Thermal performance of a phase change material-based heat sink in presence of nanoparticles and metal-foam to enhance cooling performance of electronics

Adeel Arshad^{a,*}, Mark Jabbal^a, Hamza Faraji^b, Pouyan Talebizadehsardari^c, Muhammad Anser Bashir^d, Yuying Yan^{a,*}

^aFluids & Thermal Engineering (FLUTE) Research Group, Faculty of Engineering, University of Nottingham, Nottingham, NG7 2RD, UK

^bPhysics Department, LPMMAT Laboratory, Faculty of Sciences Ain Chock, Hassan II University, Casablanca, Morocco

^cDepartment of Mechanical and Aerospace Engineering, Institute of Energy Futures, Brunel University London, Uxbridge, Middlesex UB8 3PH, UK

^dDepartment of Mechanical Engineering, Mirpur University of Science & Technology (MUST), Mirpur 10250, AJK, Pakistan

Abstract

The present study explores the parametric investigation of a heat sink filled with composite of pure phase change material (PCM), nanocomposite phase change material (NCPCM), metal-foam (MF) by employing the numerical approach for effective passive thermal management of electronics. The combinations of heat sink are varied by filling PCM, NCPCM, MF+PCM and NCPCM+MF. Different parameters such as MF materials, porosities, pore densities (PPI-pores per inch), volume fractions of nanoparticles in NCPCM, power levels and combination of MF+NCPCM by varying different porosities and nanoparticles volume fractions. Copper (Cu) nanoparticles of 1%, 3% and 5% volume fraction were dispersed in RT-35HC, used as a PCM, and copper, aluminium (Al) and nickel (Ni) MFs were embedded inside the heat sink. Transient simulations with conjugate heat transfer and melting/solidification schemes were formulated using finite-volume-method (FVM). The thermal performance and melting process of the NCPCM filled heat sink were evaluated through melting time, heat storage capacity, heat storage density, rate of heat transfer and rate of heat transfer density. The results showed that with the addition of Cu nanoparticles and MF, the rate of heat transfer was increased and melting time was reduced. The melting time was reduced by -1.25%, -1.87% and -2.34%; and rate of heat storage is enhanced

^{*}Correspondence authors

Email addresses: adeel.arshad@nottingham.ac.uk, adeel_kirmani@hotmail.com (Adeel Arshad), yuying.yan@nottingham.ac.uk (Yuying Yan)

by 1.35%, 0.76%, and 0.19% with the addition of 1%, 3% and 5% volume fraction of Cu nanoparticles, respectively. The composite of MF+NCPCM showed the lower heat sink temperature and higher liquid-fraction were obtained. The latent-heating phase duration was decreased with the increase of Cu nanoparticles volume fraction. Additionally, the lower reduction in melting time of -18.10% and higher rate of heat transfer of 8.12% were obtained with 1% Cu nanoparticles, 95% porosity and 10 PPI Cu MF based heat sink. *Keywords:* Thermal management, Nanocomposite phase change material based heat sink, Phase change material, Electronics cooling

1 1. Introduction

The reliability of smart electronic devices is purely based on their ability to perform 2 the preprogrammed functions under the safe operating conditions. With the advancement 3 in electronics and telecommunications industries, the electronics products have shifted to-4 wards lightweight, low power consumption, faster and smaller [1]. Gradually, the growing 5 miniaturization and advanced multi-features of smart electronic devices have led towards 6 the more complexity in thermal management (TM) solutions subjected the medium to 7 high-heat-flux. Internal overheating and rise of the temperature as a result of this, lead 8 to damage or failure of electronic devices around 55% compared with the other failures [2]. 9 Modern TM technologies including active and passive, the active cooling methods such air 10 or liquid cooling, adopted to cool the portable electronic devices have some limitations such 11 as high-power consumption, acoustic noise production, bulk in volume and size [1, 3]. In 12 such instances, a novel technology is needed which can ensure the passive TM in the best 13 possible way. Significant outcomes of cooling through PCMs with heat sink, used as passive 14 heat exchanger, in electronics equipment have been achieved as they emanate heat during 15 heavy usage due to their high latent-heat of fusion with isothermal phase transformation 16 [4]. To facilitate the reliable TM of electronic devices, a latent-heat storage unit (LHSU) 17 containing high thermal conductivity enhancers in form of extruded fins [5–8], metal-foams 18 [9, 10] and nanoparticles [11–13] with PCMs embedded in a heat sink are invariably used 19 [14, 15].20

Many researchers have introduced the applications on PCM with porous medium using ei-21 ther metallic foams or carbon material such as expanded graphite integrated with heat sink, 22 heat pipe and shell and tube heat energy storage system. Zhao et al. [16] conducted a 23 experimental study to investigate the heat transfer enhancement through a PCM/MF heat 24 storage system. RT–58 and Cu MF were embedded and melting and solidifying processes 25 were analysed. The results found that using the Cu MF as heat transfer enhancement 26 media, increased the overall heat transfer rate by 3–10 times while phase–change process. 27 Further, Tian and Zhao [17] conducted a numerical study to effect of MFs on heat transfer 28 enhancement using 2D LTNE model and conduction and convection heat transfer modes 29 were considered at phase-transition and liquid-phase. The key findings were reported that 30 heat conduced rate was improved by using the MF because of high thermal conductivity 31 and natural convection effects were suppressed because of the large flow resistance in MF. 32 Sundarram and Li [18] investigated the pore size and porosity of aluminium MFs infiltrated 33

with paraffin wax as a PCM using a three-dimensional finite element model. The results 34 found that at a constant porosity, the smaller pore size reduced the more heat source tem-35 perature for a longer duration compared with higher pore size. In addition, the effective 36 thermal conductivity of MF+PCM system was doubled by reducing the pore size from 100 37 to 25 μ m. Chen et al. [19] conducted a experimental and numerical study using aluminium 38 made MF integrated with paraffin wax at pore scale. Authors studied the temperature 39 field and melting evolution MF+PCM and found that MF had the capability to enhance 40 the phase-change heat transfer during solid-liquid phase transformation due to the thermal 41 conduction in the metal matrix. Nada and his co-authors [20] conducted the comprehensive 42 parametric study by using carbon–foams of varying porosities and thermal conductivities 43 and PCMs of different latent-heat of fusions and melting temperatures at different input heat 44 fluxes. The effects of module thickness and power densities were investigated and the results 45 showed that by decreasing carbon-foam and PCMs thermal conductivities, increasing the 46 module height and carbon-foam porosity increased the module temperature and delayed the 47 steady-state temperature time. Further, the authors used the nano-carbon tubes along with 48 carbon-foam and PCM (RT-65) and the effect of pure carbon-foam, carbon-foam+RT-65 49 and carbon-foam+RT-65/nano-carbon tubes of different porosities were investigated nu-50 merically [21]. The results revealed that carbon-foam+RT-65/nano-carbon tubes reduced 51 the 11.5% module surface temperature with less then 75% carbon–foam porosities and a 52 7.8% reduction was obtained with 88% porosity. Alipanah and Li [22] proposed a numerical 53 study for TM of Li-ion battery by using octadecane, gallium and octadecane+aluminium 54 MF. Three different porosities of 0.88, 0.925 and 0.97 and three heat fluxes of 400, 600, and 55 800 W/m^2 were varied. The results revealed that octadecane+aluminium MF of 0.8% led 56 to the 7.3 times longer discharge time compared to the pure octadecane case. Furthermore, 57 the addition of aluminium MF increased the uniformity in battery surface temperature. 58 Zhang et al. [23] conducted the experimental and numerical study consisting of Cu MF 59 and paraffin wax in a square cavity for thermal energy storage and TM applications. The 60 phase-change heat transfer and melting phenomenon were studied and found that there was 61 a quite significant difference between the ligament of Cu MF and paraffin wax because of 62 the non–equilibrium thermal effects in heat transfer between the paraffin wax and Cu MF. 63 Mahdi and Nsofor [24, 25] conducted the numerical studies to study the melting and solid-64 ification processes in a triplex-tube thermal energy system using Al_2O_3 nanoparticles and 65 Cu MF combination under three heat transfer fluid temperatures. Three different nanopar-66

ticles volume fractions of 0.01, 0.03 and 0.05 and two MF porosities of 0.95 and 0.98 were 67 varied for each inlet fluid temperature. The authors found that melting and solidification 68 processes were improved with the addition of MF and nanoparticles further improved the 69 melting and solidification processes with the increase of volume fraction. In addition, the 70 authors reported that with the increase of MF porosity, the potential of nanoparticles for 71 enhancing the thermal energy storage was decreased. Further, the combination of NCPCM 72 and MF was numerically investigated by Bernardo et al. [26] by adding 1% and 5% of Al₂O₃ 73 nanoparticles in RT–58, as a PCM, and aluminium MF having 80% and 90% porosities. The 74 authors revealed that MF improved the charging or discharging rate more significantly of the 75 system instead of the nanoparticles. The nanoparticles could be used to only to adjust the 76 charging or discharging time moderately during phase-transformation processes. Chamkha 77 et al. [27] conducted a numerical study using a L-shape enclosure heat sink consisting of 78 Cu MF and paraffin wax under the pulse heat flux conditions to study the flow and heat 79 transfer phenomenon. A constant and uniform efficiency were obtained by the MF+PCM 80 heat during the pulse heating. The results revealed that the higher pulse heat flux showed 81 that higher heat sink efficiency. The efficiencies of 1.75 and 2.4 were obtained by providing 82 the element heat flux to fourfold and sixfold of the steady heat flux, respectively. Recently, 83 Li et al. [28] conduced a numerical study by using porous/NCPCM system in a counter 84 current triple-tube to explore the effect of MF and NCPCM under different heat transfer 85 fluid temperatures and directions. The results showed that by 5% of Cu nanoparticles, the 86 melting/solidification period was reduced by 25.9%/28.2%. In addition, with 95% porosity 87 of MF, the melting/solidification period was reduced by 83.7%/88.2% which revealed that 88 embedding the MF with PCM had the more advantage than the adding nanoparticles into 89 the PCM. 90

The aforementioned literature reveals that a few studies have explored the effect of adding 91 nanoparticles into the PCM and MF+PCM specially in an application of latent-heat ther-92 mal energy storage system. Therefore, the present study aim to explore the effect of four 93 different heat absorbing mediums such as PCM, NCPCM, MF+PCM and MF+NCPCM 94 filled in a heat sink for passive thermal cooling of electronic devices. The RT–35HC is used 95 as a PCM and copper (Cu) nanoparticles of varying volume fractions are dispersed into the 96 PCM. Three different MFs made of Cu, aluminium (Al), and nickel (Ni) are investigated 97 to find the best heat conductive MF with PCM. Three different porosities, five different 98 pore densities and four different input power levels of 5, 6, 7, and 8 W are varied. Firstly, 99

the MF of constant porosity is embedded into NCPCM of different volume fractions and 100 secondly, the NCPCM of constant volume fraction is varied with different porosities of MF 101 to explore the effect of two heat conductive mediums. The melting phenomenon and tem-102 perature distribution of of pure PCM, NCPCM, MF+PCM and NCPCM+MF based heat 103 sink is studied. Moreover, thermal cooling performance is presented using four different 104 performance evaluation parameters such as heat storage capacity, heat storage density, rate 105 of heat transfer, and rate of heat transfer density along with the total melting time. This 106 will eventually provide a better picture to select the optimum heat storage medium filled in 107 a heat sink for efficient solution of passive thermal cooling of electronic devices. 108

¹⁰⁹ 2. Geometric and Mathematical description

110 2.1. Physics of the problem

In present study, a two-dimensional (2D) heat sink is considered filled up with the 111 composite of PCM, NCPCM, MF+PCM and MF+NCPCM, as shown in Fig. 1 and the 112 configuration of each case is presented in Fig. 2 investigated in current study. The heat sink 113 with width (W = 70 mm) and height (H = 25 mm) is heated with a heat source of volumetric 114 heat generation (q''') with sizes of l = 50 mm and t = 2 mm. All the sides of heat sink are 115 adiabatic except top surface which undergoes with natural convection. The internal cavity 116 of the heat sink has width of w = 60 mm and height h = 20 mm. The heat sink is made 117 of Cu and numerically modelled to investigate the thermal performance for passive cooling 118 of electronic devices. The effect of different parameters such as different MF materials, 119 porosities, pore densities, volume fractions of nanoparticles in NCPCM, combination of 120 NCPCM+MF by varying different porosities and nanoparticles volume fractions, and power 121 levels. The RT–35HC is used as PCM having melting temperature of 35 °C. Three different 122 volume fractions of $\varphi = 1\%$, 3%, and 5% of Cu nanoparticles are dispersed in RT-35HC to 123 study the effect of NCPCM based heat sink. Three different porosities of 95%, 90%, and 80% 124 and five different pore densities or PPI of 10, 20, 30, 40 and 50 are varied of MF embedded 125 inside the PCM heat sink to study the effect of MF+PCM heat sink. The MFs made of 126 Cu, Al and Ni are investigated with PCM. Five different power levels of 4, 5, 6, 7, and 8 W 127 are provided at the base of heat sink to analyse the temperature distribution and melting 128 phenomenon MF+PCM based heat sink. The 95% porosity and 10 PPI of Cu MF is kept 129 constant with pure PCM and NCPCM of $\varphi = 1\%$, 3%, and 5% heat sink cases. Similarly, 130 NCPCM of 5% volume fraction of Cu nanoparticles is kept constant with pure PCM and MF 131

of 95%, 90%, and 80% porosities to investigate the optimum volume fraction and porosity of composite NCPCM+MF filled heat sink. The current system is designed based on the average dimensions of portable hand-held electronic devices to investigate passive thermal performance using pure PCM, NCPCM, MF+PCM and MF+NCPCM based heat sink.



Figure 1: Schematic diagram of the computational domain used in current study.

136 2.2. Mathematical formulation

The governing equations are applied based on the PCM and NCPCM based heat sink 137 embedded with MF, shown in Fig. 1. During melting process of pure PCM and NCPCM, 138 heat is transferred through conduction mode while in solid–phase and then conduction and 139 natural convection modes contribute the heat transfer while generation of liquid-phase due 140 the temperature gradient. Since, the pure PCM has low thermal conductivity, a solid porous 141 medium is embedded inside the PCM and NCPCM which enhances the heat transfer rate 142 by conduction rather than convection because of the high flow-resistant effect of MF struc-143 ture. Since, the presence of MF increases the volume of heat sink having a similar PCM or 144 NCPCM mass. Moreover, due to the flow-resistant of MF structure, the effect of natural 145 convection is negligible in MF+PCM and NCPCM+MF cases. By adding the nanoparticles 146 into the pure PCM, the thermophysical properties of PCM are changed. The purpose to 147 add the nanoparticles is to modify the thermal conductivity of PCM. The effective density 148 is increased, whereas latent-heat of fusion and specific heat capacity are reduced. 149

¹⁵⁰ In numerical modelling, the enthalpy-porosity method is adopted to model the effect of ¹⁵¹ phase transformation of pure PCM and NCPCM inside the MF embedded heat sink, where ¹⁵² the porosity is equal to the liquid-fraction of each cell. Due the presence of MF, the vis-



Figure 2: Different configurations of heat sink investigated in current study.

cous and inertial losses are overcome by introducing the pressure drop effect in momentum
equation. The MF as a porous medium is modelled by considering the Darcy–Forchheimer–
Brinkman model (DFBM) and a source term is introduced in momentum equation. The
following assumptions are considered to define the continuity, momentum and energy equations [29–31]:

- An open cell MF is assumed homogeneous and isotropic.
- An incompressible, transient, laminar and Newtonian fluid is considered of liquid PCM
 and NCPCM inside the MF.
- Local thermal equilibrium model is assumed between the PCM, NCPCM and MF in
 energy equation.
- Volume expansion of PCM and NCPCM is negligible during phase transformation process.

- Viscous dissipation is negligible and no-slip boundary condition are assumed for velocities.
- Constant thermophysical properties are considered for nanoparticles, PCM and NCPCM
 expect the density for PCM and NCPCM.
- The heat sink is considered as solid-state with homogeneous and isotropic properties and thermal conduction heat transfer exists.
- The Boussinesq approximation is assumed to simulate the buoyancy driven effect under natural convection as $\rho = \rho_m / [\beta(T - T_m) + 1]$, where $T_m = (T_s + T_l)/2$.
- The NCPCM is considered as colloid suspension which exhibits as a Newtonian fluid.
 The liquid NCPCM flow regime is 2D, laminar, unsteady and incompressible.
- The dispersion of nanoparticles in PCM is assumed homogeneous, no agglomeration is considered.
- The nanoparticles and PCM are in local thermal equilibrium and there is no-slip between them.
- The initial temperature of heat sink, PCM, NCPCM and MF are the same.
- Adiabatic boundary conditions are assumed from the surroundings.

According to the above assumption the governing conservation equations for mass, momentum and energy can be written as follow:

Mass conservation:

$$\frac{\partial u}{\partial x} + \frac{\partial v}{\partial y} = 0 \tag{1}$$

Momentum conservation:

$$\frac{\rho_{ncpcm}}{\varepsilon} \left(\frac{\partial u}{\partial t} + \frac{u}{\varepsilon} \frac{\partial u}{\partial x} + \frac{v}{\varepsilon} \frac{\partial u}{\partial y} \right) = -\frac{\partial p}{\partial x} + \frac{\mu_{ncpcm}}{\varepsilon} \left(\frac{\partial^2 u}{\partial x^2} + \frac{\partial^2 u}{\partial y^2} \right) - A_m \frac{(1 - f_l)^2}{(f_l^3 - 0.001)} \cdot u - \left(\frac{\mu_{ncpcm}}{K} u + \frac{C_F}{\sqrt{K}} \rho_{ncpcm} u \sqrt{u^2 + v^2} \right)$$
(2)

$$\frac{\rho_{ncpcm}}{\varepsilon} \left(\frac{\partial v}{\partial t} + \frac{u}{\varepsilon} \frac{\partial v}{\partial x} + \frac{v}{\varepsilon} \frac{\partial v}{\partial y} \right) = -\frac{\partial p}{\partial x} + \frac{\mu_{ncpcm}}{\varepsilon} \left(\frac{\partial^2 v}{\partial x^2} + \frac{\partial^2 v}{\partial y^2} \right) - (\rho\beta)_{ncpcm} g\varepsilon (T - T_{ref}) - A_m \frac{(1 - f_l)^2}{(f_l^3 - 0.001)} \cdot v - \left(\frac{\mu_{ncpcm}}{K} v + \frac{C_F}{\sqrt{K}} \rho_{ncpcm} v \sqrt{u^2 + v^2} \right)$$
(3)

where, the ρ_{ncpcm} , μ_{ncpcm} , β_{ncpcm} are the density, dynamic viscosity, and thermal ex-183 pansion coefficient of the NCPCM, respectively; p and q are the pressure and gravitational 184 acceleration, respectively. A_m is the mush-zone constant which represents the damping of 185 the velocity to zero during the solidification. The A_m constant does not affect the behaviour 186 of melting time of PCM or NCPCM when MF is combined in the system. In present simu-187 lation, the value of A_m is set equal to the $A_m = 10^5$ [29, 30]. The small numerical value of 188 0.001 is used to prevent division by zero. The K is the permeability and C_F is the inertial 189 coefficient of MF. 190

Since, the two models (i) local thermal equilibrium (LTE) and (ii) local thermal non– 191 equilibrium (LTNE), are employed to solve the energy equation for PCM and NCPCM in 192 presence of MF. However, the LTE model is used to solve the energy equation in present 193 to save the computational resource without affecting the solution accuracy, in which the 194 MF and PCM in each cell have the same temperature [29, 32]. Although, the LTNE model 195 provide the more accurate solution compared with the LTE model because of considering 196 the local convective heat transfer coefficient inside pores of porous between the metal-foam 197 and PCM or NCPCM. Moreover, the difference between LTE and LTNE depends on the ini-198 tial and boundary conditions which are more prominent while sensible heating mode rather 199 than phase-change process due to having an most constant temperature. In addition, it is 200 notable that, the LTNE model cannot be applied in 2D and axisymmetric models because 201 of the generation of porous boundaries at the walls between the heat sink and PCM and 202 limitation of coupled boundary condition for it in FLUENT software [31]. Thus, the energy 203 equation with LTE is modelled as follows: 204

Energy conservation:

$$\overline{(\rho c_p)}\frac{\partial T}{\partial t} + (\rho c_p)_{ncpcm} \left(u\frac{\partial T}{\partial x} + v\frac{\partial T}{\partial y}\right) = k_{eff} \left(\frac{\partial^2 T}{\partial x^2} + \frac{\partial^2 T}{\partial y^2}\right) - \varepsilon \rho_{ncpcm} L_{ncpcm} \frac{\partial f_l}{\partial t} \quad (4)$$

The product $\overline{(\rho c_p)}$ is evaluated as the weighted average of the densities of MF and NCPCM as follows:

$$\overline{(\rho c_p)} = [1 - \varepsilon](\rho c_p)_{mf} + \varepsilon (\rho c_p)_{ncpcm}$$
(5)

where, $(\rho c_p)_{mf}$ and $(\rho c_p)_{ncpcm}$ are the volumetric heat capacities of MF and NCPCM, respectively. The L_{ncpcm} represent the latent-heat of fusion of NCPCM.

The effective thermal conductivity (k_{eff}) introduced in LTE model, Eq. 4 is calculated as the volume average thermal conductivities of MF and NCPCM as follows:

$$k_{eff} = (1 - \varepsilon)k_{mf} + \varepsilon k_{ncpcm} \tag{6}$$

where, k_{mf} and k_{ncpcm} are thermal conductivity of MF and NCPCM, respectively. Additionally, f_l is the liquid-fraction during the phase-change in temperature interval of $T_s < T < T_l$ and it varies between 0 (solid) to 1 (liquid), which is defined as:

$$f_l = \frac{\Delta H_{ncpcm}}{L_{ncpcm}} = \begin{cases} 0 & \text{if} \quad T < T_s \\ \frac{T - T_s}{T_l - T_s} & \text{if} \quad T_s \le T \le T_l \\ 1 & \text{if} \quad T < T_l \end{cases}$$
(7)

where, ΔH_{ncpcm} is the fractional latent-heat of the NCPCM that gains during the phasechange process between 0 for solid and L_{ncpcm} for liquid during melting or releases during the solidification process. Thus, when all the NCPCM melts, ΔH_{ncpcm} is equal to the L_{ncpcm} and when the entire NCPCM is still solid, ΔH_{ncpcm} is equal to zero as follow:

$$\Delta H_{ncpcm} = \begin{cases} 0 & \text{if } T < T_m \\ f_l L_{ncpcm} & \text{if } T > T_m \end{cases}$$
(8)

The total enthalpy (H) of the NCPCM is the summation of sensible and latent-heat defined as:

$$H = h + \Delta H_{ncpcm} \tag{9}$$

where, h is the sensible enthalpy which is defined as follows:

$$h = h_{ref} + \int_{T_{ref}}^{T} c_{p_{ncpcm}} dT \tag{10}$$

Energy (heat sink):

$$(\rho c_p)_{HS} \left(\frac{\partial T}{\partial t}\right) = k_{HS} \left(\frac{\partial^2 T}{\partial x^2} + \frac{\partial^2 T}{\partial y^2}\right) \tag{11}$$

Energy (heat source):

$$(\rho c_p)_{hs} \left(\frac{\partial T}{\partial t}\right) = k_{hs} \left(\frac{\partial^2 T}{\partial x^2} + \frac{\partial^2 T}{\partial y^2}\right) + q^{'''}$$
(12)

where, $(\rho c_p)_{HS}$, k_{HS} $(\rho c_p)_{hs}$, and k_{hs} are the thermal capacitances and thermal conductivities of heat sink and heat source, respectively.

223 2.3. MF characterization

For laminar flow regime driven by pressure gradient, the permeability (K) is widely to develop the relationship between the pressure gradient and velocity. The permeability, K and inertial coefficient, C_F are determined using correlation proposed by Calmidi and Mahajan [33] as follows:

$$K = 0.00073(1 - \varepsilon)^{-0.0024} \left(\frac{d_l}{d_p}\right)^{-1.11} d_p^2$$
(13)

228 and

$$C_F = 0.00212(1-\varepsilon)^{-0.132} \left(\frac{d_l}{d_p}\right)^{-1.163}$$
(14)

The other basic parameters such as porosity (ε) , ligament or cell diameter (d_l) , pore diameter or pore size (d_p) , and pore density (ω) are defined to describe the structure of MF. The porosity (ε) is the ratio of the pore volume to the MF total volume of pores and ligaments. The pore density (ω) is the number of pores per linear inch (PPI). The ligament diameter (d_l) can be calculated based on the pore diameter (d_p) as follows [33]:

$$\frac{d_l}{d_p} = 1.18 \sqrt{\frac{1-\varepsilon}{3\pi}} \left(\frac{1}{1-e^{-[(1-\varepsilon)/0.04]}}\right)$$
(15)

where, d_p is calculated as:

$$d_p = \frac{0.0254(m)}{\omega(PPI)} \tag{16}$$

235 2.4. Thermophysical properties of NCPCM

With the addition of Cu nanoparticles, the thermophysical properties of pure PCM are changed by varying the volume fractions. All the effective properties of NCPCM are constant except thermal conductivity and calculated based on the volume fraction of nanoparticles. The effective density (ρ_{ncpcm}), specific heat capacity ($c_{p_{ncpcm}}$), latent-heat (L_{ncpcm}), and thermal expansion coefficient (β_{ncpcm}) of the NCPCM can be calculated using simple theoretical models of mixtures as follows [24–26, 28]:

$$\rho_{ncpcm} = \varphi \rho_{np} + (1 - \varphi) \rho_{pcm} \tag{17}$$

$$c_{p_{ncpcm}} = \frac{\varphi(\rho c_p)_{np} + (1 - \varphi)(\rho c_p)_{pcm}}{\rho_{ncpcm}}$$
(18)

$$L_{ncpcm} = \frac{(1-\varphi)(\rho L)_{pcm}}{\rho_{ncpcm}}$$
(19)

$$\beta_{ncpcm} = \frac{\varphi(\rho\beta)_{np} + (1-\varphi)(\rho\beta)_{pcm}}{\rho_{ncpcm}}$$
(20)

In above Eqs. 17–20, φ is the volume fraction of nanoparticles, the subscripts *ncpcm*, *np* and *pcm* refer to the NCPCM, nanoparticles, and PCM, respectively. The effective dynamic viscosity (μ_{ncpcm}) and thermal conductivity (k_{ncpcm}) of NCPCM are calculated using modelled by the Vajjha et al.[34]:

$$\mu_{ncpcm} = 0.983 e^{(12.959\varphi)} \mu_{pcm} \tag{21}$$

$$k_{ncpcm} = \frac{k_{np} + 2k_{pcm} - 2(k_{pcm} - k_{np})\varphi}{k_{np} + 2k_{pcm} + (k_{pcm} - k_{np})\varphi}k_{pcm} + 5 \times 10^4 \beta_k \zeta \varphi \rho_{pcm} c_{p_{pcm}} \sqrt{\frac{BT}{\rho_{np} d_{np}}} f(T,\varphi) \quad (22)$$

246

where, B is Boltzmann constant which is equal to $1.381 \times 10^{-23} \text{ J/K}$, $\beta_k = 8.4407(100\varphi)^{-1.07304}$,

²⁴⁷ and function $(f(T, \varphi))$ is defined as follows:

$$f(T,\varphi) = (2.8217 \times 10^{-2}\varphi + 3.917 \times 10^{-3})\frac{T}{T_{ref}} + (-3.0669 \times 10^{-2}\varphi - 3.91123 \times 10^{-3})$$
(23)

where, T_{ref} is the reference temperature which is equal to 273.15 K. The first part of Eq. 248 23 relates with Maxwell model to determine the thermal conductivity of solid PCM while 249 second part of Eq. 23 accounts the effects of Brownain motion of nanoparticles, nanoparticles 250 size, volume fraction and temperature dependence. Additionally, ζ is a correction factor 251 which comes in Brownian motion term, because there is no Brownian motion in solid-252 phase. Therefore, the value of ζ is defined as the same as for liquid-fraction, f_l [35]. The 253 thermophysical properties of pure PCM (RT-35HC), Cu nanoparticles, all the MFs (Cu, Al 254 and Ni) are summarized in Table 1. 255

256 2.5. Initial and boundary conditions

The initial and boundary conditions applied in current study are labelled in Fig. 1. The side walls of the heat sink are defined as an adiabatic boundary condition except the top surface which is undergoes the natural convection effect. Following are the initial and boundary conditions applied in this work to solve the governing equations as follows:

1. Initial conditions 261 $t = 0, T = T_{ini} = 296.15 \text{ K}, f_l = 0$ 262 2. Boundary conditions 263 • No-slip condition at walls: u = v = 0264 • Adiabatic walls: 265 $\begin{array}{l} -k \left. \frac{\partial T}{\partial x} \right|_{x=0,W} = 0 \\ -k \left. \frac{\partial T}{\partial y} \right|_{\substack{x=0-10,60-70\\y=0}} = 0 \end{array}$ Along vertical walls 266 At bottom surface 267 • Natural convection: 268 $-k\frac{\partial T}{\partial y}\Big|_{y=H} = h(T - T_{\infty})$ At Top surface 269 • Volumetric heat generation provided from heat source: 270 $-k \left. \frac{\partial T}{\partial y} \right|_{\substack{x=10-60\\y=0-2}} = q^{\prime\prime\prime}$ 271

272 2.6. Numerical methodology

The commercial package of CFD ANSYS-FLUENT 19.1 is used to solve the unsteady 273 simulations. The governing equations of continuity, momentum and energy are solved are 274 discretized by finite volume method (FVM) with double precision. The "melting/solidification" 275 model in connection with local thermal equilibrium porous model is adopted to study the 276 phase-change phenomenon of PCM or NCPCM. The melting/solidification model is based 277 on the enthalpy-porosity method in which the solution is based on a fixed-gird. The gov-278 erning equations are modified in such a manner that they are valid for both solid and liquid 279 phases. The mush-zone where these both phases coexist is treated as "pseudo" porous 280 medium where porosity behaves according to the liquid fraction. A User-defined function 281 (UDF) is written in C++ language to account the temperature dependent k_{ncpcm} of NCPCM 282 due to dispersion of Cu nanoparticles. The PRESSURE-BASED method is selected which 283 is recommended for incompressible flow with high–order Quadratic Upstream Interpolation 284 for Convective Kinematics (QUICK) differencing scheme presented by Leonard [37] to en-285 hance the accuracy of the numerical method. The Semi-Implicit Pressure-Linked Equation 286 (SIMPLE) algorithm was adopted for pressure–velocity coupling by Patanker [38]. The 287 PRESTO (PREssure STaggering Option) scheme was adopted for pressure correction equa-288 tion. The gravitational effect is also considered and second-order upwind difference scheme 289 is selected to discretize convective terms in momentum and energy equations. The under-290 relaxation factors for pressure, velocity, energy and liquid-fraction are set to 0.3, 0.3, 0.8 291 and 0.5, respectively. The convergence criteria are set to 10^{-4} , 10^{-6} and 10^{-8} for continuity, 292 momentum and energy equations, respectively. 293

The grid independence test is also carried out using different gird size of 43753, 48305, 294 54087 and 60796 to avoid its effects on numerical accuracy. The results of melting time and 295 total energy of PCM are summarized in Table 2. The maximum deviation in melting time 296 and total energy is obtained of 0.38% and 0.06% between the elements size of 48305 and 297 54087, respectively. Thus, the grid with the size of 54087 elements is selected for further 298 simulation. Three different time-steps of 0.05, 0.1 and 0.2 s are varied for mesh size of 54087 299 elements and no significance variation is observed. The reason is that PCM upfront velocity 300 and thermal front movement are low, reflecting a low Peclet number and Courant number 301 situation. Therefore, the mesh-size and time-step are 54096 elements and 0.1s, respectively, 302 considered are set in current study. 303

304 2.7. Performance evaluation parameters

To estimate the thermal performance of pure PCM, NCPCM, MF+PCM and NCPCM+MF 305 based heat sink, four different performance evaluation parameters such as heat storage ca-306 pacity (Q), heat storage density (q), rate of heat transfer (Q), and rate of heat transfer 307 density (\dot{q}) along with the total melting time (t_{melt}) . The total Q is defined as the total 308 thermal energy storage capacity during the pre-sensible heating, latent-heat of fusion, and 309 post-sensible heating of pure PCM or NCPCM. Whereas, q indicates the total thermal en-310 ergy storage capacity per unit mass of the pure PCM or NCPCM. Since, the pre-sensible 311 heating and latent-heat are the most significant parameters to determine the Q of pure PCM, 312 NCPCM, MF+PCM and NCPCM+MF based heat sink whiling charging mode. Therefore, 313 Q and q can be defined by Eqs. 24 and 25, respectively, as follows [39]: 314

$$Q = m_{mf} \int c_{p_{mf}} dT + m_{ncpcm} \left(\int_{solid} c_{p_{ncpcm}} dT + \lambda L_{ncpcm} + \int_{liquid} c_{p_{ncpcm}} dT \right)$$
$$\approx m_{ncpcm} [c_{p_{ncpcm}} (T_m - T_i) + f_l L_{ncpcm}] \quad (24)$$

315 and

$$q = \frac{Q}{m_t} = \frac{m_{mf} \int c_{p_{mf}} dT + m_{ncpcm} \left(\int_{solid} c_{p_{ncpcm}} dT + f_l L_{ncpcm} + \int_{liquid} c_{p_{ncpcm}} dT \right)}{m_{mf} + m_{ncpcm}} \approx \frac{m_{ncpcm} [c_{p_{ncpcm}} (T_m - T_i) + f_l L_{ncpcm}]}{m_{mf} + m_{ncpcm}}$$
(25)

Since, the Q and q can only evaluate the storage capacity of pure PCM, NCPCM, 316 MF+PCM and NCPCM+MF based heat sink relative to the mass of PCM or NCPCM. 317 However, there is no relationship of total t_{melt} of PCM or NCPCM with Q and q. Thus, the 318 overall thermal performance of heat sink cannot be evaluate only with Q and q. Therefore, 319 the effect of t_{melt} , m_{mf} , m_{ncpcm} , and Q are combined together to define the rate of heat 320 transfer (Q) and rate of heat transfer density (\dot{q}) . The Q indicates the total thermal energy 321 storage capacity per unit melting time and \dot{q} is defined as total thermal energy storage 322 capacity per unit melting time and per unit mass of MF ad PCM or NCPCM, by Eqs. 26 323 and 27, respectively, as follows: 324

$$\dot{Q} = \frac{Q}{t_{melt}} = \frac{m_{mf} \int c_{p_{mf}} dT + m_{ncpcm} \left(\int_{solid} c_{p_{ncpcm}} dT + f_l L_{ncpcm} + \int_{liquid} c_{p_{ncpcm}} dT \right)}{t_{melt}} \approx \frac{m_{ncpcm} [c_{p_{ncpcm}} (T_m - T_i) + f_l L_{ncpcm}]}{t_{melt}} \quad (26)$$

325 and

$$\dot{q} = \frac{Q}{t_{melt}.m_t} = \frac{m_{mf} \int c_{p_{mf}} dT + m_{ncpcm} \left(\int_{solid} c_{p_{ncpcm}} dT + f_l L_{ncpcm} + \int_{liquid} c_{p_{ncpcm}} dT \right)}{t_{melt}.m_{ncpcm}} \approx \frac{m_{ncpcm} [c_{p_{ncpcm}}(T_m - T_i) + f_l L_{ncpcm}]}{t_{melt}(m_{mf} + m_{ncpcm})}$$
(27)

326 3. Model validation

327 3.1. Experimental validation with PCM based heat sink

The current model is validated with experimental results of a no fin heat sink filled with PCM having the dimension of similar dimensions used in Ashraf et al. [5]. The RT– 35HC was used as a PCM having melting temperature of 35 °C at a input power level of 5W. The results of average temperature of heat sink was compared of both numerical and experimental cases, as shown in Fig. 3a. The results of both numerical and experimental results showed a good agreement and revealed that the present numerical model can be used for further simulations.

335 3.2. Experimental and numerical validation MF/PCM based thermal energy storage

A comprehensive validation is carried with the previous experimental and numerical 336 studies reported by Zhao et al. [16] and Tian and Zhao [17], respectively, and numerical 337 study by Liu et al. [40] by using both LTE and LTNE models of MF+PCM system. The 338 2D system of MF+PCM is adopted for validation proposed in experimental and numerical 339 results of Zhao et al. [16] and Tian and Zhao [17] and as well as Liu et al. [40]. A rectangular 340 geometry having the dimensions of $200 \times 50 \text{ mm}^2$ is selected and a constant heat flux of 1600 341 W/m^2 is provided at the bottom. The RT-58 is used as a PCM and Cu MF having 95% 342 porosity and 10 PPI. Natural convection effects are considered at the side and top walls as 343 considered in Refs. [16, 17] and [40] to get the more better results. The result of temperature 344 variations are presented at a height of 8 mm in comparison with the Zhao et al. [16] and 345

Tian and Zhao [17], as shown in Fig. 3b. A excellent agreement can be seen between the 346 LTNE model of present study with experimental and numerical results with Refs. [16, 17] 347 and numerical results of Ref. [40]. The results of LTE model also show a good agreement 348 with LTE model results reported by Liu et al. [40]. The variations in results of numerical 349 and experimental studies reported in Zhao et al. [16] and Tian and Zhao [17], respectively, 350 are because of considering a constant melting temperature in numerical study, as reported 351 in Liu et al. [40]. Therefore, in present study and the study reported by Liu et al. [40] use 352 the different solidus (T_s) and liquidus (T_l) temperatures for the simulations. The maximum 353 deviation of ± 4.2 °C is obtained between the current study results and experimental results 354 of Zhao et al. [16]. In addition, it is always difficult to justify the discrepancy in results, 355 shown in Fig. 3b, because the data is taken out from the experimental and numerical 356 results of electronic copies of Refs [16, 17, 40]. Although, the LTNE model can predict 357 the T_{HS} and f_l more accurately compared to the LTE model. However, due the to higher 358 computational cost of LTNE model in porous–PCM medium, especially in 3D solid–liquid 359 cases, LTE model is employed for further analysis. It has been revealed in previous studies, 360 the difference between LTE and LTNE models is very small due to the primary domination 361 of heat conduction rather than the natural heat convection effect [41]. 362

³⁶³ 4. Results and discussion

364 4.1. Effect of MF materials

The average heat sink temperature (T_{HS}) and melt-fraction (f_l) variations for PCM, 365 Cu, Al and Ni MF+PCM cases are shown in Fig. 4a. The uniform and non–uniform 366 melting of PCM can been seen clearly for PCM and MF+PCM embedded heat sinks cases. 367 At a constant input power level, the transient variation of T_{HS} for each case shows that 368 MF+PCM embedded heat sinks have the significant reduction in heat sink temperature, as 369 shown in Fig. 4a. The reduction in T_{HS} is due to uniform heat transfer from the heat sink 370 base towards the ambient due to conjugate heat transfer mode. This reveals that a heat 371 sink embedded with MF+PCM has the better heat transfer performance. Thus effective 372 cooling performance is achieved compared to a heat sink of pure PCM filled. A closer look of 373 temperature variation between the different materials of MF reveals that Cu MF+PCM heat 374 sink shows the better reduction in T_{HS} compared to the Al and Ni MF+PCM embedded 375 heat sinks. The reduction in heat sink temperature by Cu MF heat sink is because of the 376 higher thermal conductivity of Cu, followed by Al and Ni. The variations in f_l of all cases, 377



Figure 3: Validation of present simulation (a) with experimental results based on PCM case and (b) experimental results by Zhao et al. [16] and Tian and Zhao [17], and numerical results by Liu et al. [40] with MF case.

shown in Fig. 4a, reveals that the higher rate of f_l is obtained for the MF cases compared to the pure PCM case because of the higher heat transfer area by MF+PCM which strengthen the conduction heat transfer. The melting rate is different between the PCM and MF+PCM cases. The variation in f_l of MF embedded heat sinks is slightly varying and the highest f_l is obtained by Cu MF heat sink, followed by Al and Ni MF heat sinks. This trend is expected since Cu has the higher thermal conductivity compared to Al and Ni. The evolution of t_{melt} ,

Q and \dot{Q} of all cases is shown in Fig. 4b. It can be seen that the t_{melt} of PCM is decreased 384 by embedding the MF+PCM and lower time of 44.17 min is obtained with Cu MF+PCM 385 compared to PCM, Al and Ni MF+PCM filled heat sink cases by obtaining 53.42, 44.92, 386 and 47.08 mins, respectively. The reduction in t_{melt} is obtained of -17.32%, -15.91% and 387 -11.86% for Cu, Al and Ni MF+PCM cases compared to the pure PCM case. The similar 388 trend is obtained in Q for pure PCM and MF+PCM cases. The Q of 279.78, 266.15, 266.27 389 and 270.76 kJ are obtained for PCM, Cu, Al and Ni MF+PCM heat sinks, respectively. The 390 reductions in Q due to embedding the MFs are achieved of -4.87%, -4.83% and -3.22%391 for Cu, Al and Ni MF+PCM heat sinks, respectively. However, the enhancement in \dot{Q} is 392 obtained with MF+PCM compared to the pure PCM based heat sink. The \dot{Q} for pure 393 PCM, Cu, Al and Ni MF+PCM heat sinks is obtained of 80.28, 86.84, 86.31 and 84.30 W, 394 respectively. It can be revealed that highest enhancement of 8.16% in \dot{Q} is obtained with 395 Cu MF+PCM compared to the Al and Ni MF+PCM of 7.51% and 5.01%, respectively, 396 compared to the pure PCM filled heat sink, because of the higher thermal conductivity. 397 Thus, it can be suggested that a PCM filled heat sink with Cu MF+PCM shows the better 398 thermal cooling performance. Since, the Fig. 4 presents thermal cooling performance in 399 terms of T_{HS} , f_l , t_{melt} , Q and \dot{Q} , however the further comparison of melting phenomenon 400 of pure PCM and all MF+PCM embedded heat sinks is shown in Fig. 5. The f_l contours 401 are presented at different flow times of 900, 1500, 2100, 2700 and 3300 s. It can be seen 402 clearly that by using MF+PCM, the melting pattern of PCM melting changes significantly. 403 In addition, the uniform and faster melting of PCM is obtained with the case of MF+PCM 404 embedded heat sinks especially in case of Cu MF+PCM followed by Al and Ni MF+PCM 405 heat sinks. Initially, at 900 s, the layers of solid-liquid interface (i.e. heat transfer area) 406 can been seen clearly through blue and red colours, respectively, for a PCM heat sink case. 407 Whereas a uniform melting patterns can be seen with MF+PCM heat sinks by showing no 408 visible solid-liquid interface. With the increase of melting time, the gradually solid-liquid 409 interface can be seen showing the effect of buoyancy and gravitational forces. A uniform 410 melting is observed in case of MF heat sinks due the conduction and natural convection 411 heating modes. At 3300 s, the complete melting of PCM is observed in case of MF+PCM 412 whereas there is still small tiny portion of solid PCM, which shows that the addition of MFs 413 with PCM improves the heat transfer rate and melting process uniformly. 414



Figure 4: Effect of different MF materials: (a) T_{HS} and f_l , and (b) t_{melt} , Q and \dot{Q} .

415 4.2. Effect of power levels

Fig. 6 presents the effect of different power levels of a Cu MF+PCM embedded heat 416 sink. The T_{HS} and f_l results are presented in Fig. 6a. Whereas, the comparison of t_{melt} and 417 Q at different power levels are presented in Fig. 6b. It is can be seen that latent-heating 418 phase duration of PCM is reduced and T_{HS} is increased as the input power level increase, 419 as expected. In addition at lower input power of 4 W, there is no phase changing occurred 420 even after 3600 s and supplied heat is absorbed by the PCM due to its higher latent-heat 421 enthalpy. The maximum T_{HS} and f_l of 309.62 °C and 0.941 are obtained at lower power 422 level of 4 W. The comparison of t_{melt} and \dot{Q} of Cu MF+PCM embedded heat sink shows 423 that with the increase of power level, the t_{melt} of PCM is decreased whereas Q increases, as 424 shown in Fig. 6b. The t_{melt} of PCM is obtained of 44.17, 36.83, 31.58, and 27.75 mins for 5, 425 6, 7, and 8 W, respectively. Similarly, the \dot{Q} is obtained of 86.84, 104.34, 121.81, and 139.19 426

Liquid Fraction



Figure 5: Variation of f_l at various t of PCM, Cu, Al and Ni MF+PCM embedded heat sinks.

⁴²⁷ W for 5, 6, 7, and 8 W, respectively. The higher \dot{Q} is because of the lower t_{melt} of PCM ⁴²⁸ filled in MF heat sink. Furthermore, it can be suggested that at lower power level, the T_{HS} ⁴²⁹ does not rise enough to melt the PCM. Therefore, a MF+PCM embedded heat sink has less ⁴³⁰ advantage for passive thermal management of electronic devices at lower power level.



Figure 6: Effect of different power levels of MF+PCM filled heat sink: (a) T_{HS} and f_l , and (b) t_{melt} and \dot{Q} .

431 4.3. Effect of PPI

Fig. 7 shows the effect of changing the PPI or pore density of MF embedded in a PCM filled heat sink. The five different values of 10, 20, 30, 40 and 50 PPI are investigated of Cu MF+PCM heat sink a constant porosity of 95% and input power level of 5 W. A slight variation in t_{melt} of PCM is obtained by changing the PPI of MF. The t_{melt} of PCM is obtained of 44.17, 44.08, 44.0, 43.92, and 43.83 mins for 10, 20, 30, 40 and 50 PPI, respectively. It

can be revealed that a small variation in t_{melt} and temperature distribution is achieved by 437 changing the PPI of MF under local thermal equilibrium heat transfer conditions. Further-438 more, a closer look of t_{melt} result reveals that a MF with lower PPI enhances the t_{melt} of 439 MF+PCM filled heat sink resulting in reduces the base temperature of the heat sink. The 440 higher number of PPI means the unit cell is divided into higher number of small pore which 441 reduce the effect of low thermal conductivity of the PCM. In addition, a higher value of PPI 442 contains a smaller size of a pore which possesses the higher surface area per unit volume. 443 Thus, higher heat transfer rate exists between the PCM and MF ultimately reduces the t_{melt} 444 of the PCM. Contrarily, the higher PPI suppresses the natural convection effect of PCM 445 melting with the MF because of the high thermal resistance in the PCM flow direction. 446 Thus, the influence of PPI is negligible for a MF+PCM embedded heat sink under local 447 thermal equilibrium heat generation conditions which has also been reported in literature 448 [40, 42]. As shown in Fig. 7 that negligible effect of PPI is achieved on the variation of f_l . 449 The higher f_l is obtained by increasing the PPI from 10 to 50. 450



Figure 7: Effect of different PPI: comparison of PCM t_{melt} .

451 4.4. Effect of Porosities

Fig. 8 illustrates the effect of different porosities (95%, 90%, and 80%) of Cu MF+PCM embedded heat sink compared with the pure PCM filled heat sink with 10 PPI at 5W. The result of T_{HS} and PCM f_l as a function of time are presented in Fig. 8a for MF+PCM and PCM cases. It can be seen clearly that T_{HS} reduces significantly by embedding the MF

inside the heat sink. In addition, a close look of the T_{HS} and f_l results of the porosities 456 95%, 90%, and 80% show that a Cu MF+PCM based heat sink of 95% porosity has lower 457 T_{HS} and f_l . The minimum T_{HS} of 323.09 °C is obtained with 95% MF+PCM heat sink 458 compared with 90%, 80% and pure PCM heat sink cases. More further, a lower f_l is obtain 459 with 95% MF+PCM heat sink compared with 90% and 80% MF+PCM cases. Since, the 460 complete meting of PCM occurs after 3065 s. By increasing the porosity of MF, as a result 461 reduced the amount of higher thermal conductivity MF inside the heat sink, thus higher 462 PCM melting time is achieved due to lower rate of heat transfer, as expected. 463

Fig. 8b shows the results of t_{melt} , the amount of Q and Q of all cases of PCM and MF+PCM 464 filled heat sinks. The t_{melt} and Q are reduced with the reduction of porosity since higher 465 amount of MF inside the heat sink exhibits the higher effective thermal conductivity of the 466 PCM result in increases the heat transfer rate. The reduction in t_{melt} and Q are obtained of 467 -17.32% and -4.87%, -20.28% and -5.44%, and -22.93% and -5.46% for 95%, 90%, and 468 80% MF+PCM, respectively, compared with PCM filled heat sinks. Whereas, in comparison 469 with PCM filled heat sink, the \dot{Q} is increased of 8.16%, 10.77% and 13.22% with 95%, 90%, 470 and 80% MF+PCM cases, respectively. Therefore, it can be suggested that a heat sink with 471 constant mass of PCM, the higher porosity of 95% is favourable for passive thermal cooling 472 of electronics, since it prolongs the PCM t_{melt} duration and lower the T_{HS} . 473

474 4.5. Effect of NCPCM

Fig. 9 reveals the effect of nanoparticles dispersion in PCM filled heat sink by varying 475 the loading contents of 1%, 3% and 5%. The addition of higher thermal conductivity 476 nanoparticles improve the thermal conductivity of pure PCM on the cost of decreasing the 477 latent-heat of fusion and specific heat capacity which is not preferable for passive cooling 478 of electronic components. The results of T_{HS} and f_l , as shown in Fig. 9a, reveal that T_{HS} 479 is decreased initially but reaches higher as the volume fraction of nanoparticles increases. 480 Further, the higher f_l is obtained with the increase of nanoparticles loadings which improve 481 the PCM melting rate. Since the amount of PCM increases with the increase of nanoparticles 482 loadings, the time of complete PCM melting is almost constant. The t_{melt} of PCM is obtained 483 of 52.75, 52.45 and 52.17 mins for 1%, 3% and 5% volume factions of Cu nanoparticles, 484 respectively, as shown in Fig. 9b. The t_{melt} is reduced of -1.25%, -1.87%, and -2.34%485 by 1%, 3% and 5% concentrated NCPCM heat sink, respectively, compared with PCM 486 filled heat sink. Similarly, the results of Q and \dot{Q} are almost negligible. The Q and \dot{Q} are 487



Figure 8: Effect of different porosities: (a) T_{HS} and f_l , (b) t_{melt} , Q and \dot{Q} .

increased initially at 1% of Cu nanoparticles concentration, however, these are decreased for 488 later on 3% and 5% NCPCM cases which is due to the increase of PCM mass by adding the 489 nanoparticles. Since, the system has a close boundary so the heat sink volume is constant 490 however, by the addition of nanoparticles the thermophysical properties such as density, 491 latent-heat of fusion, thermal expansion coefficient, specific heat capacity, viscosity and 492 volumetric heat capacity of PCM are changed which also effect the thermal performance 493 of heat sink. The \dot{Q} is enhanced by 1.35%, 0.76%, and 0.19% by adding 1%, 3% and 5% 494 volume concentration of Cu nanoparticles, respectively, compared with the PCM filled heat 495 sink case. Similarly, the variations in \dot{Q} are obtained of 0.34%, -0.69%, and -1.68% for 1%, 496 3% and 5% NCPCM cases, respectively, compared with PCM filled heat sink. It can been 497 reveal that Q and best Q are obtain at 1% of Cu nanoparticles concentration. Therefore, 498 it is recommended that by adding 1% loading of nanoparticles has the optimum results in 499

terms of thermal cooling performance because the higher concentrations can lead toward 500 the agglomeration and sedimentation of nanoparticles in real time system. Additionally, 501 these are also reduced the latent-heat fusion and increase t_{melt} of PCM. A rapid decreasing 502 trend can observed in results of q and \dot{q} because of increase of the PCM mass with the 503 increase of Cu nanoparticles concentrations, as shown in Fig. 9b. By adding the 1%, 3%504 and 5% loadings of Cu nanoparticles the q is decreased of -8.63%, -23.27%, and -34.04%, 505 respectively, compared with PCM filled heat sink case. In similar manner, the \dot{q} for NCPCM 506 filled heat sinks is reduced by -7.70%, -22.16%, and -32.79%, respectively, compared with 507 PCM filled heat sink. 508

For better understanding the effect of NCPCM heat sinks of different volume fractions, 509 Figs. 10a and 10b illustrate the results of k_{eff} and k_{eff} enhancement, respectively, for 510 PCM and NCPCM filled heat sinks. It can be seen that k_{eff} increases with the increase 511 of Cu nanoparticles concentration with respect to time. More further, it has been reported 512 that k_{eff} also increases and decreases with respect to temperature during melting and 513 solidification processes, respectively, with the increase of nanoparticles loading by using 514 Equation 22 [28]. After 3600 s, the maximum enhancement in k_{eff} are obtained of 17.99%, 515 24.33%, and 31.50% for 1%, 3% and 5% loadings of Cu nanoparticles. 516

Further, the distribution of T_{HS} at the surface of heat sink in vertical direction is shown in 517 Fig. 11 by representing the isotherm contours at different time intervals for different φ of 518 Cu nanoparticles. The T_{HS} is increased gradually with the increase of operating time and φ 519 during the melting process. Initially, at the 1200 s, the lower to higher variation in T_{HS} can 520 be seen for all cases of PCM and NCPCM filled heat sinks over the surface because of the 521 conduction and natural convection modes. A clear observation of conduction heat transfer is 522 reflected at the boundaries of PCM and heat sink due to the temperature gradient. During 523 the heating process, a uniform the distribution of T_{HS} isotherms is seen between the solid-524 liquid interfaces of both PCM and NCPCM filled heat sinks which that the conduction heat 525 transfer mode is more dominant than convection mode. With the increase of time from 1800 526 to 3000 s, the both conduction and natural convection heat transfer modes are observed from 527 the isotherms for all heat sink cases by seeing the deformation in PCM shape, which shows 528 the initiating of natural convection inside the heat sink. In addition, the circulating pattern 529 of isotherms are observed at the bottom of the heat sink because of the buoyancy effects 530 and gravity force. The more dominant convection patterns of isotherms are obtained with 531 the increase of time, developed by the temperature gradient, because of the growing role of 532

533 convection heat transfer in melt zone.

The further visualization of melting phenomenon of PCM and NCPCM filled heat sink 534 cases is illustrated using f_l contours presented in Fig. 12. Initially, at 1200 s, there are 535 clear zones of solid and liquid PCM representing by blue and red colours, respectively, and 536 layers of PCM melting can be observed at the bottom and sides walls for all heat sink cases. 537 With the increase of melting time, at 1800 and 2400 s, the clear observation of circulating 538 patterns of liquid PCM is found at the bottom of the heat sink due the effect of buoyancy 539 and gravity forces. Since, the addition of nanoparticles enhance the thermal conductivity of 540 NCPCM as the well as the viscosity of NCPCM which enhance the heat transfer rate and 541 also affects the melt movement of PCM. Therefore, conduction heat transfer mode dominates 542 over convection mode. In addition, the significance of natural convection is noticeable by 543 appearing the more deformation and size of rotating circles of melted PCM during melting 544 process of NCPCM. Moreover, a regular decrease in circulating patterns is observed with 545 the increase of nanoparticles volume fraction. A closer look reveals that relative cold PCM 546 moves downward from solid-liquid interface because of gravitational effect which improves 547 the complete melting of PCM. This movement of melted PCM enhances the rate of PCM 548 melting at the bottom half of the heat sink compared to the upper half. Later on during 549 the melting process, at 3000 s, the higher rate of f_l of NCPCM is obtained in most of 550 the part of heat sink domain which shows the dominant contribution of natural convection 551 heat transfer because of the influence of buoyancy effects. There is still movement of cold 552 or relative less melted PCM towards the bottom because of gravity effects. At 3600s, the 553 complete melting of NCPCM is obtained for 1%, 3%, and 5% φ because of conduction and 554 natural convection contribution. Since, the addition of nanoparticles improves the thermal 555 conductivity of PCM, thus, it improves the conductive heat transfer rate within the PCM 556 and faster melting is achieved. 557

558 4.6. Effect of different nanoparticles concentrations with constant MF

Fig. 13 presents the effect of varying Cu nanoparticles volume concentrations in presence of MF+PCM filled heat sink. The constant porosity and PPI of 95% and 10 are selected, respectively, and three different volume fractions of 1%, 3% and 5% of Cu nanoparticles are investigated individually. Since it has been explored that the MF having 95% and 10 porosity and PPI, respectively, has the best thermal cooling performance. Thus, effect of nanoparticles concentrations are investigated further with MF+PCM embedded heat sink.



Figure 9: Effect of PCM and NCPCMs filled heat sink: comparison of (a) T_{HS} and f_l , (b) t_{melt} , Q, \dot{Q} , q and \dot{q} .

Fig. 13a shows the results of T_{HS} and f_l for PCM, MF+PCM and MF+NCPCM heat sink 565 cases at a constant input power level of 5 W. The lower T_{HS} and higher f_l are obtained 566 for both MF+PCM and MF+NCPCM heat sink cases. Furthermore, a closer look reveals 567 that MF+NCPCM heat sink has the higher f_l and less latent-heat phase-change duration 568 result in increases the T_{HS} after complete melting of PCM. The melting durations of PCM 569 are obtained of 53.42, 44.17, 43.75, 42.92, and 42.08 mins for the case of PCM, MF+PCM, 570 MF+NCPCM (1%), MF+NCPCM (3%), and MF+NCPCM (5%), respectively, as shown 571 in Fig. 13b. A reduction in PCM t_{melt} was obtained of -17.32%, -18.10%, -19.66%, 572 and -21.22% for MF+PCM, MF+NCPCM (1%), MF+NCPCM (3%), and MF+NCPCM 573 (5%), respectively, compared with PCM filled heat sink. Since, both MF and nanoparticles 574 have the higher thermal conductivity which results in enhance the heat transfer rate within 575 PCM and reduced the t_{melt} . Similarly, a decreasing trend in results of Q are obtained and 576



Figure 10: Results of (a) k_{eff} (b) k_{eff} enhancement of NCPCMs.

a reduction of -4.87%, -5.68%, -7.19%, and -8.70% for MF+PCM, MF+NCPCM (1%), 577 MF+NCPCM (3%), and MF+NCPCM (5%), respectively, compared with PCM filled heat 578 sink. It can be seen that a slight variation in \dot{Q} of MF+PCM and MF+NCPCM heat sinks 579 cases. The \dot{Q} sharply increases by adding the MF however, by adding the Cu nanoparti-580 cles of 1%, 3%, and 5% it is slightly decreased which is less significant in overall thermal 581 performance. The enhancement in \dot{Q} is obtained of 8.16%, 8.12%, 7.99%, and 7.86% for 582 MF+PCM, MF+NCPCM (1%), MF+NCPCM (3%), and MF+NCPCM (5%), respectively, 583 compared with PCM filled heat sink. It can be seen from Fig. 13b that sharp decreasing 584 trend is obtained in results of q and \dot{q} for the case of MF+NCPCM heat sinks which is due 585 to increase of the total mass of the PCM by adding the different concentrations of nanopar-586 The q is reduced by -4.87%, -14.11%, -28.30%, and -38.75% for MF+PCM, ticles. 587 MF+NCPCM (1%), MF+NCPCM (3%), and MF+NCPCM (5%), respectively, compared 588 with PCM filled heat sink. The result of \dot{q} presents that \dot{q} increases by 8.16% adding the MF 589



Figure 11: Variation of isotherms contours at various t of PCM filled heat sink and various nanoparticles concentrations of NCPCM based heat sinks.





Figure 12: Variation of f_l at various t of PCM filled heat sink and various nanoparticles concentrations of NCPCM based heat sinks.

⁵⁹⁰ however it is decreased by -1.54%, -16.57%, and -27.64% by adding the Cu nanoparticles ⁵⁹¹ of 1%, 3%, and 5% concentration, respectively, in MF because of the increase of the total ⁵⁹² mass of PCM.

The comparison of MF+PCM and MF+NCPCM isotherms are further presented in Fig. 593 14 for different time intervals of 1200, 1800, 2400 and 3000 s. It can be seen that the 594 temperature increases with the increase of time and with the increase of Cu nanoparticles 595 φ . The MF provides a constant heat transfer path from bottom to top of the heat sink 596 due to interconnected MF structure. Thus, the main contribution of heat transfer is be-597 cause of heat conduction. In addition, nanoparticles have the higher thermal conductivity 598 which further improve the heat transfer enhancement with the increase of added amount. 599 A closer look reveals that in combination of nanoparticles and MF, the significance heat 600 transfer contribution is because of the MF. Further more, the comparison of NCPCM and 601 MF+NCPCM isotherms, shown in Figs. 11 and 14, respectively, reveals that the addition of 602 MF in PCM provide a more uniform heat distribution compared to the NCPCM heat sink. 603 Furthermore, Fig. 15 shows the f_l contours of MF+PCM and MF+NCPCM heat sink at 604 different time periods of 1200, 1800, 2400 and 3000 s. The melting behaviour of PCM shows 605 the uniform melting because of the addition of MF with increase of time. However, the 606 higher f_l contours can be seen the addition of nanoparticles for each time step especially at 607 2400 s. The PCM melting improves as the φ increases from 1% to 5% because of the higher 608 thermal conductivity of Cu nanoparticles. Since, the both MF and nanoparticles have the 609 thermal conductivity, thus, improves the melting of the PCM resulting in lower t_{melt} and 610 latent-heating phase is obtained. However, the higher heat transfer rate is achieved with 611 the addition of MF. Therefore, by comparing the results, shown in Figs. 13b, 11 and 15, it 612 can be suggested the nanoparticles having 1% volume fraction is enough in combination of 613 MF embedded heat sink for effective thermal cooling performance. 614

⁶¹⁵ 4.7. Effect of different MF porosities with constant nanoparticles concentration

Fig. 16 illustrates the effect of varying MF porosities (85%, 90% and 95%) with a constant nanoparticles concentration of 5% and PPI of 10 at 5 W. The results of T_{HS} and f_l of PCM, NCPCM and MF+NCPCM heat sinks are presented in Fig. 16a. It can be revealed that lower T_{HS} is achieved initially before the complete melting of PCM for the cases of NCPCM and MF+NCPCM heat sinks which is due the effective heat transfer from the heat sink base towards ambient in presence of high thermal conductive nanoparticles



Figure 13: Effect of different nanoparticles concentrations with constant MF+PCM based heat sink: (a) T_{HS} and f_l , (b) t_{melt} , Q, \dot{Q} , q and \dot{q} .

and porous medium with PCM. However it can be seen that MF+NCPCM heat sink of 622 5% Cu concentration and 95% porosity has the lowest T_{HS} of 326.11 °C and higher latent-623 heating phase completion duration after 3600 s compared with the NCPCM (5%) heat sink. 624 This reveals that a MF+PCM heat sink has the better thermal performance compared with 625 PCM-only and NCPCM heat sinks. Similarly, higher f_l is obtained for MF+NCPCM heat 626 sinks compared with PCM and NCPCM heats sink. However, a closer analysis presents that 627 NCPCM heat sink has the higher f_l compared with PCM-only heat sink results in shorter 628 the melting time of PCM. Moreover, a 95% porosity and 5% nanoparticles concentration 629 heat sink has the lower f_l which takes the higher time to melt the PCM completely compared 630 with 90% and 85% porosities MF+NCPCM heat sinks. The t_{melt} for PCM, NCPCM and 631 MF+NCPCM of 95%, 90% and 85% porosities is obtained of 53.42, 52.17, 41.50, 40.08 and 632 38.83 mins, respectively, as shown in Fig. 16b. The reduction in t_{melt} is obtained of -2.34%, 633



Figure 14: Variation of isotherms at various t of MF+PCM and MF+NCPCM heat sink of different nanoparticles volume fractions.

Liquid Fraction



Figure 15: Variation of f_l at various t of MF+PCM and MF+NCPCM heat sink of different nanoparticles volume fractions.

-22.31%, -24.96%, and -27.30% for NCPCM, MF+NCPCM (95%), MF+NCPCM (90%), 634 and MF+NCPCM (85%), respectively, compared with PCM filled heat sink. The results of 635 Q and q reveal the decreasing trend for both NCPCM and MF+NCPCM filled heat sink 636 compared with PCM filled heat sink. However, it can be observed that the variations in 637 Q and q for MF+NCPCM heat sink cases are very less significant. The reduction in Q 638 is achieved of -1.68%, -9.23%, -9.52%, and -9.64% for NCPCM, MF+NCPCM (95\%), 639 MF+NCPCM (90%), and MF+NCPCM (85%), respectively, compared with PCM filled 640 heat sink. Similarly, the reduction in q is obtained of -34.04% for NCPCM heat sink, how-641 ever, a slight decrease is obtained of -39.10%, -39.30%, and -39.38% for MF+NCPCM 642 heat sink of 95%, 90%, and 85% porosities, respectively, compared with PCM filled heat 643 sink case. Fig. 16b, a increasing trend is observed in \dot{Q} for NCPCM and MF+NCPCM 644 heat sinks. A slight enhancement of 0.19% for NCPCM heat sink, however, a sharp en-645 hancement of 7.97%, 10.44%, and 12.85% is obtained for MF+NCPCM heat sink of 95%, 646 90%, and 85% porosities, respectively, compared with PCM filled heat sink. Comparably, 647 the reduction in \dot{q} results is obtained of -32.79%, -27.57%, -25.91%, and -24.28% for 648 NCPCM, MF+NCPCM (95%), MF+NCPCM (90%), and MF+NCPCM (85%), respec-649 tively, compared with PCM filled heat sink. The result illustrates that only NCPCM heat 650 sink reduced the q and \dot{q} more significantly as compared to MF+NCPCM heat sink. Since, 651 the q and \dot{q} depend on the mass of the PCM which means that more the PCM mass more 652 the decrease in q and \dot{q} or vice versa. 653

⁶⁵⁴ 5. Concluding remarks and suggestions

The present study explores the parametric investigation of a two-dimensional (2D) heat 655 sink filled up with the composite of PCM, NCPCM, MF+PCM and MF+NCPCM through 656 numerical approach. The influence of Cu nanoparticles is evaluated by varying the volume 657 fraction with PCM and MFs. The effect of different parameters such as different MF mate-658 rials, porosities, pore densities, volume fractions of NCPCM, combination of MF+NCPCM 659 by varying different porosities and nanoparticles volume fractions, and power levels. The 660 contours of liquid-fraction and isotherms are presented at different time periods and volume 661 fractions of Cu nanoparticles. Thermal performance of heat sink is evaluated using different 662 performance indicators such as melting time, heat storage capacity, heat storage density, 663 rate of heat transfer, and rate of heat transfer density. The key findings from results are 664 identified as follows: 665



Figure 16: Effect of different MF porosities with constant nanoparticles concentration NCPCM based heat sink: (a) T_{HS} and f_l , (b) t_{melt} , Q, \dot{Q} , q and \dot{q} .

- A uniform heating and melting of PCM is observed b by embedding the MF with PCM filled heat sink. The lower heat sink temperature and enhanced melting rate is obtained by Cu MF followed by Al and Ni MFs. The higher heat transfer rate of 86.84 W and lowest melting time of 44.17 min are obtained Cu MF+PCM embedded heat sink.
- The melting time and latent-heating phase duration were decreased with the increase
 of power level. However, the rate of heat transfer increased with the increase of power
 level.
- The pore size showed a slight variation in reduction of heat sink temperature and
 melting of PCM. However, the 10 PPI showed the lower heat sink temperature and
 higher melting time. In addition, the comparison of porosities revealed the highest
 porosity of 95% showed the lowest heat sink temperature and liquid-fraction of PCM.

The higher reduction melting time and heat storage capacity were obtained of -17.32%and -4.87%, respectively, however, lower rate of heat transfer was increased of 8.16%with 95\% porosity and 10 PPI Cu MF+PCM based heat sink.

• There are less significance effects observed by adding the nanoparticles inside the PCM 681 in reduction of heat sink and PCM temperatures. The lower heat sink and NCPCM 682 temperatures were achieved with the increase of volume fraction of nanoparticles. 683 The addition of nanoparticles improved the thermal conductivity and viscosity of 684 PCM with the increase of volume fraction and improved the uniformity in melting 685 process. The melting time was reduced by -1.25%, -1.87% and -2.34% with the 686 addition of 1%, 3% and 5% volume fraction of Cu nanoparticles. Increasing trend was 687 observed in heat storage capacity and rate of heat transfer at 1% volume fraction of Cu 688 nanoparticles. Later on, they were decreased. A rapid decrease in heat storage density 689 and rate of heat transfer density was obtained. The rate of heat storage is enhanced 690 by 1.35%, 0.76%, and 0.19% by adding 1%, 3% and 5% volume concentration of Cu 691 nanoparticles, respectively, compared with the PCM filled heat sink case. 692

- The addition of nanoparticles enhanced the melting rate and thermal conduction mode of PCM with the increase of volume fraction because of the enhancement in effective thermal conductivity and viscous effects of NCPCM. Addition of nanoparticles improved the uniformity in melting process.
- With the addition of the NCPCM and MF, lower heat sink temperature and higher 697 liquid-fraction were obtained. Additionally, the latent-heating phase duration was 698 decreased with the increase of Cu nanoparticles volume fraction. The reduction in 699 melting time was obtained of -18.10%, -19.66% and -21.22% by adding the 1%, 3%700 and 5% volume fraction of Cu nanoparticles with 95% porous Cu MF. Higher rate of 701 heat transfer of 8.12% was obtained at 1% volume fraction of Cu nanoparticles. How-702 ever, a sharp decrease in heat storage density and rate of heat transfer density because 703 of increase in total mass of the PCM due to addition of nanoparticles. A uniform tem-704 perature distribution and melting phenomenon was observed through isotherms and 705 liquid-fraction contours. 706
- The different porosities (85%, 90% and 95%) of MF with constant volume fraction of 5% revealed that lower heat sink temperature and higher latent-heating phase

completion duration. The lower liquid-fraction was obtained with 95% porous of MF 709 and 5% volume fraction of Cu nanoparticles. The reduction in melting time was 710 obtained of -2.34%, -22.31%, -24.96% and -27.30% for for NCPCM, MF+NCPCM 711 (95%), MF+NCPCM (90%), and MF+NCPCM (85%), respectively, compared with 712 PCM-only case. The slight improvement of 0.19% for NCPCM heat sink, however, a 713 sharp enhancement of 7.97%, 10.44%, and 12.85% is obtained for NCPCM/MF+PCM 714 heat sink of 95%, 90%, and 85% porosities, respectively, compared with pure PCM 715 filed heat sink. 716

After carefully analysis of the pure PCM, NCPCM, MF+PCM and MF+NCPCM based heat sink, it can been reveal that optimum heat transfer rate and heat absorption capacity are obtained at 1% of Cu nanoparticles concentration with 95% porosity and 10 PPI Cu MF heat sink for efficient thermal cooling performance of electronic devices.

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726 Conflict of interest

The authors declare no conflict of interest regarding this research article.

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Nom	Nomenclature							
Abbreviations			Rate of heat transfer density (W/Ka)					
Al	Aluminum	a‴	Volumetric heat generation (W/m^3)					
Cu	Copper	$\frac{q}{S}$	Source term in momentum equation					
FVM	Finite volume method	\tilde{T}	Temperature (K)					
HS Heat sink		+	Time (sec)					
NCPO	M Nanocomposite phase change material	u	Velocity component in x -axis					
Ni	Nickel		(m/s)					
DFBM Darcy-Forchheimer-Brinkman model		v W	Velocity component in y -axis (m/s) Width (mm)					
LTE	Local thermal equilibrium	c_p	Specific heat capacity $(J/kq,K)$					
LTNE	Local thermal non-equilibrium	ΔH	Fractional latent-heat $(J/kq.K)$					
PCMs Phase change materials		2D	Two dimensional					
PPI	Pores per inch	Gree	reek letters					
ТМ	Thermal management	φ	Volume fraction					
UDF	User–defined function	β	Thermal expansion coefficient $(1/K)$					
Symbols		μ	Viscosity (Pa.s)					
A_m	Mushy zone	f_l, λ	Liquid fraction					
В	Boltzman constant (J/K)	ε	Porosity					
ρc_p	Volumetric heat capacity $(J/m^3.K)$	ω	Pore density					
C_F	Inertial coefficient	Subscripts						
K	Permeability	HS	Heat sink					
d_l	Ligament or cell diameter (m)	hs	Heat source					
d_p	Pore diameter or pore size (m)	ini	Initial					
g	Gravitational acceleration $({\rm m/s^2})$	l	Liquidus					
H	Height (mm)	m	Melting					
Q	Heat storage capacity (J)	mf	Metal-foam					
q	Heat storage density (J/Kg)	ncpcn	<i>ncpcm</i> Nanocomposite phase change mate-					
k	Thermal conductivity $(W/m.K)$		rial					
L	Latent heat of fusion $(J/kg.K)$	np	Nanoparticles					
m	Mass (Kg)	ref	Reference					
p	Pressure (Pa)	x	x-axis					
\dot{Q}	Rate of heat transfer (W)	y	y-axis					

Divisional proportion	RT–35HC	Nanoparticles	Metal-foam	Metal-foam	Metal-foam
r nysicar properties		(Cu)	(Al)	(Cu)	(Ni)
T_m (K)	308	-	-	-	-
T_s (K)	307	-	-	-	-
T_l (K)	309	-	-	-	-
L (kJ/kg)	240	-	-	-	-
k (W/m.K)	0.2	400	202.4	400	91
$c_p(J/kg.K)$	2000	380	871	385	440
$a \left(\frac{l_{rg}}{m^3} \right)$	880 (solid)	8920	2719	8960	8002
p (kg/m)	770 (liquid)				8902
$\beta (1/K)$	0.0006	-	-	-	-

Table 1: Thermophysical properties of PCM, nanoparticles, and MFs [10, 28, 36].

Table 2: Grid independence analysis.

Number of elements	Melting Time	Deviation $(\%)$	Total energy (kJ/kg)	Deviation $(\%)$
43753	1290	0.00	242.47	0.00
48305	1300	0.78	242.34	0.05
54087	1295	0.38	242.49	0.06
60796	1295	0.00	242.39	0.04