Granular Activated Carbon Produced from Asphalt


Summary: A process for preparing granular activated carbon has been developed, utilizing petroleum residues such as asphalt. Successful pilot plant operations have been made. The process consists of three steps: They are char production, char activation, and exhaust gas treating. Granular char, hard and uniform, has been produced by reacting asphalt with fuming sulfuric acid or elemental sulfur. A co-current, two-step fluidized bed furnace with unique features developed by the authors has been used for activating char above obtained. The product has indicated some excellent properties such as large specific surface area, extensive methylene blue adsorption, characteristic pore distribution, and high mechanical strength. Some practical applications have shown promising results.

1 Introduction

Activated carbon, especially granular activated carbon, is one of the essential materials used for prevention of air and water pollutions, and its demand is constantly increasing. Fresh water supply in Japan is limited, and its shortage will be a serious problem in the near future. The Ministry of Construction has estimated that in 1985 water demand in Japan will be 170% greater than that in 1965.

Up to now, woody materials such as coconut shells and sawdust have been valued as sources of activated carbon of good quality. But supplies of these raw materials are limited. Processes for production of activated carbon from coal have recently been developed, but they are only in limited commercial operation. Also these processes involve complicated steps, and they are not easily adaptable to large scale operations. Moreover, supply of coals with desirable properties for activated carbon production is limited.

The various materials mentioned above cannot be considered as dependable sources. On the other hand, the heavy fractions of petroleum, such as asphalt, are readily available, and they can be important sources of activated carbon. To develop a new process for large scale production of good quality activated carbon from these fractions, we have made studies using asphalts since 1970, and we are now operating a pilot plant using the process successfully.

2 Process Description

2.1 Process Flow Scheme

Figure 1 shows a flow diagram of the new process for producing activated carbon from asphalt. The process consists essentially of the following three steps: (1) char producing step, (2) char activating step, and (3) exhaust gas treating step.

Two different processes, as shown in Fig. 1, were used for the char producing step.

2.1.1 Char Producing Procedure

Petroleum residue or asphalt as a raw material was treated with sulfuric acid or fuming sulfuric acid in Process A. The purpose was to obtain a char with high melting point by sulfonation, oxidation, and dehydrogenation reactions. These reactions were carried out in a paraffinic or halogenated hydrocarbon solvent under vigorous agitation at 70~100°C for 1~3 hours.

In Process B, the raw material was reacted with elemental sulfur to obtain a char by dehydrogenation. The reaction was carried out in kerosene or gas oil used as solvent at 200~230°C for 2~4 hours.

Solvents in both processes were recovered by distillation, and the grainy chars of varying
sizes from about 1 to 5 mm in diameter remained. Such gases as \( \text{SO}_2 \) (from Process A) and \( \text{H}_2\text{S} \) (from Process B) produced in the reaction were subjected to gas treating, and the \( \text{SO}_2 \) recovered could be converted to sulfuric acid and \( \text{H}_2\text{S} \) to elemental sulfur. The yield of char from either process was from 120 to 160% by weight of the raw material used.

2.1.2 Char Activating Step

The char thus obtained was activated with steam at 850–900°C in our co-current, two-step fluidized bed furnace\(^1\) where steam was also used for fluidization, and activation could be done continuously and efficiently. Both steam and char were charged at the lower zone of the fluidized bed, and activated carbon was obtained from the upper zone. This is quite possible, because the density of activated carbon particles is smaller than that of the char. This is one of the unique features of our process.

The yield of activated carbon from the char was from 30 to 40% by weight. The yield based on asphalt was from 30 to 45% by weight.

2.1.3 Exhaust Gas Treatment

In Process A, the exhaust gas generated from the char producing step and the char activating step was substantially \( \text{SO}_2 \), which could be oxidized catalytically to sulfuric acid. \( \text{H}_2\text{S} \) was generated from Process B and it was converted to elemental sulfur by the Claus process. Both sulfuric acid and elemental sulfur thus recovered could be reused.

2.2 Process Characteristics and Economy

Although a number of reports are available for reactions between petroleum residues and sulfuric acid or elemental sulfur, there are only few reports concerned with the production of activated carbon\(^2\)–\(^4\). Moreover, the char produced by these reactions is usually lumpy and brittle, and additional steps such as pulverizing and pelleting are required for manufacturing granular activated carbon of high quality.

In our process, however, the reaction was carried out in the liquid phase and granular char was precipitated from the reaction mixture. So, it was possible to obtain a hard char of almost uniform particle size.

The char was so easy to handle and it could be activated smoothly in the fluidized state, without the need for pulverizing or pelleting, to produce high quality granular activated carbon.

Since the co-current, two-step fluidized furnace for activation is used in our process, activation can be done continuously with high thermal efficiency. This makes the commercial plant compact and simple that enhances the possibility of mass production of activated carbon.

Generally, the yield of activated carbon from coal used as the raw material is about 20%\(^5\)–\(^6\), while the yield from asphalt by our process is from 30 to 45% by weight of asphalt used.
Since the solvents, sulfuric acid or elemental sulfur used can easily be recovered and reused, the process can be operated in a closed system. About 85% or more of the sulfuric acid used as reactant can be recovered and reused in Process A. About 75% or more of elemental sulfur used as reactant can be recovered and reused in Process B.

Table 1 gives construction costs and utilities requirements for Process B. It should be noted that activated carbon can be produced quite economically from asphalt by our process.

3 Properties of Activated Carbon

3.1 Basic Properties

Table 2 shows some of the properties of activated carbon A (product of Process A), B (product of Process B), C (market product), and D (market product). A and B have higher specific surface areas and high methylene blue adsorption. Pore distributions are also shown in Fig. 2.

The existence of many micropores (under 15Å pore radius) is remarkable in A, while the existence of transitional pores (from 15Å to 1,000Å pore radius) is remarkable in B. It is presumed that this is the reason why the products from Process A and B indicate such high specific surface areas. Figures 3, 4, 5 and 6 show the surfaces of four species of activated carbon, enlarged 4,500 times by a scanning electron microscope. The surface of A or B differs from that of the market products, and the presence of small pores is clearly shown on the surfaces of A and B.

Mechanical strength of the activated carbons were measured by the modified method of JIS K 1412. B was found to be superior in mechanical strength.

Table 1 Typical Economics

<table>
<thead>
<tr>
<th>Plant Capacity</th>
<th>5,000 t/yr</th>
</tr>
</thead>
<tbody>
<tr>
<td>Investment Cost</td>
<td>1,860 × 10⁶ Yen</td>
</tr>
<tr>
<td>Raw Materials (per product ton)</td>
<td></td>
</tr>
<tr>
<td>Asphaltic Petroleum Residue</td>
<td>2.2 t</td>
</tr>
<tr>
<td>Heavy Gas Oil</td>
<td>0.3 t</td>
</tr>
<tr>
<td>Sulfur</td>
<td>0.8 t</td>
</tr>
<tr>
<td>Utilities</td>
<td></td>
</tr>
<tr>
<td>Electricity</td>
<td>620 kW·hr</td>
</tr>
<tr>
<td>Steam</td>
<td>5.9 t</td>
</tr>
<tr>
<td>Industrial Water</td>
<td>89 t</td>
</tr>
<tr>
<td>Fuel</td>
<td>1.5 × 10⁶ kcal</td>
</tr>
</tbody>
</table>

Table 2 Characteristics of Typical Activated Carbons

<table>
<thead>
<tr>
<th></th>
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</tr>
</thead>
<tbody>
<tr>
<td>Specific Surface Area (m²/g)</td>
<td>1,250</td>
<td>1,190</td>
<td>1,040</td>
<td>775</td>
</tr>
<tr>
<td>Pore Volume (m³/g)</td>
<td>1.29</td>
<td>1.31</td>
<td>0.99</td>
<td>0.85</td>
</tr>
<tr>
<td>Adsorption of Methylene Blue (mg/g)</td>
<td>356</td>
<td>359</td>
<td>324</td>
<td>134</td>
</tr>
<tr>
<td>Bulk Density (g/ml)</td>
<td>0.40</td>
<td>0.44</td>
<td>0.42</td>
<td>0.42</td>
</tr>
<tr>
<td>Mechanical Strength* (%)</td>
<td>70</td>
<td>91</td>
<td>68</td>
<td>65</td>
</tr>
<tr>
<td>Size Distribution (wt%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>8 mesh over</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>8~12 mesh</td>
<td>1</td>
<td>1</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>12~16 mesh</td>
<td>4</td>
<td>5</td>
<td>2</td>
<td>3</td>
</tr>
<tr>
<td>16~24 mesh</td>
<td>23</td>
<td>29</td>
<td>50</td>
<td>38</td>
</tr>
<tr>
<td>24~32 mesh</td>
<td>35</td>
<td>40</td>
<td>29</td>
<td>42</td>
</tr>
<tr>
<td>32 mesh under</td>
<td>28</td>
<td>20</td>
<td>13</td>
<td>10</td>
</tr>
</tbody>
</table>

* Mechanical strength was measured by the modified method of JIS K 1412 as follow: 17~18 g of activated carbon sieved into 24~32 mesh range were charged in the steel-bound mill, and the carbon was crushed by 10 steel-bound balls (10 mmϕ) with rotating the mill at 100 rpm for 60 minutes. Then the activated carbon was sieved into 24~32 mesh range, and its weight was determined.

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3.2 Adsorption Characteristics

The adsorption characteristics of activated carbons were examined by determining the adsorption isotherms and rates.

The change in the amount of adsorption with time and the amount of adsorption at equilibrium in both vapor phase and liquid phase were determined. The relationship between the equilibrium pressure or concentration and the amount of adsorption was summarized according to Freundlich's formula of adsorption (1)\(^9\),\(^10\).

\[
\log Q_\infty = \frac{1}{n} \log q + K
\]  

(1)

\(Q_\infty\) is the amount of adsorption at equilibrium, \(q\) is the equilibrium pressure or concentration, and \(1/n\) and \(K\) are constants. A small \(1/n\) means a slight change in adsorption capability with change of equilibrium pressure or concentration, and a large \(K\) indicates a large amount of adsorption.

The change in the amount of adsorption with time was summarized according to Boyd's equation (2)\(^11\).

\[
\frac{Q_t}{Q_\infty} = \frac{6}{R} \sqrt{\frac{D_i \cdot t}{\pi}}
\]  

(2)

\(R\) is the average radius of activated carbon particles, \(D_i\) is the internal diffusion constant, and \(t\) is time.

Generally, the rate of diffusion within the pores is considered to be the rate-determining step\(^11\).

As an example of the results in vapor phase adsorption, the adsorption isotherms and adsorption rates of SO\(_2\) are shown in Fig. 7 and 8. As an example of liquid phase adsorption, the results of methylene blue adsorption are shown in Fig. 9 and 10.

In the vapor phase adsorption, A and B have characteristics similar to those of market product C or D. On the other hand, in the liquid phase adsorption, A and B show better adsorption characteristics than those of C or D. Especially for high molecular weight substances, A and B incline to give higher adsorption rates than those of the market products.

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3.3 Performance Test

Two experiments were carried out to test the performance of our activated carbon in the vapor and liquid phase applications. As an example of vapor phase application, capability for prevention of motor gasoline vaporization was determined by Toyota Motor's Test Method, which consists of vapor phase adsorption and desorption of benzene on activated carbon. A 150 g sample of activated carbon was introduced into a U-tube to form a layer, and benzene vapor was passed through it at the rate of 400 ml/min at 25°C until the weight of the adsorbent layer became constant. The results are shown in Fig. 11 (1).

In the desorption experiment, air was passed through the activated carbon layer, which had adsorbed benzene, at the rate of 400 ml/min at 25°C, and the decrease in the weight of the adsorbent layer with time was determined. The results are shown in Fig. 11 (2).

As for the activated carbon used to prevent motor gasoline vaporization, it is desirable that the activated carbon has a high rate of and capacity for adsorption of hydrocarbons (benzene) and a high rate of desorption of hydrocarbons (benzene). The market products E and F are used industrially for this purpose. The products A and B are expected to be more useful, because they have better adsorption and desorption characteristics than those of E and F.

As an example of liquid phase applications, the removal of phenol from FCC waste water was tested with activated carbon. The FCC waste water contained about 100 ppm of phenols and about 200 ppm of H₂S. Since the latter might interfere with the adsorption of phenols, it was removed prior to the adsorption experiment by bubbling air through the waste water, which was then charged to the 150 g activated carbon layer at the rate of 6 l/hr. The
break-through curves are shown in Fig. 12. The break-through point is defined as the point at which the phenol content in the effluent from the activated carbon column attains a value of 0.3 ppm.

4 Conclusion

We have developed a new and unique process for economical production of granular activated carbon from asphalts.

To produce a char, which can withstand high activating temperatures such as 850~900°C, from petroleum fractions of low softening points, they were reacted with sulfuric acid, fumic sulfuric acid or elemental sulfur.

In the first step, granular char is produced by precipitation from the liquid phase in which asphalt is made to react with one of the above sulfurous substances.

In the next step, the char is activated continuously using the unique co-current, two-step fluidized furnace developed by us.

The product from our process shows good properties as regards specific surface area, methylene blue adsorption, pore distribution and mechanical strength. The vapor and liquid phase adsorption characteristics of our product are superior to those of market products.

We, Maruzen Oil Co., Ltd. are planning to commercialize this new process for production of high quality activated carbon.

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