Modelling, simulation and comparison of phase change material storage based direct and indirect solar organic Rankine cycle systems

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Abstract

The thermodynamic performance of a novel direct solar organic Rankine cycle system and conventional indirect solar organic Rankine cycle system is compared in this study. The working fluid is vaporized directly in the solar collectors in direct solar organic Rankine cycle system while heat transfer fluid is used to vaporize the working in indirect solar organic Rankine cycle system. The evacuated flat plate collectors array covering a total aperture area of 150 m^2 is employed as a heat source and a phase change material tank having a surface area of 25.82 m^2 is used as thermal storage for both configurations. R245fa and water are chosen as heat transfer fluids for direct and indirect solar organic Rankine cycle systems, respectively. However, R245fa is used as a working fluid for both configurations. The performance of both configurations is compared by carrying out weekly, monthly and annual dynamic simulations in MATLAB by using hourly weather data of Islamabad, Pakistan. The direct solar organic Rankine cycle system outperforms the indirect solar organic Rankine cycle system in terms of thermal efficiency and net power. The annual system efficiency and an annual average net power of the direct solar organic Rankine cycle system are 71.96% and 64.38% higher than indirect solar organic Rankine cycle system respectively. However, average annual heat stored by phase change material during charging mode of indirect solar organic Rankine cycle system is 4.24 MJ more than direct solar organic Rankine cycle system. Conversely, direct solar organic Rankine cycle system has provided annual daily average power of 33.80 kW extra to heat transfer fluid during the discharging mode of phase change material storage. Furthermore, with phase change material storage, the capacity factor is increased by 17 % and 21.71 % on annual basis for direct and indirect solar organic Rankine cycle systems, respectively.

Keywords: Organic Rankine Cycle; Direct and Indirect solar ORC system; Phase Change Material; Capacity Factor; Thermodynamic, Comparison

1 Introduction

Solar energy has emerged as one of the most rapidly growing renewable sources of electricity. It has a minimum time of replenishment and maximum capacity among all available energy resources [1]. Furthermore, it is an attractive option for coupling with a low-medium temperature organic Rankine cycle (ORC) system. A temperature of 100 ^{o}C or slightly higher is enough to run a solar ORC system. Hence, these solar ORC systems are able to work efficiently within a temperature range of 100 to 150 ^{o}C. They are beneficial in terms of power, all the way down to small unit sizes, low technical demand in heat storage, co-generation close to the usage point and suitability in regions with less direct solar radiation resources.

To date, most of the studies investigated solar ORC systems use heat transfer fluid (HTF) to transfer energy from collectors to the organic fluid [2–5]. Heat transfer irreversibility largely occurs in the evaporator as explained by Jing et al. [6]. Furthermore, investment cost also increases while using HTF. Extra power is required for the pumping of HTF that can decrease the system's net power output specifically for small-scale solar ORC systems. The direct solar organic Rankine cycle system (DSOS) is one in which the evaporator is replaced with the solar collectors. Hence, all of the aforementioned problems can be resolved by adopting the novel DSOS.

To date, the solar ORC system with direct vapor generation (DVG) is studied by different researchers. Few have conducted experimental work [7,8] while others have done theoretical studies [6,9,10]. All of them focused on working fluid selection and performance of the system. This kind of system is found to be promising. In DVG solar ORC system vapor is generated directly in the collectors, the working fluid influences not only the heat to power conversion but also the solar energy collection.

Jing et al. [6] explained heat transfer irreversibility largely occurs in the evaporator. Moreover, they found 10% relative increment in overall electricity efficiency of DVG based solar ORC over conventional solar ORC system at solar radiation of 1000 W/m^2. Furthermore, extra power is required for the pumping of HTF in conventional solar ORC that can decrease the system's net power output and overall system efficiency specifically for small-scale solar ORC systems. Freeman et al. [23] calculated that almost 58% of the exergy loss occurs in the evaporator.

The thermal energy storage is a core component in the development of a solar power system. Thermal energy storage for solar thermal applications can be divided into Latent Heat Storage (LHS) and Sensible Heat Storages (SHS) [11]. Phase Change Materials (PCMs) lies into the LHS group and are one of the promising technology for the development of efficient thermal storage. Moreover, 5–14 times additional energy per volume can be stored by using PCM as compared to sensible-heat storage materials [12]. PCM is also advantageous compared with SHS systems because the process of phase

change is nearly isothermal for pure substances, and takes place over a finite temperature range for composite materials [13,14]. In contrast, past research had shown some disadvantages of PCM related to low thermal conductivity which results in lower charging and discharging rates [15]. PCMs are classified into three different transition phases such as liquid to solid, solid to gas and solid to liquid. The solid-liquid transition can be further subdivided into organic, inorganic and eutectics [12,16]. The selection of PCM to make the best latent heat storage system is critical for the specific application since the operating conditions are widely variable. The melting point temperature is a key parameter in the selection process of PCM. An overview of the PCM properties and related applications are studied by Agyenim et al. [14]. The applications are subdivided into low temperature (1-65 ^{o}C), medium temperature (80-120 ^{o}C) and high temperature (>150 ^{o}C). Medium temperature range PCMs are suitable for solar ORC applications.

Previously, researchers have proposed and analyzed different configurations of solar ORC systems. However, few of them have focused on integrating the systems with heat storage. Li et al. [17] analyzed solar organic Rankine cycle with thermal energy storage. A dynamic model of the solar ORC system was developed. The effect of storage capacity, solar fluctuation and evaporation temperature on the solar ORC system were evaluated. It was concluded that a proper thermal energy storage capacity should be selected in order to cater the solar fluctuations of a given area. Wang et al. [3] investigated the off-design performance of the solar ORC system with the compound parabolic collector and sensible thermal storage unit. The system performance was observed under time-varying conditions and changing ambient temperature. The system could obtain maximum exergy efficiency in December while maximum power output in June. Freeman et al. [18] had studied domestic-scale distributed solar combined heat and power system consisting of an organic Rankine cycle for the UK climate. The system comprised of 15 m^2 solar collectors, 1 m^3 thermo-chemical storage, and ORC engine. Two staged solar collectors were considered. It was found that the proposed system could meet 32 % electricity demands of a UK home. The author suggested that future studies will be focused on providing a proper solution of a finite-sized thermal storage system. Moreover, the effectiveness of the storage system will be assessed for load profile matching over seasonal time-scales. Wang et al. [19] developed a mathematical model to simulate the solar ORC system under steady-state. It was concluded that by employing a heat storage unit into the system, uninterrupted and steady operation of the solar-driven regenerative organic Rankine cycle could be achieved over a long period of time.

Pantaleo et al. [20,21] carried out two different studies on ORC coupled with heat storage. In the first study, the heat was recovered from exhaust gasses of gas turbine via thermal energy storage. Two molten salts storage tanks of (one cold at 200 ^{o}C and one hot at 370 ^{o}C) and three different plant locations were selected. The thermodynamic modelling was performed assuming two CSP sizes, and consequently two thermal energy storage sizes. The thermodynamic performance indicated higher global energy conversion efficiencies while using CSP integration. The second study was focused on

thermodynamic modelling and chosen two different CSP sizes, storage levels, and operation modes. Two molten salt storage tanks were considered which provided the 4.8–18 MWh energy storage. The plant capacity factor was increased and operating hours increased from 5500–6000 to 8000 h per year.

The integration of ORC with the PCM storage unit has gained attention in the recent past. Gang et al. presented two different configurations [22,23] of the solar ORC system with PCM storage unit. The first study was focused on the comparison of a regenerative ORC with the solo cycle. The system consists of non-tracking solar collectors and ORC engine integrated with a PCM storage tank. The research was focused on the effect of the regenerative cycle on system efficiency and collector efficiency. However, the second study was focused on the arrangement of the two-stage solar collectors. Flat plate collectors with corresponding PCM storage were used for preheating. Compound parabolic concentrators with corresponding PCM storage were used to achieve a higher temperature. Furthermore, collector efficiency and overall cycle efficiency were calculated.

Freeman et al. [24] proposed different thermal energy solutions. Authors applied various combinations of PCMs storage, water storage and solar collectors in a small-scale solar organic Rankine cycle combined heat and power system. Their performance was evaluated for UK and Cyprus climates, respectively. Furthermore, PCMs resulted in a 20% higher total daily electrical output per unit storage volume as compared to water storage.

Marcello et al. [25] analyzed and compared different PCMs coupled with the ORC system. It was found that the amount of energy stored and the thermal efficiency of the system increased with increasing heat source temperature. Conclusively, it was observed that the amount of energy stored by PCM increased dramatically by using a metal foam. This happened due to the higher thermal conductivity of the foam which resulted in the faster melting process. Manfrida et al. [26] and Sagar et al. [27] integrated PCM storage with solar ORC system. Their studies were focused on developing a mathematical model of PCM storage. Then dynamic simulations of overall systems were carried out for seven and ten days, under time-varying weather conditions (solar radiation, ambient temperature), for first and second study respectively. In case of first study, the model gave time-dependent HTF temperature profile of PCM storage tank outlet. It was then applied to solar ORC system. Weekly average energy efficiency achieved by the PCM storage was 83% during charging mode and 93% during discharging mode, respectively. Two different diameters and lengths of PCM storage tank were selected in the second study. It was found that smaller diameter and longer length showed the overall better performance of solar ORC system. In contrast, the pressure drop was significantly high for aforementioned system.

However, previous studies mainly focused on the design and short-time simulations. Detailed modeling and simulation of integrated direct solar organic Rankine cycle system and direct solar organic Rankine cycle system (ISOS) for a whole year had not been reported yet. The novelty of this work lies in the thermodynamic performance assessment and comparison of the performance of DSOS and ISOS based

on the thermodynamic model. The contribution includes

- Development and validation of the 1-dimensional PCM model for integrated solar ORC system. Temperature profiles of both HTF and PCM are demonstrated and compared for the first time under time-varying weather conditions.
- The weekly, monthly and annual dynamic simulations of the integrated solar ORC system in relation to the charging and discharging characteristics of the PCM storage system. Furthermore, increment in the capacity factor of both configurations by using PCM storage is also analyzed and compared for the first time.
- Thermodynamic performance and comparative assessment of DSOS and ISOS based on system efficiencies and net power output.

The present study is divided into five sections. Section 1 of the paper provides an overview of the current research status in the area of solar organic Rankine cycle (ORC) system with a focus on novelty and originality of the present work. The layout, working principle, and control of the integrated solar ORC system presented in Section 2. The modelling and simulation approach is described in detail in Section 3. The results and discussion of the DSOS and ISOS are provided in Section 4. Finally, the concluding remarks are added in Section 5 of the paper.

2 System configurations and control

The schematic diagrams of the DSOS and ISOS are shown in Figure 1 and Figure 2, respectively. Present work has been carried out to compare the performance of both configurations on the basis of hourly weather data. Both configurations are comprised of evacuated flat plate collectors (EFPCs), phase change material storage tank and basic ORC plant (evaporator, expander, condenser, and pump) coupled with a generator. PCM storage works in two operating modes (charging mode & discharging mode). In the case of DSOS, solar collectors work as an evaporator during charging mode as shown in Figure 1. However, during the discharging mode, the PCM storage tank works as an evaporator. One variable flow organic fluid flow pump namely P_1 is employed to regulate fluid flow in the system. HTF is replaced by organic fluid in this system. Moreover, R245fa is used as a working fluid.

In the case of ISOS, water works as HTF in the solar loop while R245fa is used as a working fluid ORC loop as shown in Figure 2. Moreover, two variable flow pumps entitled P_1 and P_2 are employed to control the flow of HTF & working fluid in solar and ORC loop respectively. Five control valves are also used to properly regulate the flow of fluid in both configurations. The valves open and close based on the operation and control conditions, which are discussed in detail in the section below. Both configurations integrated with PCM storage are simulated in MATLAB programming environment under time-varying solar radiation conditions.

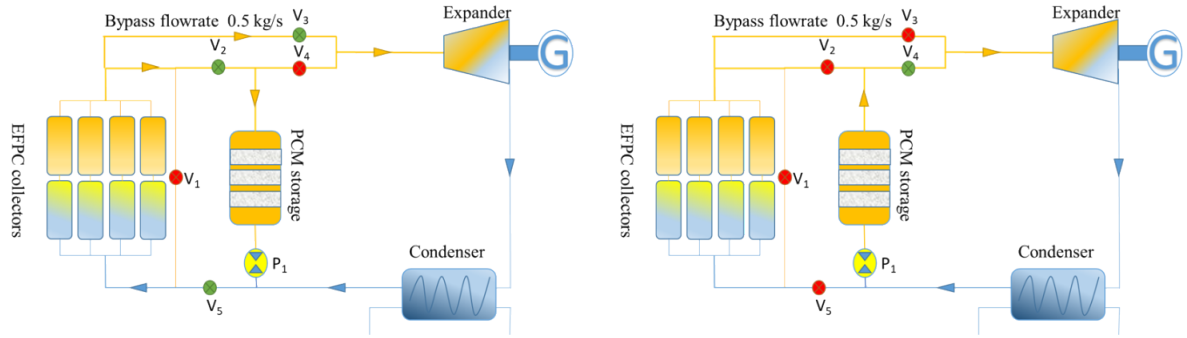

Figure 1: Layout diagram of direct solar organic Rankine cycle system (DSOS) during (a) charging mode and (b) discharging mode

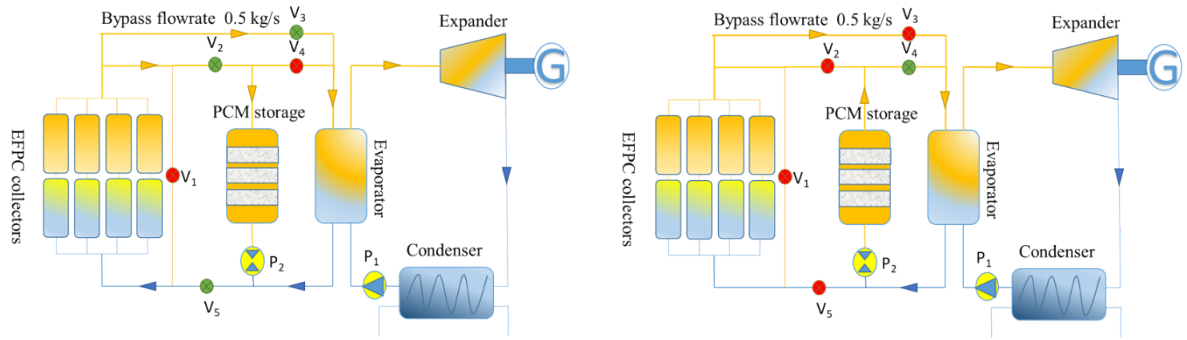

Figure 2: Layout diagram of indirect solar organic Rankine cycle system (ISOS) during (a) charging mode and (b) discharging mode

Hourly based climatic data of Islamabad-Pakistan has been used for the current study. Typical meteorological year (TMY) data of Islamabad is obtained by using Meteonorm software [28]. Islamabad represents cold winter and very hot and hot-humid summer. The monthly average ambient temperature along with the solar radiation falling on the solar collector surface for the whole year is shown in Figure 3. It can be observed that June is the hottest month with the maximum amount of solar radiation and maximum ambient temperature while January is the coldest month of the year along with low solar radiation and minimum ambient temperature. The first week of January and the second week of June are considered to be the coldest and hottest weeks of the whole year, respectively.

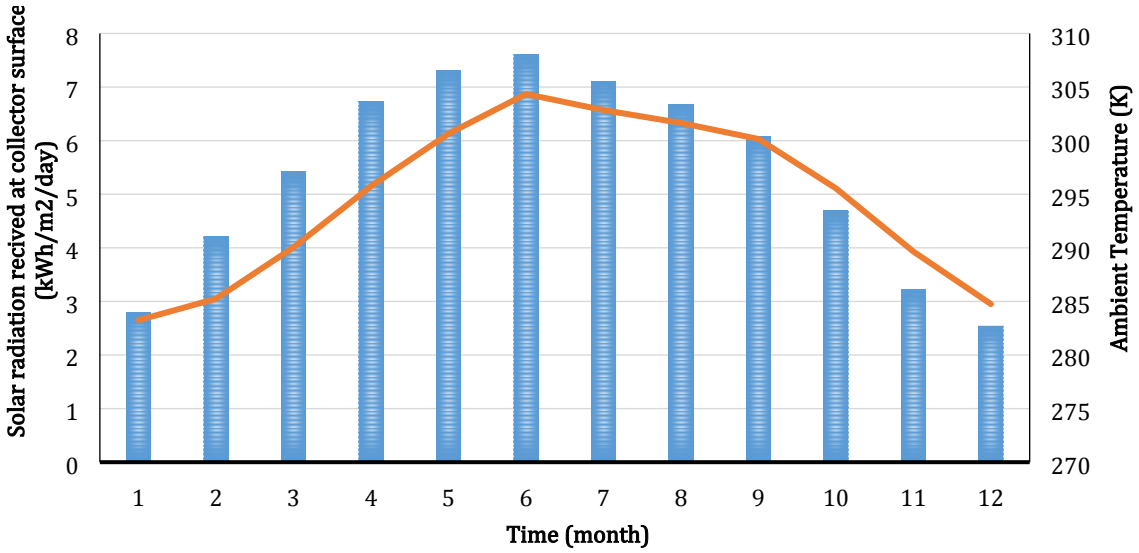

Figure 3: Climatic data of Islamabad- Pakistan (daily average monthly)

To apply the desired logical control system, every component for both configurations is controlled and turned on and off by logical functions depending on several simulation parameters. The values of TMY data of the Islamabad is imported in MATLAB from the metronome software. Both configurations have been evaluated at the same operation and boundary conditions. It is assumed that the system starts working when the solar radiation received at the surface of the collector goes above 400 W/m^2. On the contrary, the system stops or undergoes to discharging mode.

There are two modes of operation of the storage system namely charging and discharging mode. The initial temperature of PCM is assumed to be 373.15 K. This shows that PCM is not charged and in the solid phase at the beginning of the simulation process. Five control valves are employed in both configurations which are opened and closed depending upon the mode of operation. During charging mode, valves V2, V3 and V5 are opened while V1 and V4 remain close. In contrast, during the discharging mode, V4 is opened while the rest of the valves remains close. The heat storage system is designed to work at melting point temperature of the PCM. It means that major part of the energy is released or absorbed at the melting point of PCM.

The maximum temperature at the outlet of collector array is selected to be 390 K that is slightly higher than the PCM melting point temperature. In this regard, when the collector outlet temperature rises above the limit imposed, V1 is opened and the rest of valves are closed and the HTF mass flow rate is increased by 10 % at every iteration until it reaches below the limit imposed. To avoid the supercritical condition, Initial temperature of HTF mass flow rate during charging mode is selected to be 3 kg/s and it increases with increment in collector outlet temperature. However, to achieve consistent power generation, HTF bypass mass flow rate is kept at a constant rate of 0.5 kg/s in both charging and discharging mode. The discharging limit of the storage tank is maintained to 370 K, which means that the system is allowed to discharge the storage in sensible heat region.

3 Thermodynamic modelling

3.1 Solar radiation

Solar radiation received by the sloped surface of the solar collectors is calculated by [29].

$$I_t = \left(I_b + I_d \frac{I_b}{I_h} \right) R_b + I_d \left(1 - \frac{I_b}{I_h} \right) \left(\frac{1+\cos\beta}{2} \right) \left(1 + \sqrt{\frac{I_b}{I_h}} \mathrm{Sin}^3 (\frac{\beta}{2}) \right) + I_h \rho_g \left(\frac{1-\cos\beta}{2} \right) \quad (1)$$

3.2 Solar collectors

For the solar collector's field, evacuated flat plate collectors are selected. Due to vacuum inside the collector, this type of collector is highly efficient at high temperatures up to 200°C. Moreover, they are non-tracking, non-concentrating collectors. Therefore, they do utilize both beam and diffused radiation [30]. The efficiency of the solar collector is calculated as a function of collector inlet temperature, the ambient air temperature and total solar radiation received at collector surface.

$$\eta_{cl} = a_0 - a_1 \frac{(T_i - T_{amb})}{I_t} - a_2 \frac{(T_i - T_{amb})^2}{I_t} \quad (2)$$

Where a_0, a_1 and a_2 are the optical efficiency of the collector, linear heat loss coefficient, and quadratic heat loss coefficient, respectively. A total number of solar collectors were assumed to be 75 in collector array and each solar collector has a size of 2 m^2. Amount of energy received by the solar collector's array is computed by:

$$q_{cl} = \eta_{cl} \times I_t \times n \times A_{cl} \quad (3)$$

Furthermore, the temperature at the outlet of the solar collector array is calculated by

$$T_o = T_i + \left(\frac{q_{cl}}{\left(m_f C_p \right)} \right) \quad (4)$$

3.3 Phase change material storage

Figure 4 shows a typical configuration of the PCM storage tank. A cylindrical shape tank completely filled with PCM is considered. Moreover, it has a coiled shape pipe containing HTF passes through the tank. The walls of the PCM storage tank are assumed to be adiabatic.

The tank operates in charging and discharging mode depending upon operation and boundary conditions. During charging mode, the heat is transferred from HTF to the PCM. The PCM temperature rises from solid-phase until it reaches the melting point temperature. After that, the temperature of PCM

remains constant during the melting process.

After completion of the melting phase during which all of the PCM changes into the liquid phase, the temperature of the liquid PCM further rises up to the limit imposed by HTF. However, during a discharging phase, thermal energy stored by liquid PCM is extracted by cold HTF [26]. The well-known enthalpy method is used to solve the governing equations for HTF and PCM as shown in equation (5) [31][32]. In order to determine the heat transfer in PCM while solving the enthalpy method few assumptions are made as follows:

- Conductive heat transfer is considered to be the dominant mechanism.
- Only One-dimensional heat transfer is contemplated.
- The thermos-physical properties of PCM remain constant for each state.
- Natural convection is neglected in this model which may occur due to the density difference in the PCM [33].

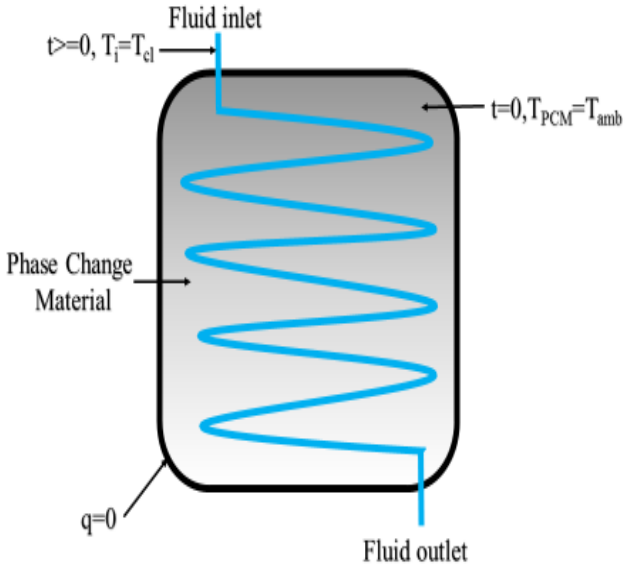

Figure 4: Layout diagram of PCM storage Tank

$$\rho\frac{\partial H}{\partial t} = \kappa_{pcm}\frac{\partial^2 T_{pcm}}{\partial y^2} \tag{5}$$

Where "H" is the total volumetric enthalpy, which consists of both the latent heat and the sensible heat of PCM at a given temperature. Therefore, the total volumetric enthalpy of PCM at any given temperature is calculated using the following relation:

$$H = \int_{T_m}^{T} \rho_{pcm} C_{pcm} \Delta T_{pcm} + \rho_{pcm} L F(\lambda) \tag{6}$$

In the above formulation equation (6), the latent heat of the PCM is related to the liquid fraction of the

PCM "LF". To compute the latent heat of the PCM, the liquid fraction LF needs to be defined that is calculated as given by the Equation (7)

$$LF = \begin{cases} 0 & for \quad T_{pcm} < T_m \quad Solid\ region \\ 1 & for \quad T_{pcm} > T_m \quad Liquid\ region \end{cases} \tag{7}$$

From equation (6) and (7) enthalpy of PCM can be calculated as:

$$H = \begin{cases} \rho_{pcm} C_{pcm} (T_{pcm} - T_m) & for \quad T_{pcm} < T_m \quad Solid\ region \\ \rho_{pcm} C_{pcm} (T_{pcm} - T_m) + \lambda \rho_{pcm} & for \quad T_{pcm} > T_m \quad Liquid\ region \end{cases} \tag{8}$$

The above correlation depicts, if the temperature of PCM is less than its melting point temperature, it only contains sensible heat. Conversely, if the temperature of PCM is more than or equal to its melting point temperature, total volumetric enthalpy is the combination of latent heat and sensible heat. The temperature of the PCM "T_{pcm}" is further derived from the volumetric enthalpy of the PCM as follows:

$$T_{pcm} = \begin{cases} T_m + \dfrac{H}{\rho_{pcm}.C_{pcm}} & for \quad H < 0 \\ T_m & for \quad 0 < H < \rho_{pcm}.\lambda \\ T_m + \dfrac{H - (\rho_{pcm}.\lambda)}{\rho_{pcm}.C_{pcm}} & for \quad H > \rho_{pcm}.\lambda \end{cases} \tag{9}$$

In equation (9) λ is the latent heat of the PCM while ρ_{pcm} is the density of PCM. The thermophysical properties of PCM used in the current study are given in Table 1.

Table 1: Thermo-Physical properties of PCM used for the current study [26]

Commercial Name		Salt hydrate
PCM category		Inorganic
Melting point (K)		389.85
Latent heat (kJ/kg)		160
Length of PCM tube (m)		24
Diameter of PCM tube (m)		0.35
Thermal conductivity (W/m^2-K)	both phases	0.7
Specific heat capacity (kJ/kg-K)	both phases	2.61

The amount of energy stored by PCM during the charging mode is calculated by multiplying the mass

of the PCM with the difference in latent heat between a final and initial node of the PCM storage tank.

$$Q_{st,c} = M_{pcm}(\lambda_{\mathrm{mx}} - \lambda_{in}) \tag{10}$$

Power transferred to HTF by PCM during discharging mode is calculated by equation (11)

$$P_{tr,d} = m_{HTF} C_{HTF} (T_{HTF,o} - T_{HTF,i}) \tag{11}$$

3.4 Validation of the Computational Model

Currently used PCM computational model has been validated from the experimental results of Zivkovic et al. [33]. Results are presented for the melting case of the PCM. Zivkovic et al. [33] using $CaCl_2.6H_2O$ PCM encapsulated in the rectangular container made of stainless steel. The length and breadth of the container are both 100 mm, while the thickness is 20mm, respectively. The container with solid PCM was placed in a bath having a constant temperature of 333.15 K. The PCM container was well insulated on the lateral sides. The experimental results were reproduced through the current computational model. To reproduce the result the convective heat transfer coefficient between air and the container wall is determined by h_{conv}=16 W/m^2-K *and* $T_\infty$was 333.15 K [33]. Thermophysical properties of the $CaCl_2.6H_2O$ are tabulated in Table 2.

Table 2: Thermo-Physical properties of CaCl2. 6H2O

Melting Point (K)		303.05
Latent heat (kJ/kg)		187
Density *(kg/m^3)*	Solid	1710
	Liquid	1530
Thermal conductivity *(W/m^2-K)*	Solid	1.09
	Liquid	0.53
Specific heat capacity *(kJ/kg-K)*	Solid	1.4
	Liquid	2.2

A comparison between the experimental results from Zivkovic et al. [33] and the current computational model, for the PCM temperature at the center of the container, is presented in Figure 5.

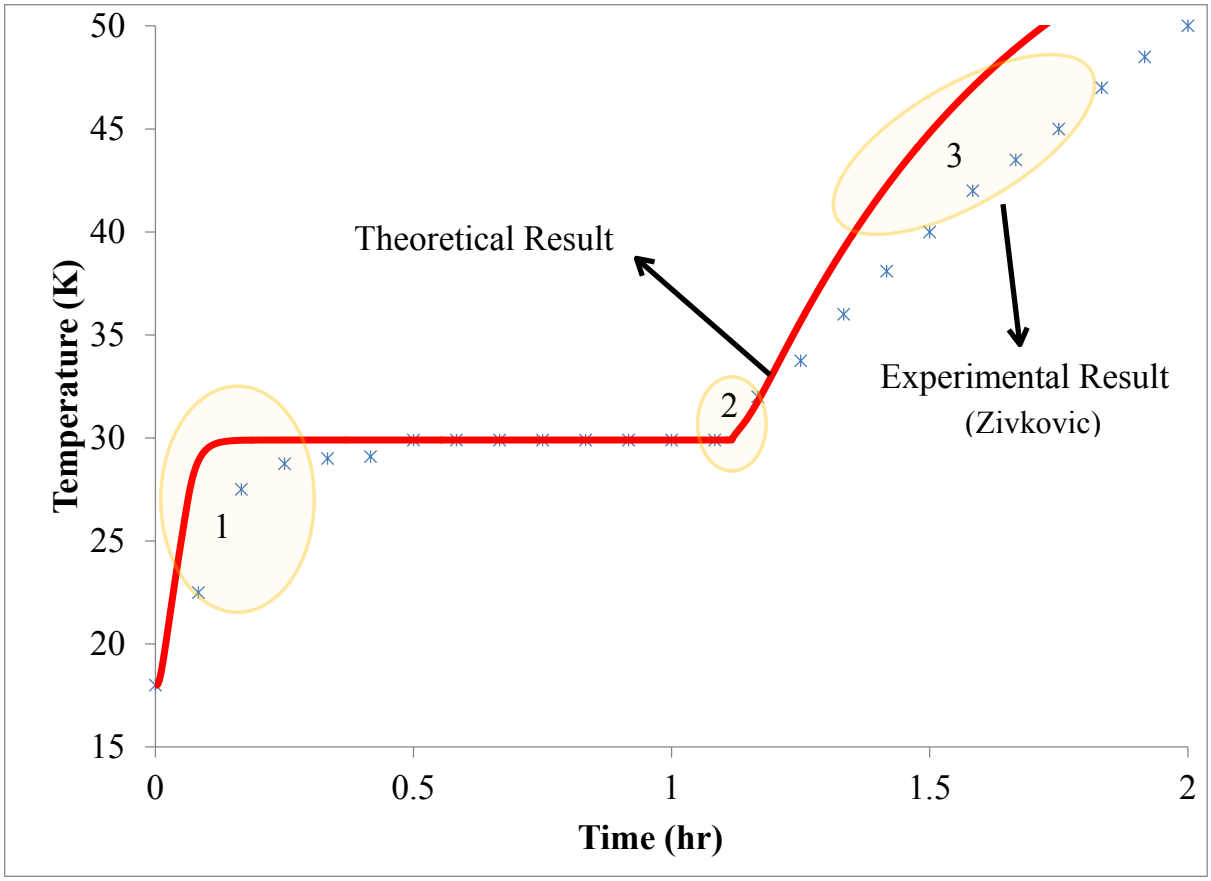

Figure 5: Comparison of the PCM modeling results with the experimental results of Zivkovic and Fujii [33] *for the melting of PCM.*

It has been observed from Figure 5 that agreement is well within the uncertainties indicated by Zivkovic et al. [33]. Three major observations and discrepancies (mentioned in Figure 5) between the predicted results and the experimental results are observed that are explained below:

1. In the beginning, sensible heat released by the PCM was predicted faster compared to the experimental result. This is mainly due to the assumption considered in the theoretical model which neglects the conduction between the container wall and PCM. This point is indicated as 1 in Figure 5.
2. Calculated PCM melting time is observed slightly higher compared to the experimental result which is mainly due to the reason that the natural convection currents within the PCM are not incorporated in the theoretical model. This point is indicated as 2 in Figure 5.
3. The theoretical model predicted the higher temperature in the liquid region compared to the experimental results at any time. This discrepancy is indicated as 3 in Figure 5. This mainly due to the reasons indicated by Zivkovic et al. [33] that the calculated heat transfer coefficient is a little bit higher compared to the real one.

Therefore it can be assumed very carefully that avoiding the natural convection and thermal conduction of the container wall at this stage may not produce a significant error in the prediction of the PCM

temperature.

3.5 **Organic Rankine cycle**

The basic Organic Rankine Cycle (ORC) configuration is chosen for the current simulation study due to lower capital investment for low-medium temperature applications [34][35]. The operating conditions and assumptions for the design of the ORC system are listed in Table 3.

Table 3. Assumptions of the boundary conditions for ORC system design

Parameter	Value
Hot water (heat source) mass flow rate	0.5 kg/s
Pinch point temperature difference in evaporator and condenser [36]	5 K
Turbine efficiency for initial cycle design [37]	80%
Pump efficiency for initial cycle design [38]	60%
Generator efficiency	85%
Degree of superheating at turbine inlet [39]	3 K
Condensation temperature	303.15 K

Properties of the working fluid R245fa are shown in Table 4.

Working fluid	R245fa
Critical temperature	154.01°C
Critical pressure	3.651 MPa
ODP	0
Net GWP	4
Flammability	High

The power generated by the expander is calculated by equation (12).

$$W_t = m_{wf}(h_{t,i} - h_{t,o}) \tag{12}$$

The total power consumed by the pump is computed by equation (13) and (14) for ISOS and DSOS respectively.

$$W_p = W_{p1} + W_{p2} \tag{13}$$

There is only one pump used in DSOS. Therefore, pump power is given by equation (14)

$$W_p = W_{p1} \tag{14}$$

Pump power within the ORC cycle is calculated by enthalpy difference across the organic fluid pump.

$$W_{p1} = m_{wf}(h_{p,o} - h_{p,i}) \tag{15}$$

However, in the case of ISOS, the power consumed by the pump in the solar loop is calculated by equation (16) using a fixed value of 65 % for solar pump efficiency ε_{sp}.

$$W_{p2} = \frac{m_{HTF}\Delta P}{\varepsilon_{sp}\rho_{HTF}} \tag{16}$$

The pressure drop ΔP comprises the pressure loss in the system's pipe network ΔP_L and the pressure loss across the solar collector array ΔP_{cl}. The total length of the pipe network is assumed to be 200 m and diameter 20 mm. The Darcy friction factor f associated with pressure drop is computed according to correlations based on the Reynolds number given in Incropera et al. [40]

$$f = 0.316\,\mathrm{Re}^{-1/4}; \quad \mathrm{Re} < 2\times10^4; \tag{17}$$

$$f = 0.814\,\mathrm{Re}^{-1/5}; \quad \mathrm{Re} \geq 2\times10^4; \tag{18}$$

$$\frac{\Delta P_L}{L} = \frac{8\,m^2{}_{HTF} f}{\pi^2 \rho_{HTF} D^5} \tag{19}$$

The pressure drop across the solar collector array is calculated by an empirical correlation given by Freeman et al. [18].

$$\Delta P_{cl} = A_{cl}(21.77 m_{HTF}^2 + 3.54 m_{HTF}) \tag{20}$$

The isentropic efficiency of the expander and the pump is defined by equation (21) and (22)

$$\varepsilon_{t} = \frac{h_{t,i} - h_{t,o}}{h_{t,i} - h_{t,os}} \tag{21}$$

$$\varepsilon_{p} = \frac{h_{p,os} - h_{p,i}}{h_{p,o} - h_{p,i}} \tag{22}$$

Where os represents the isentropic process. The energy required in the heating process of the ORC is calculated by the enthalpy increment of the organic fluid from the pump to the expander.

$$W_{net} = W_t \varepsilon_g - W_p \tag{23}$$

$$Q_{ORC} = m_{wf}\left(h_{t,i} - h_{p,o}\right) \tag{24}$$

The ORC efficiency for both configurations is defined by the ratio of the net power output to the heat supplied [6].

$$\eta_{ORC} = \frac{W_{net}}{Q_{ORC}} \tag{25}$$

The system efficiency of both solar ORC systems is expressed by

$$\eta_{sys} = \eta_{ORC}.\eta_{cl} \tag{26}$$

The increment in a capacity factor of both configurations is calculated by a relative increment in working hours by use of PCM storage.

$$CF_{inc} = \frac{Wh_{\mathrm{w,pcm}} - Wh_{wo,pcm}}{Wh_{\mathrm{w,pcm}}} \tag{27}$$

4 Results and Discussions

Results obtained from dynamic MATLAB simulation of solar ORC systems are presented, analyzed and discussed in this section. A time step of 1 hour is selected for the whole simulation process. Islamabad is chosen as a reference location having coordinates: 33.7294° N, 73.0931° E. Whole year simulations are carried out to compare the performances of both DSOS and ISOS. Firstly, dynamic simulations are carried out for the coldest and hottest weeks of the year. The performance of PCM storage is also compared for both DSOS and ISOS during charging and discharging mode.

Results of system efficiencies of both configurations and net power output are also compared and discussed. Secondly, the performance of both configurations is compared for the whole year. Moreover, it is also analyzed how much heat is stored by PCM during charging mode and how much power is delivered to the PCM by HTF in discharging mode. Finally, an increase in the capacity factor by using PCM storage for both configurations are presented and analyzed.

4.1 Performance of the hottest week

4.1.1 Variation in temperature profiles of HTF and PCM during charging mode

During charging mode, hourly average daily temperature profiles of both PCM and HTF for ISOS and DSOS during the hottest week (2nd week of June) of the year are shown in Figure 6 and Figure 7 respectively. It can be seen that HTF temperature shows an increasing and decreasing trend with respect to the rise and fall of solar radiation. However, it remains constant initially with an initial decrement in solar radiation. By comparing Figure 6 and Figure 7, it is observed that the number of charging hours for ISOS and DSOS is found to be 9 and 7 hours, respectively. The number of charging hours is higher for ISOS because, within the selected range of parametric conditions, HTF temperature cannot reach

up to the imposed limit of 390 K. Increment in temperature of PCM for ISOS and DSOS is observed to be 7.7 K and 18.5 K, respectively. However, in the case of HTF, these values are found to be 14.8 K and 18 K. Moreover, at maximum HTF temperature, the temperature difference between HTF and PCM is found to be 10 K and 0 K for ISOS and DSOS, respectively. Hence, it shows that heat transfer between HTF and PCM is higher for DSOS in comparison with ISOS. Conclusively, in case of DSOS, there is a steep rise in temperature of HTF until it reaches up to the melting point of the PCM under selected operating and boundary conditions. Conversely, in case of ISOS, HTF temperature doesn't reach up to that limit. This can happen because water has 2.5 times higher value of specific heat capacity as compared to R245fa within the given temperature range.

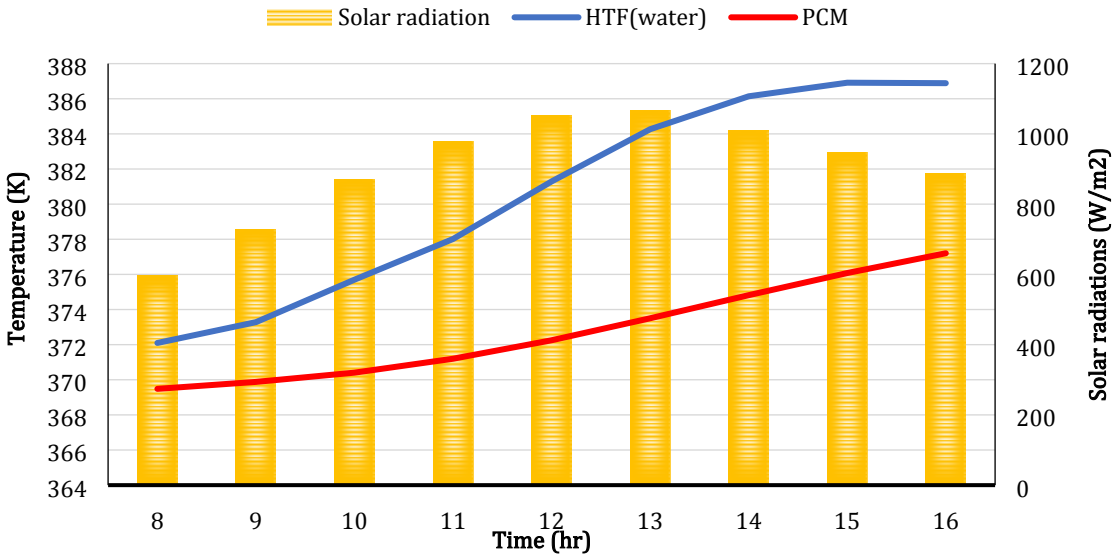

Figure 6: Variation in temperature profiles of HTF and PCM with solar radiation for ISOS during charging mode in the hottest week

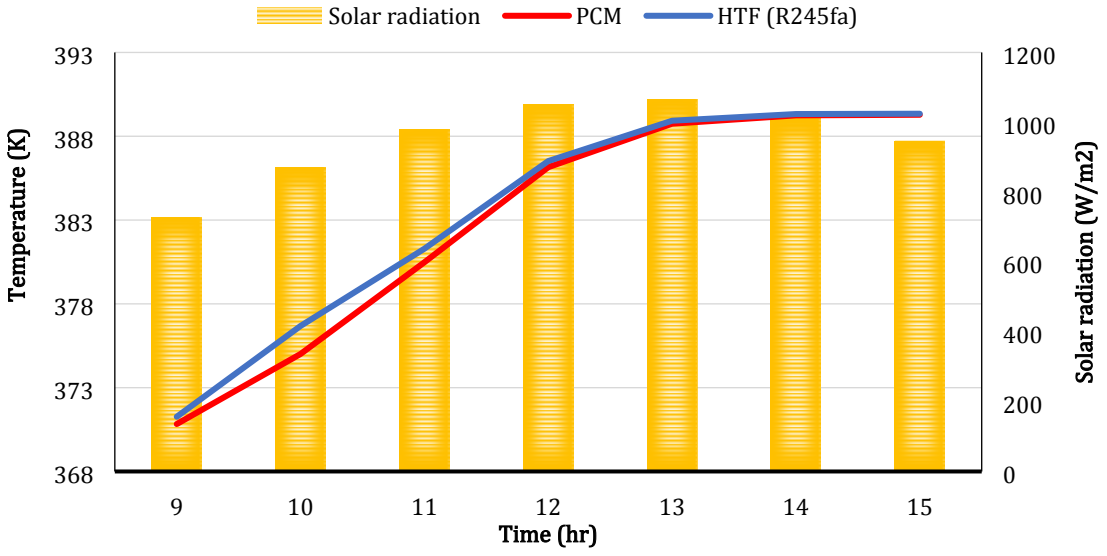

Figure 7: Variation in temperature profiles of HTF and PCM with solar radiation for DSOS during charging mode in the hottest week

4.1.2 *Variation in temperature profiles of HTF and PCM during discharging mode*

A numerical simulation model of the PCM storage tank is developed using MATLAB. The finite difference method is used to discretize time and space (along one dimension). Therefore, the length of the PCM storage tank is divided into 105 equally spaced nodes. The simulation time step is selected to be 1 hour. The model has calculated the temperature of PCM and HTF at each node for every hour during the whole simulation process. Figure 8 and Figure 9 show variation in hourly average weekly temperature of PCM and HTF in the hottest week of the year during discharging mode for ISOS and DSOS respectively. It is observed that the temperature of PCM and HTF generally increases along the length of the heat storage tank for both ISOS and DSOS. This might happens because the temperature difference between PCM and HTF is large at the initial node. Therefore, heat transfer from PCM to HTF is very high at the beginning of discharging process.

Therefore, the drop in PCM temperature is larger at that point. Conversely, there is the least temperature difference between HTF and PCM at the final node. Therefore, heat transfer between HTF and PCM is comparatively lesser at the final node. Therefore, the drop in PCM temperature is lesser at that point. Moreover, the temperature difference between a final and initial node of PCM is found to be 14.01 K and 1.19 K for DSOS and ISOS respectively. Furthermore, an increase in the temperature of HTF along the PCM storage tank for DSOS is found 7.4 times higher than ISOS. This might happen due to the higher heat capacity of water as compared to R245fa. However, the relative increment in PCM temperature with respect HTF decreases along the length of the PCM storage tank.

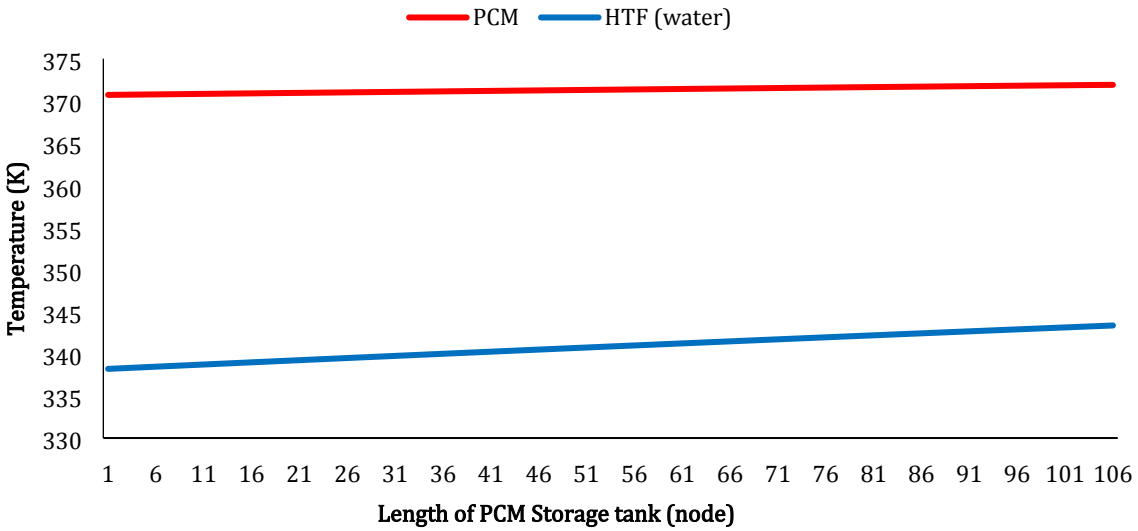

Figure 8: Variation in temperature of PCM & HTF during discharging mode for ISOS in the hottest week

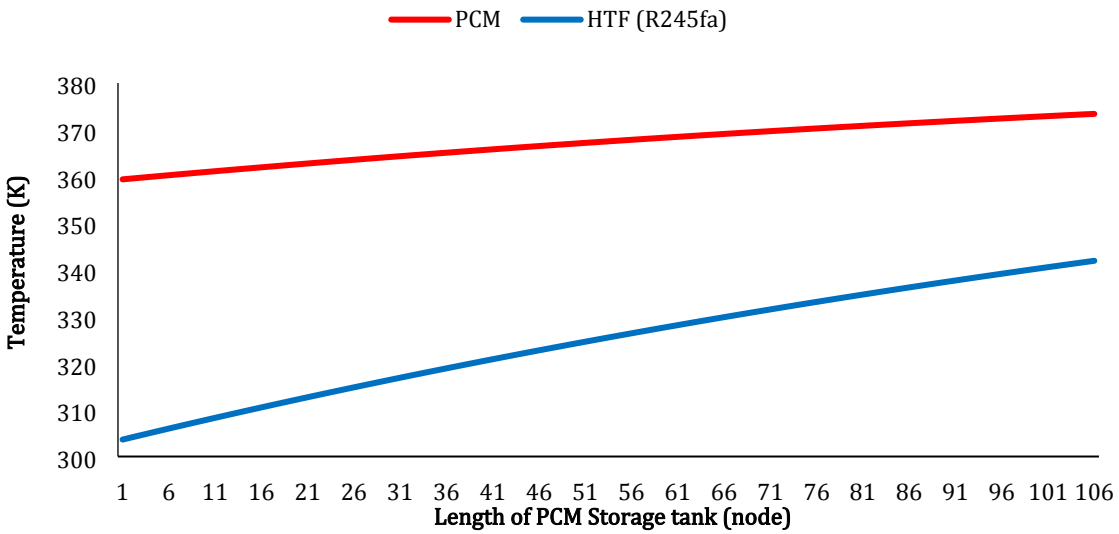

Figure 9: Variation in temperature of PCM & HTF during discharging mode for DSOS in the hottest week

4.1.3 Variation in system efficiencies and net power output

System efficiency and net power output are the two major parameters that generally describe the performance of thermal systems. Therefore, these parameters are calculated for both ISOS and DSOS. Figure 10 depicts the results for hourly average daily system efficiency and net power output of ISOS and DSOS during the hottest week of the year. In the case of DSOS, the system works from 8:00 till 21:00 hours, however, for ISOS it works between 7:00 to 20:00. DSOS has resulted in an extra working hour than ISOS on an average daily basis. The reduction in working hours for ISOS is due to the weaker thermal match between HTF and PCM as shown in Figure 6 and Figure 8. The system efficiency and the net power output for DSOS are found to be 5% and 2.4 times higher than ISOS respectively. The maximum values of the aforementioned parameters are found at the time instant of 15:00 hours. It happens due to the maximum charging of the PCM storage, which occurs at this time instant, as shown in Figure 6 and Figure 7. Further, DSOS has achieved 7.5 % higher system efficiency and 6.5 kW extra power than ISOS on an average daily basis. Hence, DSOS has demonstrated better thermal performance in comparison with ISOS.

This might be because of two major reasons: firstly, higher thermal losses occur across evaporator in ISOS (research has shown that more than 50 % of exergy losses occur in the evaporator section [18]). Secondly, extra power is required to operate the solar pump.

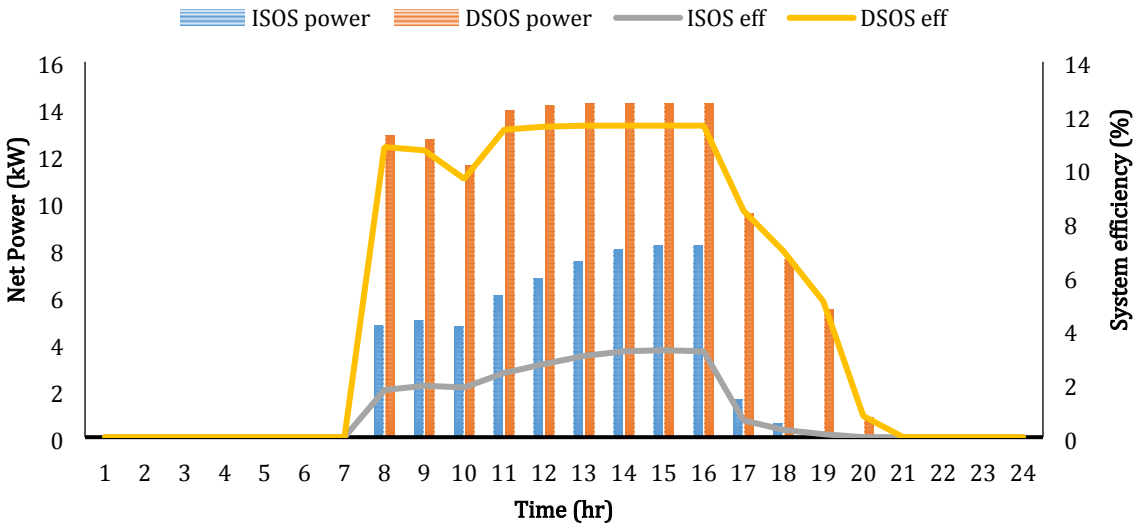

Figure 10: Variation in system efficiencies and net power output of DSOS and ISOS during the hottest week

4.2 Performance of the coldest week

4.2.1 Variation in system efficiencies and net power output

Figure 11 shows that the hourly average daily system efficiencies and net power output for ISOS and DSOS during the coldest week of the year (1st week of January). There are weak solar radiation and very low ambient temperature during the coldest week. Both of the systems work between 11:00 to 15:00. Hence, these weather conditions are not enough to rise the HTF temperature at the outlet of the storage tank up to the designed value of 370 K. Therefore, the charging process does not take place in both systems. However, both systems have demonstrated an identical trend with the hottest week. DSOS has shown higher 4.5 times system efficiency and 1.4 times larger net power output as compared to ISOS on an average daily basis. Moreover, DSOS has achieved 1.09 % higher system efficiency and 0.32 kW extra power than ISOS on an average daily basis. Furthermore, both systems have shown maximum system efficiencies and net power at the time instant of 14:00 hour because largest solar radiation and highest ambient temperature occur at that time.

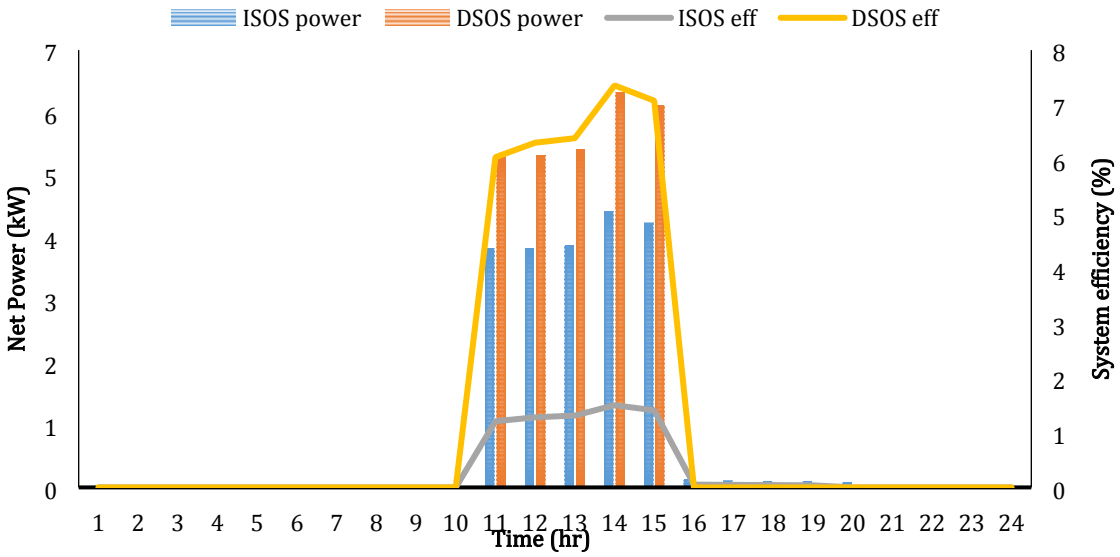

Figure 11: Variation in system efficiencies and net power output of DSOS and ISOS during the coldest week

4.3 Performance over the month

4.3.1 Variation in system efficiencies and net power output

Figure 12 shows the daily average monthly system efficiencies and net power output of ISOS and DSOS, respectively. By comparing Figure 3 and Figure 12, it is observed that both parameters (system efficiency and net power output) increase and decrease with rising and fall in solar radiation & ambient temperature. Therefore, both systems have followed similar behavior as seen in the case of weekly simulation. The maximum value of system efficiencies and net power output are observed in June being hottest month. Conversely, these values are found to be minimum in January that is the coldest month. The average system efficiency achieved and daily average net power output delivered by ISOS on annual basis is observed to be 1.71 % and 34.02 kW, respectively. While DSOS has shown 4.5 times higher system efficiency and 2.8 times higher net power output on annual basis. Furthermore, DSOS has shown 6.1% higher system efficiency and 61.5 kW higher average daily net power output than ISOS on annual basis. Hence, DSOS has shown much better thermal performance than ISOS.

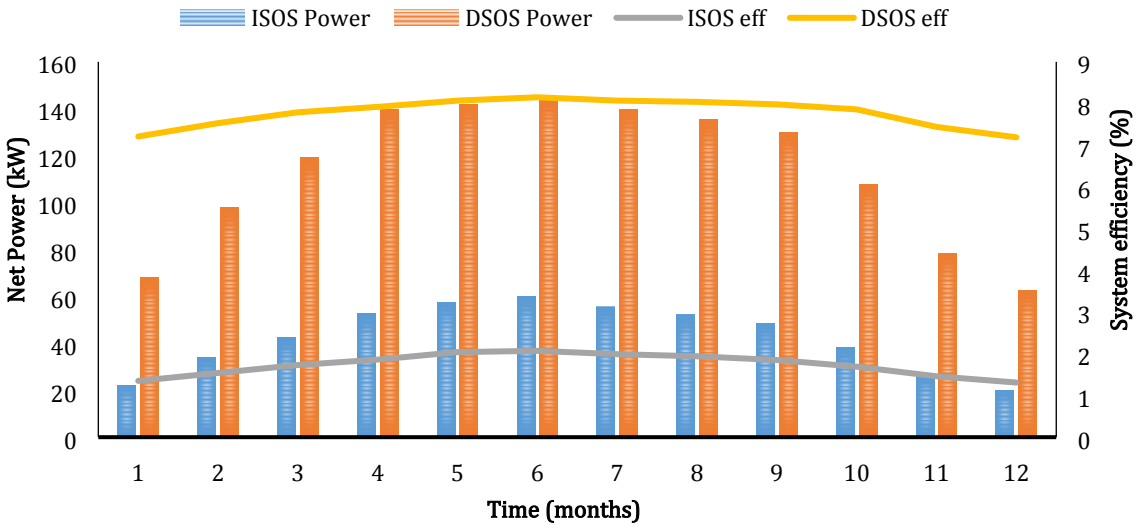

Figure 12: Variation in system efficiencies and net power output of DSOS and ISOS on daily average monthly basis

4.3.2 ***Heat stored by PCM during charging mode***

The daily average monthly amount of heat stored by the PCM storage tank during charging mode for DSOS and ISOS is shown in Figure 13. The PCM storage system is designed to work at PCM melting point temperature. However, in the current study, the total amount of heat stored by PCM is the sum of both sensible and latent heat. It is observed that the amount of heat stored for both systems increase and decrease with respect to rise and fall in solar radiation and ambient temperature as shown in Figure 3 and Figure 13. Moreover, maximum and minimum amount of heat stored by PCM is observed during June and January, respectively. However, a larger amount of heat is stored by ISOS as compared to DSOS. This can happen because the larger amount of heat transfer occurs across ISOS due to a higher number of charging hours. The daily average amount of heat stored per annum by PCM storage for ISOS is found to be 4.24 MJ more than DSOS. Furthermore, the maximum difference in the average daily amount of heat stored for ISOS and DSOS is observed to be 23 MJ that occurs in June. However, minimum difference in daily average amount heat stored for both systems is found to be 6 MJ that occurs in January.

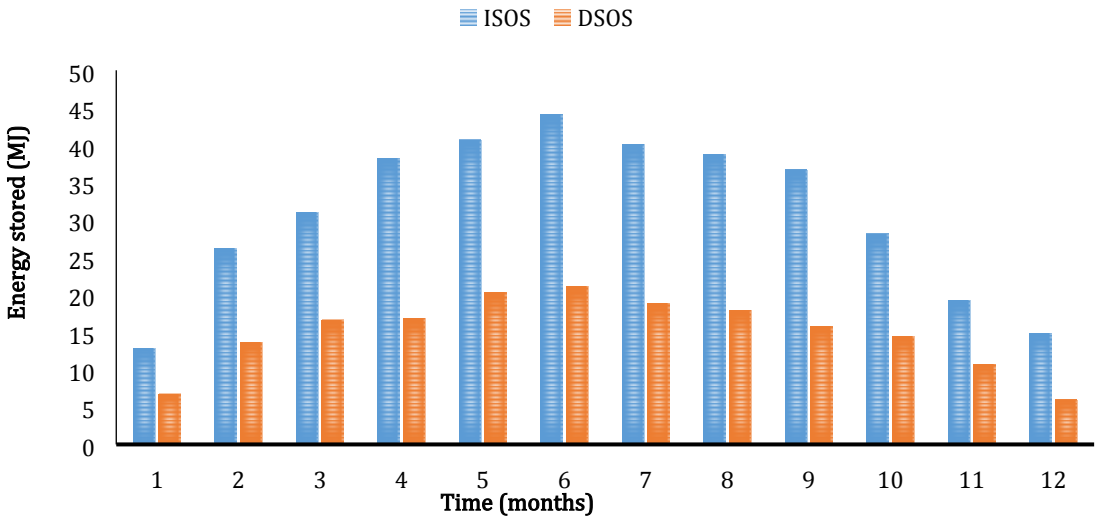

Figure 13: Variation in the amount of heat stored by PCM storage tank during charging mode for DSOS and ISOS on daily average monthly basis

4.3.3 ***Power Transferred by PCM to HTF during discharging mode***

When solar radiation fall below the imposed limit (400 W/m^2), both systems undergo discharging mode. Power transferred to the HTF from PCM for both systems on a daily average monthly basis during discharging mode is shown in Figure 14. It is observed that the amount of power transferred by PCM to HTF for both systems increase and decrease with respect to rise and fall in solar radiation and ambient temperature as shown in Figure 3 and Figure 14. Furthermore, the maximum and minimum amount of power transferred by PCM is observed during June and January, respectively. However, a larger amount of power is transferred by PCM for ISOS as compared to DSOS. This might be because of the stronger thermal match between PCM and HTF for DSOS as compared to ISOS as shown in Figure 8 and Figure 9. Furthermore, the maximum difference in the amount of power transferred by PCM for ISOS and DSOS is observed in summer months.

However, a minimum difference in power transferred by PCM for both systems is found in the winter months. Furthermore, power transferred by PCM to HTF is 4.8 times higher for DSOS as compared to ISOS on an annual basis.

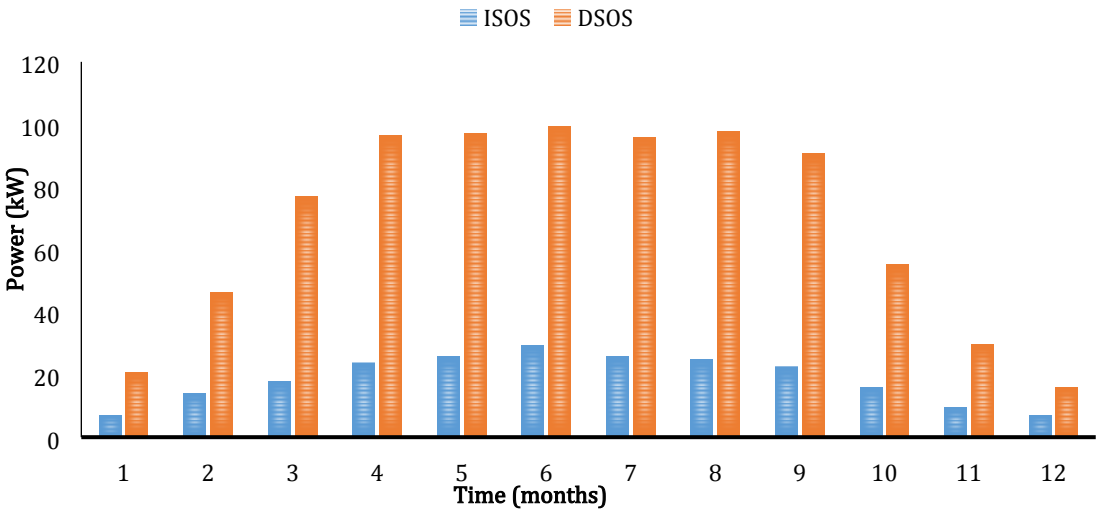

Figure 14: Variation in power transferred by PCM to HTF during discharging mode for DSOS and ISOS on daily average monthly basis

4.3.4 ***Increment in capacity factor by employing PCM storage***

The capacity factor of both systems increases by employing PCM storage in the systems. Figure 15 compares the daily average monthly increment in capacity factor by using PCM storage for both systems. The increment in capacity factor usually depends upon the size of the heat storage unit as well as operating and boundary conditions. It is observed that it generally increases with increment in solar radiation and ambient temperature as shown in Figure 3 and Figure 15. However, in the case of DSOS, it has shown higher increment for the summer months having higher solar radiation and ambient temperature. Moreover, DSOS has shown higher increment in capacity factor as compared to ISOS. Furthermore, with PCM storage, the capacity factor is increased by 17 % and 21.71 % on annual basis for ISOS and DSOS respectively.

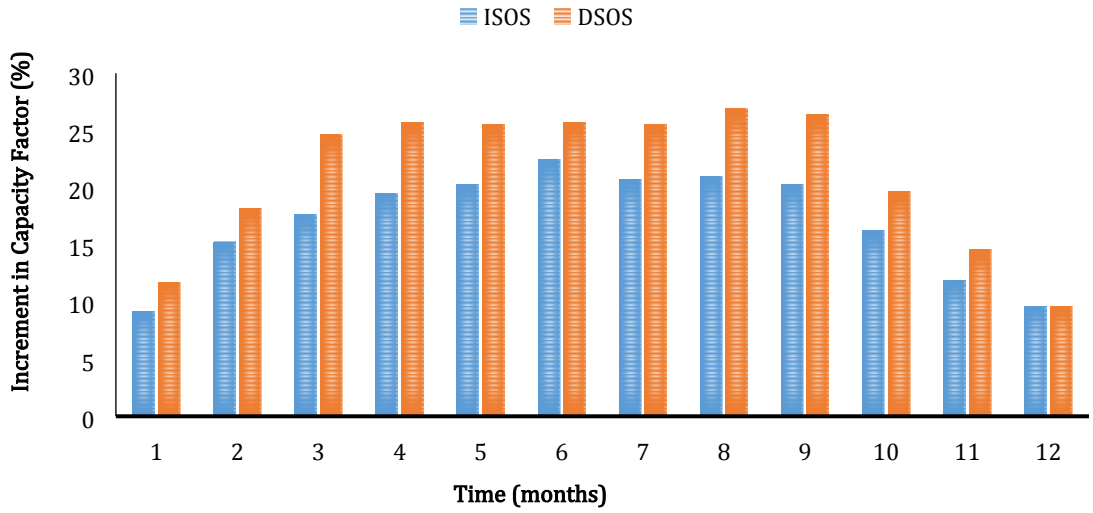

Figure 15: Variation in increment in Capacity Factor by using PCM storage for DSOS and ISOS on daily average monthly basis

5 Conclusions

The thermodynamic performance of DSOS and ISOS is compared in this study. A PCM storage tank is employed in both systems to increase the capacity factor and to assure the stability of power generation. A numerical mathematical model of PCM storage is developed. The model resulted in temperature profiles of HTF and PCM that varies with time. Annual dynamic simulations are carried out under time-varying conditions for both systems. It is concluded from results that DSOS has shown overall better thermal performance as compared to ISOS. The thermal match between HTF and PCM is stronger in the case of DSOS in comparison with ISOS. Hence, the temperature difference between HTF and PCM is found to be 0 K for DSOS and 10 K for ISOS at the maximum HTF temperature. Under given operating and boundary conditions, DSOS has shown 6.1% higher system efficiency and 61.5 kW higher daily average net power output than ISOS on an annual basis. Although the annual amount of heat stored by PCM for ISOS is 1.46 times higher than DSOS in charging mode. This happens might be due to specific heat capacity of water is almost 3 times higher than R245fa at selected operating and boundary conditions. However, during discharging mode, annual amount power transferred by PCM to HTF is 4.8 times higher for DSOS as compared to ISOS because of stronger thermal match of DSOS. Furthermore, DSOS has achieved 195 working hours higher than ISOS on annual basis. However, the performance of both systems can be improved significantly by applying more complex logical control. The future study involves the optimization of latent heat thermal energy storage systems for given constrains.

Acknowledgment

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Nomenclature

Symbols

A	Area [m^2]
a	Heat transfer coefficient
C	Specific heat [J/kg/K]
CF	Capacity factor

D	diameter [m]
F	Friction factor
H	Volumetric enthalpy [J/m^3]
h	Specific enthalpy [kJ/kg]
I	solar radiation [W/m^2]
m	Mass flow rate [kg/s]
L	Length [m]
M	Mass [kg]
P	Pressure [bar]
Q	Heat transferred [J]
q	Energy stored [J]
R	Reynolds number
T	Temperature [K]
W	Work [J]

Abbreviations

P	Pump
V	Valve
CHP	Combined heat and power
CPC	Compound parabolic concentrator
CSP	Concentrated solar power
DSOS	Direct solar organic Rankine cycle system
DSG	Direct steam generation
DVG	Direct vapor generation
EFPC	Evacuated flat plate collector
GWP	Global warming potential
HTF	Heat transfer fluid
ISOS	Indirect solar organic Rankine cycle system
LHS	Latent Heat Storages
ODP	Ozone depletion potential
ORC	Organic Rankine cycle
PCM	Phase change material
RMB	Ren Min Bi
SCHP	Scale distributed solar combined heat and power
SEGS	Solar electricity generation system
SHS	Sensible heat storage
TMY	Typical meteorological year
UK	United Kingdom

Subscript

amb	Ambient
cl	Collector
g	Generator
i	Inlet
mx	Maximum
m	melting point
HTF	Heat transfer fluid
ORC	Organic Rankine cycle
o	outlet
os	The isentropic process
p	Pump
PCM	Phase change material
sp	Solar pump
t	Turbine
th	thermal
w	with

wo	without
wf	Working fluid

Greek letters

β	Angle
λ	Latent heat [J/kg/K]
Δ	Change
ρ	Density [kg/m^3]
ε	Efficiency
η	Thermal efficiency

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