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Improved CO$_2$/CH$_4$ Separation Properties of Cellulose Triacetate Mixed–Matrix Membranes with CeO$_2$@GO Hybrid Fillers

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Abstract: The study of the effects associated with the compatibility of the components of the hybrid filler with polymer matrix, which ultimately decide on achieving mixed matrix membranes (MMMs) with better gas separation properties, is essential. Herein, a facile solution casting process of simple incorporating CeO$_2$@GO hybrid inorganic filler material is implemented. Significant improvements in material and physico-chemical properties of the synthesized membranes were observed by SEM, XRD, TGA, and stress-strain measurements. Usage of graphene oxide (GO) with polar groups on the surface enabled forming bonds with ceria (CeO$_2$) nanoparticles and CTA polymer and provided the homogeneous dispersion of the nanofillers in the hybrid MMMs. Moreover, increasing GO loading concentration enhanced both gas permeation in MMMs and CO$_2$ gas uptakes. The best performance was achieved by the membrane containing 7 wt.% of GO with CO$_2$ permeability of 10.14 Barrer and CO$_2$/CH$_4$ selectivity 50.7. This increase in selectivity is almost fifteen folds higher than the CTA-CeO$_2$ membrane sample, suggesting the detrimental effect of GO for enhancing the selectivity property of the MMMs. Hence, a favorable synergistic effect of CeO$_2$@GO hybrid fillers on gas separation performance is observed, propounding the efficient and feasible strategy of using hybrid fillers in the membrane for the potential biogas upgrading process.

Keywords: gas separation; cellulose triacetate; CeO$_2$@GO hybrid fillers; mixed-matrix membrane

1. Introduction

Membrane systems have become an accepted gas treating technology. This technology plays a key role in biogas upgrading and offshore natural gas treatment processes, especially to remove CO$_2$ [1]. Depending on the materials used for the synthesis, inorganic and polymeric membranes are in use. Inorganic membranes show significantly higher diffusivity and selectivity of gas molecules due to discerning ability based on pore size and shape, possessing high thermal and chemical stability, mechanical strength, and longer life span [2]. However, these membranes face the limitation of poor scalability, high cost and complicated fabrication procedure [3].

On the other hand, polymer membranes still dominate the current market in the separation process. Despite this, pristine polymeric membranes always suffer from the trade-off between permeability and selectivity [4]. The high demand for membrane technology in gas separation and rapid effort toward seeking membranes with higher permeability and selectivity has motivated the development of MMMs. The addition of the porous/nonporous inorganic fillers into the polymer matrixes combine the processability of polymeric membranes with the good gas separation performance of the inorganic membranes [5], synergistically contributing to the enhancement in membrane separation performance, thus minimizing the trade-off limit. Furthermore, highly absorptive inorganic
nano-fillers benefits from their large surface area and abundant active sites [6]. However, these inorganic fillers occasionally exhibited limited compatibility with the polymer matrices, poor dispensability due to their strong intermolecular van der Waals interaction and aggregation in membranes [7,8]. The resulting difficulty in achieving homogeneous fillers dispersion in a continuous polymer phase and consequent loss of selectivity thus represent the major limitation of MMMs preparation. Furthermore, the functionalization of fillers is often conducted to improve their compatibility and dispersion properties in the polymer matrix [9]. However, this technique is laborious and could disrupt the structure and lose the intrinsic properties of the native fillers particles [10]. Much research into potential new materials for gas separation membranes is thus driven by the limitation and disadvantages which is in existence in the current available MMMs.

Coronas group reported a significant improvement in the CO$_2$/CH$_4$ and CO$_2$/N$_2$ selectivity with hybrid MOF (HKUST-1)@zeolite (silicate-1) and polysulfones system [11]. They also conducted experiments to investigate the synergistic effect of two fillers with different natures [11–13]. Since then, the use of the two different filler particles into a polymer matrix to enhance the gas separation performance of the resultant MMMs has attracted much attention. Jamil et al. fabricated the MMMs by incorporating reduced graphene oxide (rGO) and ZIF-8 nanofillers into the PES matrix and observed high permeability and low selectivity for CO$_2$/CH$_4$ [14]. Wong et al. synthesized the highly dispersed CNT into a thin-film nanocomposite via the addition of an amphiphilic GO nanosheet onto a PSF support layer. They found 30% and 60% improvement in the gas permeability and selectivity, respectively [15]. Ahmed et al. prepared the PVDF membrane incorporated with POSS and SAPO-34 zeolite and observed the detrimental increment in CO$_2$ flux compared to the pristine PVDF membrane [16]. It was revealed that the interactions between the two different fillers with distinct properties, morphologies, and surface chemistry promote polymer-fillers interaction, thereby reducing the probability of the fillers agglomeration and enhancing the gas separation efficiency leading to synergy to acquire non-linear effects [13]. Nevertheless, only a countable number of such studies can be found in the literature, suggesting that such hybrid filler MMMs are still in the initial stage of development, and further research is essential.

In this study, we investigate the synergy between CeO$_2$ and GO nanosheets in MMMs. GO was selected as a filler because the sheeted morphology of GO can improve the dispersion of the fillers co-existed in the nanocomposite. Such behavior is assumed due to the structural and property-wise resemblance of GO as polyelectrolytes in a 2D configuration. Their surfactant-like characteristics at the interface thus can serve as a special class of dispersion agents [17]. Likewise, these nanosheets act as a potential support to deposit inorganic oxides nanoparticles on their surface due to the presence of a large number of polar groups that play an important role in participating in a wide range of bonding interactions [18]. Similarly, the charge transferability between GO and CeO$_2$ due to the difference of their work function create stronger interaction between them [19]. Due to the diverse reactive groups like epoxide, hydroxyl, and carboxylic acid at the edge and basal plane of GO nanosheets, there is high compatibility among the GO nanosheets with the polymer matrix via covalent or non-covalent connections [20]. Moreover, they tend to be parallel to the membrane surface. Such sheeted GO morphology gave rise to a strong steric effect and prevented the aggregation of other fillers [21]. Furthermore, the randomly distributed GO sheet may act as a barrier in the polymer matrix due to the hindered diffusion pathway through the nanocomposite, thus enhancing the selectivity [14,15]. GO, which retains the lamellar structure of graphite, contains several unpaired π electrons and oxygenated functional groups, making GO highly selective towards small and polar molecules like CO$_2$ [22]. On the other hand, Cerium oxide (CeO$_2$), one of the most reactive rare earth metal oxides, has received attention as a promotor or catalyst even in industrial processes due to its oxygen storage capacity [23]. Similarly, owing to the high adsorption affinity of both oxidized/reduced forms of CeO$_2$ with CO$_2$ forming bridged, monodentate, bidentate carbonates, and bicarbonates, and leading to the improved permeability when blended
with polymer matrix [24,25], CeO$_2$ was selected as another material for the formation of composite nanofillers. The synthesized membrane’s affinity was then determined by evaluating the CO$_2$ uptake and CO$_2$ and CH$_4$ permeability capacity. To the best of our knowledge, there is no evidence of using CeO$_2$@GO hybrid filler to fabricate MMMs in conjunction with polymer for gas separation purposes yet.

2. Materials and Methods

2.1. Chemicals

CTA (acetyl content 43–44%) was obtained from Acros Organics (Waltham, MA, USA). Cerium nitrate hexahydrated (Ce(NO$_3$)$_3$·6H$_2$O, 99.99%), sodium hydroxide (NaOH, 98.99%), and 1-methyl-2-pyrrolidinone (NMP, ACS reagent > 99.0%) were purchased from Sigma-Aldrich (St. Louis, MO, USA). Ethanol (C$_2$H$_5$OH, 99.89%) was purchased from VWR Chemicals (Radnor, PA, USA). All the chemicals were of analytical grade and were used as received without any further purification. Similarly, GO was synthesized in the laboratory. The detail of the synthesis process is explained in suplementry information section.

2.2. Preparation of CeO$_2$@GO Hybrid Fillers

CeO$_2$ nanoparticles were prepared using the facile hydrothermal process [25]. 0.1 g of as-prepared CeO$_2$ nanoparticles were mixed with 130 mL ethanol, and different dopamine functionalized GO concentrations (3 wt.%, 5 wt.%, 7 wt.%), and 10 wt.% with respect to the wt. of CeO$_2$ nanoparticles). The mixture was sonicated for 30 min. (VWR ®Ultrasonic cleaner, USC-THD (Lutterworth, UK) stirred for 4 h. followed by hydrothermal treatment using Teflon lined vessel at 110 °C for 18 hrs. Afterwards, the mixture was dried at 80 °C in an oven.

2.3. Preparation of CTA-CeO$_2$@GO Mixed-Matrix Membrane and Its Mechanism of Formation

0.045 g of the hybrid nanoparticles (CeO$_2$@GO) was mixed with 26 mL of NMP. The mixture was sonicated for 30 min. and stirred for 3 h afterwards. 1.62 g of CTA polymer was added to the initial mixture and stirred for 18 h to obtain the optimal dispersion of nanoparticles in the polymer solution. The mixture was further sonicated for 30 min. and left undisturbed for 4 h. The membrane was then cast in a glass plate using an applicator (Elcometer 3580, Manchester, UK). The casted film was left undisturbed at ambient conditions till complete evaporation of the solvent took place. The membrane was then further kept in a vacuum oven at 60 °C for overnight for complete drying. The membrane with 3 wt.%, 5 wt.%, 7 wt.%, and 10 wt.% of GO are referred to as CTCeGO3, CTCeGO5, CTCeGO7, and CTCeGO10, respectively, throughout the manuscript. The physico-chemical properties and the gas sorption/permeation behavior of CTA-CeO$_2$ and pristine CTA membranes were reported previously [25].

Among various techniques of membrane synthesis, the dry-casting method was used. This technique is a suitable method for preparing asymmetric membranes with dense skin applicable for the gas separation process. This synthesis process is characterized by nonsolvent and/or solvent evaporation from an initially homogeneous single-phase polymer solution. During solidification, polymer from the polymer-rich phase precipitates to form a solid matrix which enfolds the polymer–lean phase. The final membrane thickness is a fraction of the initial cast film thickness owing to both nonsolvent and/or solvent loss and excess volume of mixing effects [26]. Similarly, membrane morphology and performance are also strongly influenced by external conditions such as the casting and evaporation temperature and air circulation during the evaporation step. Since NMP is a very slow evaporating solvent, the composition change in the evaporation process is rather slow. The evaporation of NMP from the surface of the casting solution can be assumed to be compensated by NMP diffusion from the casting solution interior to the surface. This slow solvent evaporation at ambient temperature and ambient airflow results in the membranes with the thin dense skin layer desirable for gas separation [27].
2.4. Materials Characterization

The transmission electron microscopy (TEM) analysis of the nanoparticles was performed on a JEM-2200FS (Jeol, Kyoto, Japan) instrument, maintaining an accelerating voltage of 200.00 kV in TEM imaging mode. Similarly, the morphology of prepared membranes was determined using scanning electron microscopy (SEM, Oxford Aztec, 80 mm², High Wycombe, Abingdon, UK) for the detailed analysis of element distributions within the materials and chemical microanalysis of elements present. The 3D non-contact optical surface profiler New View 9000 (ZYGO, Middlefield, CT, USA) was used for the no-contact surface roughness measurement. X-ray photoelectron spectroscopy (XPS) analysis was carried out in ESCAProbeP Spectrometer, Omicron Nanotechnology (Uppsala, Sweden). The primary X-ray beam was monochrome radiation of an Al lamp with energy of 1486.7 eV. Powdered X-ray diffraction (XRD) measurement was performed at a temperature of 273.5 K using a 2nd-Generation D2 Phaser X-ray diffractometer (Bruker, Billerica, MA, USA) with Cu Kα radiation (λ = 0.15418 nm), SSD (1D mode) detector, coupled 2θ/θ scan type, and continuous PSD fast scan mode. The range of measured Bragg angles was 5°–80°. Fourier-transform infrared spectroscopy (FTIR) measurements were performed using a NicoletTM iS50 FTIR Spectrometer (Thermo Fischer Scientific, Waltham, MA, USA) in absorbance mode. The spectra were taken in the wavenumber range of 400–4000 cm⁻¹. The thermogravimetric analysis (TGA) of synthesised MMMs was done by Setsys Evolution (Setaram Lyon, France). The samples were heated in an aluminium crucible from 40 °C to 800 °C at a rate of 10 °C·min⁻¹ under an N₂ atmosphere with a flow rate of 60 mL·min⁻¹. Tensile stress-strain curves were measured using an Instron Universal Testing Machine 3365 (Instron, Norwood, MA, USA) equipped with pneumatic grips, rubber-coated sample gauge length (initial sample length between the grips) of 10 mm, sample width of 4 mm (on average), and a crosshead speed of 5 mm·min⁻¹ until specimen break. The measurement was carried out at an ambient temperature.

2.5. Gas Sorption and Gas Permeation Measurements

CO₂ and CH₄ sorption experiments were performed gravimetrically at 25 °C in a pressure range from 0.1 to 1.5 MPa using a self-developed sorption apparatus equipped with a calibrated (McBrain) quartz spiral balance. A detailed description of the apparatus and the experimental procedure is described elsewhere [28,29]. The gas permeation affinity of all synthesized membranes was determined by using single gases (H₂, CO₂, O₂, N₂, and CH₄) in a custom-built time lag permeation setup. The membrane for gas permeation measurement was cut and tightly enclosed in a circular membrane permeation cell with an effective membrane surface area of about 2.14 × 10⁻⁴ m². Similarly, the permeation cell was made air-free by using a vacuum at both ends. In addition, after each gas experiment, the permeation cell was continuously evacuated with a vacuum to remove the gases present therein. The synthesized membranes were subjected to test gases, and the data were collected using the SWeTr version 1.13 (2003, Neovision, Prague, Czech Republic) data acquisition software. All permeation data were collected when the steady-state was reached. Each gas was measured three times, and the average value was recorded to minimize the experimental error [25].

The increase in the pressure in the fixed permeate volume was monitored as a function of time as soon as the membrane was exposed to feed gas at a pressure of 1.5 bar. For the given setup, the transient permeation curve, describing the pressure increase on the permeate side after exposure of the membrane to the feed gas, takes the following form [30,31].

\[
P_t = P_0 + \left( \frac{dP}{dt} \right)_0 t + \frac{RTA}{V_p V_m} P_f \sum_{n=1}^{\infty} \frac{(-1)^n}{n^2} \exp \left( \frac{-Dn^2 \pi^2 t}{4t^2} \right) \]

(1)
where $P_t$ is the permeate pressure (bar) at time $t$ (s); $P_0$ is the initial pressure, which is usually less than 0.05 mbar; $(dp/dt)_0$ is the baseline slope, which is normally negligible for a defect-free membrane; $R$ is the universal gas constant ($8.314 \times 10^{-5} \text{ m}^3 \text{ bar mol}^{-1} \text{ K}^{-1}$); $T$ is the absolute temperature (K); $A$ is the active membrane area ($\text{m}^2$); $V_p$ is the permeate volume ($\text{m}^3$); $V_m$ is the molar volume of a gas at standard temperature and pressure ($22.4 \times 10^{-3} \text{ m}^3 \text{ STP mol}^{-1}$ at 0 °C and 1 atm); $p_f$ is the feed pressure (bar); $S$ is the gas solubility ($\text{m}^3 \text{ STP m}^{-3} \text{ bar}^{-1}$); $D$ is the diffusion coefficient, and $l$ is the membrane thickness.

In the steady-state permeation condition, the exponential term approaches zero; hence, the equation becomes:

$$P_t = P_0 + \left( \frac{dp}{dt} \right)_0 t + \frac{R T A}{V_p V_m} \times \frac{p_f P}{l} \left( t - \frac{l^2}{6D} \right)$$  \hspace{1cm} (2)

A plot of $P_t$ versus $t$, after a long time, produces a straight line, which, upon extrapolation, intersects the time axis at $t = \frac{l^2}{6D}$, which describes the time lag ($\theta$) in permeation. These equations thus allow for the calculation of diffusion and permeability coefficients. Assuming the validity of the solution-diffusion model, the solubility coefficient was then determined indirectly by a simple relation: $P = S \times D$.

The gas permeability ($P$, 1 Barrer = $10^{-10} \text{ cm}^3 \text{ (STP) cm}^{-2} \text{ s}^{-1} \text{ cmHg}^{-1}$) is expressed by the following equation:

$$P = \frac{l Q_i}{A \Delta P_i}$$  \hspace{1cm} (3)

where $l$ refers to the thickness of the membrane ($\mu\text{m}$), $Q$ represents the volume flow rate ($\text{cm}^3 \text{ s}^{-1}$, STP) of gas $i$, $A$ is the effective membrane area ($\text{cm}^2$), and $\Delta P_i$ is the partial pressure difference across the membrane (cmHg). Similarly, the selectivity is expressed as:

$$\alpha_{ij} = \frac{P_i}{P_j}$$  \hspace{1cm} (4)

where $P_i$ and $P_j$ are the permeability of two pure gases ($P_i > P_j$), respectively.

3. Results

3.1. Physico-Chemical Characterizations

Structural characterization using TEM provided insights into CeO$_2$@GO composite filler. Figure 1 shows the typical TEM images of the CeO$_2$@GO composite filler. Figure 1A displays a low magnification TEM image, where GO nanoparticles show a darker colour contrast on the CeO$_2$ surface. Figure 1B exhibits a high-resolution TEM (HRTEM) image. It can be seen that the CeO$_2$ and GO particles were uniformly dispersed one into another. The distinct lattice spacing of 0.362 ± 0.001 nm in the enlarged section (Figure 1E) corresponds to the (111) plane of CeO$_2$, indicates the formation of the CeO$_2$@GO matrix. The SAED pattern (Figure 1C) were assigned to the reflection of the CeO$_2$ and (002) of GO which agrees well with the XRD pattern. The EDS results (Figure S1) demonstrate the presence of Ce, O and C on the composite matrix. The corresponding elemental mapping confirms the homogeneous distribution of elements Ce, O and C on the matrix.
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Figure 1. (A) TEM images of CeO$_2$@GO composite; (B) HRTEM image shows both CeO$_2$ and GO region; (C) SAED of the CeO$_2$; (D,E) Enlarged HRTEM images displaying the corresponding GO and CeO$_2$ lattice structure in the selected area in B.

SEM images of the surface and cross-section of synthesized MMMs are shown in Figure 2. The bright spots in the images represent the embedded CeO$_2$@GO nanoparticles. As can be seen, CeO$_2$ nanoparticles are distributed over the membrane surface with good dispersion. The tendency of CeO$_2$ particles to aggregate seems to decrease with increasing GO concentration. The cross-section view showed a dense structure with few voids. The thick membranes (25 ± 10 µm) was obtained due to delayed mixing in the phase separation with the solvent evaporation [25]. Figure S2 and Table S1 depict the surface roughness analysis results of the synthesized membranes. The hybrid membrane exhibit a smoother surface with the increase in GO concentration in the blends. This decrease in surface roughness is likely attributed to the reduction in the aggregation tendency of the nanoparticles. Similarly, with the increase in the GO concentration, the membrane structure also becomes smoother. It suggests the homogeneous dispersion ability of the GO with a good polymer filler contact preventing the aggregation of the CeO$_2$ nanoparticles. Furthermore, it also indicated that CeO$_2$ particles were dispersed among GO nanosheets and loosely entangled in the polymer matrix. The EDS mapping of Ce over the cross-section of the CTCeGO7 sample in Figure S3 shows the homogeneous distribution of CeO$_2$ throughout the membrane cross-section. The overlaid EDS image further confirms the homogeneous distribution of the fillers throughout the membrane volume, suggesting a contribution of GO to overcome the aggregation tendency of the inorganic fillers in the membrane matrix.
The XRD patterns of pristine GO, pristine CTA, and the CeO$_2$@GO incorporated CTA MMMs are displayed in Figure 3. XRD pattern of GO showed a major peak at $2\theta = 10.5^\circ$ that corresponds to (002) plane while for pristine CTA, a broad peak centered at 19° 20 value is seen. For MMMs samples, all the peaks related to CTA, CeO$_2$ and GO compared with pristine CTA, pristine GO, and CeO$_2$ JCPDS 34-0394, respectively. This suggests the homogeneous intermixing of the hybrid filler with the polymer matrix without losing its properties and conformation.

![Figure 3. XRD patterns of the synthesized MMMs with different concentrations of GO.](image-url)
The chemical composition of the CeO$_2$@GO composites was evaluated by XPS analysis. The overall XPS survey spectrum in Figure S4 confirmed the presence of C1s, O1s, and Ce3d elements in the CeO$_2$@GO composite. The high-resolution spectrum of Ce3d$_{5/2}$ and Ce3d$_{3/2}$ levels in the binding energy range between 875–894 eV and 895–925 eV were resolved into eight peaks after deconvolution (Figure 4A). These peaks confirm the presence of trivalent and tetravalent oxidation states of Ce on the given composite. The presence of Ce$^{3+}$ ions introduces oxygen vacancies in the crystal. Such high oxygen vacancies show the charge transfer effect from CeO$_2$ to GO implying the covalent interaction between CeO$_2$ and GO [32,33]. The C1s spectra for GO in CeO$_2$@GO hybrids with characteristic peaks at 284.7 eV, 286.3 eV and 288.8 eV are attributed to the C-C, C-O and C=O groups, respectively (Figure 4B). The binding energy at 284.7 eV is a typical peak position of graphitic carbon and demonstrates the sp$^{2}$- hybridized carbon in the graphite state [34].

![Figure 4. High-resolution XPS spectra of; (A) Ce3d and (B) C1s in CeO$_2$@GO composite.](image)

Figure 5 represents the FTIR spectra of the CeO$_2$@GO-CTA MMMs. The characteristic peaks at 3420 cm$^{-1}$ refer to the O-H stretching vibration band. The peak at around 2942 cm$^{-1}$ corresponds to the C-H bond in methyl and methylene groups. The band at 1740 cm$^{-1}$ is associated with the stretching vibration of the C=O group, whereas the band at 1687 cm$^{-1}$ can be attributed to the COO$^-$ group. The peak at 1431 cm$^{-1}$ is associated with the asymmetric deformation in the plane of a methyl group. The band at 1369 cm$^{-1}$ corresponds to the C-H bending vibration of CH$_3$ in the acetyl group. The absorption peaks at 1219 cm$^{-1}$ and 1033 cm$^{-1}$ are related to the C-O-C asymmetric and symmetric stretching band of CTA, respectively. The band at 907 cm$^{-1}$ is attributed to C-O-C stretching at the β1-4 glycosidic linkage [25,35,36]. The spectra of pristine CTA, as well as the modified membranes, show no significant differences. Regarding the FTIR spectra of pristine GO (Figure S5), the change in intensity of peaks at 1000–1750 cm$^{-1}$ in MMMs can be attributed to different concentrations of GO in the membrane samples.

TGA was carried out to study the influence of the hybrid nanofillers addition on the thermal stability of the MMMs. The resulting TGA and its corresponding DTA curves are presented in Figure 6. Two-step weight loss can be observed. The major weight loss in the first step occurs at around 165 °C, which is assumed due to loss of moisture, evaporation of volatile impurities within the membrane pores, and shift in the crystallinity of the glass transition temperature of the CTA matrix. The loss was around 9–14%. A significant weight loss of 85–90% was noticed at 365 °C as a result of the decomposition of the polymer matrix [37]. No significant difference in weight loss was observed for different concentrations of CeO$_2$@GO loading, suggesting no detrimental effect on the overall thermal stability of the MMMs with the incorporation of the synthesized nanofillers in the low loading percentage. However, a noticeable improvement in the thermal stability effect is seen compared to the pristine CTA membrane [25].
Representative strain-stress curves of the synthesized MMMs showing the loading effect is depicted in Figure 7. The stress-strain curves of each sample were chosen from their corresponding data sets that are closest to their respective averages reported in Table 1. Young’s modulus increases with an increase in GO concentration, indicating the significant effect of GO on mechanical strength. The tensile strength, which characterizes sample stiffness, increases with an increase in the concentration of GO in the matrix up to 7 wt.% then after decrease at higher concentration of GO whereas elongation at maximum stress decreases. Agglomeration in the high loading of GO can be the reason for the reduction in tensile strength. Compared to pristine CTA [25], the elongation at break decreased with the addition of the hybrid fillers attributed to the rigidification of the polymer chain mainly due to favourable interaction between the polymer matrix and the nanofillers [38]. The decrease in elongation at break and increase in Young’s modulus with the increase in GO concentration manifests the role of GO in fortifying interfacial interaction and improving interfacial quality of CeO$_2$@GO hybrid fillers containing MMMs.
sorption capacity on the hybrid matrix membrane occurs at energetically uniform adsorption sites with the formation of a monolayer of CO2 on the adsorbent surface.

Furthermore, the sorption data were fitted to the Langmuir model \[39\]; where $q_e = \frac{C_e K_L q_m}{1 + C_e K_L}$ (5)

Table 1. Mechanical properties of the synthesized membranes determined at 25 °C.

| #       | Stress (MPa) | Elongation at Maximum Stress (%) | Young’s Modulus (GPa) |
|---------|--------------|---------------------------------|-----------------------|
| CTCeGO3 | 36.3 ± 4.5   | 26.2 ± 4.5                      | 1.02 ± 0.14           |
| CTCeGO5 | 50.7 ± 3.3   | 18.9 ± 3.8                      | 1.14 ± 0.15           |
| CTCeGO7 | 99.5 ± 10.2  | 20.2 ± 6.1                      | 1.31 ± 0.20           |
| CTCeGO10| 34.5 ± 7.1   | 19.5 ± 3.0                      | 1.52 ± 0.11           |

3.2. Gas Separation Performances Evaluation

In order to explain the elevated selectivity, high-pressure adsorption (0–1.5 MPa) tests were conducted on MMMs possessing different GO concentrations. CO2 uptakes for all the MMMs considerably improved with increasing pressure because of the nanofillers’ specific chemical affinity for quadrupolar CO2 molecules. Similarly, an increase in the concentration of the GO increases the CO2 adsorption amount (Figure 8). This can be attributed to the rise in the number of polar groups on the GO surface, which can uptake more CO2. Moreover, the CH4 uptake was very low, which was below the sensitivity of the measuring system. This observation is related to the high condensability of CO2 relative to CH4 and potentially to the specific interaction of CO2 molecules with the membrane matrix [25]. Furthermore, the sorption data were fitted to the Langmuir model [39];

$$q_e = \frac{C_e K_L q_m}{1 + C_e K_L}$$ (5)

where $q_e$ (mg·g$^{-1}$) is the equilibrium adsorption capacity, $C_e$ (mg·L$^{-1}$) is the equilibrium adsorbate concentration, $K_L$ is the Langmuir constant (L·mg$^{-1}$), and $q_m$(mg·g$^{-1}$) corresponds to the maximum adsorption capacity of the adsorbent and gives the amount of adsorbent adsorbed after forming a complete monolayer (mg·g$^{-1}$). The fitting parameters are enlisted in Table S2. The Langmuir model presented the best fit ($R^2 = 0.997$) to describe the equilibrium data of adsorption on the given MMMs samples suggesting that the adsorption capacity on the hybrid matrix membrane occurs at energetically uniform adsorption sites with the formation of a monolayer of CO2 on the adsorbent surface.
In order to evaluate the performance of the prepared hybrid MMMs, single gas (CO₂ and CH₄) permeation tests were performed under the constant pressure of 1.5 bar and temperature of 25 °C. Adding a small amount of the GO into CeO₂ to form hybrid fillers incorporating into the CTA matrix significantly increases the selectivity maintaining almost similar or slightly higher permeability than CTA-CeO₂ MMMs (Figure 9). The gas permeation results revealed that the hybrid nanofillers significantly affect the gas separation performance of the resultant MMMs. The incorporation of hybrid CeO₂@GO into the CTA polymer matrix causes the CO₂ permeability and CO₂/CH₄ selectivity to be improved due to the high CO₂ sorption potential and strong interfacial interaction between the fillers and polymer chains. CO₂/CH₄ ideal selectivity was enhanced from 3.35 to 50.7 by increasing the GO concentration up to 7 wt.%. Similarly, the gas separation efficiency of the synthesized membranes was compared with the single CTA-GO based membrane. The CO₂ and CH₄ permeabilities of the CTA-GO membrane was 11.29 Barrer and 0.33 Barrer, respectively, with CO₂/CH₄ selectivity of 34.22. A notable increment in the gas selectivity (almost 1.5 times) can be observed in the hybrid fillers based MMMs (CTCeGO7) compared to the CTA-GO membrane. The results proved that hybrid fillers caused a synergistic effect enhancing the gas separation performance compared to the single filler. Furthermore, comparing the permeability affinity of different gases H₂, O₂, CO₂, N₂ and CH₄ were performed, taking CTCeGO7 membrane as a reference (Figure S6). The permeability follows the order as; CO₂ > H₂ > O₂ > CH₄ > N₂. However, the kinetic diameter of CO₂ (3.30 Å) is higher than that of H₂ (2.89 Å) but shows higher permeability affinity, probably because of the interaction affinity with the polymer and the fillers matrix. Similarly, the selectivity of different gases pairs observed are as follows; CO₂/N₂ > CO₂/CH₄ > O₂/N₂ > CO₂/H₂. The higher selectivity of CO₂/N₂ can be attributed to more condensable properties and more interaction of CO₂ molecules with the membrane matrix compared to N₂ molecules. These results thus suggested the CeO₂@GO hybrid fillers based MMMs to have a potential affinity for biogas/natural gas upgrading.

Figure 8. CO₂ sorption affinity of the synthesized CTA-CeO₂@GO membranes with different concentrations of GO.
Due to the difference in physical properties and density between the nonporous fillers (like silica, TiO$_2$, CeO$_2$) and polymer matrix, the aggregation of fillers are frequently encountered. It can disrupt the polymer chain packing and increase the membrane fractional free volume, increasing gas permeability [40]. For the CTA-CeO$_2$ membrane, it can thus be presumed that an increase in diffusivity and permeability (Table 2) is associated with the substantial change in the free volume due to disruption of the chain packing provides a more specious pathway for gas transport. This increase in free volume is also advantageous for large gas molecules like CH$_4$, favouring increased permeability, ultimately decreasing selectivity. GO nanosheets with good dispersion properties can be easily homogeneously dispersed in a polymer matrix with good polymer fillers contact [41]. The addition of such GO sheeted morphology is presumed to give rise to a strong steric effect and prevented the aggregation of CeO$_2$ fillers. These nanosheets also acted as a selective barrier to render high selectivity through the hydroxyl and carboxyl groups on the GO surface of MMMs. The interactions between the CO$_2$ molecules and the polar hydroxyl and carboxyl groups on the GO surface play an important role in facilitating the transport of the CO$_2$ molecules and the polar hydroxyl and carboxyl groups on the GO surface of MMMs as compared to CH$_4$. Similarly, the high affinity of CeO$_2$ towards CO$_2$ upsurges the solubility coefficient of CO$_2$ thus resulting in an overall increase in selectivity.

**Table 2.** Permeability, selectivity and their corresponding solubility and diffusivity values of the synthesized MMMs.

| #     | Permeability (Barrer) | Selectivity (CO$_2$/CH$_4$) | Solubility Coefficient ($10^{-4}$ mol m$^{-3}$ Pa$^{-1}$) | Diffusivity Coefficient ($10^{-12}$ m$^2$ s$^{-1}$) |
|-------|-----------------------|------------------------------|----------------------------------------------------------|---------------------------------------------------|
|       | CO$_2$ | CH$_4$ | CO$_2$ | CH$_4$ | CO$_2$ | CH$_4$ |
| CTA-CeO$_2$ | 9.67    | 2.89 | 3.35 | 1.33 | 0.35 | 1.6 |
| CTCeGO3 | 5.81 | 0.42 | 13.83 | 31.71 | 0.89 | 0.61 | 1.57 |
| CTCeGO5 | 6.33 | 0.17 | 37.23 | 42.99 | 0.01 | 0.49 | 1.70 |
| CTCeGO7 | 10.14 | 0.20 | 50.7 | 79.66 | 0.72 | 0.44 | 1.20 |
| CTCeGO10 | 10.87 | 0.26 | 41.81 | 14.85 | 0.47 | 2.29 | 1.42 |

Compared with CTA-CeO$_2$, the hybrid fillers incorporated membranes show a lower CO$_2$ diffusivity coefficient resulting from the stronger interfacial interaction and the larger gas transport resistance at the interface. The diffusivity coefficient of CO$_2$ decrease with an increase in the concentration of GO. This decrease in diffusivity coefficient is the indication...
of narrowing of the pore size of the MMMs, the decreased mobility of the interfacial polymer chains and increase in tortuosity [42] which generate the larger transport resistance for the gas with a larger molecular diameter and, therefore, higher selectivity for CO\textsubscript{2}/CH\textsubscript{4} separation. The incorporation of the hybrid fillers thus significantly improve the selectivity due to the intensification of the diffusion process. At a higher concentration of GO increase in diffusivity and decrease in solubility is observed, which is assumed that the aggregation of the fillers reduces the active sites of the fillers for interaction with the CO\textsubscript{2}. It also might result in the formation of microvoids, resulting in higher permeance of gases but ultimately leading to a decrease in selectivity.

In order to compare the results with the literature data, the gas separation performance of the synthesised hybrid MMMs was compared with the Robeson upper bound plot, as shown in Figure 10. Although the gas separation performance cannot cross Robeson’s upper bound limit, the hybrid fillers’ use significantly increases the CO\textsubscript{2}/CH\textsubscript{4} separation performance. Therefore, the simultaneous modification of the CeO\textsubscript{2} filler with a small amount of GO leads to the detrimental improvement in the membrane efficiency exhibiting high potential value in the bio/natural gas separation field.

**Figure 10.** A comparison of CO\textsubscript{2}/CH\textsubscript{4} gas separation efficiency of studied MMMs with Robeson 2008 upper-bound plot [4].

Furthermore, the gas separation performance of reported hybrid fillers based MMMs compared to those in this study is shown in Table 3. From this comparison, we deduce that blending the CeO\textsubscript{2}@GO fillers into the CTA matrix presented an attractive prospect for biogas/natural gas upgrading. Therefore, we believe that the development of high-performance MMMs can be realised by optimizing the membrane preparation conditions and the gas separation parameters.
Table 3. Gas separation performance of the reported MMMs in comparison with this study.

| Membrane Type                                                                 | Permeability (Barrer)  | Selectivity (-) | References            |
|-------------------------------------------------------------------------------|------------------------|-----------------|-----------------------|
| Mixed porous fillers MOFs and zeolite silicate-1 blended with polysulfone     | $P_{CO_2} = 8.9$     | CO$_2$/CH$_4$ = 22.4 | Zornoza et al. [11] |
| [HUKUST-1/S1C-PSF (16 wt% filler mixture)]                                    |                        |                  |                       |
| Ordered mesoporous silica and layered titanosilicate fillers with 6FDA-based copolyimide [MMMs (MCM-41(8Wt.%)+JDF-L1(4 wt.%)) with 6FDA-4MPD/6FDA-DABA] | $P_{H_2} = 440$     | H$_2$/CH$_4$ = 32.0 | Galve et al. [12]    |
| GO and ZIF-8 blended with Polyethersulfone matrix followed by Pebax coating [2 rGO-ZIF-8-M] | $P_{CO_2} = N/A$     | CO$_2$/CH$_4$ = 35.0 | Jamil et al. [14]    |
| MOF(UiO-66-NH$_2$)@COF (TpPa-1) hybrid fillers in polysulfone matrix [5 wt% of MOF@COF fillers] | $P_{CO_2} = 7.1$     | CO$_2$/CH$_4$ = 46.7 | Cheng et al. [7]     |
| CNT/SiO$_2$ composite core incorporated into Pebax-1657 matrix               | $P_{CO_2} = 148.3$   | CO$_2$/N$_2$ = 66.5 | Wang et al. [5]      |
| Ordered mesoporous silica(MCM-41) and MOF (NH$_2$-MIL-53(AL) blended with polysulfones [8/8 wt.% of each MCM-41 and MOF] | $P_{H_2} = 19.5$    | H$_2$/CH$_4$ = 67.3 | Valero et al. [13]  |
| Cellulose-based poly-ionic liquid membranes [P[CA][Tf2N]]                    | $P_{CO_2} = 8.9$    | CO$_2$/CH$_4$ = 22.3 | Nikolaeva et al. [43]|
| Poly(butylene succinate)-cellulose triacetate blends [CTA + 10 wt% PBS]     | $P_{CO_2} = 3.5$    | CO$_2$/CH$_4$ = 35.0 | Cihal et al. [44]   |
| PVA grafted on UiO-66-NH$_2$ incorporated into polyvinyl amine matrix [24 wt% MOF] | $P_{CO_2} = 76.13$  | CO$_2$/N$_2$ = 45.6 | Ashtiani et al. [45]|
| CeO$_2$@GO blended CTA membrane [7 wt.% GO with respect to CeO$_2$ concentration] | $P_{CO_2} = 10.14$  | CO$_2$/CH$_4$ = 50.7 | This work            |

4. Conclusions

In summary, we have successfully synthesized hybrid fillers (CeO$_2$@GO) incorporated CTA MMMs for the first time. The CTA-CeO$_2$@GO MMMs were prepared by varying the GO concentration, and its effect on the gas separation performance was evaluated. An SEM-EDS image of the membrane cross-section shows uniform dispersion of the fillers in the polymer matrix. In contrast, the other physico-chemical analysis (XPS, FTIR, SEM and TEM) reveals the interfacial interaction between constituent components of the membrane. The fabricated hybrid MMMs demonstrated superior gas separation performance at an optimum GO loading concentration of 7 wt.% at 1.5 bar feed pressure reaching CO$_2$/CH$_4$ selectivity 50.7. This increment is almost 15 folds higher than the CTA-CeO$_2$ MMMs and 1.5 fold higher than the CTA-GO membrane. The CO$_2$/CH$_4$ separation of the resulting MMMs was relatively near the upper bound region of the 2008 Robeson plotline. Thus, the gas permeation properties of the MMMs can be tailored efficiently even with the incorporation of extremely low loading of the fillers. Compared to other fillers requiring high filler loading to demonstrate enhancement of permeability/selectivity, using a low loading of fillers is advantageous given its possibility of mitigating particles aggregation and ensuring uniform distribution of the fillers in the polymer matrix. Thus, CTA membranes with a combination of two distinct nanofillers (CeO$_2$@GO) have improved physico-chemical properties and thus show great potential for CO$_2$ gas separation due to their synergistic effect.

Supplementary Materials: The following are available online at https://www.mdpi.com/article/103390/membranes1100777/s1, Methodology for the synthesis of GO. Figure S1: EDS mapping of hybrid matrix confirming the presence of CeO$_2$@GO. Figure S2: Surface roughness measurement of the synthesized membranes via 3D non-contact optical surface profiler; (A) CTCeGO3, (B) CTCeGO5,
Figure S3: EDS mapping of Ce throughout the cross-section in CTCeGO7 MMMs. Figure S4: XPS survey spectrum of CeO$_2$@GO composite. Figure S5: FTIR spectra of a pristine GO. Figure S6: Permeability and their corresponding selectivity of different gases pair in CTCeGO7 MMMs. Table S1: Arithmetic mean height, root mean square height and a maximum height of the synthesized membranes determined by the 3D optical non-contact profiler. Table S2: Langmuir sorption isotherm fitting parameters for CO$_2$ sorption in the CTA- CeO$_2$@GO membrane matrix.

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References

1. Ugo Moretti, I. Chapter 9—Polymeric membrane-based plants for biogas upgrading. In Membrane Engineering for the Treatment of Gases: Volume 1: Gas-Separation Issues with Membranes, 2nd ed.; The Royal Society of Chemistry: London, UK, 2018; Volume 1, pp. 242–255.

2. Vinoba, M.; Bhagiyalakshmi, M.; Alqaheem, Y.; Alomair, A.A.; Pérez, A.; Rana, M.S. Recent progress of fillers in mixed matrix membranes for CO$_2$ separation: A review. Sep. Purif. Technol. 2017, 188, 431–450. [CrossRef]

3. Park, H.B.; Kamcev, J.; Robeson, L.M.; Elimelech, M.; Freeman, B.D. Maximizing the right stuff: The trade-off between membrane permeability and selectivity. Science 2017, 356, eaab0530. [CrossRef]

4. Robeson, L.M. The upper bound revisited. J. Membr. Sci. 2008, 320, 390–400. [CrossRef]

5. Wang, D.; Zheng, Y.; Yao, D.; Yang, Z.; Xin, Y.; Wang, F.; Wang, Y.; Ning, H.; Wu, H.; Wang, H. Liquid-like CNT/SiO$_2$ nanoparticle organic hybrid materials as fillers in mixed matrix composite membranes for enhanced CO$_2$-selective separation. New J. Chem. 2019, 43, 11949–11958. [CrossRef]

6. Gangu, K.K.; Maddila, S.; Mukkamala, S.B.; Jonnalagadda, S.B. A review on contemporary metal–organic framework materials. Inorg. Chim. Acta 2016, 446, 61–74. [CrossRef]

7. Cheng, Y.; Ying, Y.; Zhai, L.; Liu, G.; Dong, J.; Wang, Y.; Christopher, M.P.; Long, S.; Wang, Y.; Zhao, D. Mixed matrix membranes containing MOF@COF hybrid fillers for efficient CO$_2$/CH$_4$ separation. J. Membr. Sci. 2019, 573, 97–106. [CrossRef]

8. Pfeifer, S.; Bandaru, P.R. A methodology for quantitatively characterizing the dispersion of nanostructures in polymers and composites. Mater. Res. Lett. 2014, 2, 166–175. [CrossRef]

9. Li, S.; Liu, Y.; Wong, D.A.; Yang, J. Recent advances in polymer-inorganic mixed matrix membranes for CO$_2$ Separation. Polymers 2021, 13, 2539. [CrossRef] [PubMed]

10. Goh, P.S.; Ismail, A.F.; Sanip, S.M.; Ng, B.C.; Aziz, M. Recent advances of inorganic fillers in mixed matrix membrane for gas separation. Sep. Purif. Technol. 2011, 81, 243–264. [CrossRef]

11. Zornoza, B.; Seoane, B.; Zamaro, J.M.; Tellez, C.; Coronas, J. Combination of MOFs and zeolites for mixed-matrix membranes. ChemPhysChem 2011, 12, 2781–2785. [CrossRef]
12. Galve, A.; Sieffert, D.; Staudt, C.; Ferrando, M.; Güell, C.; Tellez, C.; Coronas, J. Combination of ordered mesoporous silica MCM-41 and layered titanosilicate JDF-L1 fillers for 6FDA-based copolyimide mixed matrix membranes. J. Membr. Sci. 2013, 431, 163–170. [CrossRef]

13. Valero, M.; Zornoza, B.; Tellez, C.; Coronas, J. Mixed matrix membranes for gas separation by combination of silica MCM-41 and MOF NH2-MIL-53(Al) in glassy polymers. Microporous Mesoporous Mater. 2014, 192, 23–28. [CrossRef]

14. Jamil, N.; Othman, N.H.; Alias, N.H.; Shahruddin, M.Z.; Roslan, R.A.; Lau, W.J.; Ismail, A.F. Mixed matrix membranes incorporated with reduced graphene oxide (rGO) and zeolitic imidazol framework-8 (ZIF-8) nanofillers for gas separation. J. Solid State Chem. 2019, 270, 419–427. [CrossRef]

15. Wong, K.C.; Goh, P.S.; Taniguchi, T.; Ismail, A.F.; Zahri, K. The role of geometrically different carbon-based fillers on the formation and gas separation performance of nanocomposite membranes. Carbon 2019, 149, 33–44. [CrossRef]

16. Ahmad, N.A.; Mohd Noh, A.N.; Leo, C.P.; Ahmad, A.L. CO2 removal using membrane gas absorption with PVDF membrane incorporated with POSS and SAPO-34 zeolite. Chem. Eng. Res. Des. 2017, 118, 238–247. [CrossRef]

17. Tian, L.; Meziani, M.J.; Lu, F.; Kong, C.Y.; Cao, L.; Thorne, T.J.; Sun, Y.-P. Graphene oxides for homogeneous dispersion of carbon nanotubes. ACS Appl. Mater. Interfaces 2010, 2, 3217–3222. [CrossRef] [PubMed]

18. Pant, B.; Park, M.; Park, S.-J.; Kim, H.-Y. One-pot synthesis of CdS sensitized TiO2 nanorods. J. Appl. Polym. Sci. 2014, 132, 415–416, 801–809. [CrossRef]

19. Xu, L.; Huang, W.-Q.; Wang, L.-L.; Huang, G.-F. Interfacial Interactions of Semiconductor with Graphene and Reduced Graphene Oxide: CeO2 as a Case Study. ACS Appl. Mater. Interfaces 2014, 6, 20350–20357. [CrossRef] [PubMed]

20. Ahmed, D.F.; Isawi, H.; Badway, N.A.; Elbayaa, A.A.; Shawky, H. Graphene oxide incorporated cellulose triacetate/cellulose acetate nanocomposite membranes for forward osmosis desalination. Arab. J. Chem. 2021, 14, 102995. [CrossRef]

21. Bhattacharya, M. Polymer nanocomposites-A comparison between carbon nanotubes, graphene, and clay as nanofillers. Materials 2016, 9, 262. [CrossRef]

22. Cui, Y.; Kundalwal, S.J.; Kumar, S. Gas barrier performance of graphene/polymer nanocomposites. Carbon 2016, 98, 313–333. [CrossRef]

23. Ojha, G.P.; Pant, B.; Park, S.-J.; Park, M.; Kim, H.-Y. Synthesis and characterization of reduced graphene oxide decorated with CeO2-doped MnO2 nanorods for supercapacitor applications. J. Colloid Interface Sci. 2017, 494, 338–344. [CrossRef]

24. Trovarelli, A. Catalytic properties of ceria and CeO2-containing materials. Catal. Rev. 1996, 38, 439–520. [CrossRef]

25. Regmi, C.; Ashtiani, S.; Sofer, Z.; Horlicka, Z.; Prusa, F.; Vopička, O.; Friess, K. CeO2-blended cellulose triacetate mixed-matrix membranes for selective CO2 separation. Membranes 2021, 11, 632. [CrossRef] [PubMed]

26. Shojaie, S.S.; Krantz, W.B.; Greenberg, A.R. Dense polymer film and membrane formation via the dry-cast process part I. Model development. J. Membr. Sci. 1994, 94, 251–280. [CrossRef]

27. Macchione, M.; Jansen, J.C.; DiRoi, E. The dry phase inversion technique as a tool to produce highly efficient asymmetric gas separation membranes of modified PEEK. Influence of temperature and air circulation. Desalination 2006, 192, 132–141. [CrossRef]

28. Friess, K.; Hynek, V.; Šipek, M.; Kujawski, W.M.; Vopička, O.; Zgažar, M.; Kujawski, M.W. Permeation and sorption properties of poly(ether-block-amide) membranes filled by two types of zeolites. Sep. Purif. Technol. 2011, 80, 418–427. [CrossRef]

29. Vopička, O.; Friess, K.; Hynek, V.; Šysel, P.; Zgažar, M.; Šipek, M.; Pilnáček, K.; Lanč, M.; Jansen, J.C.; Mason, C.R.; et al. Equilibrium and transient sorption of vapours and gases in the polymer of intrinsic microporosity PIM-1. J. Membr. Sci. 2013, 434, 148–160. [CrossRef]

30. Jansen, J.C.; Friess, K.; DiRoi, E. Organic vapour transport in glassy perfluoropolymer membranes: A simple semi-quantitative approach to analyze clustering phenomena by time lag measurements. J. Membr. Sci. 2011, 367, 141–151. [CrossRef]

31. Friess, K.; Jansen, J.C.; Bazzarelli, F.; Izák, P.; Jarмаровá, V.; Kačirková, M.; Schauer, J.; Clarizia, G.; Bernardo, P. High ionic liquid content polymeric membrane interfaces: Correlation of membrane structure with gas and vapour transport properties. J. Membr. Sci. 2012, 415–416, 801–809. [CrossRef]

32. Joung, D.; Singh, V.; Park, S.; Schulte, A.; Seal, S.; Khondaker, S.I. Anchoring ceria nanoparticles on reduced graphene oxide and their electronic transport properties. J. Phys. C. 2011, 115, 24494–24500. [CrossRef]

33. Wang, Z.; Zhao, P.; He, D.; Cheng, Y.; Liao, L.; Li, S.; Luo, Y.; Peng, Z.; Li, P. Cerium oxide immobilized reduced graphene oxide hybrids with excellent microwave absorbing performance. Phys. Chem. Chem. Phys. 2018, 20, 14155–14165. [CrossRef]

34. Rajendran, R.; Shrestha, L.K.; Minami, K.; Subramanian, M.; Jayavel, R.; Ariga, K. Dimensionally integrated nanorocheticonics for a novel composite from 0D, 1D, and 2D nanomaterials: RGO/CNT/CeO2 ternary nanocomposites with electrochemical performance. J. Mater. Chem. A 2014, 2, 18480–18487. [CrossRef]

35. Lee, J.M.; Nguyen, D.Q.; Lee, S.B.; Kim, H.; Ahn, B.S.; Lee, H.; Kim, H.S. Cellulose triacetate-based polymer gel electrolytes. J. Appl. Polym. Sci. 2010, 115, 32–36. [CrossRef]

36. El Nemr, A.; Ragab, S.; El Sikaily, A.; Khaled, A. Synthesis of cellulose triacetate from cotton cellulose by using NIS as a catalyst under mild reaction conditions. Carbohydr. Polym. 2015, 130, 41–48. [CrossRef]

37. Sasikumar, B.; Arthanareswaran, G.; Sankaranarayanan, K.; Jeyadeepan, K. Synthesis and formation of phase-tuned TiO2- /ionic liquid-incorporated polymer membranes for ammonia sensing at room temperature. ACS Sustain. Chem. Eng. 2019, 7, 15884–15895. [CrossRef]
38. Fan, Y.; Li, C.; Zhang, X.; Yang, X.; Su, X.; Ye, H.; Li, N. Tröger’s base mixed matrix membranes for gas separation incorporating NH2-MIL-53(Al) nanocrystals. *J. Membr. Sci.* 2019, 573, 359–369. [CrossRef]
39. Langmuir, I. The adsorption of gases on plain surfaces of glass, mica and platinum. *J. Am. Chem. Soc.* 1918, 40, 1361–1403. [CrossRef]
40. Cheng, Y.; Wang, Z.; Zhao, D. Mixed matrix membranes for natural gas upgrading: Current status and opportunities. *Ind. Eng. Chem. Res.* 2018, 57, 4139–4169. [CrossRef]
41. Ehsani, M.; Rahimi, P.; Joseph, Y. Structure-function relationships of nanocarbon/polymer composites for chemiresistive sensing: A review. *Sensors* 2021, 21, 3291. [CrossRef]
42. Wu, X.; Tian, Z.; Wang, S.; Peng, D.; Yang, L.; Wu, Y.; Xin, Q.; Wu, H.; Jiang, Z. Mixed matrix membranes comprising polymers of intrinsic microporosity and covalent organic framework for gas separation. *J. Membr. Sci.* 2017, 528, 273–283. [CrossRef]
43. Nikolaeva, D.; Azcune, I.; Tanczyk, M.; Warmuzinski, K.; Jaschik, M.; Sandru, M.; Dahl, P.I.; Genua, A.; Lois, S.; Sheridan, E.; et al. The performance of affordable and stable cellulose-based poly-ionic membranes in CO2/N2 and CO2/CH4 gas separation. *J. Membr. Sci.* 2018, 564, 552–561. [CrossRef]
44. Čihal, P.; Vopička, O.; Lanč, M.; Kludský, M.; Velas, J.; Hrdlička, Z.; Michalcová, A.; Dendisová, M.; Friess, K. Poly(butylene succinate)-cellulose triacetate blends: Permeation, pervaporation, sorption and physical structure. *Polym. Test.* 2018, 65, 468–479. [CrossRef]
45. Ashtiani, S.; Khoshnamvand, M.; Regmi, C.; Friess, K. Interfacial Design of Mixed Matrix Membranes via Grafting PVA on Uio-66-NH2 to Enhance the Gas Separation Performance. *Membranes* 2021, 11, 419. [CrossRef]