Thermal performance of carbon-based microencapsulated phase change materials

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ABSTRACT – The aim of this study is to investigate the effect of carbon-based materials for the thermal performance of microencapsulated phase change material (μ PCM). The sample was prepared separately by mixing 5 wt.% of Multiwall Carbon Nanotube (MWCNT) and 5 wt.% Expanded Graphite (EG) with μ PCM using a powder metallurgy technique. The mixed powder was then compacted into a disc with a diameter of 45 mm and thickness of 5 mm using a hot compaction technique. The thermal performance was tested according to the ASTM standard. It was found that the addition of MWCNT into μ PCM can absorb heat effectively as compared to pure μ PCM and μ PCM/EG composite.

1. INTRODUCTION

Phase change material (PCM) is suitable to be applied as energy storage, thermal protection and passive cooling system [1]. This is because of its desirable characteristic such as high latent heat of fusion, low vapor pressure during melting, no super cooling and chemically inert and stable [2-3]. In principle, the heat generated by any system could be absorbed by the PCM by attaching it on the system. PCM acts as heat sink for the system to eliminate the heat produced. However, the PCM could melt when the temperature of the system exceeds the melting point and the high latent heat of PCM. The melting PCM causes complexity and limits the full use of its function. In order to solve the limitation, micro-encapsulated technique was employed to PCM to solve the problem. Microencapsulated phase change material (µPCM) products are very small bi-component particles consisting of a core material, the PCM and an outer shell or capsule wall that hold the melted PCM from leaking. Therefore, whether the PCM is in the liquid or solid state, the capsule itself will remain as a solid particle.

The μ PCM has disadvantage of low thermal conductivity to meet required heat exchange rates. Many techniques were proposed to improve thermal conductivity of PCM such as inserting fin, incorporating carbon based materials and adding metallic and nonmetallic materials [4]. Most researchers use core

PCM, typically Paraffin wax instead of μ PCM, therefore the matrix lead to melt compare to μ PCM.

Based on the above review, there are a limited number of studies [1-4] that investigate the effect of carbon-based materials on the thermal performance of μ PCM. This study aims to investigate further the thermal performance between μ PCM/EG and μ PCM//MWCNT composites.

2. METHODOLOGY

The μ PCM type of PX52 was supplied by Rubitherm Technologies, while for MWCNT and EG was supplied by Microtek Laboratories. Table 1 shows the thermal properties of the materials.

Table 1 Thermal properties of the materials			
^a Properties	μPCM	MWCNT	EG
Melting temperature, °C	52	3625	3000
Thermal conductivity, W/m.k	0.1	3500	200
Latent heat capacity, kJ/kg	100	-	-
Density, g/cm ³	0.65	0.03	2.2

Table 1 Thermal properties of the materials

^aFrom manufacturer

The samples were prepared separately by mixing 5 wt.% of MWCNT and 5 wt.% EG with μ PCM using a ball mixer. The mixed powder was then compacted into a disc with a diameter of 45 mm and thickness of 5 mm using a hot hydraulic press. The samples are show in Figure 1.



Figure 1 Photo of samples of (a) µPCM, (b) µPCM/EG composite, and (c) µPCM/MWCNT composite.

The thermal performance of the prepared sample was tested according the modified method in correlation to the ASTM C518-10 [6]. The measurement stack of the apparatus is shown in Figure 2. The sample is placed between the hot and cold copper plates. The hot plate above the electric heater assures unidirectional heat flow from the heater up through the sample. The cold plate above the sample provides uniform temperature distribution on its surface. Unidirectional heat flow is achieved by adjusting the electric power supply to such a level that temperature of the heater is equal to the hot plate temperature. The sample was covered by aluminum. The aluminum casing was heated at 12V of nominal voltage and 5 Watt of power. All tests were performed at room temperature. The temperature of aluminum casing was recorded by using an infrared thermometer for each minute. The result was obtained in the form of the temperature profile of aluminum casing over time.



Figure 2 Measurement stack in plate apparatus.

3. RESULTS AND DISCUSSION

Figure 3 shows the temperature profiles of aluminum with and without µPCM composites. It was clearly shown that the temperature of aluminum increases over time. However, the magnitude of temperature significantly decreases when the µPCM is present. Besides, the combination of µPCM and MWCNT effectively reduce the temperature of aluminum casing to 91% over the time. This shows that the composite absorbs some amount of heat from the aluminum casing, and hence decrease the temperature. The present of MWCNT into µPCM can absorb more heat from the aluminum casing as compared to the pure µPCM and µPCM/EG composite. This is due to the fact that the MWCNT has higher thermal conductivity and larger surface area compared to EG. M. Lajvaerdi et al. [5] reported that the thermal conductivity of paraffin increases to 31% with addition of 2 wt.% of MWCNT.

4. CONCLUSIONS

In conclusion, the thermal performance of μ PCM can be improved by employing carbon-based materials as an additive. However, the μ PCM/MWCNT

composites show a great improvement in terms of heat absorption due to the fact that the MWCNT has higher thermal conductivity and larger surface area compared to EG.



Figure 3 Temperature profiles of aluminum casing over time with and without μ PCM composites.

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6. **REFERENCES**

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