Thermal Diffusivity of Carbon Pellets (CPs) Treated with KOH

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Abstract: In this work thermal diffusivity of carbon pellets (CPs) treated with different percentage level of KOH has been studied. Thermal diffusivity measurements were carried out at room temperature by using photoflash technique. The technique consists of a camera flash having approximately 5 ms pulse duration for heating and a thin film of polyvinylidene difluoride (PVDF) attached to the back of the samples for signal detection. Eight carbon samples treated with different percentage level of KOH (0 to 7 mole percent) were studied and the thermal diffusivity result was compared with SEM which showed that the sample with highest thermal diffusivity had distinctly lower porosity and better grain alignment.

Keyword: Thermal Diffusivity, KOH, PVDF, Flash Technique

INTRODUCTION

The porosity effect on thermal properties of material has been studied by many workers since 1972. Since that time a lot of work has been done on porosity effect on thermal conductivity of materials. In comparison, there was less work done on porosity effect on thermal diffusivity. The grain size and grain alignment in microstructure of materials are two other important parameters that can affect thermal properties of materials as has been reported by Haydari et al. In this work the effect of porosity and grain alignment on thermal diffusivity was studied on the samples of carbon pellets (CPs) treated with KOH. The photoflash technique was used to measure thermal diffusivity whereby camera flash absorption at the front surface induced heat into the sample while the temperature history of the rear surface of the sample was detected by using polyvinylidene difluoride (PVDF) film.

MATERIAL AND METHODS

Theory: One-dimensional heat propagation in a system consisting of four layers of gas, sample, PVDF sensor and backing, assuming that the sample is optically opaque and the backing has similar thermal properties to the PVDF sensor was considered. The theoretical calculation of the signal as detected by PVDF sensor that thermally attached to the sample was obtained by approximating the long camera flash temporal profile with a finite square pulse.

Sample preparation: Carbon pellets (CPs) precursor (cotton cellulose) was first pre-carbonized at low carbonization temperature, milled for 20 h and sieved to produce self-adhesive carbon grains (SACG) that can pass through a 53 microns sieve. The SACG was treated for 16 hours with KOH having concentration from 0 to 7 moles percent and then dried in oven at 100 °C before converted into pellets (CPs) by applying 15 metric tones compression pressure on 2 g of samples in a mould. These CPs were carbonized from 500 °C to 1000 °C in nitrogen environment using a multi-steps heating profile (Vulcan Box Furnace 3-1750). The CPs were then washed with distilled water to remove impurities until a pH of 6 and dried in an oven at 100 °C for 4 hours.

Experimental setup: The experimental setup used for the thermal diffusivity measurements consisted of the pyroelectric detector, a 52 µm thick PVDF film attached to the samples with a thin layer of thermal grease. A thick vulcanized rubber layer was used as sample backing. The excitation source was a camera flash (Minolta 5400HS) with pulse duration of about 5 ms (that is much shorter than the pyroelectric response obtained from the samples) and of temporal shape that may be approximated with a finite square pulse. The signal from the PVDF transducer is recorded by a 400 MHz digital oscilloscope (LeCroy 9310A). The measurements were conducted at room temperature by averaging a total of 25 signals. The samples were polished to optimum measurement thickness of ~1 mm.

RESULTS AND DISCUSSION

Figure 1 shows PVDF response collected from the samples. As shown in the figures the model based on square pulse has reasonably good agreement with the experimental data with slight mismatches at the peak and the tail. The sources of the mismatch are from the heat loss that starts to dominate at the long time behavior at the tail and the deviation of the theoretical
Fig. 1: Thermal wave response from carbon sample with different KOH (% mole) fitted with theoretical model (continuous line)

Fig. 2: Thermal diffusivity as a function of KOH mole percent. The continuous line is for added clarity
The morphologies of the fractured surface for the KOH (0, 2, 5 and 7%) treated carbon pellet are shown in Figure 3. The SEM micrograph shows that the grain size and inter-grain distance is the most distinguishable for the particular case of KOH (2%) treatment as compare to the rest which display almost similar features. Visually it is the most compact i.e. the least porous, smaller grain size and the best aligned among others. As evident in the Figure 2 the thermal diffusivity for the case of KOH (2%) is much higher than the others justifies better grain alignment and smaller porosity that aid to more efficient heat diffusion in the sample.

CONCLUSION

The positive effect of lower porosity and better grain alignment on the thermal diffusivity for a set of carbon pellets (CPs) treated with KOH has been described. This made thermal diffusivity value of KOH (2%) higher than others.

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square pulse from the actual experimental temporal profile of the camera flash. Therefore, the mismatch is less obvious in the data from the sample with higher thermal diffusivity as expected whereby heat loss is much lower when the signal is stronger and the temperature history is shorter. The thermal diffusivity for a set of carbon treated with KOH obtained from the curve fitting is summarized in figure 2. The thermal diffusivity peaked itself at about 0.46×10⁻⁶ m²s⁻¹ for the sample treated with 2% moles of KOH. At other mole percentages the thermal diffusivity has slightly an upward trend towards higher mole percent.
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