a membrane, basing the response mechanism on disparities in 
surface energy, polymer entropy and segmental interactions [15]. For instance, surface energy 
is the driving force of the surface responsive reorientation as systems fundamentally minimise 
the interfacial energy between the surface of the responsive material and its immediate 
environment. Numerous researches were reported by utilising MOF [24, 37] and membranes 
[38, 39] for the purpose of gas separation, focusing on the interfacial energy between the 
responsive material and its immediate environment to control the pore’s opening of the 
separation system to the specific size of gas molecule. Since many gas mixtures to be 
separated are similar in size, common sieving mechanisms cannot efficiently cater to the 
specific size of gases. This is where stimuli-responsive materials, with the ability to fine-tune 
their properties with external stimulus, come into the picture.

In example, Ma and co-workers [40, 41] reported a mesh-adjustable molecular sieve 
(MAMS) based on a MOF, whose mesh can be adjusted between 2.9 and 5.0 Å, which covers 
the size range of almost all common gas separations. Size adjustment is based on the degree 
of dehydration of MAMS at various temperatures, which can show temperature-triggered 
capture and release of specific gas penetrant in a narrow temperature range through designed 
stimuli-responsive gating effects. To make a MAMS, Ma and co-workers used an amphiphilic 
ligand 5-tert-butyl-1,3-benzenedicarboxylate (BBDC), [40]) and 4’-tert-butyl-biphenyl-3,5-
dicarboxylate (BBPDC, has one more phenyl ring than BBDC), [41] that consists of 
hydrophobic and hydrophilic ends. The hydrophilic end of the ligand will bind with the metal 
clusters, where the structure will propagate into a 2D layer. Two layers of ligands will 
sandwich a metal-cluster layer, thus giving rise to a trilayer packed through van der Waals 
interaction.
In the study [40], MAMS exhibited highly selective uptake of H₂ over CO, N₂, or O₂. The molecular sieving effect most likely comes from the BBDC gates. At 77 K, MAMS gate opening was around 3.0 to 3.4 Å, with all hydrophilic channels and hydrophobic chambers connected, giving rise to a 3D gas container with space continuity. Due to the larger kinetic diameters of 3.46, 3.64, and 3.76 Å for O₂, N₂, and CO, respectively, MAMS only allows H₂ to enter the hydrophobic chambers with one-dimensional gated channels, which accounts for the high H₂ uptake using MAMS at 77 K. When the temperature was raised to 87 K, MAMS can take up a significant amount of O₂. In fact, if temperatures can be tuned continuously and precisely, gas separations such as those of N₂/O₂, which is difficult to achieve because the size difference of these two gases is very small, would be readily attainable by using this temperature responsive MAMS. Ma et al. [40] has successfully demonstrated a linear relationship between the mesh gate opening (D) and the temperature (T); D=D₀ + αT, where D₀ refers to the mesh size at 0 K and α is a constant, to predict the best temperature for the separation of a gas mixture. D₀ and α are affected by applied tert-butyl group in MAMS. The above temperature responsive MAMS was constructed of weakly bonded sheets via van der Waals interactions with one-dimensional gated channels [40, 41]. The inter-sheets bonding interactions may not stay stable in harsh flue gas applications. Thus, Wriedt and co-workers [42] had proposed an improved design of MAMS by introducing strong coordination bonds between the layers. PCN-200 (PCN=porous coordination network) was used instead of the amphiphilic ligand, to fit exactly one CO₂ molecule in a short interlayer distance of the elastic polar pocket of PCN-200. The developed high density PCN-200-ac is stable up to 218°C with high CO₂ uptake capacity of 129 gL⁻¹, which meets the temperature requirements for physical sorbents in a flue gas separation process. More importantly, PCN-200-ac showed no loss in capture capacity even after running a series of thirty adsorption–desorption cycles.
at 313 K. By alternating heating and cooling of PCN-200-ac between 423 K and 296 K, respectively, CO₂ loading capacity of 1.13 mmolg⁻¹ was realised by flowing pure CO₂ over the sample and 0.65 mmolg⁻¹ in CO₂/N₂ (15:85) mixture gas as flue gas simulant. This result emphasises the capabilities of the thermal-responsive PCN-200 as a sorbent to remove CO₂ from flue gas [42].

Stimuli-responsive materials for gas separations commonly utilise materials sensitive to temperature, as shown in the examples given above, yet another stimulus may prove to be a more practical choice. In particular, photo-responsive materials for gas separation is highly appealing, the most popular amongst the reported external stimuli (redox, temperature, pH, light, magnetic field etc.) without any by-product generation [37]. Photo-irradiation that brings about reversible changes in physicochemical properties generally works by isomerisation of the photo-chromic groups integrated in the bulk or surface of a membrane, where the membrane separation selectivity and regeneration efficiency can be tuned upon light irradiation [15, 20]. Generally, photochemical reactions are induced by ultraviolet, visible and infrared radiation, where molecules absorb the light energy through excitation of electrons. Efficiency of this process depends on the structure transformations of the photo-chromic groups (Fig. 1) as well as the energy of the light absorbed. Light absorption causes the molecules to become excited, thus initiating various photo-chemical reactions [20]. The changes in photo-chromic groups’ structures cause the changes in macroscopic properties of a membrane, including the charge, wettability and mass transport properties [15].

Park and co-workers [37] have demonstrated an instant CO₂ capture and release platform by introducing an azobenzene (AZB) functional group to the organic linker of an MOF. Before UV exposure, the freshly made MOF sample manifested significant CO₂ uptake of 22.9 cm³/g at room temperature and 1 bar, with a negligible adsorbed amount of N₂. Up to
53.9% adsorbed CO$_2$ can be released after 5h exposure to UV irradiation as a result of the isomerisation of the AZB groups inside the pores of the MOF. The MOF adsorbent was able to return to its original state upon gentle heating at 60°C for 30 minutes in the dark. Potentially, the findings showed a possible efficient separation of gas molecules (uptake of CO$_2$ and negligible adsorption of N$_2$) with fast regeneration ability (release of CO$_2$ to return to its original state through heating process) [37]. The working mechanism of photo-responsive materials, as well as the use of AZB and MOF in gas separation systems will be discussed in more detail and depth in later sections.

Release triggers for adsorbed gases, such as thermal heating, tend to be very energy intensive [20], consuming a large amount of energy from the power plant. Despite this, most research works have used temperature as stimuli for the purpose of creating stimuli-responsive gas separation systems. Therefore, there is a strong motivation for the development of release triggers that do not require intensive energy from the plant, for example photo-responsive material triggered upon light irradiation, drawing from renewable energy resources such as solar energy. Unfortunately, utilisation of stimuli-responsive materials have not been focused on gas separations, especially with light as the stimuli. Next sections are focused on photo-isomerisation in polymer and integration of photo-responsive material in gas separation.

3. Photo-Responsive Systems for Gas Separation

Light is a very intriguing source of stimulus for a responsive trigger with the potential to drastically reduce energy costs if used with stimuli-responsive material that strongly absorbs light whilst desorbing adsorbed components [43-45]. In addition, stimulation by light is interesting as photo-excitation can be relayed in a very selective and speedy manner without
imposing additional changes to the chemical potential of the system [36, 45, 46]. However, as photo-irradiation is an external stimulus, it requires the membrane housing to be optically transparent [15].

In a photo-reaction mechanism, two types of reactive entities can be distinguished, which are photo-sensitizers and photo-reactive molecules or moieties. Photo-sensitizers selectively absorb light and trigger a chemical reaction of another substance as a result. They function as “antennas” for light and from the excited state, transfer the absorbed energy through electron or radiative transfer [20]. Ding et al. [47] reported the usage of metallophthalocyanine based conjugated microporous polymers (MPc-CMPs, M=Co, Ni, Cu, Zn) as photo-sensitizer to generate singlet oxygen ($^1\text{O}_2$). In the study, an oxygen-saturated solution of 1,3-diphenylisobenzofuran (DPBF) was irradiated with light in the presence MPc-CMPs to produce $^1\text{O}_2$, for application fields such as photodynamic therapy. Results demonstrated that at different photo-catalysis capability of MPc-CMPs, the central metal species (M=Co, Ni, Cu, Zn) in the MPc will probably influence the triplet states, triggering the activation of photo-activity to generate $^1\text{O}_2$. In the report, ZnPc-CMP and CuPc-CMP have demonstrated high efficiency for the generation of $^1\text{O}_2$ over other reported organic porous materials with lower energy required for light excitation.

As for gas separation, a novel strategy to utilise stimuli-responsive materials to trigger gas adsorption and desorption through the formation of radical triplets has been designed by An et. al [46]. In the work, MOF containing photo-reactive zwitterionic pyridinium 4-carboxylate units was exposed to UV light irradiation, forming radical triplets through intramolecular electron transfer. As this reversible light-responsive generation of radicals occurs at the major adsorption sites of the MOF, the CO$_2$ adsorption could be controlled, where the light irradiation will cause the adsorption capacity of CO$_2$ to decrease. Even though
existing works are still quite scarce, it is evident that there is increasing interest for photo-responsive materials to be used for gas separation. In order to pursue the effort, a deeper understanding of the materials’ properties, their interacting mechanisms with gas penetrants and integration approaches into the separation systems are of interest. More details of MOF in photo-responsive gas separation systems will be presented in the following sections.

### 3.1. Photo-Reaction Mechanisms

After absorption of light, photo-reactive molecules or moieties can undergo either irreversible or reversible reaction, which is typically known as isomerisation. Photo-isomerisation to one isomer and back is either both reversible photo-reaction processes, and one is a photo-chemical reaction and the other is a thermal reaction. These two reactions allow the photo-chromic unit to switch between two chromophoric moiety states \[48\], which brings about molecular changes such as group polarity, charge, and size \[24, 49\]. These molecular changes would show responses that can be visualized, that is, changes in macroscopic properties \[15\].

Fig. 2 shows an example of visual representation of these changes, which can affect the mass transport through the porous and non-porous barriers of a membrane separation layer.

Weh and co-workers \[50\] had produced a photo-responsive polymethacrylate membrane by chemically binding AZB which is capable of reversible photo-isomerisation (trans-to-cis states) to the side chain of polymethacrylate. Mass transport of gases through membranes is controlled by solubility and diffusivity, which depends on the relative size of the gas molecules and free volume within the polymer. When UV light was exposed to AZB, the spheroidic cis structure of AZB within the membrane matrix would fill the pores within the polymer, hence lowering the free volume of the membrane compared to when AZB is in the trans form. In the study, the cis-form of AZB hinders diffusivity more effectively compared to
the trans isomer.

In another work carried out by Weh et al [38], two zeolites were used to host AZB groups to control the gas permeation of a photo-responsive membrane. This intra-zeolitic photo-isomerisation was found to be capable of altering the molecular size, geometry and dipole moment of the AZB, resulting in changes of the vacant pore space for specific gas molecule to transport through the membrane. In their work, Weh et al [38] had summarized a few factors that contribute to the controlled permeation of gases. First, AZB isomers in the trans- and cis-forms can reversibly occupy different positions in the framework, which can present the AZB unit as a diffusion obstacle with different efficiencies depending on the position. Second, different configurations of the AZB isomers also means that the AZB unit can alter the free volume available in the membrane, depending on the space occupied by the isomers, to control the gas permeation and selectivity of a membrane [25, 50]. Third, the electrostatic interactions between AZB and gas penetrant could also play a role, especially between the polar cis-AZB and the more polar permeant gases. Dipole-dipole interactions between cis-AZB and polar gas molecules will increase the barrier to gas transport, in addition to the fact that AZB is bulkier in its cis-state compared to its trans-state [38].

However, there is another interpretation for MOFs containing photo-sensitive units affecting gas separation processes. Instead of undergoing photo-isomerisation from trans-to-cis state, light irradiation functions to increase the surface energy of the photo-sensitive MOF, thus, weakening the intermolecular interactions between adsorbed gas molecules and the MOF surface [44]. Although most azo-MOF reacts to UV irradiation by isomerisation, the same trigger caused structural oscillations due to the C-C-N bending movement of the azo-containing ligand instead [51]. Similarly, Lyndon et al. [45] found that the UV irradiation of an azo-containing ligand in MOF resulted in C-C-C and C-C-N bending movements due to
the suppression of isomerisation. They observed that the bending modes resulted in a release of adsorbed CO\textsubscript{2} molecules, due to the higher surface energy on the pore surface of MOF when irradiated. Indeed, this light-responsive functionality was used to trigger the reversible adsorption and release of CO\textsubscript{2}, finding that the exposure to UV light triggered an instantaneous release of up to 64% adsorbed CO\textsubscript{2} with broadband radiation or concentrated solar sources.

While understanding the underlying mechanisms responsible for the observed behaviour is key to design the photo-stimulated CO\textsubscript{2} capture and release process, studies addressing the issue are rarely reported because it is difficult to investigate the molecular-level mechanisms through experiments. Huang et al. [25] investigated the mechanism of CO\textsubscript{2} capture and release using AZB functionalised MOF, calling into question the hypothesis that the drop in CO\textsubscript{2} adsorption capacity is caused by the steric effects from the changes in pore morphology of the stimuli-responsive membrane, blocking the stronger metal oxide binding sites. Two MOF with AZB functional group attached to their ligands were investigated in their work. Even though isomerisation of the AZB groups to the cis-state reduced the pore diameter of the frameworks, they observed that the change in CO\textsubscript{2} absorption capacity when photo-isomerisation occurred is actually caused by the alteration in local interactions from the accumulation of charge on metal oxides.

Through calculations of radial distribution functions between the CO\textsubscript{2} centre of mass and two specific atoms of the framework in both trans- and cis-configurations of AZB, it was suggested that the trans-AZB isomer provided extra interactions that led to the increased CO\textsubscript{2} uptake [25]. While accessible surface area, free volume, pore morphology and adsorption sites can influence the uptake of CO\textsubscript{2}, adsorption enthalpy plays the largest role in controlling the capacity of CO\textsubscript{2} adsorption in a porous structure. In the same work [25], it was found that the
conventional CO$_2$ binding sites were still available even when the AZB groups were in cis-state, not shielded as was suggested before. This further proves that the extra interactions between CO$_2$ molecules and the MOF when AZB was in the trans-state was the main factor that led to the higher CO$_2$ molecule adsorption capacity, as compared to when the AZB group was in the cis-state. This finding is in line with the CO$_2$ binding energies when the AZB groups were in trans- and cis-configurations at -31.3 and -27.5 kJ/mol, respectively.

Another photo-reaction process called photo-induced surface relief (PSR) formation is exhibited by azopolymer thin films, where surface modulation on the submicron scale occurs under light irradiation. Kameda et al. [52] found that the azopolymer in the trans-form supported by a base membrane were in the form of many localised particulate domains. However, after being exposed to irradiation, these particulate domains spread out to cover the surface of the base membrane. Based on the observation, they suggested that upon irradiation, the azopolymers were fluidised and went through plastic deformation during the relief-forming process, known as PSR, which is reversible. Gas permeability of a rubbery polymer is significantly higher than that of glassy polymer. The initially glassy azopolymer particulate domain, when irradiated, was plasticised and fluidised. Consequently, the change in state from glassy to rubbery could increase the gas permeability. Thus, the photo-responsive permeability of the membrane is caused by the thin layer of azopolymer spread on the surface of the base membrane. The photo-induced change in the azopolymer properties such as the increase in free volume was shown through the change in permeability under irradiation. Permeability of helium (He) gas increased by approximately 5% under irradiation with blue light, then returned to its initial value when irradiation was stopped.

While the PSR formation described above is very intriguing, other research suggested that the switchable properties of photo-responsive materials are derived from other causes.
Many works explained that the changes in permeability or selectivity of the photo-responsive materials upon irradiation were due to the structure alteration when exposed to light. This morphology change could decrease the free volume of the material or membrane, hindering the gas diffusivity and thus becoming a diffusion obstacle. Another suggestion is that the polarity of the photo-chromic units is affected by isomerisation, consequently increasing the ability to attract polar gases. Yet a different proposed mechanism is that the irradiation of photo-responsive material could enhance its surface energy, which could trigger the release of adsorbed gases or increase its binding energy, thus enhancing the adsorption of target molecules. Either way, the effect of irradiation on photo-responsive systems depends on the properties of the photo-chromic molecules, which would be discussed in the next section.

3.2. Photo-Chromic Units

Based on the working mechanisms of photo-responsive materials in gas separation described in Section 3.1, this section focuses on the prevailing photo-responsive materials that have been researched for gas separation. Among the popular photo-chromic units used to design stimuli-responsive systems for the purpose of gas separation, AZB and diarylethene (DAE) are the most thoroughly researched. The changes in conformation of these photo-chromic units upon irradiation can be exploited to control the absorption properties or adsorption capacity of the microporous structures hosting these molecules [25, 53].

The planar trans-isomer of AZB has a greater distance between the aryl termini than the non-planar cis-isomer [54]. The distance between the para-carbon atoms in an AZB molecule in the trans-state is 9 Å, which changes to 5.5 Å when isomerised into the cis-isomer [35, 37, 55]. This geometrical disparity between AZB isomers can be translated into bulk materials, altering their size. As for its reversible transformation capability, the
thermodynamically stable AZB trans-isomer can be switched to the cis-isomer via irradiation
with UV light (\(\lambda=365\) nm), and is able to switch back to the trans-isomer through irradiation
using visible light (\(\lambda=455\) nm) [36, 56]. Because the trans-AZB is thermodynamically more
stable, the recovering of cis- to trans-isomer can also be done through thermal relaxation [53,
57]. In addition, the isomerisation also affects the polarity, dipole moment and required space,
leading to a change in the sorption behaviour.

Through the reversible photo-stimulated changes of the photo-chromic units in a
membrane, it can reversibly switch from a non-binding to a binding site for a target molecule
by alternating photo-irradiation. The AZB isomers are significantly different in terms of their
molecular geometries and electronic properties, rendering the feasibility of the AZB-
containing membrane as a light-switchable CO\(_2\) valve for remote controllable gas permeation
[53, 56]. Such a photo-responsive function using AZB derivatives was demonstrated by
Modrow et al. [57] through the incorporation of the derivatives into MOF. Irradiation with
UV light to produce cis-isomers caused the CH\(_4\) adsorption capacity to increase, while
reverting back to the trans-isomer by irradiation with visible light resulted in a reduction of
the adsorbed CH\(_4\). Cis/trans isomerisation brought significant structural changes upon the host,
leading to a substantial change in the sorption properties.

The MOF synthesised by Prasetya and Ladewig [39] containing AZB ligands showed
photo-response towards CO\(_2\) adsorption in both static and dynamic conditions. Even though
the AZB-containing MOF had demonstrated smaller surface area compared to the parent
MOF, incorporation of the AZB units enhanced the affinity between the CO\(_2\) molecules and
the MOF. This AZB-containing MOF was used as filler inside the Matrimid (MI) polymer
and the polymer of intrinsic microporosity (PIM) to produce AZB-MOF/MI and AZB-
MOF/PIM mixed matrix membranes (MMMs). Both the AZB-MOF/MI and AZB-MOF/PIM
MMMs demonstrated significantly better permeability and CO\(_2\)/N\(_2\) selectivity compared to their pristine polymers. It is worth noting that the performance of the AZB-MOF/PIM membrane had surpassed the 2008 Robeson Upper Bound.

Another photo-chromic unit that is very popular in the development of stimuli-responsive material is the DAE derivatives. When triggered by UV and visible light, DAE will undergo isomerisation involving ring-opening or ring-closing reactions due to their fatigue-resistant and thermally irreversible properties [24, 58, 59]. Both states typically show distinct absorption spectra that can be addressed by specific individual wavelength [49]. The reversible cyclisation and cyclo-reversion of DAE are enabled by a central n-electron sextet. The ring-closing conformation after light-induced cyclisation allows an extended conjugation of the electrons due to the aromatic character of the thienyl rings. This leads to the higher electronic conductivity in a ring-closing state compared to the ring-opening state. DAE isomerisation brings about two decisive changes within the molecule; different electronic properties and a more rigid structure after cyclisation when the thienyl moieties become interconnected, thus reducing the mobility [21]. Moreover, DAE derivatives have very high photo-stability with over 104 photo-induced colouration/decolouration cycles and a lifetime exceeding a thousand years at 30°C [58]. These properties and characteristics make DAE derivatives attractive for use in low-energy gas separation by altering the separation properties such as the gas adsorption capacity, permeability and selectivity.

Fan and co-workers [24] attempted to overcome the reported trade-off associated with MOFs and membranes where high selectivity often comes with low adsorption capacity and vice versa. They approached the matter by using DAE units to demonstrate increased C\(_2\)H\(_2\)/C\(_2\)H\(_4\) separation that can be switched with light as an external stimulus. A larger alteration in the DAE unit was required for the C\(_2\)H\(_4\)-loaded structure compared to the C\(_2\)H\(_2\)-
loaded structure. Moreover, for the C_2H_2-loaded structure, other than the opening movement
of the DAE unit, the whole DAE ligand was maintained, while an intense alteration on the
entire DAE ligand would be necessary for C_2H_4. Such an intense overall alteration within the
MOF skeleton would be completely rejected by the MOF due to its rigidity. The preferential
adsorption of C_2H_2 over C_2H_4 with a selectivity of 47.1 was also reflected in the isosteric heat
of adsorption (Q_s). The stronger host-guest interactions between C_2H_2 and the MOF were
demonstrated by its larger Q_s compared to C_2H_4.

From the above examples, it could be concluded that AZB in particular had garnered a
lot of attention, in part due to its larger geometrical change compared to DAE. As steric
effects could potentially affect the gas separation performance, a bigger change in the
properties when isomerised could give a more pronounced change in performance. In addition,
a study showed the performance of DAE-incorporated gas separation system decreasing after
several cycles [21]. Further research needs to be done to understand the reason behind this so
that measures can be taken to prevent or delay the occurrence. Another interesting avenue for
future research is the integration of other photo-chromic molecules for gas separation, which
is something that is severely lacking. However, the few works that have utilised other photo-
chromic units to separate gases are given below.

Spiropyran is a photo-chromic unit that so far has rarely been used in gas separation.
However, Healey et al. [60] had successfully incorporated spiropyran into MOF through a
two-step post-synthesis modification of Zr-oxo nodes in MOF-808. Exposing spiropyran to
UV irradiation would trigger an interconversion from the colourless spiropyran to its coloured
merocyanine (MC) state [55, 61, 62]. The resulting MOF exhibited photo-sensitive response
towards pore volume and CO_2 uptake, due to the modified MOF having a markedly higher
affinity towards CO_2 than the parent MOF [60].
In another work carried out by Huang and co-workers [63], they had utilised another photo-sensitive unit, anthracene to design and synthesise covalent organic frameworks (COFs) to observe its ability to adsorb gas. Irradiation by light caused structural transformations that consequently resulted in changes of properties, including porosity, luminescence and $\pi$-electronic absorption. This thermally reversible reaction of the COF also showed a change in gas storage capability, showing potential for photo-responsive gas adsorption.

While research on AZB and DAE are relatively extensive, other photo-chromic units such as spiropyran and anthracene have not been studied much in comparison. This may due to limitations that is not clear as of yet. However, these very lacks of research done into other photo-chromic units makes these units more interesting, and give motivation to look into their prospects in gas separation systems.

3.3. Techniques to Integrate Photo-Responsive Materials into the Gas Separation Systems

MOF is already an attractive topic of research for gas storage and separation. More recently, it has been used to host photo-chromic units that could change properties as a response to external stimulus, which could then produce a change in gas separation performance. While photo-chromic guest molecules can be simply loaded or dangle from the MOF side chains, incorporating the molecules directly onto the backbone of the framework has the highest potential for drastic transformations, even though it is the most difficult to achieve. One of the common methods to integrate photo-responsive units is by incorporating them into MOF or membrane matrix, which can be classified into three generations [64, 65], as shown in Fig. 3. MOF is a class of microporous crystalline material comprised of metal centre and organic ligands, periodically joined to give a crystalline porous array [45]. It
comprises of very large surface area as well as free volume with low density [66]. In terms of chemistry properties, MOF is chemically and thermally robust, in addition to having the ability to retain structural rigidity during reversible adsorption and desorption [22, 36]. Hence, it is a promising host for photo-chromic units, which sparked many investigations into energy-efficient gas capture and release that can be externally triggered [25].

The first generation of MOF is loaded with photo-responsive guest molecules, lending them an extrinsic photo-responsive property [32, 39]. However, this approach is not applicable in all cases as the isomerisation of the guest molecule can be blocked by the host framework [35, 50]. For instance, Hermann and co-workers [67] observed that the loading of AZB units into MIL-53(Al) has not only produced densely packed MOF with reduced pore volume, but also prevented AZB trans-to-cis isomerisation.

By depositing UiO-67 layer on α-Al2O3 support, Knebel et al. [35] loaded AZB molecules into the matrix of UiO-67 membrane. Results showed that AZB had completely filled the pores of UiO-67, inhibiting isomerisation of AZB necessary for photo-switching. After in situ desorption of AZB was done to reduce the amount of AZB loaded inside the membrane, only then could gas permeation be observed. In this case, H2 and CO2 permeances decreased when the AZB units were in trans-state, with CO2 being more significantly affected due to its size. This observation was attributed to the gating mechanism where AZB in the trans-state reduced the aperture of the gate for gas molecules to diffuse through.

The second generation of MOFs differs from the first by having the photo-responsive side chains directly within the framework organic linkers [39]. Of the three generations of photo-responsive MOFs, this generation is the most researched, with researchers commonly linking the photo-chromic unit covalently to the organic linker. Several materials have shown the ability to allow the AZB side-chain to retain its photo-isomerisation ability [57], resulting
in a change in pore dimensions when irradiated and subsequently altering its gas separation capabilities [37].

For example, Li and co-workers [68] investigated the capability of PCN-250 with AZB ligands as second generation MOF in light-induced swing adsorption (LISA) to capture and release CO\textsubscript{2} using solar energy instead of electricity. The metal clusters of the MOF gave it great structural stability while the flexibility was provided by the AZB-containing ligands. Exposing the MOF to a UV light source would cause the ligand to undergo a suppressed bending in a non-periodic fashion which would release the adsorbed CO\textsubscript{2} through the LISA process and regenerating the MOF. Intermittent light exposure demonstrated a reversible CO\textsubscript{2} capture and release, or dynamic adsorption. An interesting observation was that reversing the cis-state AZB back to the trans-state AZB did not require exposure to visible light or heat treatment. The significantly accelerated relaxation of AZB-containing ligands in the MOF happened in only 5 minutes at room temperature in the dark. Reusability tests after five cycles showed only a 7.3% and 4.9% loss in adsorption and LISA capacity, respectively, confirming the robustness of the MOF and its promising function for stimuli-responsive capture and release applications.

In another work carried out by Luo et al. [58], they have constructed a porous diarylethene MOF (DMOF) using a DAE derivative with two N-donor sites. N- and O-donor ligands were used to interact with dicarboxylate ligand of biphenyl-4,4’-dicarbocxylic acid (H\textsubscript{2}bpdc). The large skeletal structure provided by the photo-responsive DAE group allows for the construction of porous DMOFs. They found that this DMOF adsorbed CO\textsubscript{2} selectively over N\textsubscript{2} due to the quadruple moment of CO\textsubscript{2}. After UV irradiation, the uptake of CO\textsubscript{2} increased significantly; 4 times the amount without UV irradiation.

While the third generation of photo-responsive MOFs is the most difficult to achieve,
it is also the most promising of the three classes of MOFs. This generation of photo-responsive MOFs incorporates a photo-chromic unit directly in the “backbone” on the linker, not as a side chain [39, 69]. Using this method, the light stimuli will directly affect the framework itself, providing the potential to demonstrate the most dramatic transformations [65].

For an example, Tang et al. [70] had utilised AZB as the base in the backbones in conjunction with Lewis bases by covalently bonding them to design novel photo-switchable sorbents for CO$_2$ capture. Results demonstrated that the difference in binding strength between the designed molecules and CO$_2$ was hardly affected by electronic and steric effects. However, by substituting various groups into the ortho position with respect to the carbon atom of the benzene group of the azo moiety, CO$_2$ molecules were expected to form intramolecular H bonds with the substitution groups. These bonds would then enlarge the difference of CO$_2$ binding strengths of the cis- and trans-isomers so that the CO$_2$-sorbent interaction can change from strong in one isomer to weak in the other by using light as stimulus, leading to a significant decrease in energy needed for sorbent regeneration.

DAE moieties had also been used as a building block for the construction of PIM by Becker et al. [21]. The properties of the DAE-PIM were studied during cyclisation and cyclo-reversion with light as the trigger. Porosity in PIMs is derived from the interstitial voids that are produced by the inefficient space packing of the polymer chains due to the building blocks possessing an inherent kink to their structure. Even though the geometrical change of DAE as a result of isomerisation is not as dramatic as AZB, DAE has a rigid and essentially planar structure in the ring-closing form and a slightly bent geometry in the ring-opening form. In this case, the stiff but kinked structure of PIMs is key to the microporosity of the polymer, as the inefficient chain packing determines the percolated free volume that is accessible for
target adsorption molecules. Isomerisation of DAE altered the distribution of electrons, but when stored in the dark at room temperature, DAE-PIM did not revert to the ring-opening isomer even after weeks of storage. Thus, as long as not exposed to visible light, DAE-PIM would not return to the ring-opening isomer for a long time. As for the separation performances, making a composite DAE-PIM with MI, yielding DAE-PIM@MI increased the diffusion coefficient (D) for both O$_2$ and CO$_2$ by 10%. Most importantly, the isomerisation of the composite significantly reduced D value in the ring-closing state while cyclo-reversion into the ring-opening state enhanced the D value. Even though cycling between the ring-opening and ring-closing states was reversible through photo-induced switching, however, after five cycles, the intensity of the band at 610 nm only retained 80% of the original value. This might due to the degradation of DAE moieties, hence preventing more comprehensive cyclo-reversion [21].

Porous coordination polymers also have highly regular nanopores useful for storage and separation as well as being responsive to guest molecules such as photo-chromic groups while retaining high regularity. In a work demonstrated by Yani et al. [32], AZB was incorporated into a flexible host compound, [Zn$_2$(terephthalate)$_2$(triethylenediamine)]$_n$ whose pores become deformed due to the incorporation of the aromatic guest molecules. UV irradiation triggered a partial expansion of the host framework when AZB was isomerised from trans- into cis-form. The composite did not initially adsorb N$_2$, possibly due to the pore blockage of the host by AZB, but upon UV irradiation, the adsorption amount increased by 8.3 fold. The significant increment of the adsorption capability of the composite is attributed to the large structural change of the host framework that was triggered by the isomerisation of AZB, as the tetragonal form in cis-AZB expanded the host framework to allow diffusion of gas molecules into the pores.
While a lot of research on photo-chromic materials for the purpose of light-triggered gas separation used MOF as the host, a considerable amount of work has also been done on introducing this switchable function into polymers and membranes for gas separation.

Ulbricht [71] stated that even though strides have been made in the development of microporous inorganic membranes, subtly fine-tuning the properties of the barrier, a necessity for molecule-selective separations, is only possible using polymeric structures. Membrane processes for gas separation has been extensively researched, but they have fixed permeation properties. In gas adsorption processes, regeneration of the adsorbent is usually costly and energy intensive [43]. Therefore, interest in producing membranes with switchable function to develop a multifunctional membrane has been growing. A change in the photo-responsive guest molecule by external stimuli would simultaneously transform the membrane according to the alteration of the guest molecules [32].

For example, Weh et al. [50] attempted to prepare photo-responsive membranes using two methods; physically loading AZB in poly(methylmethacrylate) (PMMA) matrix and by chemically binding AZB to the side chain. For the membrane with AZB chemically bound to its side chain (PMAZB), the flux rates of gases permeating through the membrane appeared in decreasing order of $\text{H}_2 > \text{n-C}_4\text{H}_{10} > \text{CH}_4 > \text{SF}_6$. They noted that the permeability of gases in the cis-state of the membrane was lower compared to the trans-state, due to the spheroidic cis-AZB being a more effective diffusion obstacle compared to the elongated trans-AZB. On the other hand, the permeability of PMMA membrane with AZB physically dissolved in the membrane matrix has shown insignificant influence when switching in between the trans- and cis-states. The flux rates of gases for this membrane (physically loading AZB in PMMA matrix) is significantly lower compared to the pure PMMA membrane and the PMAZB.

Authors concluded that the physically loaded AZB in PMMA matrix caused AZB to occupy
the free volume in the membrane, hence reducing the transport capacity of the membrane 
drastically as a result. In addition, the membrane in cis-state had a higher dipole moment 
compared to the membrane in trans-state, leading to the high permeation rate for a polar 
permeate compared to the non-polar gases [24, 38].

Another type of membrane called surface-mounted MOFs (SURMOFs) were 
fabricated by growing MOF thin films continuously in layers to obtain monolithic, crystalline 
MOF films, as reported by Wang and co-workers [72]. The fabricated membranes have a 
pillared layer structure of Cu$_2$(AzoBPDC)$_2$(AzoBiPyB) containing AZB on asymmetric 
mesoporous aluminium oxide support. Switching the AZB-groups in the membrane from 
trans- to cis-state only reduced the H$_2$ permeance slightly while the CO$_2$ permeability was 
significantly affected. Fig. 4 shows the reversible photo-switching of permeability and 
separation factors of H$_2$:CO$_2$ and N$_2$:CO$_2$ mixtures through the membrane. The cis-AZB with 
higher polarity had a stronger attraction with the fairly strong quadruple-moment of CO$_2,$ 
resulting in a decrease in the CO$_2$ diffusion rate. On the other hand, the interaction between 
H$_2$ and the membrane was not affected by the switching of isomers. This led to an increase of 
trans/cis permselectivity from 3.0 in the trans-state to approximately 8.0 in the cis-state of the 
membrane.

In the same work [72], the separation switching of N$_2$:CO$_2$ mixture was found to be 
reversible as well, with a selectivity increase from 5.5 to 8.5 when the AZB was switched 
from trans-state to cis-state, as N$_2$ permeability was hardly affected by the trans-to-cis 
switching. Amazingly, not only the separation factor of the photo-responsive membrane can 
be switched between a minimal and a maximal values, but any values between the 
aforementioned ranges can be achieved by adjusting the irradiation times or by manipulating 
the relative intensities of the multi-component light irradiation. Therefore, the separation
factor of the permeate gases can be continuously tuned. One of the advantages in being able
to tune the separation factor of the membrane can be demonstrated by the ability to control the
flammability and the safety of the pre-combustion mixture of H₂:CO₂ in the presence of air or
oxygen. Using this system, each composition in between the ranges can be tuned so that the
hazardous or non-hazardous condition of the gas mixture can be controlled remotely [72].
Other than MOF and polymeric membranes, COFs, a new class of crystalline porous
polymers have also been constructed with AZB as part of their building block. Having
permanent porosity, structural periodicity and thermal stability, COFs have been garnering
interest recently, especially for gas absorption and storage owing to their large pore volume,
high surface area and low density [63]. As COFs are constructed through atomically precise
integration of building blocks, the variations of pore sizes, functions and structures of COFs
are dependent on the diversity of the functional building blocks [69]. In fact, numerous groups
reported the excellent gas pair separation using two-dimensional layered-stacking COF-COF
composite membrane to separate H₂/CO₂, H₂/N₂, H₂/CH₄ [73], and CO₂/N₂ [74, 75] gas
mixtures, together with an enhancement in permeability to surpasses the Robeson upper
bounds. By integrating photo-responsive AZB to control the conformational changes of COF
composite membrane, a more precise gas pair separation is expected [63].
Zhu and Zhang [54] utilised a porous organic polymer (POP) to investigate whether
the change in AZB polarity would be adequate to change the bulk material properties. POP is
a porous material that is lightweight with intrinsic porosity, excellent physicochemical stability
as well as structural tenability. It was observed that a bulky substituent on the AZB moiety
hindered the photo-responsiveness of the frameworks due to the steric hindrance that
suppressed the isomerisation. These polymers demonstrated reversible changes in pore size
distribution and CO₂ uptake using UV light irradiation and thermal treatment. Even after
several cycles, the absorption properties were similar, indicating robustness and good
switchability of the pore structures of the POPs. While photo-responsive MOFs with AZB
units could induce the release of CO$_2$ by isomerisation from trans-to-cis, the POPs showed a
unique inverse alteration by absorbing more CO$_2$ upon trans-to-cis isomerisation. This is
because the dipole-quadrupole interaction between polar CO$_2$ molecules and more polar cis-
AZB was enhanced.

So far, the integration strategies of photo-chromic units into MOFs and membranes
are similar. Most photo-chromic research for gas separation has focused on the second
generation incorporation method. Even though the results of the first and second generation
methods have shown promise, researchers have started to venture into the third generation of
photo-chromic integration, which, while difficult, has the best chances of showing prominent
results.

4. Practicability of Photo-Responsive Membranes for Gas Separation

Membrane adsorbers for gas separation have vital advantages compared to conventional
porous adsorbers due to the membrane pore structure which allows directional convective
flow through the pores, greatly reducing the distance for pore diffusion. In membrane
adsorption, separation occurs based on the reversible binding of molecules on the
functionalised pore walls. Therefore, development of membrane adsorbers needs to progress
towards optimising the pore structure as well as surface layer functionality so that a maximum
number of binding sites with optimum accessibility can be obtained [71]. Conventional
methods to regenerate adsorbents such as vacuum and heating are not energy efficient [66],
providing incentive towards research on photo-responsive adsorbents that can be regenerated
using less energy at a reduced cost [51]. Various polymers and membranes have also been
investigated for incorporation of photo-chromic units to manipulate gas separation performance [21, 39]. Utilising a photo-responsive membrane could open possibilities for switchable, multi-functional membranes, in addition to potentially reducing gas separation costs. While many attributed the change in gas separation performance to the change of the conformation of photo-chromic groups, some also mentioned that the changes were brought by the alteration of dipole moment of the photo-chromic groups and their interaction with target molecules upon isomerisation [24].

Park et al. [37] chose to synthesise MOF with AZB as a dangling group as CO$_2$ adsorption on MOF at room temperature is relatively high. There was a delay in the change in gas adsorption isotherms when exposed to UV irradiation due to the slow isomerisation in the PCN-123 MOF caused by the steric hindrance among AZB groups. Initially, the AZB groups were predominantly in the trans-state, which is a relatively stable state. Upon being exposed to UV irradiation, only some of the AZB groups changed into cis-state right away, causing a decrease of CO$_2$ uptake by 26.6%, while more trans-AZB later became cis-AZB as they were triggered by the initial conformational changes, leading to the gradual reduction of gas uptake. The overall reduction of CO$_2$ uptake after irradiation was 53.9%. The change in position of the dangling benzene group in PCN-123 caused the reduction in gas adsorption capacity. The electron density of the cis-isomer around the Zn$_4$O cluster is higher than the trans-isomer, making it harder for CO$_2$ molecules to approach and occupy the metal clusters in the cis-state, which are the main adsorption sites for CO$_2$ molecules.

PCN-123 was used in addition to another MOF, Cu$_2$(AzoBPDC)$_2$(AzoBiPyB) by Yang et al. [43] as photo-responsive adsorbents for CO$_2$ adsorption. Similar to Park et al [37], they arrived to the conclusion that the decreased uptake of CO$_2$ for both AZB-containing MOFs upon trans-to-cis isomerisation due to irradiation was attributed to the blocking of
metal nodes of the MOF by cis-AZB, which function as the adsorbing sites. As both MOFs operate under the same mechanism, they hypothesised that this mechanism would apply for materials with exposed metal centre.

Another MOF combining AZB and acrylamide ligands was synthesised and reported by Dang and co-workers [66]. Results showed excellent selectivity for CO₂ adsorption over CH₄, O₂, CO and N₂ gases. Exposure to UV light caused the AZB in the MOF to isomerise rapidly from trans- to cis-state, subsequently reducing the size of the pore accommodating adsorbed CO₂ to incite CO₂ release, thus regenerating the adsorbent. Dynamic irradiation caused instantaneous CO₂ release with 21% desorption capacity, showing promise for a low energy CO₂ release in adsorption process.

A concern with using UV light to trigger trans-to-cis photo-isomerisation of AZB is that the UV light can destroy and reduce the lifespan of materials and hinder applications. To counter the problem, advanced AZB derivatives that can be reversibly switched without UV irradiation were developed [76]. AZB is particularly popular in the development of visible-light-activated photo-responsive materials as it can be tailored to demonstrate complete addressability in the visible region. Castellanos et al. [48] fabricated two MOFs; aluminium-based MIL-53(Al) and zirconium-based UiO66(Zr) that were responsive to visible light using the same ortho-fluoroazobenzene linker with different inorganic secondary building blocks (SBU)s to observe the impact of the framework structure.

Both MOFs have pore dimensions that can accommodate the AZB group within the large voids in the crystalline network in a disordered orientation. The AZB units forced the MIL-53(Al) lattice into the form with large pores, as the benzene rings of the terephthalate linkers allow a certain rotational degree of freedom, opening to the possibility of the bulky AZB units to cause rotation of the terephthalate rings away from parallel orientation. It is
suggested that the isomerisation efficiency of AZB in F-azo-MIL-53(Al) was drastically reduced due to congestion of AZB in the MOF. On the other hand, F-azo-Uio-66(Zr) responded readily to both green and blue light, with green light triggering isomerisation much more efficiently compared to UV radiation [48].

In situ photo-irradiation experiments where samples were irradiated during CO$_2$ adsorption showed that for both frameworks, the pure MOFs without AZB moieties could adsorb more CO$_2$ compared to their photo-responsive counterparts as the pores were partially occupied by the pendant AZB moieties. Even though F-azo-Uio-66(Zr) showed good photo-chromic response under visible light, the CO$_2$ adsorption capability was not affected by green light, concluding that the geometrical alteration of the side group of the linker in this MOF had no influence on the total volume of CO$_2$ molecules that can be hosted within the pores. Conversely, the CO$_2$ adsorption capability of F-azo-MIL-53(Al) dropped by 10% during exposure to green light due to the hindered isomerisation events. As AZB is not fluorescent, the excess of light energy that was absorbed by the photo-chromic units must dissipate via radiationless photo-physical deactivation, where the electronically excited azo moieties relaxed through intramolecular vibrations that were transferred to the framework, thus increasing the surface energy and locally producing heat. It was measured that the decrease of 10% in the CO$_2$ adsorption ability was comparable to that produced by an increase in overall temperature by 10°C. Although hindered isomerisation is typically undesirable in photo-chromic applications, in this case it led to the change in CO$_2$ adsorption capacity [48].

Both AZB and DAE units were used to synthesise MOF by Fan et al. [51] to observe adsorption of CO$_2$, C$_2$H$_2$ and C$_2$H$_4$ gases. Exposure of the MOF to UV light resulted in C-C-N bending of the AZB-containing ligand rather than trans-to cis-isomerisation [45, 68], while UV irradiation at 320 nm exhibited partial transformation of DAE units from ring-opening to
CO₂ adsorption occurred closer to the photo-sensitive centre of the MOF, leading to a larger effect on CO₂ adsorption upon UV irradiation compared to C₂H₂ and C₂H₄ adsorption. This gave the MOF ability to alter the selectivity of the gas mixtures as well as adsorption capacity [51].

While light is an attractive trigger for gas capture and release, Li et al. [64] raised the issue of high volume of materials needed for light to pass in a large scale operation. Therefore, they developed a magnetic induced swing adsorption (MISA) strategy by incorporating magnetic nanoparticles into the MOF to generate heat when exposed to magnetic field. This field can fully penetrate the MOF, unlike light to release the adsorbed gas from MOF. MISA process, which requires electricity is combined with LISA process obtain a low-energy and efficient approach for gas adsorption. Integrating Fe₃O₄ magnetic nanoparticles into PCN-250 MOF that contains AZB linkers produced a dual stimuli-responsive MOF that can enhance the gas release up to 96.8% of CO₂ desorption.

In an absorption process through a dense polymeric membrane, the effective flux rate represents the net transport through the membrane. This is governed by both the diffusivity and solubility below the glass transition temperature of the polymer [50]. By incorporating photo-chromic molecules inside the absorption system, the isomerisation state of the photo-chromic groups can be manipulated by light, consequently providing switching control over the flux and the separation factor of the gas permeating through the membrane. Therefore, the permeability of gases as well as the separation factor or permselectivity can be continuously and remotely tuned by adjusting the ratio of trans- and cis-isomers of the photo-chromic groups [72].

Most works investigating photo-responsive materials necessitates the use of UV light to activate the photo-sensitive moieties, which comes with several disadvantages such as poor
selectivity, low penetration into most media and potentially damaging. Therefore some
research has been done to utilise visible light as a trigger instead, which can be beneficial
especially in the aspect of exploiting solar energy as visible light is predominant in the solar
spectrum [48]. Müller et al. [76] used o-fluorozobenzene moieties to functionalise a
nanoporous, photo-switchable MOF thin film that can undergo isomerisation with visible light.
Another advantage to avoiding UV light is that the ligand-to-metal-charge transfer, which is
excited by UV light in Cu-paddle-wheel MOF structures would not be stimulated using
visible light. Therefore, very high isomerisation yields of AZB groups can be achieved, which
is unprecedented in MOFs. Green and violet light were used to control the isomerisation state
of the photo-chromic groups, consequently controlling the gas uptake and separation factors
of H₂:C₂H₄ and H₂:C₃H₆ mixtures through MOF membranes.

In the same work [76], it was found that H₂ permeability through a membrane of thin
SURMOF film grown on a mesoporous Al₂O₃ substrate was hardly affected by the trans-to-
cis switching. However, the permeability of C₂H₄ and C₃H₆ increased by approximately 25
and 30%, respectively when the membrane was irradiated with green light. The original
permeance and selectivity could be recovered by irradiating with violet light. The H₂:C₂H₄
selectivity can be switched between 6.6 and 8.2 while the C₃H₆ selectivity can be switched
between 8.8 and 12.6. Tuning these selectivities between the aforementioned ranges can be
done continuously by adjusting the irradiation times or by mixing the green and violet light.
As the separation of H₂ and CO₂ showed no significant changes upon isomerisation and the
polarity difference between trans- and cis-AZB was unable to influence the adsorption of the
apolar hydrocarbons, the conclusion was that the separation by the photo-responsive
Cu₂(F₂AzoBDC)₂(dabco) membrane was driven by steric effects instead of changes in the
dipole moment. In the trans-state, the SURMOF had smaller pore diameter, resulting in a
lower hydrocarbon permeability compared to the cis-state. This finding is in line with the larger switching impact observed for the larger hydrocarbon molecules while the permeability of the smaller H₂ molecules was not affected. It was believed that the steric effect superimposed the polarity switching effect of AZB, leading to no switching impact on the H₂:CO₂ separation [76].

MMMs incorporating light responsive MOFs into MI polymer matrix were synthesised by Prasetya et al. [77] to observe the light-responsive gas separation. JUC-62 and PCN-250 are both generation-3 light responsive MOFs with significantly reduced CO₂ adsorption capacity when exposed to UV light. On the other hand, the permeation of N₂ hardly changed with photoswitching. A maximum switchability of 8.5% was obtained using 15wt% JUC-62-MI, which was lower than expected. This was caused by the rigid structure of MI polymer matrix that limited the ligand bending of the photo-responsive MOFs. While PCN-250-MI had higher adsorption capacity than JUC-62-MI as PCN-250 had more active sites for CO₂ permeation, both MMMs had superior CO₂/N₂ ideal selectivity compared to pristine MI membranes. This was due to the good interaction and dispersion of MOFs inside the polymer matrix as well as the good affinity between the light-responsive AZB units in the MOFs with CO₂ [24, 54]. Long-term performance showed negligible changes for 15wt% JUC-62-MI and a slight decrease in 10wt% PCN-250-MI, thus showing promise for the construction of photo-responsive anti-ageing material for CO₂/N₂ separation.

In adsorption, the change in position of the photo-chromic dangling group can occupy or block the adsorption sites, which would reduce the adsorption capacity. On the other hand, the conformation change of the photo-chromic units can alter their efficiency as diffusion obstacles in absorption by changing the free volume. The polarity of the photo-chromic isomer is another factor that can affect both the permeability in absorption and adsorption.
capacity when polar target molecules are involved. There are many possibilities for photo-responsive materials in the gas separation industry, not only to tune the separation factor remotely to change the permeability and gas adsorption capacity. For instance, the triggering using external stimuli can be used to manipulate the selectivity of gas mixtures to control the flammability of the mixture and thus the safety of the system. A summary of the research available on photo-sensitive materials for gas separation so far is listed in Table 1 and the basic points covered in this review are condensed into Fig. 5.

5. Conclusions and Future Direction

This review has covered in detail the emerging use of photo-responsive materials for the specific purpose of gas separation. To help gain a better understanding of the research that has been done so far, the research to date has been explored from the aspects of photo-chromic units, their mechanism, their integration into the system, and the practicability for gas separation processes. While some work on gas separation through absorption or adsorption processes using photo-responsive materials are available, from this review it is evident that the investigation has mostly focused on two photo-chromic groups; AZB and DAE. In addition, most of the research found incorporated the photo-chromic units to MOF instead of membranes, and attached said groups to the dangling side chains instead of the backbone of the framework. It is clear that the research done so far have barely scratched the surface of the almost infinite possibilities those photo-responsive materials have to offer for gas separation processes.

Future direction of gas separation incorporating photo-responsive materials could be focused on overcoming the limitations of the process, such as the degradation of photo-chromic units, and potentially damaging effects of the UV lighting. Even though the results of
photo-responsive materials has been promising so far to separate gases, the fact that so little research has been done on photo-chromic units other than AZB and DAE shows that the scope of this topic can be expanded. By pushing beyond the current boundaries placed on the photo-responsive materials for gas separation research, various improvements and innovations can be discovered.

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Nomenclature

AZB Azobenzene
COF Covalent organic framework
DAE Diarylethene
LISA Light-induced swing adsorption
MISA Magnetic induced swing adsorption
MOF Metal-organic framework
MAMS Mesh-adjustable molecular sieve
MI Matrimid
MMM Mixed matrix membrane
PCN Porous coordination network
PCP Porous coordination polymer
PIM Polymer of intrinsic microporosity
PMMA Poly (methylmethacrylate)
**POP** Porous organic polymer

**PSR** Photo-induced surface relief

**SURMOF** Surface-mounted MOF

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Tables and Figures

Table 1. Research on Photo-sensitive Materials for Gas Separation to Date

| Photo-chromic moiety | Process       | Investigated gas | Host for the photo-chromic units | Ref.          |
|----------------------|---------------|------------------|----------------------------------|--------------|
| AZB                  | Adsorption    | CO₂              | MOF                              | [25, 37, 43, 45, 48, 68] |
| AZB                  | Adsorption    | CO₂              | Dual stimuli-responsive MOF      | [64]         |
| AZB                  | Adsorption    | CH₄              | MOF                              | [57]         |
| AZB                  | Adsorption    | CO₂, CH₄, O₂, CO, N₂ | MOF                              | [66]         |
| AZB                  | Adsorption    | CO₂              | Lewis bases                      | [70]         |
| AZB                  | Adsorption    | N₂               | PCP                              | [32]         |
| AZB                  | Adsorption, absorption | CO₂, N₂       | MOF, MI                          | [39, 77]     |
| AZB                  | Absorption    | H₂, CO₂, N₂, CH₄, C₂H₆, C₃H₈ | MOF                              | [35]         |
| AZB                  | Absorption    | H₂, n-C₄H₁₀, CH₄, SF₆, CH₃OH | PMMA membrane                    | [50]         |
| AZB                  | Absorption    | CH₄, CO₂, N₂     | Zeolite-AZB membranes            | [38]         |
| AZB                  | Absorption    | He               | Azopolymer membrane              | [52]         |
| AZB                  | Absorption    | CO₂              | POP                              | [54]         |
| AZB     | Absorption | H₂, CO₂ | SURMOF | [72] |
|---------|------------|---------|--------|------|
| AZB     | Absorption | H₂, C₂H₄, C₃H₆ | SURMOF | [76] |
| AZB, DAE| Adsorption | CO₂, C₂H₂, C₂H₄ | MOF    | [51] |
| DAE     | Adsorption | CO₂, N₂  | MOF    | [58] |
| DAE     | Adsorption | C₂H₂, C₂H₄ | MOF    | [24] |
| DAE     | Absorption | N₂      | PIM    | [21] |
| Spiropyran | Adsorption | CO₂  | MOF    | [60] |

**Fig. 1.** Examples of photo-chromic conjugates used in photo-responsive materials for gas separation: azobenzene, spiro-benzopyran and diarylethene, with their reversible photo-reactions. Reproduced with permission from Wandera et al. [15]; Copyright (2010) Elsevier and He et al. [20]; Copyright (2008) Elsevier.
Fig. 2. Isomerisation of photo-chromic azobenzene (AZB) unit resulting in changes of conformation (size) and properties (polarity or charge) (Authors’ own constructed figure).
**Fig. 3.** Integration of photo-responsive units into metal-organic frameworks. Reproduced with permission from Coudert [65]; Copyright (2012) American Chemical Society.

**Fig. 4.** The separation of $\text{H}_2$:CO$_2$ and $\text{N}_2$:CO$_2$ gas mixtures (a) and (b), respectively through the membrane under dynamic photo-switching. Red spheres with the scale on the left-hand side represent the separation factors. $\text{H}_2$ and CO$_2$ (a) as well as $\text{N}_2$ and CO$_2$ (b) permeances are represented by black solid squares and black open squares, respectively, and the logarithmic scale is shown on the right-hand side. Reproduced with permission from Wang et
Fig. 5. Summary of photo-responsive materials for gas separation (Authors’ own constructed figure).