

# Multi-layered solid-PCM thermocline thermal storage concept for CSP plants. Numerical analysis and perspectives.

*Short title: MLSPCM thermocline concept for CSP*

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## Abstract

Thermocline storage concept has been considered for more than a decade as a possible solution to reduce the huge cost of the storage system in concentrated solar power (CSP) plants. However, one of the drawbacks of this concept is the decrease in its performance throughout the time. The objective of this paper is to present a new thermocline-like storage concept, which aims at circumventing this issue. The proposed concept consists of a storage tank filled with a combination of solid material and encapsulated PCMs, forming a multi-layered packed bed, with molten salt as the heat transfer fluid. The performance evaluation of each of the prototypes proposed is virtually tested by means of a detailed numerical methodology which considers the heat transfer and fluid dynamics phenomena present in these devices. The virtual tests carried out are designed so as to take into account several charging and discharging cycles until periodic state is achieved, i.e. when the same amount of energy is stored/released in consecutive charging/discharging cycles. As a result, the dependence of the storage capacity on the PCMs temperatures, the total energy and exergy stored/released, as well as the efficiencies of the storing process are compared for the different thermocline, single PCM, cascaded PCM and multi-layered solid-PCM (MLSPCM) configurations. The analysis shows that the multi-layered solid-PCM concept is a promising alternative for thermal storage in CSP plants.

*Keywords:* Thermal Energy Storage, CSP, Phase Change Materials, Thermocline, Multi-Layered Solid-PCM, Numerical Analysis

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## NOMENCLATURE

$A$	Surface area
$A_t$	Transversal area of tank
$A_w$	Internal surface area of tank's lateral wall
$C_p$	Specific heat at constant pressure
$d_p$	Diameter of filler PCM capsule/solid particle
$e_{cap}$	Capsule's shell width
$ex$	Exergy
$f$	Mass liquid fraction (PCM)
$g$	Gravity acceleration
$h$	Specific total enthalpy
$h_{conv}$	Convection coefficient
$k$	Thermal conductivity
$k_{eff}$	Effective thermal conductivity
$L$	specific latent enthalpy
$m, \dot{m}$	Mass and mass flux
$n_{fm}$	Number of filler particles/capsules in tank section
$Nu$	Nusselt number
$N_r$	Number of control volumes of one filler particle/capsule
$N_x$	Number of tank sections
$p$	Pressure
$Pe$	Péclet number
$Pr$	Prandtl number
$r$	Radial direction
$R_{cond}$	Thermal conduction resistance of capsule shell
$R_{conv}$	Convection resistance between fluid and capsule/solid filler
$Re$	Reynolds number
$t$	Time
$T$	Temperature
$U_{amb}$	Global heat transfer convection coefficient between the fluid and the ambient
$v$	Velocity magnitude
$V$	Volume

$\Delta t$	Time step
$\Delta x$	Tank section height
$\epsilon$	Volume liquid fraction (porosity)
$\mu$	Dynamic viscosity
$\rho$	Density

*Superscripts and subscripts:*

<i>amb</i>	ambient
<i>cap</i>	PCM capsule shell
<i>f</i>	fluid flow
<i>fm</i>	filler material (PCM or solid)
<i>i</i>	Index of tank section/control volume
$i \pm 1/2$	Index of tank section's face limiting $i$ and $i \pm 1$
<i>in</i>	Tank inlet
<i>j</i>	Index of capsule/solid filler control volume
$j \pm 1/2$	Index of filler control volume's face limiting $j$ and $j \pm 1$
<i>l, liq</i>	Liquid phase
<i>out</i>	Tank outlet
<i>s, sol</i>	Solid phase

*Abbreviations:*

CSP	Concentrated Solar Power
HTF	Heat Transfer Fluid
LCOE	Levelized cost of Electricity
MLSPCM	Multi-Layered Solid-PCM
PCM	Phase Change Material
TES	Thermal Energy Storage

## 1. Introduction

Thermal energy storage (TES) systems are an essential feature to make a major profit of solar energy. These systems allow using the thermal energy stored in hours of high solar radiation in times of lower radiation and higher energy demands, reducing the mismatch between the supply and demand. In solar power generation stations, the incorporation of TES systems produce an increase in system reliability and generation capacity, and a decrease of the levelized cost of electricity (LCOE) [1, 2].

For concentrated solar power plants (CSP) the current standard for thermal energy storage is the

8 two-tank molten salt system [3, 4], which make profit of the sensible energy changes of a heat transfer  
9 fluid (molten salt) under a temperature difference. In the search for investment costs reduction,  
10 different designs which result in lower container volumes or in the use of less and/or cheaper storage  
11 media have been proposed as alternatives. Some of these, making use of the materials sensible energy  
12 capacity, are the thermocline tanks [5, 6] and the concrete storage designs [7].

13 Thermocline storage system consists of a single tank, with a volume somewhat higher than one of  
14 the two-tank system, filled with a solid material forming a porous packed bed through which the heat  
15 transfer fluid (HTF) flows. Most of the tank volume is occupied by the solid, which acts as a sensible  
16 energy storage medium, and therefore less of the more costly HTF is needed when compared to an  
17 equivalent two-tank system. Different solid materials have been considered, such as quartzite rocks,  
18 granite, sand [5], asbestos-containing wastes [8], etc.

19 Thermocline tanks rely on the principle of thermal stratification, which occurs in a fluid having  
20 temperature gradients under the action of the gravitational force. The hot fluid, having a lower density  
21 than the cold fluid, is pushed upwards by the buoyancy force while the low temperature fluid is  
22 displaced downwards. Therefore, the hot fluid is placed in the upper part of a tank, while the colder  
23 fluid stays at the bottom. As a consequence, a vertical temperature gradient is formed which is called  
24 “thermocline”. The filler material helps in maintaining the thermal gradient, preventing possible  
25 mixing flows that may be present due to effects such as cooling through the walls (see for instance  
26 [9, 10]) or strong inlet flow currents [11].

27 Phase change materials (PCM) can also be used to store energy, using less storage material than  
28 would be used with a sensible energy storage medium, taking advantage of the latent energy changes  
29 during a phase change. The resulting storage device should be more compact, and hopefully cheaper,  
30 than one that only makes use of the sensible energy changes. Thermal storage devices using encapsu-  
31 lated PCMs have been studied as a form of thermal storage devices for CSP applications by several  
32 authors. Liu et al. [12] perform an extensive review of PCMs suitable to be used in TES for CSP  
33 plants and of heat transfer enhancement methods. Michels and Pitz-Paal [13] studied, experimentally  
34 and numerically, the performance of storage systems using vertical shell and tube heat exchangers  
35 with different PCMs enclosed between the shell and the tubes, with different melting points (cascaded  
36 PCM), for parabolic trough plants. Shabgard et al. [14] studied cascaded latent heat storage with  
37 gravity-assisted heat pipes for CSP. They performed numerical simulations using a thermal network  
38 model and evaluated thermal performance of the different designs after a single charge/discharge cycle.  
39 Nithyanandam et al. [15] numerically analyzed packed bed thermal storage with single encapsulated  
40 PCMs by studying their performance after a single charging/discharging cycles and after a periodic

41 cyclic state is reached. Parametric studies were performed and guidelines for designing latent ther-  
42 mocline storage systems for CSP were established. Flueckiger et al. [16] studied thermocline storage  
43 for solar power stations augmented with latent heat. They integrated their numerical model of the  
44 thermocline into a system-level model for the CSP plant and evaluated the effect of the increase of  
45 the storage capacity with latent heat. When compared against solid-filled thermocline, limitations in  
46 the thermal performance of designs including a single PCM were observed, while some improvement  
47 was obtained with some of the cascaded PCM configurations.

48 Moreover, combination of latent and sensible storage devices has been studied for CSP plants  
49 with direct steam generation (DSG) [17], where a PCM storage unit is intended for vapor generation  
50 (evaporation) and the sensible energy units for absorbing the sensible energy of the heat transfer fluid  
51 (preheating and superheating).

52 One of the configurations that have received the most attention is that of packed beds. Many  
53 numerical investigations of thermal storage in packed beds can be found in the literature. Ismail and  
54 Stuginsky [18] performed a comparative analysis of different packed bed models used for sensible and  
55 latent heat storage. Flueckiger et al. [19] reviewed different experimental and numerical studies on  
56 thermocline tanks for solar thermal storage. On one hand, models for packed beds of solid materials  
57 [6, 19, 20, 21] usually disregard the temperature gradients inside the particles. On the other hand, if  
58 the bed consists of PCM capsules [15, 16, 22, 23], thermal gradients inside them may be significant  
59 and are generally, but not always, taken into account. Karthikeyan and Velraj [24] performed a  
60 comparison of three one-dimensional models for packed beds of PCM spherical capsules, where the  
61 effect of considering the radial variation of temperature inside the capsules was tested. All these  
62 models are based on discretizing the conservation equations for the heat transfer fluid and the filler  
63 bed, and usually, several simplifying assumptions are made. Depending on the scope of the numerical  
64 code, one-, two- or three-dimensional simulations can be performed. However, due to the significantly  
65 higher computational costs associated with two- (e.g. [6, 20]) and three-dimensional (none found in  
66 the literature) models, one-dimensional analysis is usually chosen for studying several working cycles  
67 of TES systems [15, 16, 22, 23, 24].

68 In this work, a new concept of thermocline-like storage system is proposed, consisting in com-  
69 bining low-cost solid and PCM filler materials, appropriately chosen and placed inside the tank in a  
70 multi-layered manner. This concept, initially presented in a congress paper [25], has been called Multi-  
71 Layered Solid-PCM (MLSPCM). The main idea behind MLSPCM configurations is the inclusion of  
72 high and low melting-point PCMs as filler materials at the ends of the tank, close to the inlet/outlet  
73 ports. In [25], the thermal performance of different MLSPCM designs was tested by means of a sim-

74 plified numerical model. Preliminary results indicated that MLSPCM configurations may reduce the  
75 thermocline degradation occurring in the single-solid filled tanks and thus achieve a higher efficiency  
76 in the use of the storage capacity.

77 Zanganeh et al. [26] recently studied a similar configuration using air as the heat transfer fluid,  
78 in which a single PCM layer of high melting point was placed at the top of a packed bed of rocks.  
79 They concluded that the PCM layer contributed to the stabilization of the outflow temperature but  
80 did not have an effect on the thermal efficiency of the system. The evaluation was performed after  
81 several charge/discharge cycles of the same duration.

82 In this work, a numerical model for evaluating the thermal behavior and optimizing the design  
83 of packed bed systems is presented. The model considers axial thermal conduction within the heat  
84 transfer fluid and an special treatment for the convective term of the energy equation, diminishing  
85 the artificial numerical diffusion. Since the intention is to test both solid and PCM filler materials,  
86 radial variation of temperature within particles/capsules is taken into account. Numerical experiments  
87 are carried out in order to compare the thermal performance of the MLSPCM prototypes against  
88 different existent thermocline-like designs, such as solid-filled thermocline, single encapsulated PCM  
89 and cascaded PCM concepts. Several configurations are here analyzed, and the results are discussed in  
90 detail. Thermal performance is evaluated after running several charge/discharge cycles, until a periodic  
91 state is reached. Time operation of the processes (charge and discharge) are not fixed beforehand but  
92 depend on the temperature of the fluid coming out of the TES, which is limited by some temperature  
93 thresholds. The intention with this is to mimic the operating conditions of a real CSP plant, where  
94 the solar receivers and power block impose certain limits to the temperature of the HTF. Thermal  
95 evaluation is mainly based on terms of total energy storage, efficiency in the use total storage capacity,  
96 proportion of PCM effectively changing phase and exergy outputs.

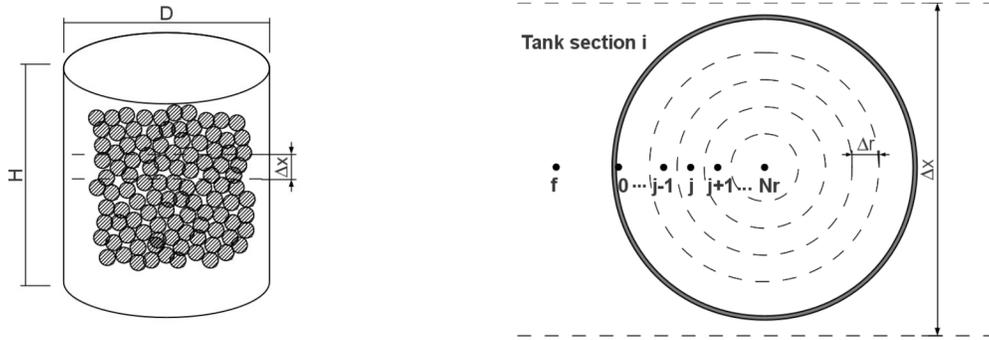
## 97 **2. Mathematical modeling and numerical implementation**

98 Mass, momentum and energy conservation equations have to be solved in order to be able to  
99 simulate the thermal behavior of a thermocline-like tank. Some simplifying assumptions are made  
100 and empirical correlations are used. The most relevant assumptions are:

- 101 1. One-dimensional fluid flow and temperature distribution (in the flow direction).
- 102 2. One-dimensional heat transfer in filler particles/capsules (radial direction).
- 103 3. Spherical shape of filler particles/capsules.
- 104 4. Constant density of both fluid and filler bed materials (solid and PCM).
- 105 5. Natural convection and contact melting inside the PCM capsules are neglected.

- 106 6. Heat conduction between different filler material particles/capsules is not considered.  
 107 7. Negligible radiation heat transfer.

108 The energy conservation equations are discretized using the Finite Volume Method (FVM). The  
 109 tank is divided in  $N_x$  transversal cylindrical sections of height  $\Delta x$  (see Fig. 1a). In each tank  
 110 section, a single representative particle/capsule needs to be simulated, as all are affected by the same  
 111 fluid temperature due to the one-dimensionality assumption. This filler particle/capsule, assumed as  
 112 spherical, is discretized in the radial direction in  $N_r$  control volumes, as shown in Fig. 1b.



(a) Sketch representing the cylindrical container with the PCM capsules packed in a random fashion

(b) Discretization details of the tank and of a representative particle/capsule, indicating the sub-indices used for tank sections (i) and capsule control volumes (j)

**Figure 1:** Domain and discretization

## 113 2.1. Energy

### 114 2.1.1. Heat transfer fluid (HTF)

115 The semi-discrete energy conservation equation of the fluid in the  $i^{\text{th}}$  tank section ( $i = 1 \dots N_x$ )  
 116 results in:

$$\begin{aligned} \rho_f \epsilon_i V_i C_{p,f} \frac{\partial T_{f,i}}{\partial t} = A_t \left( k_{eff} \frac{\partial T_f}{\partial x} \right) \Big|_{i-1/2}^{i+1/2} - \dot{m} C_{p,f} (T_{f,i+1/2} - T_{f,i-1/2}) \\ - n_{fm,i} \frac{T_{f,i} - T_{i,0}}{R_{conv,i} + R_{cond,i}} - U_{amb} A_{w,i} (T_{f,i} - T_{amb}) \end{aligned} \quad (1)$$

117 where  $T_{i,0}$  is the temperature of the internal surface of the particles/capsules (boundary node in fig.  
 118 1b). In the advective term (second in the right hand side) the fluid is assumed to be coming from  
 119 section  $i - 1$  and going to section  $i + 1$ .  $R_{cond}$  stands for the thermal resistance in the PCM capsules  
 120 due to the capsule shell. The mass of the shell is disregarded here and is not considered to add any  
 121 thermal inertia. The calculation of the thermal resistance due to convection between the HTF and the

122 filler material ( $R_{conv}$ ) requires the fluid-to-bed Nusselt number, which is calculated using the following  
 123 correlation, obtained from [27]:

$$Nu = 2.0 + 1.1Re^{0.6}Pr^{1/3} \quad \text{where } Re = \frac{\rho_f v_f d_p}{\mu_f} \quad (2)$$

124 As stated in [27], the correct use of Eq. (2) should take into account the effects of solid-phase  
 125 conduction and thermal dispersion in the diffusive term [first term on the right hand side of Eq. (1)]  
 126 and not only molecular diffusion. Therefore, the effective thermal conductivity is evaluated as follows:

$$k_{eff} = k_{eff}^0 + k_{eff}^{disp}$$

127 where the stagnant effective thermal conductivity ( $k_{eff}^0$ ) is determined here as in [28]:

$$k_{eff}^0 = \left( \frac{k_{fm}}{k_f} \right)^{0.280 - 0.757 \log_{10}(\epsilon) - 0.057 \log_{10}(k_{fm}/k_f)} \quad (3)$$

128 and the effective conductivity due to thermal dispersion ( $k_{eff}^{disp}$ ) is determined according to the following  
 129 correlation [29]:

$$\frac{k_{eff}^{disp}}{k_f} = 0.00232 Pe^2 \quad \text{where } Pe = RePr$$

130 In this work, some of the studied cases include one or several PCMs as filler materials. Since solid  
 131 and liquid phases may have different thermal conductivities, a criterion is needed to determine the  
 132 value of  $k_{fm}$  used in Eq. (3). The criterion adopted here consists in using the thermal conductivity  
 133 calculated at the radius of capsule that divides the sphere in two parts of the same volume.

### 134 2.1.2. Filler material

135 The energy balance for the inner nodes ( $j = 1 \dots N_r$ ) of the filler material remains:

$$\rho_f V_{i,j} \frac{\partial h_{i,j}}{\partial t} = \left( k_{fm} A \frac{\partial T}{\partial r} \right)_{i,j-1/2} - \left( k_{fm} A \frac{\partial T}{\partial r} \right)_{i,j+1/2} \quad (4a)$$

136 while for the boundary node ( $j = 0$ ), in contact with the heat transfer fluid, results in:

$$\rho_f V_{i,0} \frac{\partial h_{i,0}}{\partial t} = \frac{T_{f,i} - T_{i,0}}{R_{conv,i} + R_{cond,i}} - \left( k_{fm} A \frac{\partial T}{\partial r} \right)_{i,1/2} \quad (4b)$$

137 In order to solve these equations it is necessary to define a relation between the enthalpy and the  
 138 temperature of the filler materials. Considering constant specific heats for each phase, these relations  
 139 are:

$$\begin{aligned}
h - h_0 &= C_{p,s}(T - T_0), & T &\leq T_s \\
h - h_0 &= C_{p,s}(T - T_0) + fL, & T_s &< T \leq T_{sl} \\
h - h_0 &= C_{p,l}(T - T_{sl}) + C_{p,s}(T_{sl} - T_0) + fL, & T_{sl} &< T \leq T_l \\
h - h_0 &= C_{p,l}(T - T_{sl}) + C_{p,s}(T_{sl} - T_0) + L, & T_l &< T
\end{aligned}$$

140  $T_{sl}$  indicates the temperature in the phase change range chosen as the transition temperature for  
141 the specific energy from solid to liquid, or vice versa. Mass liquid fraction ( $f$ ) values range from 0  
142 (pure solid) to 1 (pure liquid), which, in this work, are calculated as a linear function of temperature  
143 in the phase change interval:

$$f = \frac{T - T_s}{T_l - T_s} \quad (5)$$

144 By taking a very narrow temperature range ( $T_l - T_s$ ), fixed melting point PCMs can also be  
145 modeled with this approach. Hence, a unique value of  $h$  exists for each value of  $T$ , and the energy  
146 balance [Eq. (4)] may be expressed with  $T$  as the only variable. It should be noted that, since the  
147 location of the solid-liquid interface is implicitly determined by values of  $f$ , explicit tracking of interface  
148 is avoided with this strategy.

## 149 2.2. Momentum

150 To determine the pressure drop in the packed bed, the following momentum equation is solved:

$$\frac{\delta p}{\delta x} \Big|_i = \pm \left( \frac{5}{Re_{1,i}} + \frac{0.4}{Re_{1,i}^{0.1}} \right) \frac{6\rho_f v_f^2 (1 - \epsilon_i)}{d_{p,i} \epsilon_i^3} - \rho_f g \quad (6)$$

$$\text{where } Re_{1,i} = \frac{\rho_f v_f d_{p,i}}{6(1 - \epsilon_i)\mu_f} \text{ (spherical particles) and } v_f = \frac{\dot{m}}{\rho_f A_t}$$

151 Eq. (6) is the Carman correlation for packed beds, which is generally used for solid objects forming  
152 a bed [30]. In this equation  $x$  increases from the bottom to the top, and therefore, the positive sign is  
153 used in the discharge of the tank while in the charge process the negative sign is used. The last term  
154 accounts for the pressure reduction/increase due to the gravitational action.

## 155 2.3. Exergy

156 For evaluating the power generating potential of the energy delivered by the thermal storage, the  
157 exergy global balance of the heat transfer fluid is calculated in the following manner:

$$\dot{m}(ex_{out} - ex_{in}) = \dot{m}C_{p,f}(T_{out} - T_{in} - T_{ref}\ln\frac{T_{out}}{T_{in}}) \quad (7)$$

158 where  $T_{ref}$  is the temperature corresponding to the dead state, which in this work has been taken as  
 159 45°C due to being a reasonable value for the temperature at which the vapor is condensed in the power  
 160 generation block. The resulting exergy flow is the difference between the exergy exiting and entering  
 161 the tank with the fluid.

#### 162 2.4. Discretization details

163 Fully implicit schemes, such as that previously adopted in [25], may suffer from numerical diffusion  
 164 in some degree. Since an accurate modeling of the temperature gradient is needed to evaluate the  
 165 thermocline tank performance, a different method, similar to that presented by Oppel et al., [31], is  
 166 used.

167 An upwind scheme with a first order explicit time integration has been adopted for the advective  
 168 term, combined with the choice of a time step such that CFL=1 ( $\Delta t = \epsilon\Delta x/u$ ). Therefore, each tank  
 169 section is “filled” completely in each time step by the fluid coming from the upstream section with a  
 170 temperature equal to the obtained in the previous time step. With this, if no damping is present (e.g.  
 171 heat transfer to the filler material), a sharp temperature front is exactly transported from the inlet to  
 172 the outlet with the velocity of the fluid flow. In cases where different porosities are present throughout  
 173 the packed bed, different CFL numbers result for the same  $\Delta t$ . Here, CFL=1 has been enforced for  
 174 the most restrictive zone (with the lowest porosity), and thus in the rest of the domain it has been  
 175 maintained between 0 and 1, where some numerical diffusion is present. The physical diffusion in the  
 176 HTF is modeled by the diffusive term in Eq. (1). A central difference spatial discretization and an  
 177 explicit time integration have been adopted (although if the error in this term is significant, more  
 178 than one iteration per time step is performed, which results in an approximation to an implicit time  
 179 integration). To avoid instabilities and for accuracy reasons, the restriction in the time step by the  
 180 diffusive term is:  $\Delta t = 0.3(\epsilon\rho C_p\Delta x^2/2k_{eff})$ . If the refinement of the grid is high, this limitation  
 181 may be more restrictive than that of the convective term, and therefore, the CFL=1 condition is not  
 182 applied anymore. For the diffusive terms of the filler energy equations [Eq. (4)], central differencing  
 183 and fully implicit time integration have been adopted.

184 The algorithm used is similar to that indicated in [25], except that more than one iteration through  
 185 the tank sections has been necessary at times, especially at the beginning of each process, in order  
 186 to calculate more accurately the diffusive term. In each tank section ( $i$ ), fluid and filler material  
 187 temperatures have to be solved. The final matrix of coefficients derived from the system of equations

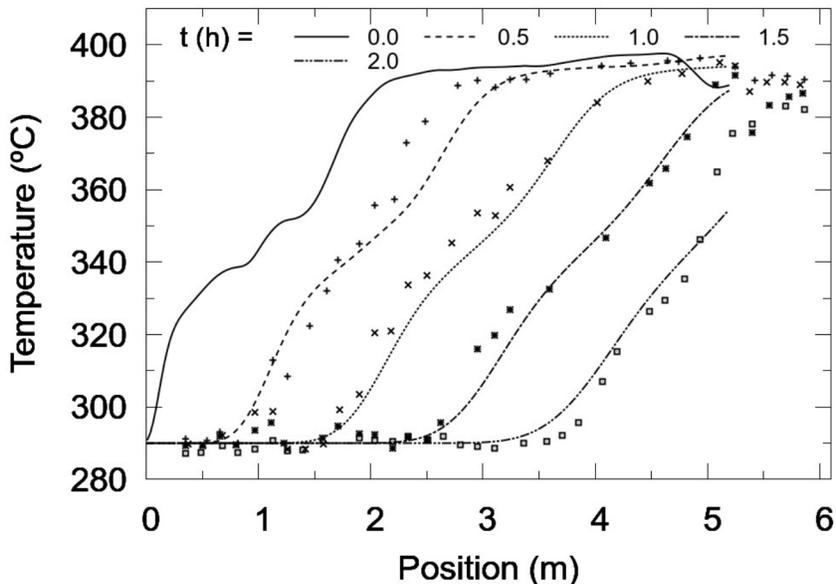
188 in each container section has a tri-diagonal pattern. This allows the usage of a TDMA algorithm to  
 189 solve the linear system.

190 *2.5. Model validation*

191 The first case used to validate the model is that coming from the experimental work of Pacheco  
 192 et al. [5]. There, a thermocline tank filled with quartzite rock and sand was tested, using molten  
 193 salt as the heat transfer fluid. This case has been adopted as the validation case in several works on  
 194 thermocline numerical modeling with solid and encapsulated PCM filler materials, see for instance  
 195 [15, 16, 19, 20, 21].

196 The thermo-physical properties of the molten salt and of the mixture of quartzite rock and sand  
 197 are indicated in Table 2. The HTF mass flow, which has been calculated following the same strategy  
 198 as in [32], is set to 5.852 kg/s. The porosity is 0.22 and the effective particle diameter is 0.015 m.  
 199 The initial state adopted in the simulations is the temperature map obtained from the experimental  
 200 measures at time 11:30 [5]. Filler material and fluid inside the tank have been assumed to be at  
 201 thermal equilibrium at this initial state.

202 The part of the tank containing solid filler material, of 5.2 m height, has been discretized axially in  
 203 208 sections ( $\Delta x = 2.5$  cm), while each simulated filler particle has been divided radially in 10 control  
 204 volumes and the time step was set to  $\Delta t \sim 12.45$  s (CFL = 1). These parameters have been verified  
 205 in the sense of producing grid independent results (a difference of  $\sim 0.1\%$  has been calculated against  
 206 a case with 416 sections). Ambient losses have been neglected.

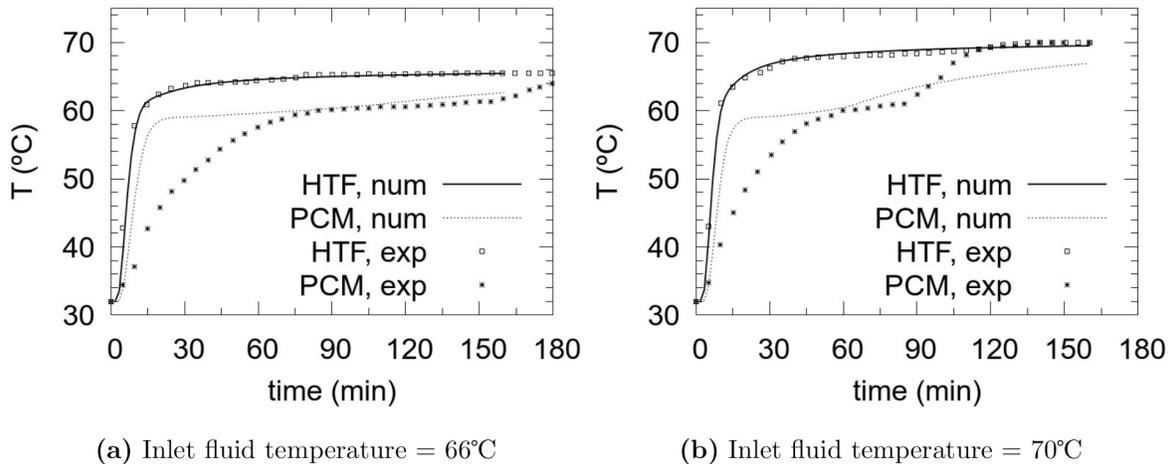


**Figure 2:** Validation. Solid lines correspond to the numerical results, while dots indicate the experimental results from Pacheco et al. [5]. The chronological order of the curves is from left to right.

207 Simulation results are presented in Fig. 2. It can be observed that the thermal gradient is well  
 208 reproduced and that it moves towards the top as the discharge proceeds. Good agreement between  
 209 experimental and numerical results is observed. The small discrepancies observed may be due to several  
 210 causes, such as: uncertainties of the experimental measurements, unavailability of all the parameters  
 211 from the original work of Pacheco et al. [5] and simplifications of the numerical model.

212 The second validation case corresponds to the charging of a tank filled with encapsulated PCM,  
 213 from the experimental work of Nallusamy et al. [33]. All the configuration parameters and properties  
 214 have been adopted from that work.

215 Figure 3 shows the results of evolution of temperature of the HTF (water) and PCM (paraffin  
 216 melting at  $60 \pm 1^\circ\text{C}$ ), measured at  $x/L = 0.5$ , where  $x$  is the axial position and  $L$  is the total height. In  
 217 [33], the position inside the capsule at which the temperature is measured is not indicated. Numerical  
 218 results of PCM temperatures shown in figure 3 are those obtained at the radius that divides the sphere  
 219 in two parts of equal volume. Grid resolution for this case is  $N_x = 92$  and  $N_r = 55$ , which has been  
 220 checked for grid independence.



**Figure 3:** Temperature evolution at  $x/L = 0.5$  of HTF and PCM. Experimental results, indicated by symbols, were extracted from [33]. Numerical results of PCM temperature are taken at the radius that divides the sphere in two parts of equal volume.

221 A good agreement is observed between experimental results (obtained from [33]) and numerical  
 222 simulations, specially for the HTF. Some discrepancies are observed in the PCM temperatures. This  
 223 is probably due to several reasons, such as not accounting for natural convection at the melting,  
 224 differences between real thermo-physical properties and those used (paraffins have been observed to  
 225 present high phase-change temperature ranges, as in [24], and thus melting probably starts at a  
 226 lower temperature than that assumed) and uncertainty about the exact position of the temperature  
 227 sensor. However, these discrepancies do not reflect significantly in the HTF temperature, for which

228 the agreement is very good.

### 229 **3. Evaluation of different solutions**

230 This section is devoted to the evaluation of thermocline tanks as thermal storage systems for CSP  
231 applications. The thermocline prototype tested by Pacheco et al. [5], is adopted as a reference case  
232 from which the operating conditions and tank global dimensions are taken.

233 Usually, in a CSP plant, the outlet temperature in the discharge process is limited by the minimum  
234 temperature that is admissible for the fluid feeding the power block. Similarly, for the charging process,  
235 the outlet temperature is limited by the restriction in the temperature of the fluid coming into the  
236 solar receivers. Therefore, both the operating time and the stored energy are determined by the level  
237 of temperatures attained by the outlet fluid in the charge and discharge processes.

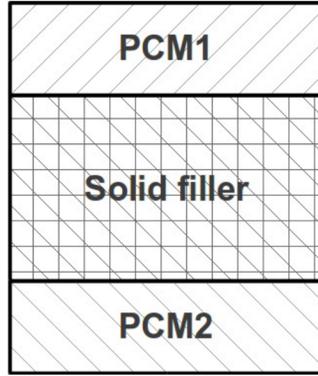
238 Since the main objective is the evaluation of the performance of single thermocline-like tanks as  
239 storage devices in the context of a CSP plant, which is intended to be operated in charge/discharge  
240 cycles (one per day) during several years, this evaluation should not rely on values of energy (and  
241 exergy) stored or released in a single charge/discharge cycle. As the performance in a single charge  
242 (or discharge) process is highly dependent on the initial temperature distribution inside the tank,  
243 numerical experiments are carried out for several cycles until a periodic state—independent of the  
244 initial conditions of the first cycle—is achieved. Final performance values are determined at this  
245 state.

246 Furthermore, the substitution of all or part of the solid filler material by encapsulated PCM is  
247 considered. Therefore, tanks exclusively filled with an encapsulated PCM are tested, where the melting  
248 point has been taken as a parameter. Results obtained by the different choices of the melting point  
249 lead to the proposal of different combinations of “hybrid” thermocline configurations, in which both  
250 solid and encapsulated PCM filler materials are included. This new concept of thermocline-like storage  
251 system is herein called Multi-Layered Solid-PCM (MLSPCM). Fig. 4 shows a sketch of a MLSPCM  
252 configuration with 3 layers. Furthermore, cascaded PCM configurations are also tested and compared.

253 The same spatial discretization has been used for all the present cases, with 416 axial tank sections  
254 sections and 10 filler capsules/particles nodes. The difference with a coarser grid having 208 axial  
255 sections is less than 5% in all the cases, and therefore, the accuracy obtained is assumed to be good  
256 enough.

#### 257 *3.1. Study cases*

258 The following operating conditions are assumed:



**Figure 4:** Sketch of a 3-layered MLSPCM configuration. PCM1 and PCM2 have high and low melting points, respectively.

- 259 • The geometry of the tank and operating conditions are the same for all the cases. Diameter  
260 of the filler particles/capsules is 15 mm. The porosity of the packed bed depends on the filler  
261 material and it is assumed to be 0.22 for the solid filler material and 0.34 for the encapsulated  
262 PCM. Moreover, a shell thickness of 0.4 mm is assumed for the PCM capsules.
- 263 • The operation time is not fixed. Instead, outlet temperature limits are imposed, which force the  
264 end of each process (charge or discharge) if the outflow temperature is not within these limits.  
265 These temperature intervals will be referred to as “admissible” temperature ranges. Here, both  
266 admissible ranges have been assumed to be 15% of the maximum temperature interval (100°C);  
267 i.e. in the charging phase, the outlet fluid temperature is allowed to be between 290°C and  
268 305°C, while in the discharge it must be between 375°C and 390°C.
- 269 • Molten salt flow is fixed to 5.852 kg/s for both processes.
- 270 • Ambient losses are neglected [ $U_{amb} = 0$  in Eq. (1)].
- 271 • Several consecutive charge/discharge cycles are simulated until a periodic thermal state is reached,  
272 i.e. when there is no variation of stored/released energy between consecutive cycles. Since ambi-  
273 ent losses are neglected, in the periodic state the same energy that is stored in the charge must  
274 be released in the discharge.

275 As the admissible temperature intervals for both charge and discharge processes are quite narrow,  
276 outlet fluid temperatures for all the cases here studied are very similar. Therefore, a higher operation  
277 time is directly related to a higher stored (or released) energy.

278 In Table 1, a code for each case/configuration is defined. The presented cases can be classified  
279 according to the filler material/s used as: “pure” thermocline (A); single PCM (B); multi-layered

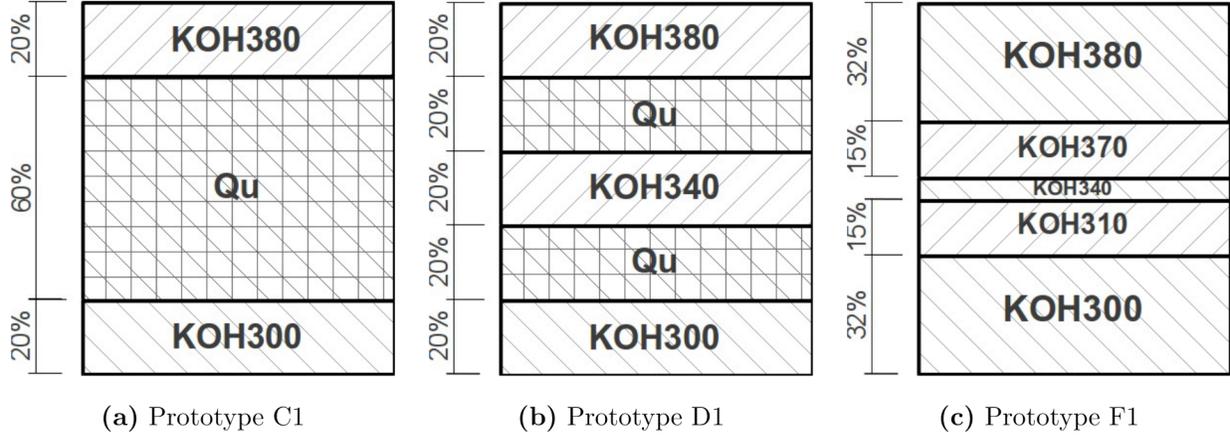
280 solid-PCM (C and D); and cascaded PCM (F). Percentages between brackets indicate the portion of  
 281 total height occupied by each filler material. It should be noted that the chosen PCMs are fictitious,  
 282 having the same thermal properties as those of potassium hydroxide (KOH) but with different fusion  
 283 temperatures. The exception to this is case B1, where KOH is considered with its actual melting point  
 284 (360°C according to [13]). This procedure has been adopted in order to account for the variations in  
 285 performance exclusively due to the change in the fusion temperature of the PCMs. Figure 5 depicts  
 286 sketches of some of the prototypes tested.

**Table 1:** Codification of cases.

Filler material <sup>1</sup>	Code
Quartzite rock & sand (Qu) (100%)	A
KOH (100%)	B1
KOH380 (100%)	B2
KOH300 (100%)	B3
<b>MLSPCM:</b> KOH380-Qu-KOH300 (20%-60%20%)	C1
<b>MLSPCM:</b> KOH380-Qu-KOH300 (40%-20%40%)	C2
<b>MLSPCM:</b> KOH380-Qu-KOH300 (10%-80%10%)	C4
<b>MLSPCM:</b> KOH380-Qu-KOH340-Qu-KOH300 (20%-20%-20%-20%-20%)	D1
<b>MLSPCM:</b> KOH380-Qu-KOH340-Qu-KOH300 (20%-25%-10%-25%-20%)	D2
KOH380-KOH370-KOH340-KOH310-KOH300 (32%-15%-6%-15%-32%)	F1
KOH380-KOH370-KOH340-KOH310-KOH300 (32%-9%-18%-9%-32%)	F2

<sup>a</sup>Materials KOHXXX (where XXX is a 3 digit number) are fictitious PCMs with fusion temperatures indicated by the number XXX (e.g. 300°C), whose thermal properties are equal to those of KOH (melting point = 360°C). The order in which the materials are indicated is the one in which they are placed inside the tank, from the top to the bottom. Between brackets, the proportion of the tank height occupied by each filler layer is indicated.

287 Table 2 shows the physical properties used in the simulations. Table 3 shows the mass of solid  
 288 filler material, PCM and HTF contained. Due to the higher porosity of the PCM layers, more confined  
 289 heat transfer fluid is contained in the configurations including encapsulated PCMs. Furthermore, as  
 290 the solid filler material is denser than the PCM, prototypes including more of the former contain a  
 291 higher total mass. The same table also presents data of the storage capacity for each configuration,  
 292 i.e. the maximum amount of energy that could be stored taking into account both sensible and latent  
 293 energy contributions, with a temperature jump equal to the difference between the hot and cold inlet  
 294 fluid temperatures of charge and discharge processes, respectively. Here, the maximum temperature



**Figure 5:** Sketches of some of the different filler configurations tested.

**Table 2:** Thermo-physical properties.

	Quartzite rock & sand (Qu) [20]	PCM (KOHXXX) [13]	Molten Salt (HTF) [34]
$\rho$ ( $kg/m^3$ )	2500	2040	1873.8
$C_{p,s}$ ( $J/kg K$ )	830	1340	-
$C_{p,l}$ ( $J/kg K$ )	-	1340	1501.5
$k_s$ ( $W/m K$ )	5.69	0.5	-
$k_l$ ( $W/m K$ )	-	0.5	$0.443 + 1.9 \times 10^{-4}T(^{\circ}C)$
$\mu$ ( $Pa.s$ )	-	-	$22.714 \times 10^{-3} - 0.12 \times 10^{-3}T +$ $2.281 \times 10^{-7}T^2 - 1.474 \times 10^{-10}T^3$
$L$ ( $J/kg$ )	-	$1.34 \times 10^5$	-

295 difference is  $100^{\circ}C(290^{\circ}C- 390^{\circ}C)$ . It can be observed that, even in the cases where the filler material is  
 296 only encapsulated PCM (cases B1-3 and F1-2), the sensible energy capacity is higher than the latent  
 297 one. This is due basically to two reasons: first, the temperature jump is relatively high, making the  
 298 sensible energy capacity of the PCMs to be equal to their latent energy capacity ( $C_p\Delta T = L$ ); and  
 299 second, the HTF confined inside the tank contributes with an extra sensible energy capacity.

**Table 3:** Mass confined inside the tank and storage capacity.

Mass data (ton)	A	B1	B2	B3	C1	C2	C4	D1	D2	F1	F2
Mass of PCM	0.0	42.0	42.0	42.0	17.0	33.9	8.5	25.0	21.0	42.0	42.0
Mass of solid filler material	71.7	0.0	0.0	0.0	42.7	13.8	57.2	28.9	35.8	0.0	0.0
Mass of confined HTF	15.2	23.4	23.4	23.4	18.5	21.8	16.8	20.1	19.3	23.4	23.4
Total mass	86.8	65.4	65.4	65.4	78.2	69.5	82.5	74.1	76.1	65.4	65.4
Storage Capacity											
Filler material (MWh)	1.65	3.13	3.13	3.13	2.25	2.84	1.95	2.53	2.39	3.13	3.13
Confined HTF (MWh)	0.63	0.98	0.98	0.98	0.77	0.91	0.70	0.84	0.80	0.98	0.98
Total (filler + HTF) (MWh)	2.28	4.10	4.10	4.10	3.02	3.75	2.65	3.37	3.19	4.10	4.10
Total sensible energy (%)	100.0	61.9	61.9	61.9	79.1	66.4	88.1	72.3	75.5	61.9	61.9
Total latent energy (%)	0.0	38.1	38.1	38.1	20.9	33.6	11.9	27.7	24.5	38.1	38.1

300 *3.2. Results and discussion*

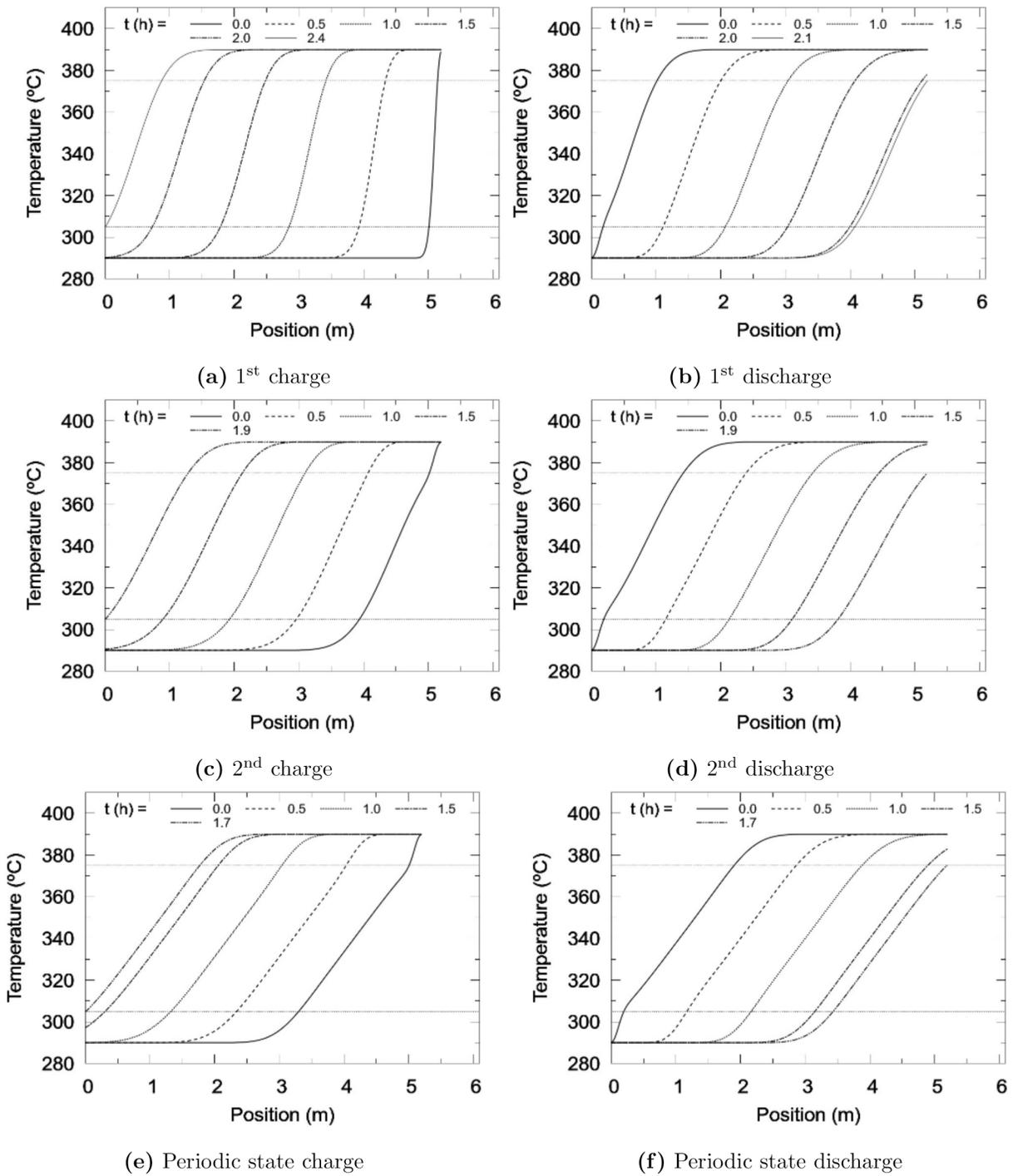
301 *3.2.1. Case A: Solid filler material*

302 Table 4 shows the thermal performance results for each case. Case A (“pure” thermocline) is shown  
303 to behave quite poorly in terms of stored energy when compared against the rest of the cases (with  
304 exception of case B1). The stored energy at thermal equilibrium is 1.45 MWh, which is around 63% of  
305 the storage capacity. This somewhat low efficiency is due to the thermocline degradation throughout  
306 the several charging-discharging cycles, enforced by the assumed temperature thresholds.

**Table 4:** Performance results for each configuration

Results	A	B1	B2	B3	C1	C2	C4	D1	D2	F1	F2
Operation time (h) <sup>1</sup>	1.67	1.16	2.61	2.62	2.86	2.99	2.68	1.82	3.00	3.28	2.03
Stored Energy in Filler material (MWh)	1.05	0.63	1.42	1.43	1.72	1.71	1.65	1.11	1.83	2.00	1.25
Stored Energy (Filler + confined HTF) (MWh)	1.45	1.00	2.19	2.20	2.32	2.42	2.22	1.48	2.43	2.66	1.64
Stored Energy / Storage capacity (%)	63.4	24.5	53.4	53.5	76.9	64.5	83.7	43.8	76.2	64.9	40.0
Sensible energy stored / Total stored (%)	100.0	96.3	90.7	90.7	80.0	80.0	86.7	69.8	73.7	64.0	60.6
Latent energy stored / Total stored (%)	0.0	3.7	9.3	9.3	20.0	20.0	13.3	30.2	26.3	36.0	39.4
Effective mass of PCM changing phase (%)	-	2.4	13.1	13.1	73.6	38.4	93.2	47.9	81.7	61.3	41.4
Exergy difference at charge (MWh)	-0.70	-0.48	-1.05	-1.06	-1.12	-1.17	-1.07	-0.71	-1.18	-1.29	-0.79
Exergy difference at discharge (MWh)	0.69	0.48	1.04	1.05	1.11	1.15	1.06	0.70	1.16	1.27	0.78
Pressure losses due to filler bed (Pa)	< 400	< 100	< 100	< 100	< 250	< 150	< 350	< 200	< 250	< 100	< 100

<sup>1</sup>In cases where the charge and discharge operation times are different, (e.g. B2 and B3) the mean value between processes is shown.



**Figure 6:** Case A. Temperature maps at various instants for the first two cycles and the periodic state. The chronological order of the curves is from left to right in the charge and from right to left in the discharge.

307 Figure 6 shows the temperature maps obtained for case A, of charge and discharge processes at  
 308 various instants for the first two cycles and the periodic state. The degradation of the thermocline  
 309 can be clearly observed by comparing the 1<sup>st</sup> and last cycles. The initial condition in the first charge  
 310 process is a uniform cold temperature throughout the tank. For the 2<sup>nd</sup> cycle, the initial condition of

311 the filler material and HTF is not anymore a uniform temperature curve, but one with the temperature  
312 gradient resulting from the last discharge process. This change in the initial conditions is inevitable  
313 due to the requirements imposed on the outgoing fluid temperature, which enforce it to lie inside the  
314 admissible range. Since the difference between the incoming fluid temperature and that of the filler  
315 material is lower than in the 1<sup>st</sup> charge process, the heat transfer rate, and thus the thermal gradient  
316 (in absolute value), are also lower in the 2<sup>nd</sup> charging.

317 Therefore, during the consecutive charging/discharging cycles, the thermal gradient tends to get  
318 “flattened” until a periodic unsteady state is reached. Due to this thermocline degradation, the  
319 stored/released energy in this periodic state is lower than those of the previous cycles, and so is the  
320 operation time, since outlet thresholds are reached earlier.

321 A useful way of estimating the stored/released energy in each process is to calculate the area  
322 between the initial temperature map and the last one, since sensible energy differences are proportional  
323 to the temperature jumps. From the comparison between figures 6a and 6e, a clear difference can be  
324 observed in the area between initial and last temperature maps of the charge processes.

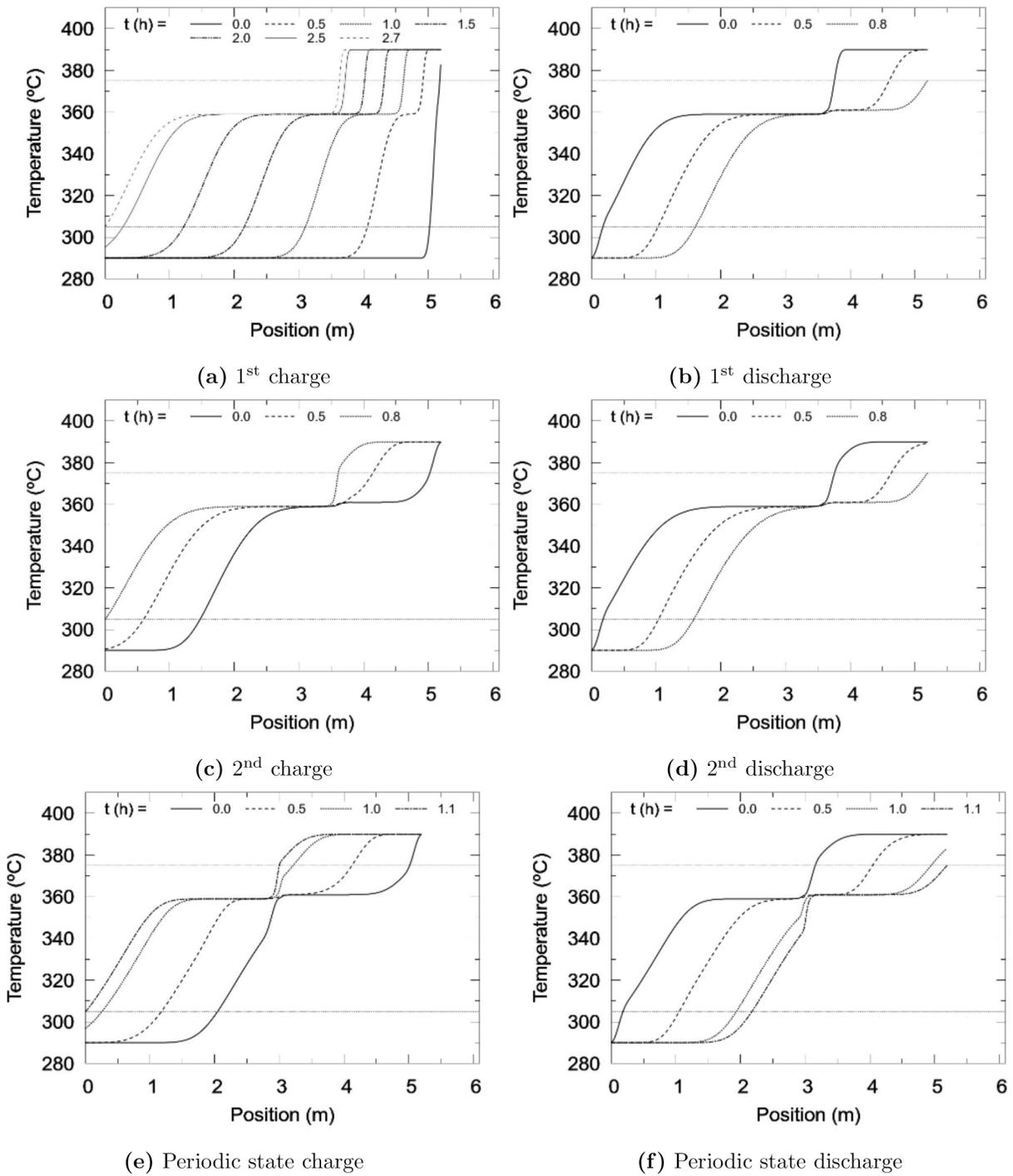
325 Once the periodic state is reached, symmetry between processes can be observed; and therefore,  
326 the energy which is stored in the charge is later delivered in the discharge. This is due to having  
327 disregarded the thermal losses to the ambient.

### 328 3.2.2. Cases B1-3: Encapsulated PCM

329 A first look into the storage capacity values of Table 3 can induce the reader to think that a storage  
330 tank filled with an encapsulated PCM should store more energy than the same tank filled with a solid  
331 filler material. However, results shown in Table 4 reveal a different situation.

332 Prototype of case B1 is filled with a single encapsulated PCM with a fusion temperature of 360°C  
333 (KOH), which is well between the operating temperatures of the storage system.

334 The temperature maps of the HTF inside the tank at various instants for the first two and periodic  
335 state cycles, are plotted in figure 7. A clear picture of the problem results from their observation. In  
336 the first charge, with an initially cold tank, the outlet temperature threshold is reached in a moment  
337 where only part of the contained PCM capsules have melted. The position inside the tank where  
338 the phase-changing capsules are located, can be identified by the location where there is a steep  
339 temperature gradient ranging from 390°C to 360°C, which for the final state is located at a distance of  
340 around 3.6 m from the bottom. Therefore, only a portion of the available latent heat has been used.  
341 Moreover, the sensible energy capacity of both the PCM and HTF is much less harnessed than in case  
342 A. Thus, the initial condition for the subsequent discharge is one where only a part of both the latent  
343 and sensible energy capacity can be exploited.



**Figure 7:** Case B1. Temperature maps at various instants for the first two and last cycles. The chronological order of the curves is from left to right in the charge and from right to left in the discharge.

344 An important observation is that the melting point of KOH lies outside the admissible temperature  
 345 ranges for both charging and discharging processes. In the figure it can be observed how the tem-  
 346 perature of the HTF passing through the phase-changing PCM capsules is kept close to their melting  
 347 point. Therefore, in the charging phase, the filler material located downstream of the phase-changing

348 capsules receive the HTF with a temperature equal to the melting point, not being able to melt. Since  
349 the threshold temperature is lower than the melting point, the charging process ends before all the  
350 PCM has melted, when the capsules located at the outlet (top) cannot bring the HTF temperature  
351 below the threshold, with the use of their sensible energy capacity alone.

352 At the periodic state, the area between the initial and last temperature curves is somewhat higher  
353 than those corresponding to the initial cycles (except from the 1<sup>st</sup> charge). Furthermore, the location  
354 of the phase-changing layers has been shifted slightly to the center of the tank.

355 Cases B2 and B3 have the common feature of using PCMs whose melting points lie inside each of  
356 the admissible temperature ranges. In B2 a melting point of 380°C has been chosen, which is inside  
357 the admissible range for the outlet fluid temperature of the discharging process (375°C- 390°C); while  
358 a melting point of 300°C has been chosen for case B3, lying inside the admissible range of the charging  
359 process (290°C- 305°C).

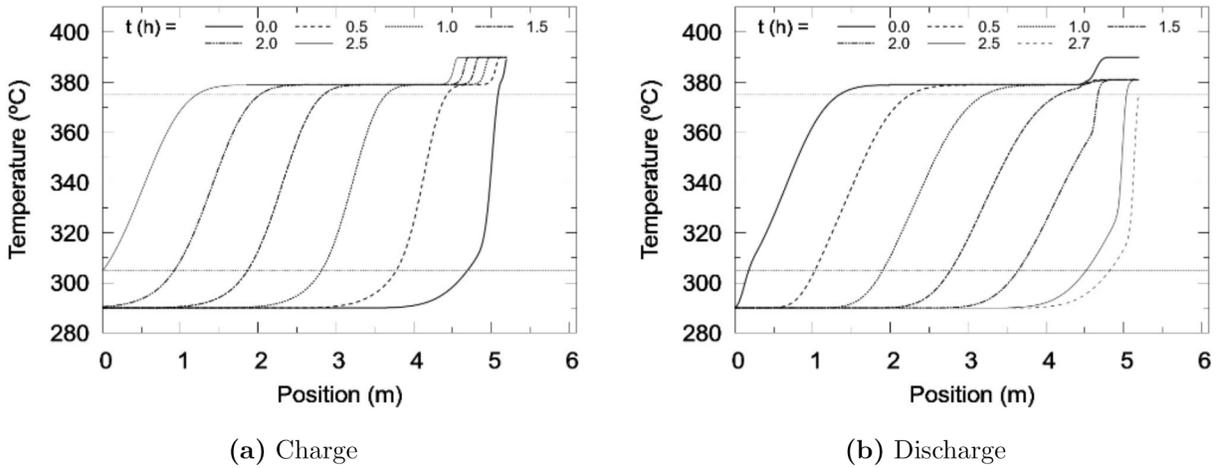
360 Temperature maps for the periodic state of both cases are shown in figures 8 and 9. A first  
361 observation is that the area between the initial and final maps for both cases is higher than that of  
362 case B1. As a result, a higher energy is stored in both cases, as can be observed in Table 4.

363 In these cases, due to the more “intelligent” choice of the melting points, a higher utilization of the  
364 whole storage capacity has been attained. For example, in case B2, as the cold fluid comes through  
365 the bottom of the tank and the thermal gradient travels to the top, the layers of PCM located at  
366 the hot zone, which have melted in the previous charging, act as thermal “buffers” for the outgoing  
367 fluid, keeping its temperature close to the melting point until almost all the PCM near the outlet has  
368 solidified. Since this temperature is inside the admissible range, the process does not stop, and the  
369 rest of the upstream filler material can be thermally discharged.

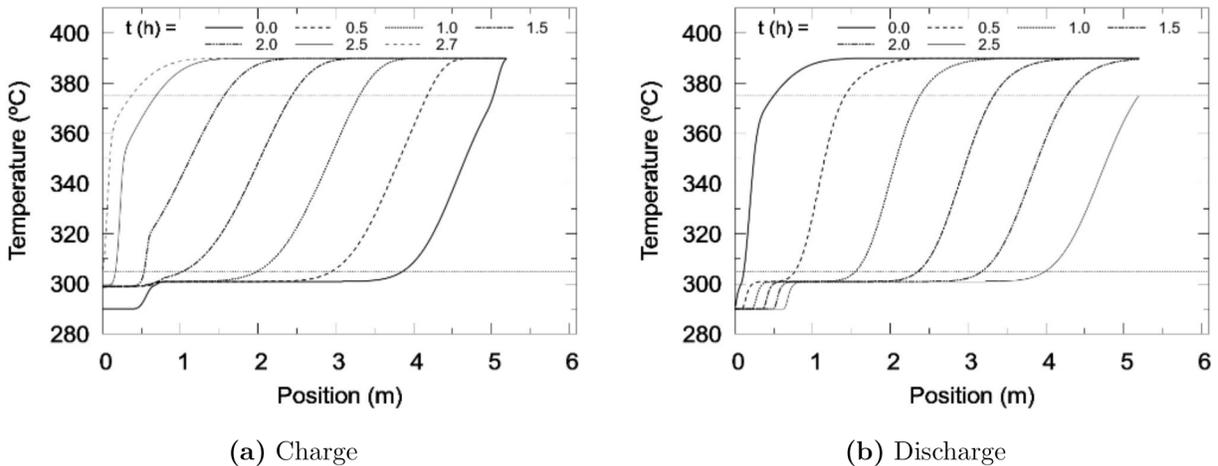
370 Due to the symmetry between key temperatures of cases B2 and B3 (melting points, thresholds and  
371 operating range), the resulting temperature maps for the periodic state are also symmetric. Charging  
372 process of case B2 looks the same as the discharging process of B3, with a shift of the sense in which  
373 both axis increase. Furthermore, the results of both cases shown in Table 4 are almost identical.

374 However, a slight difference is encountered in the value of exergy delivered in the discharging phase.  
375 Result for case B3, with a PCM melting point of 300°C is slightly higher than that of case B2, where  
376 a melting point of 380°C has been adopted. This result may seem strange, since one could think that  
377 a higher melting point of the PCM should result in a higher amount of hot fluid coming out of the  
378 tank and a higher amount of exergy delivered in the discharge. The explanation for this result can  
379 also be extracted from the temperature maps shown in figures 8 and 9. In case B2, the temperature  
380 of the outgoing fluid in the discharge is seen to be the PCM melting point during most of the process

381 (Fig. 8b); while in case B3, it is observed that during most of the discharging phase, the outlet fluid  
 382 temperature is the maximum possible (Fig. 9b). The reason for this is that the low melting point  
 383 PCM, acting as a thermal buffer during the charging phase, allows the upstream filler material to be  
 384 charged of sensible energy up to the maximum temperature, while the high melting point PCM of case  
 385 B3 is storing most of the high temperature energy (between 380°C and 390°C) in the form of latent  
 386 heat at 380°C. On the other hand, case B2 also stores less exergy than case B3, since it needs less  
 387 time to be charged (see figures 8a and 9a), and its final exergy efficiency (delivered/stored) is slightly  
 388 higher in the former.



**Figure 8:** Case B2. Periodic state. Temperature maps at various instants. The chronological order of the curves is from right to left for the charge process and from left to right for the discharge.



**Figure 9:** Case B3. Periodic state. Temperature maps at various instants. The chronological order of the curves is from right to left for the charge process and from left to right for the discharge.

389 *3.2.3. Cases C1-3, D1-2: Multi-layered solid-PCM (MLSPCM)*

390 Results obtained with cases B1 - B3 show that even in the best case, only a little amount of PCM  
391 is effectively changing phase in a cycle (less than 15%). Moreover, the results of cases B2 and B3  
392 show that an effective way of increasing the stored energy of an encapsulated PCM tank is to choose a  
393 PCM whose fusion temperature lies between an admissible temperature interval for either one of the  
394 processes (charge or discharge). Thus, the PCM capsules located at the end of the tank, where the  
395 HTF temperature is close to its melting point, act as a thermal buffer maintaining a desirable outlet  
396 temperature, while the rest of the tank is charged (or discharged) with sensible energy.

397 Therefore, a tank which is filled with PCM in such a way that most of it can effectively undergo  
398 through the phase change, together with the inclusion of a cheaper solid filler material to store the  
399 sensible energy, should be a much more efficient and cost-effective thermal storage device.

400 Hence, all the configurations studied in this section contain PCM layers at both extremes, one  
401 with a high melting point placed at the top of the tank (hot zone) and another with a low melting  
402 point placed at the bottom (cold zone), together with solid filler material placed in between.

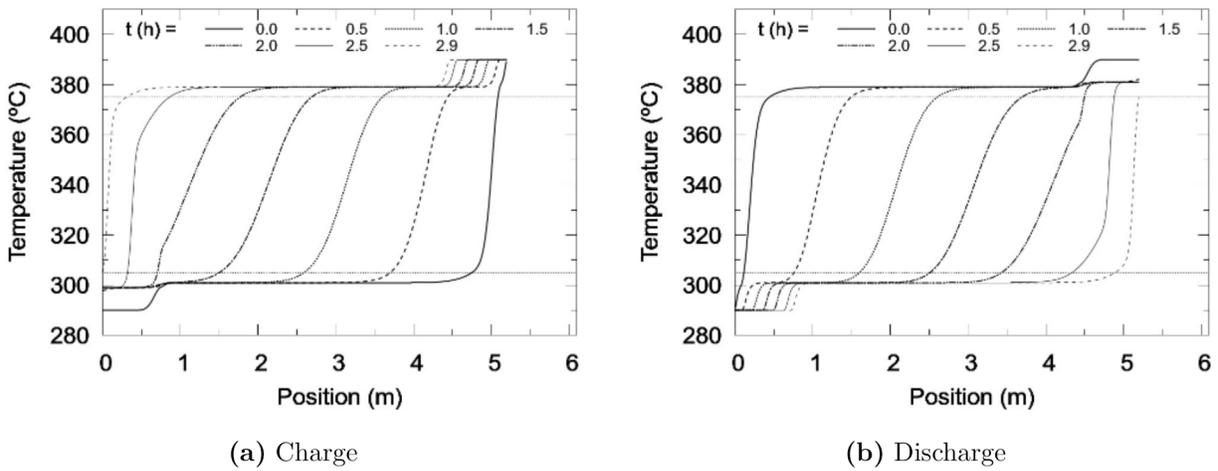
403 The symmetry of the proposed problem, given the operation conditions, induces to design the  
404 multi-layered prototypes using symmetric configurations, i.e. using the same width for the PCM  
405 layers whose melting point are at the same distance (in temperature units) from the corresponding  
406 outlet temperatures.

407 Cases C1 (see sketch on Fig. 5a), C2 and C4 are MLSPCM configurations with only two different  
408 PCMs collocated at both extremes of the tank and a solid filler material (quartzite rocks & sand) in the  
409 middle zone, forming a 3-layer arrangement, only differing in the width of the layers. PCMs used are  
410 those of cases B2 (KOH380) and B3 (KOH300), having melting points lying inside the corresponding  
411 admissible range.

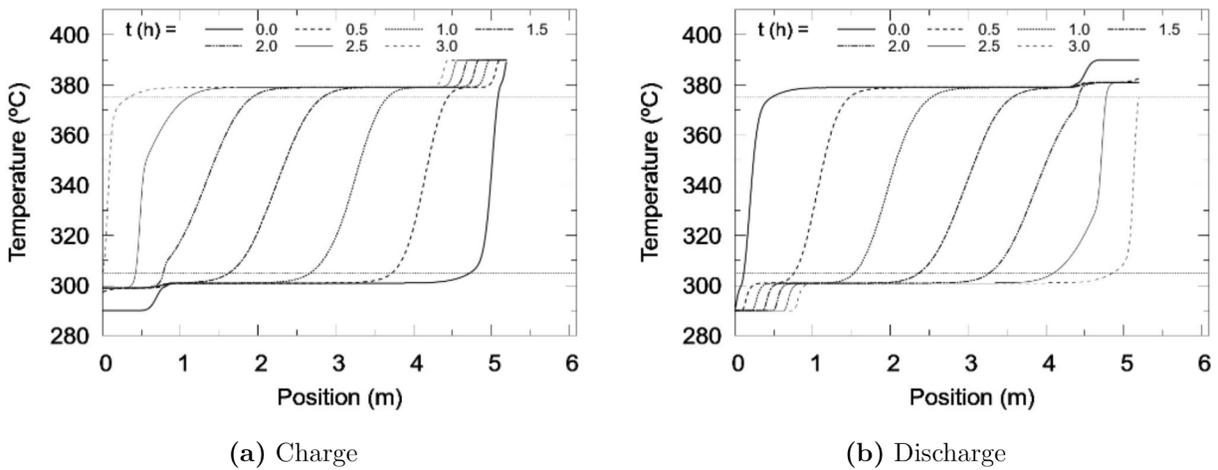
412 Performance results of the three cases, in Table 4, show a significant improvement with respect  
413 to cases A and B1-3. The “buffering” effect of the PCM at both ends can be appreciated in figures  
414 10 to 12. An increase in the amount of stored/released energy in the periodic state is observed. The  
415 efficiency in the usage of both the total thermal capacity and the latent energy capacity are also higher.

416 C2 stores the highest amount of energy of the three, but is the one with the lowest efficiencies in  
417 terms of utilization of both total and latent storage capacities. In fact, C4, with the least amount of  
418 PCM, is the best in terms of efficiency. Around 93% of the PCM is effectively changing phase between  
419 successive processes in the latter case, while 74% is the corresponding value for C1 and 38% for C2.  
420 In terms of total storage, C1 stores around 4% less than C2, while C4 is around 8% worse than C2.

421 Regarding the values of fraction of energy stored in the form of latent heat, it can be observed

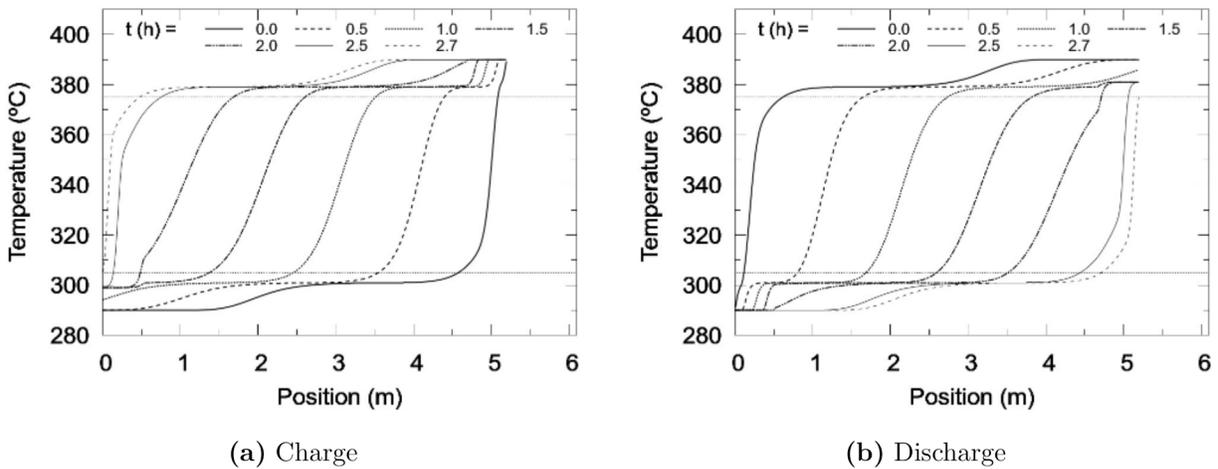


**Figure 10:** Case C1. Periodic state. Temperature maps at various instants. The chronological order of the curves is from right to left for the charge process and from left to right for the discharge.



**Figure 11:** Case C2. Periodic state. Temperature maps at various instants. The chronological order of the curves is from right to left for the charge process and from left to right for the discharge.

422 that for cases C1 and C2 they are almost exactly the same and lower for case C4. This can be  
 423 explained by the following reasoning. The available energy for storage, from the HTF, is in the form  
 424 of sensible energy and thus, it is proportional (with the approximation of having constant specific  
 425 heat) to the temperature difference. Therefore, from the point of view of the high melting point PCM,  
 426 melting at 380°C, the energy contained in the HTF which is available for being stored in the melting  
 427 process, is only that between 380°C and 390°C. This is only 10% of the energy contained in the HTF  
 428 between 290°C and 390°C, and 11.7% of that between 305°C and 390°C (considering both limits of the  
 429 admissible range of outlet HTF temperatures in the charge). This means that, at most, only 10%-12%  
 430 of the energy that needs to be extracted from the HTF can be used for melting this PCM, given the  
 431 constraint for the outgoing temperature of not surpassing 305°C.



**Figure 12:** Case C4. Periodic state. Temperature maps at various instants. The chronological order of the curves is from right to left for the charge process and from left to right for the discharge.

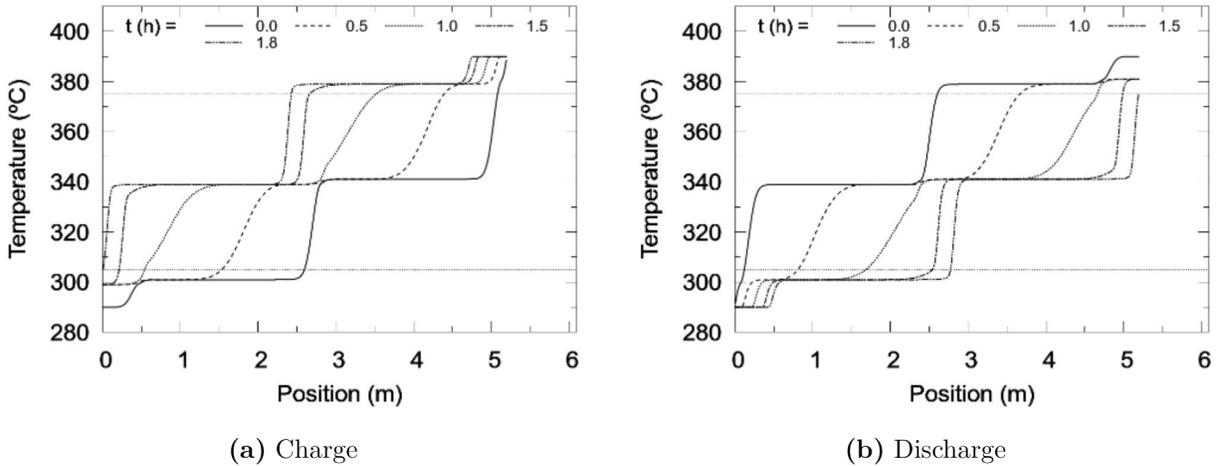
432 It could be argued that, having a second PCM placed at the outlet with a much lower melting  
 433 point (300°C), much more energy could be stored in the form of latent heat, since the energy contained  
 434 in a much broader range of temperatures is available for melting this other PCM (in the case of total  
 435 utilization of the first 10% for melting the first PCM, this range would be 300°C–380°C; or less, if the  
 436 sensible energy stored in the layers upstream from the second PCM is considered). However, this is  
 437 not the case when the periodic state has been reached. In this state, and in the absence of thermal  
 438 losses, the energy stored in the charge is the same as that delivered in the subsequent discharge.  
 439 Furthermore, the same amount of PCM that melts in the charge, is solidified in the next discharge.  
 440 For this reason, if 10% of the total available energy is used to melt the high melting point PCM, then  
 441 only this same amount will be delivered by it in the subsequent discharge. Given the symmetry of the  
 442 problem, the same reasoning can be applied to the low melting point PCM.

443 Therefore, as each PCM layer can only theoretically store around 10% - 12% of the total available  
 444 energy, both PCMs can sum up to 20% - 24% of it, at most. Case C1 and C2 both result in a latent  
 445 storage of 20%. C4, having significantly less amount of PCM, only reach to around 13%. Table 3  
 446 shows that the latent capacity of prototype C1, C2 and C4 is around 21%, 34% and 12%, respectively.  
 447 This is the reason why in case C2 only a small fraction of the PCM effectively changed phase, having  
 448 a higher latent heat capacity fraction than the theoretically possible.

449 Cases D1 (see sketch in Fig. 5b) and D2 consist of 5-layered MLSPCM tanks. Both PCM layers  
 450 at the ends are maintained, while an extra PCM layer is added in the middle zone, with a melting  
 451 point equal to the mean temperature of the operation range, i.e. 340°C. Other two solid filler material  
 452 layers are placed between the PCM layers.

453 Case D1 is seen to behave worse than the previous C1-3 cases. Figure 13 shows the temperature  
 454 maps obtained for this case, with a similar behavior to that of B1. The presence of KOH340 acts  
 455 as a thermal buffer keeping the HTF temperature close to its melting point, which is outside both  
 456 admissible temperature ranges. The thermal buffering provided by the other two PCMs is not enough  
 457 to allow it to melt and solidify completely before the end of the charge and discharge processes, causing  
 458 the melting point of 340°C to be a limit to the temperature jump of the solid filler layers and also of  
 459 much of the middle PCM layer. As a result, the stored energy and the efficiencies are much worse  
 460 than those of the previous MLSPCM prototypes.

461 On the other hand, case D2, with the only difference of containing a middle layer of half the width  
 462 compared to D1, results in a significantly different behavior. Figure 14 shows the temperature maps  
 463 obtained for this case, where a behavior similar to that of cases C1 and C2 can be observed. Here, the  
 464 thermal buffering effect of the middle layer does not last so long and the top and bottom PCM layers  
 465 are capable of bearing the “extra” exigency. In fact, the inclusion of the middle layer has resulted in  
 466 an overall increase in the stored energy with almost the same efficiency, compared to case C1, which  
 467 had a lower amount of PCM effectively changing phase.

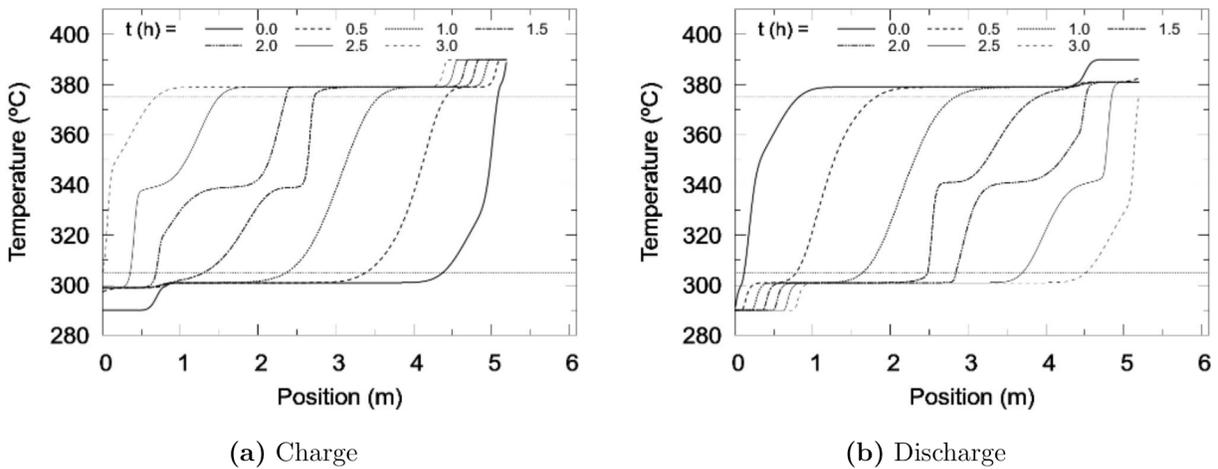


**Figure 13:** Case D1. Periodic state. Temperature maps at various instants. The chronological order of the curves is from right to left for the charge process and from left to right for the discharge.

### 468 3.2.4. Cases F1-2: Cascaded PCMs

469 In the cascaded PCM configurations considered here, as in MLSPCM cases, PCM layers located  
 470 at both ends with melting points lying inside the admissible ranges are included.

471 The difference between these prototypes and MLSPCM ones, is that no solid filler material is  
 472 included in the formers. Instead, several layers of different PCMs are placed inside the tank, with  
 473 increasing melting points from the bottom to the top. This kind of configuration has been studied by



**Figure 14:** Case D2. Periodic state. Temperature maps at various instants. The chronological order of the curves is from right to left for the charge process and from left to right for the discharge.

474 other authors, e.g. Michels and Pitz-Paal, [13]. A sketch of configuration F1 is depicted in figure 5c.

475 The difference between F1 and F2 is in the width of the three middle layers, where the PCMs with  
 476 melting points outside the admissible ranges are located. Case F1 includes more of the PCMs with  
 477 melting points more near the admissible limits (KOH370 and KOH310), while F2 includes more of the  
 478 middle PCM with a temperature equal to the operating temperatures mean value (KOH340).

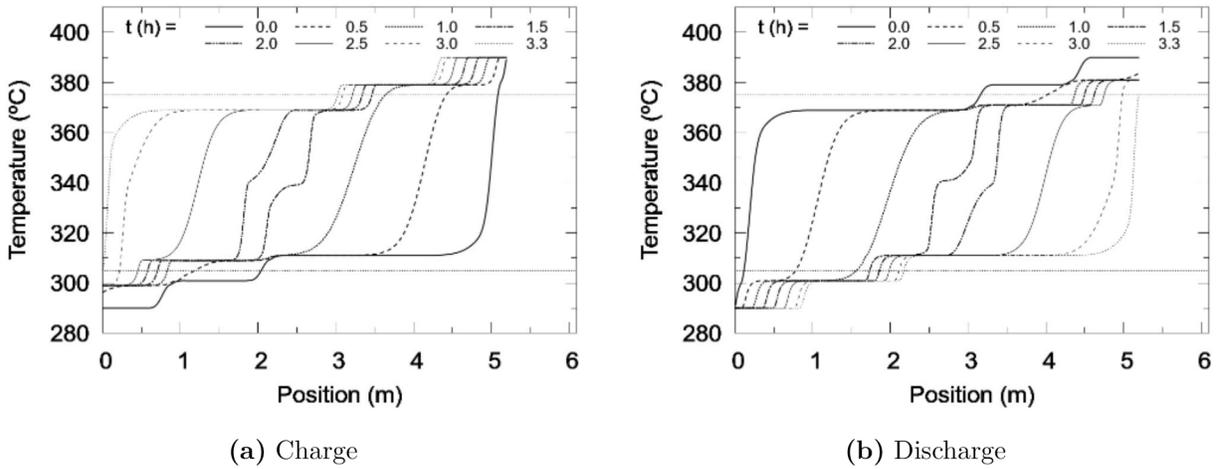
479 Results of Table 4 show that the performance of case F1 is significantly better than that of case  
 480 F2. As both cases use the same PCM layers, the difference between them is only due to the different  
 481 proportion of materials included. Temperature maps of both cases at the periodic state can be observed  
 482 in figures 15 and 16. Significant differences are encountered. In case F1, the temperature range  
 483 traversed by the fluid inside the tank is sensibly higher than in case F2. In case F1, when the  
 484 PCM at the ends have almost completely changed of phase —maintaining outlet fluid temperature  
 485 inside the admissible range— a considerable amount of the PCM layers inside the tank have also  
 486 melted/solidified. This is not what happens in case F2, for which the outlet fluid temperature cannot  
 487 stay within the admissible ranges enough time to allow a good utilization of sensible and latent  
 488 capacities. Hence, as it has been observed with MLSPCM cases, a correct design of the PCM layers  
 489 is a critical aspect in the final performance.

490 Case F1 has been the best of all the studied cases in thermal storage capacity. The energy stored  
 491 is around 15%, 10%, 20% and 9% higher than for cases C1, C2, C4 and D2, respectively. However, the  
 492 efficiency in the use of the storage capacity is not very high (65%), with 61% of the PCM effectively  
 493 changing phase.

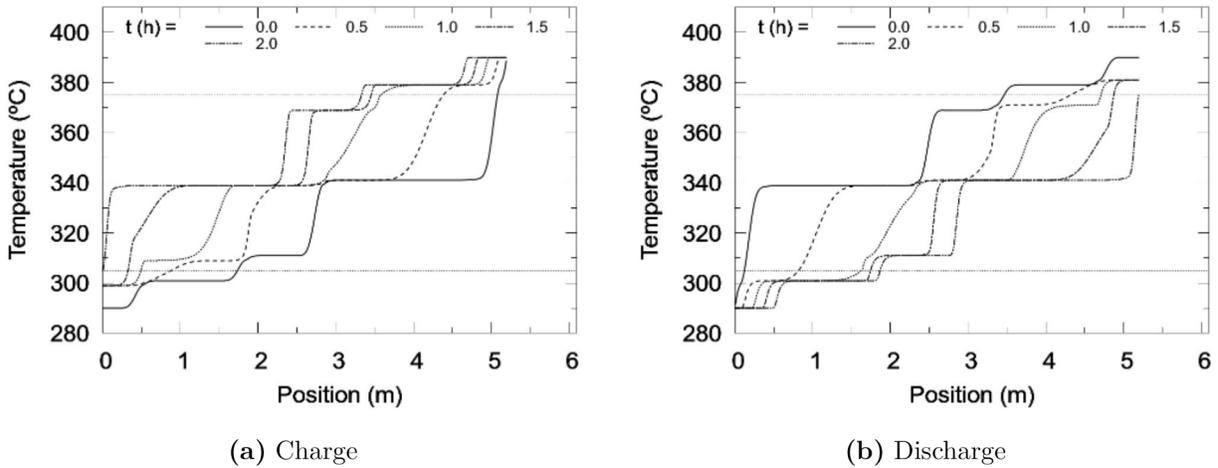
494 Some other cascaded PCM cases with different layer thicknesses have been tested (not presented),

495 with the result of F1 being the best of all. It might be possible to find other arrangements, with the  
 496 same PCM layers, with better performance, although it is the authors belief that this arrangement is  
 497 close to the best possible results.

498 Considering that an encapsulated PCM is probably much more costly than the solid filler material,  
 499 MLSPCM prototypes can be expected to be more cost-effective than cascaded PCM, also considering  
 500 the better efficiency of the former in the use of PCMs latent heat.



**Figure 15:** Case F1. Periodic state. Temperature maps at various instants. The chronological order of the curves is from right to left for the charge process and from left to right for the discharge.



**Figure 16:** Case F2. Periodic state. Temperature maps at various instants. The chronological order of the curves is from right to left for the charge process and from left to right for the discharge.

501 *3.2.5. Pressure losses*

502 Pressure losses due to the presence of the filler material [first term on the right hand side of  
 503 Eq. (6)] are below 400 Pa for all the cases studied. The highest losses are found in the thermocline

504 system filled uniquely with solid material (case A), due to the higher compactness of the solid filler  
505 material compared to those of the encapsulated PCMs. However, when compared against the pressure  
506 differences arising due to gravitational effects ( $\rho gh \sim 1 \times 10^5$  Pa) these represent less than 0.4% in all  
507 the cases, and thus, are negligible.

#### 508 **4. Conclusions**

509 A new multi-layered solid-PCM thermocline-like thermal storage concept for CSP plants has been  
510 presented. The key aspect of this new concept is the inclusion of PCM layers at both ends of the tank,  
511 whose fusion temperatures are conveniently chosen to lie inside the predefined admissible temperature  
512 ranges for the outgoing fluid in both charge and discharge processes. These admissible temperature  
513 ranges depend on the requirements of the power generation block and the solar receivers. The PCM  
514 layers act as thermal buffers, causing the outlet fluid to remain close to their melting points, and  
515 therefore inside the admissible temperature range for the corresponding process.

516 In order to design and evaluate the performance of such storage devices, as well as of the other  
517 thermocline-like systems considered, a numerical model has been developed and implemented. This  
518 model has been successfully validated against experimental data.

519 Several simulations have been carried out for different designs of thermocline tanks, where configu-  
520 rations of solid, single-PCM, multi-layered solid-PCM and cascaded PCM filler material configurations  
521 have been tested. The obtained results show that the multi-layered solid-PCM concept prevents from  
522 the high thermocline degradation presented by the pure thermocline, resulting in a much higher ef-  
523 ficiency in the use of the overall thermal capacity of the system. Furthermore, compared against the  
524 cascaded PCM concept, this new approach has the advantage of using much less encapsulated PCM  
525 for almost the same total stored energy, again with a higher thermal efficiency. For example, prototype  
526 C4 (MLSPCM) stores around 83% of the energy stored with prototype F1 (cascaded PCM), using  
527 only 20% of the mass of PCM and 72% of molten salt; which is a consequence of being more efficient  
528 in the use of the storage capacity (84% vs. 65%, respectively).

529 Therefore, the MLSPCM thermocline storage systems can be considered as a promising solution  
530 for their use in CSP plants.

#### 531 **Acknowledgments**

532 This work has been financially supported by the *Ministerio de Economía y Competitividad, Secre-*  
533 *taría de Estado de Investigación, Desarrollo e Innovación*, Spain (ENE-2011-28699), by the EIT via  
534 the KIC InnoEnergy TESCONSOL project (ref. 20\_2011\_IP16) and by the *Secretaria d'Universitats*

535 *i Recerca (SUR) del Departament d'Economia i Coneixement (ECO) de la Generalitat de Catalunya*  
536 and by the European Social Fund.

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