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Enhanced N-doped Porous Carbon Derived from KOH-Activated Waste Wool: A Promising Material for Selective Adsorption of CO$_2$/CH$_4$ and CH$_4$/N$_2$

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Received: 15 January 2019; Accepted: 12 February 2019; Published: 15 February 2019

Abstract: Separation of impurities (CO$_2$ and N$_2$) from CH$_4$ is an important issue for natural gas alternatives (such as coalbed gas, biogas, and landfill gas) upgrading. It is notably challenging to synthesize high N-doped porous carbon with an appropriate porous structure. In this work, high N content (14.48 wt %) porous carbon with micropore size of 0.52 and 1.2 nm and specific surface area of 862 m$^2$ g$^{-1}$ has been synthesized from potassium hydroxide (KOH) activated waste wool upon the urea modification. Pure component adsorption isotherms of CO$_2$, CH$_4$, and N$_2$ are systematically measured on this enhanced N-doped porous carbon at 0 and 25°C, up to 1 bar, to evaluate the gases adsorption capability, and correlated with the Langmuir model. These data are used to estimate the separation selectivities for binary mixtures of CO$_2$/CH$_4$ and CH$_4$/N$_2$ at different mixing ratios according to the ideal adsorbed solution theory (IAST) model. At an ambient condition of 25°C and 1 bar, the predicted selectivities for equimolar CO$_2$/CH$_4$ and CH$_4$/N$_2$ are 3.19 and 7.62, respectively, and the adsorption capacities for CO$_2$, CH$_4$, and N$_2$ are 2.91, 1.01, and 0.13 mmol g$^{-1}$, respectively. This report introduces a simple pathway to obtain enhanced N-doped porous carbon with large adsorption capacities for gas separation of CO$_2$/CH$_4$ and CH$_4$/N$_2$.

Keywords: enhanced N-doped; porous carbon; absorbent; CO$_2$/CH$_4$ and CH$_4$/N$_2$; selectivity

1. Introduction

In recent decades, to satisfy the rapid growth in chemical and manufacturing industries, a large amount of fossil fuels burning has increased the level of carbon dioxide (CO$_2$) in the atmosphere, which causes serious environmental issues, such as global climate change, the greenhouse effect, and ocean acidification [1–3]. A number of methods to reduce the CO$_2$ gas have been suggested. One of the potential methods might be the use of natural gas (NG) as fuel for transportation and heating purpose [3]. Compared with the fossil fuels such as coal and petroleum, NG produces less CO$_2$ per energy unit and has a high specific combustion enthalpy; therefore, it is considered a cleaner and potential substitute energy [4–6]. Additionally, NG has no threat to water or land, because of its non-toxic and no vapor pool forming on the ground as the liquefied petroleum gas does [3]. Nowadays, NG has provided a large amount of energy to satisfy business, industry, vehicle, and home needs throughout...
the world. The consumption of NG is expected to grow by 50% in the next 20 years, which makes the energy shortage situation much more serious [7]. As a result, coalbed gas, biogas, and landfill gas are deemed as attractive alternatives to NG, because they possess high methane (CH₄) content and obvious socioeconomic advantages. However, the existence of some impurities such as CO₂ and nitrogen (N₂) will hamper the application of these alternative gases by reducing the heating value and corroding the transportation or storage system [7–9]. Consequently, urgent research is required to separate impurities (CO₂ and N₂) from CH₄. At present, various technologies have been developed for gas separation and purification including cryogenic distillation, membrane separation, and pressure swing adsorption (PSA) [6,10]. Among these technologies, PSA has attracted intense interest due to its advantage of low cost, ease control, and energy effectiveness [6]. Accordingly, a lot of adsorbents have been evaluated for the adsorptive separation of CO₂/CH₄ and CH₄/N₂ such as metal–organic frameworks (MOFs) [11–14], organic zeolites [14–16], organic frameworks [17], and carbon-based materials [6–9,18,19].

Porous carbons have been considered as the most promising candidates for gas storage and separation because they are easy to synthesize and have good economical properties and strong thermal stability [20–22]. The adsorption of CO₂, CH₄, and N₂ could be described as physisorption through weak van der Waals interaction forces [22]. Current research has focused on increasing the specific surface area (SSA) and controlling the pore size distribution (PSD), since the adsorption capacity is closely related to the textural characteristics of the carbon materials [22–24]. For example, Leventis et al. [25,26] reported that microporous carbon aerogels demonstrated an exceptionally high adsorption capacity for CO₂. Meanwhile, for the purpose of improving the adsorption efficiency and further enhancing the interactions between gas molecules and carbon material, it is important to incorporate heteroatoms (i.e., N, P, S) into the carbon framework [21,22,25–28]. Recently, many efforts have been devoted to the synthesis of robust porous carbon that possesses high adsorption capacity and good selectivity, especially the N-doped porous carbon adsorbent. For example, the N-doped porous carbon synthesized from benzimidazole-linked polymers demonstrated high adsorption selectivity for CO₂/CH₄ and CO₂/N₂ [21]. The N-rich porous carbon through polyamine-incorporated metal–organic framework materials, which exhibited excellent CO₂-selective adsorption ability over CH₄ and N₂ [27]. The N-doped graphene sheets by reducing graphene oxide sheets in the presence of polyindole displayed outstanding adsorption selectivity ratios for CO₂ over N₂, CH₄, and H₂, respectively [28]. However, the synthesis processes of these N-doped porous carbons are very complex and time-consuming, and the precursor is not economical, which will enormously influence the practical application for the adsorbents. Thus, there still remains a challenge to develop porous carbon adsorbents with strong adsorption capacity, high selectivity, facile synthesis, and low cost.

Porous carbon adsorbents obtained from biomass materials are very competitive, which are not only cost-effective, easy to synthesis, abundant and sustainably renewable, but also happen to be excellent adsorbents for gas adsorption and separation [29–32]. So far, there have already been some biomass-derived porous carbons reported for CO₂/CH₄/N₂ adsorption and separation. For example, Li et al. [31] investigated starch-derived porous carbon spheres for high uptakes of CO₂, CH₄, and H₂. Niya et al. [32] synthesized N-doped carbon foams for CO₂/N₂, CO₂/CH₄, and CH₄/N₂ separation by using banana peel as a precursor and zinc complex as a template. Because of the N-rich nature, waste wool is a desirable precursor for the fabrication of N-doped porous carbon [33]. In a previous study, we have successfully synthesized N-doped hierarchical porous carbon from waste wool by potassium hydroxide (KOH) activation, exhibiting a good selective adsorption capability of CO₂/N₂ [34]. Although KOH activation is a well method for developing the porosity structure of the carbon, it unavoidably leads to a loss of N from the carbon framework [33–35].

Therefore, enhancing the N content of the porous carbon after KOH activation is an urgent task for further gas selective adsorption application. Keeping the above purpose in mind, in this work we propose a simple method to synthesize N-enhanced porous carbon from KOH activated waste wool for selective adsorption of CO₂/CH₄ and CH₄/N₂. As shown in Figure 1, first waste wool is chosen as carbon precursor to be activated by KOH, and then modified by urea. Here, waste wool is chosen
because it is largely composed of keratin amino acids and is a unique renewable natural material with high N contents. However, the KOH activation can make a significant evolution of the structure for the original wool derived activated porous carbon (WAPC), which simultaneously reduces the N groups and introduces many O groups into the carbon frameworks [34–36]. Upon the urea modification, a significant amount of N groups can be again introduced into the activated carbon matrix, because the high N content (46 wt %) urea can react with O-containing functional groups [37]. The CO2, CH4, and N2 adsorption properties are evaluated at 0 °C and 25 °C, respectively, on the as-obtained enhanced N-doped wool derived activated porous carbon (N-WAPC) The isotherm data are used to predict the separation selectivities for binary mixtures of CO2/CH4 and CH4/N2 assuming different mixing ratios by the ideal adsorption solution theory (IAST).

![Figure 1. Schematic illustration of the synthesis of enhanced N-doped porous carbon.](image)

2. Experimental Section

2.1. Materials

Waste wool was obtained from rural pastures of Henan, China. Potassium hydroxide (KOH), hydrochloric acid (HCl), and urea (NH2CONH2) were purchased from Sinopharm Chemical Reagent Co. Ltd. China (Beijing, China) and used as received without any further purification.

2.2. Sample Preparation

First, the waste wool was thoroughly washed, and then pre-carbonized at 300 °C under N2 atmosphere. The as-obtained black and crisp carbide was ground to powder and mixed with a KOH agent (the ratio of KOH/pre-carbonized sample = 3), activated under N2 atmosphere as we have previously reported [34]. The resulting waste wool derived activated porous carbon was denoted as WAPC. In the following synthesis, 0.1 g of WAPC and 0.1 g of urea were thoroughly mixed by grinding for 20 min. After that, the mixture was transferred into a quartz boat placing in a horizontal tube furnace. Subsequently, the mixture was calcinated at 600 °C for 1 h with a heating rate of 3 °C min−1 under N2 atmosphere. After cooling down, the sample was thoroughly washed with hot distilled water to remove the residual urea, and then dried at 60 °C to obtain the final product. The as-obtained N-enhanced porous carbon sample was denoted as N-WAPC.

2.3. Material Characterizations

Scanning electron microscopy (SEM) images and elemental mappings were obtained by field emission scanning electron microscopy (FE-SEM, Hitachi S-4800, Hitachi, Tokyo, Japan) with an energy dispersive X-ray spectrometer (EDS) (Hitachi, Tokyo, Japan). Transmission electron microscopy (TEM) and high-resolution TEM (HR-TEM) measurements were carried out on a JEOL-2100F transmission electron microscope (JEOL, Tokyo, Japan) operated at 200 KV. Powdered X-ray diffraction (XRD)
patterns were performed on a Bruker-D8 powder diffractometer (Bruker, Madison, WI, USA) using Cu Kα radiation (λ = 0.15406 nm). Raman spectra were collected on a Renishaw inVia (Renishaw, London, UK) with an excitation wavelength of 514 nm. Fourier transform infrared (FT-IR) spectra were collected on a Nicolet iS10 FT-IR spectrophotometer (Nicolet, Madison, WI, USA), using the KBr pellet technique. X-ray photoelectron spectroscopy (XPS) was performed on a Thermo Scientific ESCALAB250 (Thermo Fisher Scientific, Waltham, MA, USA) equipped with an Al Kα excitation source. Elemental analysis (C, H, and N) was performed on a dry basis using a VarioEL III Elemental Analyzer (Elementar, Hanau, Germany). The N2 adsorption–desorption isotherms were measured on a Quantachrome NOVA1000e volumetrically sorption analyzer (Quantachrome, Boynton Beach, FL, USA) at liquid nitrogen temperature. Prior to each adsorption experiment, the degassing procedure was carried out for 3 h at 300 °C to remove the guest molecules in the sample. The SSA was calculated by multiple points Brunauer–Emmett–Teller (BET) method and PSD was determined by a non-local density functional (NLDFT) theory using nitrogen adsorption data and assuming a slit pore model.

2.4. Adsorption Measurements

The CO2, CH4, and N2 adsorption isotherms were measured by the Micromeritics ASAP 2020 volumetrically at two temperatures of 0 °C and 25 °C, respectively. Ultrahigh-purity CO2, CH4, and N2 were used as received. As aforementioned, the degassing procedure was also carried out for 3 h at 300 °C prior to the adsorption measurement.

3. Results and Discussions

3.1. Characterizations of the As-Prepared Samples

The SEM images of the pre-carbonized sample show a bulky morphology with a smooth surface, and no pores can be observed (Figure S1). After the KOH activation process, the activated sample WAPC exhibits an obviously developed porous structure with spongy pores on the surface (Figure 2a). Further urea nitridation, the structure of the N-enhanced sample N-WAPC becomes more or less collapsed; however, the whole sponge-like porous structure still can be well kept (Figure 2b). TEM and HR-TEM measurements are also carried out to study the overall morphology and microstructure of the N-WAPC. As showed in Figure 2c, a continuous sponge-like architecture with a connected porous texture is clearly observed. Furthermore, the HR-TEM image of N-WAPC (Figure 2d) shows an amorphous curved surface with worm-like micropores, as well as the selected area electron diffraction (SAED) pattern exhibiting typical diffuse rings (inset in Figure 2d).

![Figure 2](image-url) SEM images of (a) wool derived activated porous carbon (WAPC), (b) N-doped wool derived activated porous carbon (N-WAPC); TEM image (c,d) HRTEM image of N-WAPC (inset: SAED).
XRD and Raman are employed to characterize the phase structures of the porous carbon samples. As shown in Figure 3a, the two broad low-intensity diffraction peaks around 25° and 43° correspond to the (002) and (100) diffraction patterns of amorphous graphitic carbon [38–40]. The weak peaks suggest a low graphitization of the WAPC. After urea nitridation treatment, these two peaks become much weaker, indicating a more disordered graphitic structure of N-WAPC. In Figure 3b, the two bands centered at about 1340 cm\(^{-1}\) (D band) and 1590 cm\(^{-1}\) (G band) are separated completely. The D band is closely related to the presence of disordered carbon structures, whereas the G band is associated with the sp\(^2\)-hybridized carbon atoms in a graphitic layer [38–40]. The intensity ratio of the D and G band (ID/IG) is known to be an indicator of the degree of graphitization, the higher the ID/IG, the lower the degree of graphitization [39]. In our case, the ID/IG peak ratios for WAPC and N-WAPC are 1.12 and 2.71, respectively, indicating a low degree of graphitization. The ID/IG ratio of N-WAPC is higher than that of WAPC, suggesting more defect sites are introduced in N-WAPC during the urea nitridation, which corresponds with the XRD result.

The corresponding N species are presented in Figure 4b. The peaks at ~398.1 eV, ~400.0 eV, ~400.7 eV, and ~402.8 eV are attributed to pyridinic-N (N-6), pyrrolic-/pyridonic-N (N-5), quaternary-N (N-Q), and N-oxides of pyridine (N-x), respectively [43,44,46–48]. The structure of these different N species can be assigned to the in-plane deformation vibration or C=C stretching vibration in aromatic rings [42,43]. The characteristic N–H and/or O–H stretching vibrations [41–43]. The band at about 1580 cm\(^{-1}\) is ascribed to the KOH activation treatment [34]. The high level of oxygen-containing functional groups be associated with the KOH activation treatment [34]. The high level of oxygen-containing functional groups in the WAPC makes it very reactive for subsequent urea nitridation [37]. After the urea treatment, the N contents increase from 4.14 wt % for WAPC to 14.48 wt % for N-WAPC. On the contrary, the O contents decrease from 24.79 wt % for WAPC to 12.44 wt % for N-WAPC. While the C contents for the two samples do not change significantly. Above results demonstrate that the oxygen-containing groups on the WAPC can interact with the amino groups in the urea, contributing to the incorporation of N into the carbon framework.

In order to further investigate the feature of surface states, the XPS technique is applied for the carbon samples. As shown in Figure 4a, both samples exhibit C1s, N1s, and O1s peaks, respectively. It is noteworthy that the N-WAPC has a considerably strong N1s peak compared with WAPC (Figure 4a). The corresponding N species are presented in Figure 4b. The peaks at ~398.1 eV, ~400.0 eV, ~400.7 eV, and ~402.8 eV are attributed to pyridinic-N (N-6), pyrrolic-/pyridonic-N (N-5), quaternary-N (N-Q), and N-oxides of pyridine (N-x), respectively [43,44,46–48]. The structure of these different N species

![Figure 3. Powder XRD patterns (a), Raman spectra (b), and FT-IR spectra of WAPC and N-WAPC (c).](image-url)
is shown in Figure 4c. To the best of our knowledge, the pyrrolic-N and pyridonic-N cannot be well distinguished from each other within the accuracy of XPS measurements. However, considering the oxygen presenting in the porous carbons and the fact that pyridonic-N is more stable than pyrrolic-N, the pyridonic-N is more likely to be present in the porous carbons [42]. As to pyridonic-N, the p orbital in its –OH would produce p–π conjugation effect with its p bond, which endows the N located in the ortho-position of –OH with stronger electron atmosphere and consequently enhances its Lewis basicity, resulting in the improvement of the CO2 adsorption ability of porous carbons [42]. Thus, the pyridonic-N, acting as an anchor for CO2 capture, is beneficial for CO2 capture [42,43]. Whereas, other N species have no active effect for CO2 capture [46]. Quantitative analysis calculations from the N1s peak area show that the N amounts of form N-5 is higher than those present of form N-6, N-Q and N-x, in the order of N-5 > N-6 > N-Q > N-x for both samples of WAPC and N-WAPC (Table 1). It should be noted that after urea nitridation treatment, the percentage of form N-5 in N-WAPC increases significantly. The relative percentage of N-5 in WAPC is only 33%, while in N-WAPC it is 44% (Table 1). This would be favorable for CO2 capture, because N-5 contributes more significantly to capturing CO2 than other forms of nitrogen species [46]. The high-resolution spectra of C1s and O1s for WAPC and N-WAPC are shown in Figure S2. The C1s peak can be deconvoluted to six sub-peaks at 284.5, 285.3, 285.9, 286.7, 287.3, and 288.8 eV, respectively (Figure S2a). The main peak centers at 284.5 eV corresponds to C-C [49], while the other weak peaks can be identified as C-OH at 285.3 eV [36,49], C=N at 285.9 eV [49], C-O-C at 286.4 eV [48], N-C=O at 287.4 eV [49], and -COOH at 288.8 eV [48]. Three characteristic peaks are observed at 531.8, 532.5, and 533.4 eV in the O 1s spectra (Figure S2b), which can be ascribed to C=O, C-OH, and O-C=O, respectively [48–50]. Furthermore, from the SEM energy dispersive X-ray spectroscopy (EDS) element mapping, it can be seen that N, C, and O elements are uniformly distributed on the N-WAPC surface (Figure S3).

**Table 1.** Elemental composition and relative percentages of nitrogen species of the carbon samples.

<table>
<thead>
<tr>
<th>Sample</th>
<th>Elemental Composition (wt %)</th>
<th>Relative Percentages of Nitrogen Species [%]</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>C</td>
<td>N</td>
</tr>
<tr>
<td>WAPC</td>
<td>69.65</td>
<td>4.14</td>
</tr>
<tr>
<td>N-WAPC</td>
<td>71.98</td>
<td>14.48</td>
</tr>
</tbody>
</table>

[a] Calculated by difference. [b] Calculated from the corresponding peak areas of the deconvoluted N1s spectra.

**Figure 4.** XPS for WAPC and N-WAPC (a) full-survey spectra and (b) high-resolution spectra of N1s; (c) the different N species possibly present in the porous carbons.
N₂ adsorption–desorption isotherms of WAPC and N-WAPC and the corresponding PSD curves are shown in Figure 5. The isotherms of WAPC and N-WAPC (Figure 5a) display a combination form of type I and type IV, with a rapid N₂ uptake at very low relative pressure (P/P₀) and a slight hysteresis loop at medium P/P₀ range, indicating the presence of rich micropores and a small number of mesopores in the samples [51]. However, the N₂ uptake amount at low P/P₀ region for N-WAPC is obviously less than that of WAPC, which can be attributed to the structural collapse that resulted from urea nitridation. Correspondingly, the PSD curves of WAPC and N-WAPC calculated by the NLDFT theory both exhibit the presence of micropores and mesopores (Figure 5b). The PSD curve for WAPC displays three relatively strong micropore peaks at around 0.52, 0.85, and 1.2 nm, respectively, and one very weak mesopore peak at about 2.6 nm. As can be seen clearly, the peaks intensities of micropores and mesopores for N-WAPC are much inferior to that of N-WAPC, which is in accordance with their N₂ adsorption isotherms and microstructure results. As shown in Table 2, which includes BET SSA (S_BET) that calculated from N₂ adsorption branch and total pore volume (V_{tot}) that obtained from N₂ adsorption amount at P/P₀ = 0.99, respectively. The S_BET for WAPC and N-WAPC is 1352 and 862 m² g⁻¹, respectively; and the V_{tot} for WAPC and N-WAPC is 0.74 and 0.5 m³ g⁻¹, respectively. We note that both the S_BET and the V_{tot} significantly decrease after nitridation treatment with urea, because the structure collapse and addition N-groups may block partial pore volume.

![Figure 5](image_url)

**Figure 5.** (a) N₂ adsorption–desorption isotherms and (b) pore size distribution (PSD) curves of WAPC and N-WAPC.

### 3.2. Gas Adsorption and Separation Studies

The pure component adsorption isotherms of CO₂, CH₄, and N₂ for WAPC and N-WAPC are measured at two temperatures (25 °C and 0 °C) and pressure up to 1 bar, as shown in Figure 6 and summarized in Table 2. All the isotherms show excellent reversibility without hysteresis, suggesting that the adsorbed molecules can be well removed during the desorption process. Thus, the samples of WAPC and N-WAPC can be easily regenerated by a vacuum without any heating treatment, which makes these samples superior to some MOFs and zeolites materials [6]. The adsorption isotherms for CO₂ are fitted by the Double Site Langmuir (DSL) model and the adsorption isotherms for CH₄ and N₂ are fitted by the Langmuir (L) model. The calculation details and parameters are summarized in Figure S4–S9 and Table S1–S3 in the supporting information. It is clear that the amounts of gases adsorbed varied widely for samples N-WAPC and WAPC, in the sequence of CO₂ > CH₄ > N₂. In both WAPC and N-WAPC, CO₂ is the most strongly adsorbed gas owing to its significant quadrupolar moment and polarizability [6,9,52]. CH₄ shows stronger adsorption than N₂, which is attributed to its higher polarizability than that of N₂ [6,9,52].
The adsorption capacity is an important factor to assess the gas separation capability for an adsorbent [6]. The higher CO$_2$ capacity is obtained by N-WAPC, which is measured to be 4.50 and 2.91 mmol g$^{-1}$ at 0 and 25 °C, respectively. These values are superior to some recently reported carbon-based adsorbents: sOMC (2.0 mmol g$^{-1}$), MAC (2.13 mmol g$^{-1}$), and N-HPMCs-5-0.6-973 (1.82 mmol g$^{-1}$) at 25 ºC and 100 kPa [6,7,53]. Although the $S_{\text{BET}}$ and $V_{\text{tot}}$ for the WAPC are high, the CO$_2$ capacity for WAPC (only 3.73 and 2.81 mmol g$^{-1}$ at 0 and 25 ºC, respectively) is much lower than that of N-WAPC. This can be clarified by the significantly enhanced N content in N-WAPC (14.48 wt %), especially the high percentage of form N-5 in N-WAPC (44%). Furthermore, the N-WAPC also exhibits high CH$_4$ adsorption capacities of 1.70 and 1.01 mmol g$^{-1}$ at 0 and 25 ºC, respectively, at the pressure of 1 bar. As for the N$_2$ uptake, the N-WAPC shows adsorption capacities of 0.67 and 0.13 mmol g$^{-1}$ at 0 and 25 ºC, respectively, under the pressure of 1 bar. Under the same conditions, all these values outperform the WAPC adsorption capacities on CH$_4$ (1.18 mmol g$^{-1}$ at 0 ºC and 0.64 mmol g$^{-1}$ at 25 ºC) and N$_2$ (0.37 mmol g$^{-1}$ at 0 ºC and 0.35 mmol g$^{-1}$ at 25 ºC).

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In addition to the desirable gas uptake, selectivity is also very important for potential application in gas separation. Considering the good gas adsorption capacity of N-WAPC, the gas separation capability of N-WAPC for CO\textsubscript{2}/CH\textsubscript{4} and CH\textsubscript{4}/N\textsubscript{2} are further measured. The IAST is a widely used method to predict the binary mixture gas adsorption selectivity behavior based upon single component gas adsorption results \cite{6,9,34}. Here, we use IAST to predict the selectivity of N-WAPC for the binary mixtures (CO\textsubscript{2}/CH\textsubscript{4} and CH\textsubscript{4}/N\textsubscript{2}). In a mixture, the selectivity of the preferential adsorption gas 1 over gas 2 can be formally defined as the following equation:

\[
S_{\text{ads}} = \frac{q_1}{q_2} \frac{p_2}{p_1}
\]  

(1)

where \(S_{\text{ads}}\) is the selectivity factor, \(q_1\) and \(q_2\) are the absolute adsorbed loadings at a partial pressure of \(p_1\) and \(p_2\), respectively, in the binary mixture. The IAST selectivities of the binary mixtures at different gas proportions on N-WAPC in this paper are typical components of CH\textsubscript{4}-related gases, such as coalbed gas, biogas, and landfill gas. Accordingly, the mixing ratios of 50/50, 15/85, and 5/95 for CO\textsubscript{2}/CH\textsubscript{4}, and CH\textsubscript{4}/N\textsubscript{2} are used to calculate the selectivity and the overall performance is summarized in Table 3.

**Table 3.** The selectivity of N-WAPC for binary mixtures \(^{[a]}\).

<table>
<thead>
<tr>
<th>Temp.</th>
<th>CO\textsubscript{2}/CH\textsubscript{4}</th>
<th>CH\textsubscript{4}/N\textsubscript{2}</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>50/50</td>
<td>15/85</td>
</tr>
<tr>
<td>0 °C</td>
<td>3.03</td>
<td>6.91</td>
</tr>
<tr>
<td>25 °C</td>
<td>3.19</td>
<td>5.98</td>
</tr>
</tbody>
</table>

\(^{[a]}\) Calculated from IAST(0 and 25 °C) for the binary mixtures assuming different mixing ratios at 1 bar.

The selectivities of binary mixtures for CO\textsubscript{2}/CH\textsubscript{4} and CH\textsubscript{4}/N\textsubscript{2} on N-WAPC, are plotted as a function of total bulk pressure in Figure 7. For equimolar binary mixture (50/50) of CO\textsubscript{2}/CH\textsubscript{4}, the selectivities decrease with the increased pressure, obtaining about 3.03 (0 °C) and 3.19 (25 °C) at 1 bar (Figure 7a). The CH\textsubscript{4}/CO\textsubscript{2} selectivity displayed by N-WAPC is comparable to or higher than some other reported adsorbents \cite{6,54–56}. When it comes to the CH\textsubscript{4}/N\textsubscript{2} (50/50) separation, the selectivities are also decreased with the increased pressure and obtained 3.04 (0 °C) and 7.62 (25 °C) at 1 bar (Figure 7b), which is comparable to that of ultra-microporous carbon \cite{57} and polycarbazole polymers under the same condition \cite{58}. The selectivities for the binary mixture CO\textsubscript{2}/CH\textsubscript{4} and CH\textsubscript{4}/N\textsubscript{2} at other mixing ratios are further predicted by IAST. As shown in Figure 7c, with the mixture ratio of 15/85 for CO\textsubscript{2}/CH\textsubscript{4}, although the selectivities are decreased with the increased pressure, it can maintain a relatively high level of 6.91 (0 °C) and 5.98 (25 °C) at 1 bar. Whereas in the mixture ratio for CO\textsubscript{2}/CH\textsubscript{4} is 5/95, the selectivity situation is different. As shown in Figure 7e, the CO\textsubscript{2}/CH\textsubscript{4} selectivities gradually increase with the increased pressure, finally reaching 12.26 (0 °C) and 8.66 (25 °C) at 1 bar. As for the binary mixture of CH\textsubscript{4}/N\textsubscript{2}, when the gas component is 15/85, the CH\textsubscript{4}/N\textsubscript{2} selectivity slightly decreases with the increased pressure, keeping 4.66 at 0 °C and 1 bar (Figure 7d). However, at 25 °C, the CH\textsubscript{4}/N\textsubscript{2} selectivity obviously increases with the increased pressure, reaching about 11.76 under the pressure of 1 bar (Figure 7d). For the case of binary mixture CH\textsubscript{4}/N\textsubscript{2} with mixing ratio of 5/95, the selectivities increase with the increased pressure, reaching 5.73 (25 °C) and 13.71 (0 °C) at 1 bar (Figure 7f). It is worth noting that, in the same conditions, the selectivities of CO\textsubscript{2}/CH\textsubscript{4} and CH\textsubscript{4}/N\textsubscript{2} keep increasing in the composition range (50/50 ~ 15/85 ~ 5/95), which is an attractive feature of an adsorbent. The listed selectivities indicate that the as-synthesized N-WAPC is a promising adsorbent in gas adsorptive separation.
Figure 7. IAST-predicted selectivities assuming different binary mixing ratios on N-WAPC
(a) CO₂/CH₄ = 50/50, (b) CH₄/N₂ = 50/50, (c) CO₂/CH₄ = 15/85, (d) CH₄/N₂ = 15/85, (e) CO₂/CH₄ = 5/95, and (f) CH₄/N₂ = 5/95 at 0 and 25 °C.

The isosteric heat of adsorption (Q_{st}) is the standard enthalpy of adsorption at a fixed surface coverage [59], which is an important value to evaluate the interactions between gas molecules and adsorbents and further elucidates the selectivity. The Q_{st} for CO₂ and CH₄ on N-WAPC are calculated by obtaining the CO₂ and CH₄ isotherms at two different temperatures (25 and 0 °C). The single component Q_{st} as a function of surface loading can be determined by a variant of the Clausius–Clapeyron equation as following [59–61]:

$$\ln(P_1/P_2) = \Delta H_{ads} \left(\frac{T_2}{R \cdot T_1} - \frac{T_1}{T_2}\right)$$  \hspace{1cm} (2)

where \(\Delta H_{ads}\) gives the isosteric heat of adsorption. \(P_1\) and \(P_2\) are the pressures, for the same amount of gas uptake, at two different temperatures \(T_1\) and \(T_2\) respectively. \(R\) is the ideal gas constant. As can be seen clearly in Figure 8, the Q_{st} values of CO₂ is in the range of 35.44–27.10 kJ mol⁻¹ gently dropping from low coverage to saturation, which means that the interaction between CO₂ molecules and the absorbent is stronger than that between CO₂ molecules [58]. The Q_{st} of CH₄ is almost in
the horizontal range from 22.59 to 22.88 kJ mol\(^{-1}\), which apparently exceeds standard enthalpy for interaction of CH\(_4\) with carbon (16 kJ mol\(^{-1}\)) and many activated carbon materials, such as M1273-150 (17–11 kJ mol\(^{-1}\)) [31], C1000 (17.4–16.3 kJ mol\(^{-1}\)) [62], K-PAF-1 (20.6–15.9 kJ mol\(^{-1}\)) [63]. The enhanced N-doped porous carbon with appropriate pore structure can greatly enhance CH\(_4\) adsorption through weak interactions, and can therefore achieve a high CH\(_4\)/N\(_2\) selectivity. Furthermore, the Q\(_{st}\) of CO\(_2\) is obviously higher than that of CH\(_4\) within the experiment range, indicating a high CO\(_2\)/CH\(_4\) selectivity. These high selectivities demonstrate that the enhanced N-doped porous carbon has potential in applications for CH\(_4\)-related gas upgrading, such as coalbed gas, biogas, and landfill gas.

4. Conclusions

Enhanced N-doped porous carbon with an appropriate pore structure has been synthesized from KOH activated waste wool upon the urea modification. The as-obtained porous carbon N-WAPC has a high N content of 14.48 wt%, SSA of 862 m\(^2\) g\(^{-1}\), and an appropriate micropore size of 0.52 and 1.2 nm. The gas the adsorption capacities of N-WAPC follows the order CO\(_2\) > CH\(_4\) > N\(_2\). The adsorption selectivities of binary mixtures CO\(_2\)/CH\(_4\) and CH\(_4\)/N\(_2\) were predicted by the IAST method. At 25 and 1 bar, the selectivities of N-WAPC for equimolar CO\(_2\)/CH\(_4\) and CH\(_4\)/N\(_2\) are 3.19 and 7.62, respectively. The good selective adsorption performance of N-WAPC reveals its great potential for efficient methane mixture gases upgrading.

Supplementary Materials: The following are available online at http://www.mdpi.com/2079-4991/9/2/266/s1, Figure S1: SEM image of the pre-carbonized waste wool; Figure S2: High-resolution XPS spectra of (a) C1s and (b) O1s for WAPC and N-WAPC; Figure S3: (a) SEM image for N-WAPC and the corresponding EDS element mappings of (b) N, (c) C and (d) O; Figure S4: CO\(_2\) gas adsorption for N-WAPC at 0 °C (a) and 25 °C (b). The continuous solid line corresponds to the DSL fittings of the experimental data; Figure S5: CH\(_4\) gas adsorption for N-WAPC at 0 °C (a) and 25 °C (b). The continuous solid line corresponds to the L fittings of the experimental data; Figure S6: N\(_2\) gas adsorption for N-WAPC at 0 °C (a) and 25 °C (b). The continuous solid line corresponds to the L fittings of the experimental data; Figure S7: CO\(_2\) gas adsorption for WAPC at 0 °C (a) and 25 °C (b). The continuous solid line corresponds to the DSL fittings of the experimental data; Figure S8: CH\(_4\) gas adsorption for WAPC at 0 °C (a) and 25 °C (b). The continuous solid line corresponds to the L fittings of the experimental data; Figure S9: N\(_2\) gas adsorption for WAPC at 0 °C (a) and 25 °C (b). The continuous solid line corresponds to the L fittings of the experimental data; Table S1: Langmuir parameters and coefficient of determination for adsorption of CO\(_2\) in N-WAPC and WAPC; Table S2: Langmuir parameters and coefficient of determination for adsorption of CH\(_4\) in N-WAPC and WAPC; Table S3: Langmuir parameters and coefficient of determination for adsorption of N\(_2\) in N-WAPC and WAPC.

Author Contributions: Y.L. proposed and planned the research, and supervised the experiments; R.X. performed the experiment and analyzed the data; B.W., J.W. and J.Y. improved the data analysis; L.W. polished and edited the English; M.S. prepared the porous carbon. All the authors discussed the results and participated in the writing of the manuscript.

Acknowledgments: This work was supported by Project funded by China Postdoctoral Science Foundation (2018M632775), NSFC (No. U1804156, 51574112, 51874124), Henan Province Colleges and Universities Key
Research Project (18A620002, 18A150005), State Key Laboratory Cultivation Base for Gas Geology and Gas Control (Henan Polytechnic University) (WS2018A05), Innovative Research Team in University of Ministry of Education of China (IRT_16R22), National Key R&D Program of China (2017YFC0804207), and Program for Innovative Research Team of Henan Polytechnic University (T2018-2).

Conflicts of Interest: The authors declare no conflict of interest.

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