Numerical Analysis of Water Melting and Solidification in the Interior of Tubes

Latent energy storage systems find applications in many engineering fields, including industrial refrigeration plants, air conditioning installations, recovery of heat in industrial processes, etc. To tackle the design of such systems, it is necessary to have correlations to account for the heat transfer during the melting and solidification of the phase change material (PCM). This work describes and analyzes the results obtained from the numerical simulation of pure water melting and solidification in the interior of tubes, which are typically present in ice banks of air conditioning systems. The shown results consider natural convection, accounting for the inversion in the water density. In the melting process, the considered initial conditions followed the classical Stefan and Neumann approach. The presented simulation results include the evolution of the phase change interface, and of the temperature, density and streamlines fields. Correlations for the Nusselt number and for the melted material volume as functions of time have been proposed.

**Keywords:** Phase change, melting and solidification, ice banks, finite volumes, polar geometry

Introduction

The increasing energy demand and the requirements for an environmentally sustained development have been major motivating factors for the implementation of more efficient means of energy conservation. In addition to conservation, the rational use of energy has been the focus of many studies, so that the processes are more efficient and less expensive, reducing waste, preserving the environment and the nonrenewable energy sources, and avoiding unnecessary expenses. In this sense, the employment of thermal energy storage has the objective of making the energy consume more rational. This storage is performed in three basic, distinct ways: sensible heat, latent heat and chemical. The systems that operate with sensible heat require high temperature differences with respect to the environmental, as well as a storage substance with large mass. This leads to some disadvantages, to mention the need of large spaces to accommodate the system and thick, expensive thermal insulation. The storage systems that use the latent heat present a superior amount of volumetric capacity, besides requiring smaller temperature differences with respect to the environment, which decreases the thermal insulation expenses.

According to the “ASHRAE Applications (1995)”, the use of thermal storage systems may be an economically attractive approach for heating or cooling loads when they are of short time duration and occur cyclically; when there is an incompatibility with the available energy source, or its supply is limited; when the energy costs are time-dependent; and when economical incentives are provided for the use of load-shifting equipment.

In the refrigeration field, ice banks have been employed as an energy storage solution to reduce the electrical power required during the hours of peaks in the demand, shifting the consume curve to a more favorable situation. It must be noted that in many countries, the cost of industrial electricity is time dependent during the day, having higher values in the periods of high demand. It can be also mentioned the contractual demand, resulting into a fine to the user in case the energy consumption is exceeded. Aiming at economical feasibility, it has been increasingly attempted to develop systems that employ the storage of latent heat, despite its disadvantages. Among those, it can be mentioned the inherent irreversibilities due to heat transfer, the decrease in the efficiency of the thermodynamic cycle due to the decrease in the evaporation temperature in the air-conditioning installations, the increase in the physical space that is necessary to accommodate the ice bank, and uncertainties in the evaluation of the heat transfer between the PCM and the system working fluid.

Water in the solid phase has been employed as the storage substance due to its properties and characteristics: high melting latent heat (334 kJ/kg), melting temperature of 0°C at the atmospheric pressure, accessibility, low cost, no pollutant effects, and chemical stability. As the most important disadvantage characteristic, it can be pointed the supercooling phenomenon (a metastable equilibrium) during the solidification, “Chen and Lee (1998)”. The ice bank consists of thermally insulated tanks that are inserted into a refrigeration loop. In some systems, the ice is formed on the outside of the tubes. In such systems, a limit must be impose to the ice thickness formed on the piping, since if it exceeds a given value there will be a decrease in the heat transfer during the discharge process, caused by the ice thickness and the flow obstruction. Examples of works that investigated the phase change occurring in the outside of the tubes are “Ho and Chen (1986)”, “Zhang and Faghri (1996)”, “Abudarah and Ismail (2000)”, “Stampa et al. (2002)”.

Encapsulating the ice in tubes is one of the most employed examples of latent heat storage, consisting of a tank with a bank of horizontal tubes, across which flows a fluid that works as a thermal device that promotes the charge or discharge of the system. Among the works that deal with the process of melting and solidification in the interior of tubes, it can be pointed “Rieger et al. (1983)”, “Ho and Viskanta (1984)”, and “Rieger and Beer (1986)”. The latter is one of the first experimental and numerical studies on phase change that employ water as the PCM. Another investigation worth mentioning is that of “Cao et al. (1991)”. The encapsulations in spherical, rectangular and spiral configurations are other possibilities.
Nomenclature

\(A = \text{area, m}^2\)
\(B = \text{body force, m/s}^2\)
\(c = \text{specific heat, J/kg}\)
\(E = \text{coefficient of the proposed function for the Kirchhoff temperature, coefficient of the Nusselt number correlation}\)
\(Fo = \text{Fourier number, } \alpha t/\varepsilon^2\)
\(g = \text{gravitational field, m/s}^2\)
\(h = \text{enthalpy, J/kg; convection heat transfer coefficient, W/(m}^2\text{K)}\)
\(k = \text{thermal conductivity, W/(mK)}\)
\(L = \text{representative grid size}\)
\(L_0 = \text{average thickness of the liquid ring, m}\)
\(m = \text{mass, kg}\)
\(Nu = \text{Nusselt number, } r_0 \ln(r/r_0)\partial T_{\text{adm}}/\partial r|_{r_0}\)
\(p = \text{pressure, N/m}^2\)
\(q = \text{heat transfer rate, W}\)
\(r = \text{radial coordinate, mesh relation, radius, m}\)
\(Rac = \text{Rayleigh number, } \alpha \eta g |T_{\text{adm}} - T_0| \sqrt{\varepsilon} / \nu v\)
\(Ste = \text{Stefan number, } c_p(T_w - T_0)/\lambda\)
\(T = \text{temperature, } ^\circ\text{C, K}\)
\(t = \text{time, s}\)
\(u = \text{tangential velocity, m/s}\)
\(V = \text{volume, m}^3\)
\(v = \text{radial velocity, m/s}\)

Greek Symbols

\(\alpha = \text{thermal diffusivity, m}^2/\text{s}\)
\(\beta = \text{thermal expansion coefficient, } 1/\text{K}\)
\(\Phi = \text{generic variable}\)
\(\eta = 9.297173 \times 10^{-6}, \circ\text{C}^{-\text{b}}\)
\(\lambda = \text{melting and solidification latent heat, J/kg}\)
\(\mu = \text{dynamic viscosity, Ns/m}^2\)
\(\nu = \text{kinematic viscosity, m}^2/\text{s}\)
\(\theta = \text{angular coordinate, rad}\)
\(\rho = \text{density, kg/m}^3\)

Subscripts

\(c = \text{relative to convection}\)
\(\text{conv} = \text{relative to convective process}\)
\(f = \text{relative to melting}\)
\(H = \text{relative to hydrostatic component}\)
\(\text{ini} = \text{relative to initial}\)
\(L = \text{relative to liquid}\)
\(\text{max} = \text{relative to maximum}\)
\(\text{mist} = \text{relative to mixture}\)
\(p = \text{relative to constant pressure}\)
\(r = \text{relative to radial direction}\)
\(\text{ref} = \text{relative to reference}\)
\(s = \text{relative to solid}\)
\(S = \text{relative to solid}\)
\(w = \text{relative to wall}\)
\(\text{wall} = \text{relative to tube wall or to initial}\)
\(\theta = \text{relative to angular direction, angular position}\)

Superscripts

\(* = \text{melting point}\)
\(b = 1.894816, \text{constant to } "\text{Eq. 31}"\)

Physical Model and Boundary Conditions

The focus of this work is a latent “cold storage” modulus, which consists of a bank of tubes containing pure water inside, subjected to a crossed external flow. In the present approach only one tube is considered through a symmetric, two-dimensional model, since such tubes are typically subjected to small temperature variations along its length.

The boundary conditions are now established in order to facilitate the physical model comprehension. For the thermal problem the first-kind boundary conditions are adopted, “Eq (1)”, both for the melting and solidification, as seen in “Fig.1”.

\[ T(r_o, \theta) = T_w \rightarrow t > 0 \]  

where \(u\) and \(v\) are, respectively, the angular and radial velocities, and \(r_o\) and \(r_s\) are the radius of the tube and of the PCM, as seen in “Fig. 1.” The initial condition adopted for the thermal problem during the melting is dependent on the approach that is applied. In the so called Stefan problem,

\[ \Phi(r, \theta) = \phi(r, \theta) \rightarrow t = 0 \]  

while in the Neumann problem,

\[ \Phi(r, \theta) = \phi_r, \phi_s \rightarrow t = 0 \]  

In the solidification the initial boundary condition, for the thermal problem, is:

\[ T(r, \theta) = T_{\text{in},L} \rightarrow t = 0 \]  

where \(T(r, \theta)\) is the temperature inside the tube, \(T_{\text{in}}\) is the initial temperature and the subscripts \(S\) and \(L\) stand for the solid and liquid regions, \(t\) is time, \(r\) and \(\theta\) are respectively the radial and angular coordinates.

For the hydrodynamic problem, the initial conditions for the melting and solidification are:

\[ u = v = 0 \rightarrow t = 0 \]  

The non-slip and impermeability boundary conditions are adopted during all the situations.
Governing Equations

The heat transfer from or to the encapsulated fluid originates the thermal gradients, which in turn cause the existence of gradients in the density in the interior of the fluid domain. Therefore, thermal convection may be present due to the fluid flow caused by buoyancy. For the solution of such a problem, it is necessary the knowledge of the velocity, temperature, density and pressure fields, which characterizes the coupling of the hydrodynamic and thermal problems. The equations that govern the phenomenon for the considered geometry are present below in radial coordinates. The continuity equation is given by:

\[
\frac{\partial \rho}{\partial t} + \frac{1}{r} \frac{\partial (\rho v)}{\partial r} + \frac{1}{r} \frac{\partial (\rho u)}{\partial \theta} = 0 \quad (7)
\]

where \( \rho \) is the density and \( t \) is de time. The momentum equations in the \( r \) and \( \theta \) directions are:

\[
\frac{\partial (\rho v)}{\partial t} + \frac{1}{r} \frac{\partial (\rho v^2)}{\partial r} + \frac{1}{r} \frac{\partial (\rho uv)}{\partial \theta} + \frac{\rho v^2}{r} = \frac{\partial p}{\partial r} + \frac{\mu}{r^2} \left( \frac{\partial v}{\partial r} + \frac{v}{r^2} \frac{\partial \theta}{\partial r} \right) + \mu \frac{\partial}{\partial \theta} \left( \frac{\partial v}{\partial \theta} + \frac{v}{r^2} \frac{\partial \theta}{\partial \theta} \right) + \rho B_r \quad (8)
\]

\[
\frac{\partial (\rho u)}{\partial t} + \frac{1}{r} \frac{\partial (\rho u^2)}{\partial r} + \frac{1}{r} \frac{\partial (\rho uv)}{\partial \theta} + \frac{\rho u^2}{r} = \frac{\partial p}{\partial \theta} + \frac{\mu}{r^2} \left( \frac{\partial u}{\partial r} + \frac{u}{r^2} \frac{\partial \theta}{\partial r} \right) + \mu \frac{\partial}{\partial \theta} \left( \frac{\partial u}{\partial \theta} + \frac{u}{r^2} \frac{\partial \theta}{\partial \theta} \right) + \rho B_\theta \quad (9)
\]

in which \( \rho \) is the pressure, and \( B_r \) and \( B_\theta \) are the radial and angular components of the gravitational field acceleration \( g \). The dynamic viscosity is represented by \( \mu \). The Boussinesq approximation is adopted for the buoyancy terms in the liquid phase. The references for the temperature and density are taken from the melting point: \( T_{ref} \) and \( \rho_{ref} \). From the Maxwell thermodynamic relations, one can set the state equation that relates the density to temperature and pressure for a given substance:

\[
\rho = \rho(T, p) \quad (10)
\]

For small variation in the density, it is acceptable to expand the above function in the vicinity of a point of reference by means of the Taylor series to give:

\[
\rho \approx \rho_{ref} + \left( \frac{\partial \rho}{\partial T} \right)_{p} (T - T_{ref}) + \left( \frac{\partial \rho}{\partial p} \right)_{T} (p - p_{ref}) + \ldots \quad (11)
\]

which is valid for liquids. Using now the definition of the thermal expansion coefficient \( \beta \), “Eq. (11)” becomes:

\[
\rho \approx \rho_{ref} \left[ 1 + \beta (T_{ref} - T) \right] \quad (12)
\]

inserting “Eq. (12)” into “Eq. (8)” and “Eq. (9)” results:

\[
\frac{\partial (\rho v)}{\partial t} + \frac{1}{r} \frac{\partial (\rho v^2)}{\partial r} + \frac{1}{r} \frac{\partial (\rho uv)}{\partial \theta} + \frac{\rho v^2}{r} = \frac{\partial p}{\partial r} + \frac{\mu}{r^2} \left( \frac{\partial v}{\partial r} + \frac{v}{r^2} \frac{\partial \theta}{\partial r} \right) + \mu \frac{\partial}{\partial \theta} \left( \frac{\partial v}{\partial \theta} + \frac{v}{r^2} \frac{\partial \theta}{\partial \theta} \right) + \rho g \beta_\theta (T - T_{ref}) \quad (13)
\]

\[
\frac{\partial (\rho u)}{\partial t} + \frac{1}{r} \frac{\partial (\rho u^2)}{\partial r} + \frac{1}{r} \frac{\partial (\rho uv)}{\partial \theta} + \frac{\rho u^2}{r} = \frac{\partial p}{\partial \theta} + \frac{\mu}{r^2} \left( \frac{\partial u}{\partial r} + \frac{u}{r^2} \frac{\partial \theta}{\partial r} \right) + \mu \frac{\partial}{\partial \theta} \left( \frac{\partial u}{\partial \theta} + \frac{u}{r^2} \frac{\partial \theta}{\partial \theta} \right) + \rho g \beta_r (T - T_{ref}) \quad (14)
\]

where \( p_{hy} \) is the pressure that incorporates the hydrostatic component. Since the density of the water is not a linear function of the temperature, “Fig. 2”. The behavior of the volumetric expansion coefficient, with respect to the temperature, was based on the expression of “Vasseur et al. (1983)”, for the water density, resulting in the following expression:

\[
\beta = -1.9364 \times 10^{-3} T^2 + 1.7507012 \times 10^{-5} T - 6.59759847 \times 10^{-8} \quad (15)
\]

![Figure 2. Behavior of the water density with respect to the temperature.](Image)

In problems of phase change, the domain can be divided into three distinct regions: solid, liquid and interface, where each region has its own characteristics. For the pure substances, the temperature is constant in the liquid-solid interface, and so it does not provide any information on the evolution of the process. To overcome this difficulty, it is considered the energy equation in the enthalpy form. Applying the model proposed by “Cao et al. (1989)”, in which it is employed the concept of the Kirchhoff Temperature, described and applied in “Vielmo (1993)”, results:

\[
\frac{\partial (\rho h)}{\partial t} + \frac{1}{r} \frac{\partial (\rho vh)}{\partial r} + \frac{1}{r} \frac{\partial (\rho uh)}{\partial \theta} + \frac{1}{r} \frac{\partial (\rho \beta)}{\partial T} \left( \frac{\partial (\rho h)}{\partial T} \right) + \rho^* \frac{g}{\beta} \beta \theta (T - T_{ref}) \quad (16)
\]

where \( h \) is the specific enthalpy. The remaining terms are described below:

\[
E(h) = k \left( \frac{c_p}{L} \right) \quad S(h) = 0 \quad \rightarrow h \leq 0 \quad \rightarrow \text{(Solid phase)}
\]

\[
E(h) = 0 \quad S(h) = 0 \quad \rightarrow 0 < h < \lambda \quad \rightarrow \text{(Interface)}
\]

\[
E(h) = k \left( \frac{c_p}{L} \right) \quad S(h) = \frac{\lambda}{c_p} \quad \rightarrow h \geq \lambda \quad \rightarrow \text{(Liquid phase)}
\]

in which \( c_p \) is the specific heat, \( \lambda \) is the latent heat, \( k \) is the thermal conductivity.
Numerical Methodology

For the integration of the governing equations, the finite volumes method is applied, as described in “Patankar (1980)” and “Maliska (2004)”. In the equation, the diffusion coefficient depends whether the considered region is the liquid, interface or solid. The system of equations resulting from the integration of the differential equations is solved by the TDMA scheme, with a block correction. The adopted interpolation function is the Power-Law, and the coupling between the pressure and velocity fields is based on the SIMPLEC algorithm. A computational mesh formed by 40x40 volumes was employed after grid independence tests were performed. The velocities are stored in the interfaces of the control volumes. In most of the solutions, relaxation factor of 0.8 for all the variables proved to be adequate to deal with the non-linearities of the problem. Values in the order of $10^{-8}$ were used for convergence criteria, applied to relative error of each variable between two consecutive iterations, as well as to the residual masses in each control volume. In order to obtain stable, implicit transient solutions, the time steps were varied and chosen so that the enthalpy, in the liquid domain, changed about 200 J/kg each time step (no control volumes submitted to the phase change). When there were control volumes close to the phase change region (the biggest period of time), the time steps were reduced to values that caused variation in the enthalpy in the range of approximately 10 J/kg in the liquid.

Validation of the Proposed Methodology

For the validation, the results obtained in this work were compared to the numerical and experimental results presented in “Rieger and Beer (1986)”, “Fig. 3”. The solid is kept fixed in the center of the tube. Figure 4 presents the solution obtained in this work for the same situation considered in the aforementioned work. As seen, there exists an excellent agreement between the two solutions.

For verification of the independence of the solutions with respect to the discretization, it was applied the Grid Convergence Index (GCI), according to “Roache (1998)”, and the usual procedure for estimation and reporting of discretization error in CFD applications. The melting was simulated using the Stefan approach for the initial condition $T_{ini,s} = 0 \, ^\circ C$, for the boundary condition $T_w = 10 \, ^\circ C$ and for a tube with a diameter $r_0 = 0.032$ m. The following equations were initially applied to evaluate the apparent order $p$ of the employed method:

$$ L = \left[ \frac{1}{N} \sum_{i} (\Delta A_i) \right]^{1/2} \quad (18) $$

$$ r = \frac{L_{ coarse}}{L_{ fine}} \quad (19) $$

$$ p = \frac{\ln(f_3 - f_1)}{\ln(r)} \quad (20) $$

where $L$ is the representative grid size, the $\Delta A_i$ is the area of the $i^{th}$ volume, and $N$ is the total number of volumes used. The $f$ terms are the solutions, and the subscripts 1, 2 and 3 refer from the more refined to the less refined mesh. The three different meshes tested have 40×40, 120×120 and 360×360 control volumes. Such an approach was necessary to allow geometrical agreement between the evaluation points. The temporal agreement was also observed. The enthalpy was the variable used for the inspection of the apparent order $p$ of the method, since this property is the key variable in the problem. The obtained results are presented in “Tab. 1”.

Such evaluations were carried out for three angular positions in the tube, and in each position three locations were chosen along the radius. The angles of the positions are referred to with respect to the “Fig. 1”. These positions were chosen in such a way to cover the upper, intermediary and lower regions of the tube.

For the evaluation of the relative error, $\varepsilon$, and of the GCI, “Eq’s. (21) and (22)” were employed considering the apparent order $p$ of local approximation, “Eq (20)”:

$$ \varepsilon = \frac{|f_2 - f_1|}{f_1} \quad (21) $$

$$ GCI = 1.25 - \frac{\varepsilon}{r^p} \quad (22) $$
As seen in the “Tab. (1)”, the maximum relative error, ε, reached values lesser than 1 % in the comparison of the results obtained for the meshes 40x40 and 360x360. The CGI presents a maximum error of approximately 1.5 % for the meshes 40x40 and 120x120. One can state, based on the results presented in “Tab. (1)”, that the employed grid of 40x40 control volumes proved to be adequate for the analysis presented in this work. Also it is possible to observe the appearance of a negative value for the apparent order p, indicating an oscillatory convergence in that position (close to the center of the tube). In this position low values for the relative errors also occur. In rare situations these variations, when close to zero, are indicative that the exact solution was reached. Anyway, it does not represent unsatisfactory calculations, or parameter to invalidate the results.

In all the obtained solution, for the different meshes, it was not registered any variation in the pattern of the variables fields. The verification of the presence or not of multiple solutions was investigated through alterations in the directions of the TDMA scheme. As observed, changing the directions in all directions did not cause any modification in the solutions.

Results and Discussion

The Nusselt Number

It is first necessary to define an expression for the Nusselt number in the melting of the PCM encapsulated in the interior of the tube, and for that matter it was adopted the enclosure treatment. For the geometry considered in this work, the solution of the steady-state, one-dimensional diffusion equation, with no source term, provides the below temperature profile:

\[
T(r) = \frac{T^* - T_w}{\ln \left( \frac{r}{r_0} \right)} + T_w
\]

The heat transfer by diffusion, \( q_r \), in a given position \( r \) is given by:

\[
q_r = -k A(r) \frac{dT}{dr} \left[ \frac{T^* - T_w}{\ln \left( \frac{r}{r_0} \right)} + T_w \right] = -\frac{2\pi k}{\ln \left( \frac{r}{r_0} \right)} \left( T^* - T_w \right)
\]

The convective heat transfer \( q_{\text{conv}} \) is written using the general formulation below:

\[
q_{\text{conv}} = -A(r) h \left( T^* - T_w \right)
\]

where \( h \) is the heat transfer coefficient. In a situation where the PCM is partially liquid and solid, but before the convection has started, the heat rate across \( r_0 \) is therefore transferred by diffusion to the liquid up to the phase change boundary. In this situation, it can be written:

\[
-k A(r) \frac{\partial T(r)}{\partial r} \bigg|_{r=r_0} = -2\pi k \frac{T^* - T_w}{\ln \left( \frac{r}{r_0} \right)} = -2\pi r_0 h \left( T^* - T_w \right)
\]

The Nusselt number is then defined for this problem as the relation \( q_{\text{conv}}/q_r \). For the situation of pure diffusion:

\[
Nu = \frac{q_{\text{conv}}}{q_r} = \frac{h}{k} r_0 \ln \left( \frac{r}{r_0} \right) = 1
\]
Results for the Melting Process

Next, it is presented the solutions for the wall boundary conditions of $T_0=4\, ^\circ C$ and $T_0=8\, ^\circ C$, applied to a tube with radius $r_0=0.032\, m$, and starting at the initial condition in the solid of $T_{i1}=0\, ^\circ C$. The thermal resistance imposed by the tube wall was neglected. Since the phenomenon is transient, solutions were collected at various instants $t$ for the two wall boundary conditions. The streamlines in the liquid region are plotted in the same graphs of the density. This allows a better understanding of the phenomenon associated with the inversion in the density, based on the observation of the local Nusselt number, “Eq. (29)”, and of the average Nusselt number, “Eq. (30)”.  

For $t=3000\, s$, there are variation of the Nusselt number with the angle. It can be observed the beginning of the convective effects through a small increase on the Nusselt number for the angular position $\theta<100^\circ$. For the lower region, $100^\circ<\theta<180^\circ$, there is a decrease in the value of the Nusselt number. The maximum temperature that the liquid reached in this simulation was $4\, ^\circ C$, while the minimum was $0\, ^\circ C$. The portion of liquid with the highest temperature will assume the highest density. Due to the gravitational field action, the fluid with the lowest density and temperature will be concentrated in the upper positions, and so intensifying the heat rate in that region. The fluid with the highest temperature and density will occupy the lower positions of the tube, leading to a decrease in the heat rate from the wall to the fluid. As a result, the Nusselt number presents values lower than the unity. This decrease occurs due to the assumptions taken for the derivation of “Eq. (26)”, in which it was considered that the heat rate that penetrates in the boundary by diffusion is mostly transported by advection in the interior of the tube. In “Fig. 6”, it can be observed the evolution of the phase change boundary as well as the behavior of the process parameters, temperature and density, which validate the aforementioned explanations. The temperature evolution demonstrates the concentration of liquid with higher temperatures in the lower regions of the domain. The aforementioned characteristics of the process intensify with the time evolution; that is, the Nusselt number increases in the upper positions and decreases in the lower positions.

![Figure 5. Nusselt number behavior as a function of the angle $\theta$ and time $t$ for melting with $T_0=4$.](image)

**Figure 5.** Nusselt number behavior as a function of the angle $\theta$ and time $t$ for melting with $T_0=4$.  

For $t=7000\, s$, there is another vortex structure. The vortex that spins in the clockwise direction causes the formation of two locations with peaks in the Nusselt number, existing in the initial instants, $t=100\, s$, and $t=5000\, s$, the first peak in the Nusselt number is due to the formation of a vortex in the upper part of the tube, and a second peak occurs in a intermediary angular position, and is formed by the PCM ring thickness variation in the liquid state. For $t=18000\, s$, there is another vortex structure. The vortex that spins in the clockwise direction, a negative vortex, predominates, while a positive vortex, located in the lower region of the domain, does not affect significantly the heat transfer process. In this situation, there is a peak in the Nusselt number in the upper part of the tube, a region where liquid with lower temperature and density is concentrated.

Figure 7 shows the behavior of the Nusselt number along the angular position for $T_0=8\, ^\circ C$, while “Fig. 8” presents the behaviors of the temperature and the density fields, to which streamlines are juxtaposed, for this situation. Again, the diffusive process is observed in the initial instants, $t=100\, s$, in “Fig. 7”. After some time, the convection mechanism takes place, as can be observed by the behavior of the Nusselt number. In this situation, one can observe the inversion in the behavior of the Nusselt number as compared to the previous situation. The peaks in the Nusselt number occur for angular positions $\theta=50^\circ$. For $\theta=50^\circ$, there is a decrease in the Nusselt number.
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Figure 6. On the right, the streamlines (+ means anticlockwise recirculation) are juxtaposed to the density; the temperatures fields for melting with $T_w=4 \, ^\circ C$, $T_{ini,S}=0 \, ^\circ C$, $t=3000 \, s$, $t=7000 \, s$, $t=12000 \, s$, and $t=18000 \, s$ are shown on the left.

The difference between the two solutions can be explained by the inversion in the density for temperatures close to 4 °C. For this wall temperature, there is concentration of liquid with high temperature and lower density in the upper regions. The fluid formed from the melting of solids with lower temperatures also presents a density inferior to the maximum value, and is prone to move to the upper regions. However, as the fluid starts the motion, it mixes with fluids at higher temperature, reaching the temperature of 4 °C, a situation in which the water has the maximum density, and so it moves towards the lower positions of the tube. For the previous situation, $T_w=4 \, ^\circ C$, there was an inverse behavior, although with the same driving mechanism.

The behaviors of the density and temperature fields for $T_w=8 \, ^\circ C$ can be observed in “Fig. 8”. The simulations indicate that, for situations in which $T_w<4 \, ^\circ C$, the behaviors are similar to those found for $T_w=4 \, ^\circ C$, while for $T_w>8 \, ^\circ C$ it is similar to $T_w=8 \, ^\circ C$. For $T_w=6 \, ^\circ C$, in the beginning of the process the behavior resembles the situation for $T_w=4 \, ^\circ C$; after some time, the solution presents an inversion, with a behavior similar to that for $T_w=8 \, ^\circ C$.

“Figs. 5, 6, 7 and 8” show the dynamic aspect of the heat transfer phenomenon, which makes it unfeasible to draw conclusions from local results to predict the heat transfer behavior. Instead, it is more adequate to consider the average behavior, as is in “Fig. 9”, which shows the average Nusselt number, “Eq. (30)”, for the two considered situations.
Figure 8. On the right, the streamlines (+ means anticlockwise recirculation) are juxtaposed to the density. The temperatures fields for melting with $T_w=8 \, ^\circ C$, $T_{S}=0 \, ^\circ C$, $t=3000 \, s$, $t=6500 \, s$, $t=8500 \, s$, and $t=11000 \, s$ are shown on the left.

Figure 9. Average Nusselt number behavior as a function of the relation $V_f/V_0$.

In the search for correlations that enable the evaluation of the heat transfer, and that are not prone to be affected by the “singular” behavior of the water density, it was employed an expression that has been presented in “Sasaguchi et al. (1998)”. Applying it into the dimensionless momentum equations, more specifically in the buoyancy terms, leads to the following Rayleigh number:

$$Ra = \frac{\eta g |T_w - T_s| \cdot L}{\alpha \nu}$$  \hspace{1cm} (31)
where $\eta = 9.297173 \times 10^{-6}$, $L$ is the average thickness of the liquid ring, $b = 1.894816$, $\alpha$ is the thermal diffusivity, $v$ is the kinematic viscosity, and $T_{ref}$ takes the value of $T_w$ in the melting.

This parameter allowed the identification of a pattern behavior of the $Nu_{avg}/Ra$ ratio, as a function of the melted volume $V_f$ to the initial volume $V_0$, $V_f/V_0$, and also as a function of $Fo$, as can be observed in “Fig. 10”, where logarithm scales were adopted.

Such behaviors were approximated by the following relations:

$$\log \left( \frac{Nu}{Ra} \right) = a \log \left( \frac{V_f}{V_0} \right) + b$$  \hspace{1cm} (32)

$$\log \left( \frac{V_f}{V_0} \right) = d \log (Fo) + c$$  \hspace{1cm} (33)

where $a$, $b$, $c$ and $d$ are the coefficients to be determined. This allows writing an expression for the Nusselt number in the following way:

$$Nu = 10^a \cdot Fo^b \cdot Ra^c$$  \hspace{1cm} (34)

The terms $E_1$ and $E_2$ are defined as:

$$E_1 = \begin{cases} 531.105 \cdot Ste^3 + 244.60 \cdot Ste^2 - 50.58 \cdot Ste - 1.682 & \rightarrow Ste \leq 0.1 \\ 531.105 \cdot Ste^3 + 259.85 \cdot Ste^2 - 53.95 \cdot Ste - 1.5 & \rightarrow Ste > 0.1 \end{cases}$$  \hspace{1cm} (35)

$$E_2 = \begin{cases} -0.395 \cdot Ste^2 - 0.894 \cdot Ste - 1.6418 & \rightarrow Ste \leq 0.1 \\ 5.2 \cdot Ste^2 - 2.86 \cdot Ste - 1.4924 & \rightarrow Ste > 0.1 \end{cases}$$  \hspace{1cm} (36)

$$Ste = \frac{c_{pc} \left( T_w - T^* \right)}{\lambda}$$  \hspace{1cm} (37)

$$Fo = \frac{aT}{c_T}$$  \hspace{1cm} (38)

where the $\alpha$ is the thermal diffusivity.

Based on the behaviors of the presented solutions, it is possible to establish correlations for the melted volume $V_f$ and, consequently, for the average thickness of the melted PCM:

$$V_f = 10^d \cdot Fo^e \cdot V_0$$  \hspace{1cm} (39)

$$L = r_0 \cdot \sqrt{c_T \left( 1 - Fo^{d+10} \right)}$$  \hspace{1cm} (40)

where the terms $d$ and $c$ are obtained according to:

$$d = 0.2631 \cdot Ste + 0.5304 \rightarrow Ste \leq 0.1$$

$$d = 1.3044 \cdot Ste + 0.4264 \rightarrow Ste > 0.1$$

$$c = 3.6912 \cdot Ste - 0.4531$$

The efficacy of the proposed correlations can be evaluated in “Fig. 11”, where they are compared to the results obtained with the simulation. In the values of the simulations, the percentage variations were added, allowing the visualization of the maximum existing differences between the two results. It can be observed for the Nusselt number that the maximum differences occur at the beginning of the melting process, where the diffusion is the predominant heat transfer mode. In the time periods where the convection is the dominant mode, the differences between the two solutions decrease. Indeed, in literature, convection correlations with uncertainties as high as 20% occur.

**Results for the Melting Process Followed by Resolidification**

As a consequence of the variable ambient conditions, which cause variations in the thermal load, it is common the partial use of the stored energy in the ice banks along the time. Therefore, simulations considering this situation also must be carried through, allowing a better knowledge of the heat transfer mechanisms. The results for the partial melting process followed by resolidification are shown in “Figs. 12 and 13”, considering that the PCM in the solid phase is initially supercooled (Neumann approach), $T_{int,L}$=5 °C. For $t=0$ s, the tube wall was exposed to $T_w$=8 °C, initiating the melting. After a period of 7760 s, the surface temperature on the external surface changes to $T_{ext}$=-5.5 °C, initiating the resolidification. The behaviors of the enthalpy and density in the domain are also presented. The dashed line represents the enthalpy in the region where the liquid reaches its maximum density, for a temperature of 4 °C. On the left corner on the bottom, the Rayleigh number for the considered instant is presented. Together with the relative density, the streamlines are presented, enabling the visualization of the liquid behavior during the phenomenon evolution. The instants of time for which the simulations were run are presented on the bottom corner.

The behavior of the melting process is first shown for the time instant $t=200$ s. It can be observed that the sensible energy associated with the solid supercooling rapidly disappears, and in this instant diffusion predominates. For $t=3000$ s, there is no longer traces of sensible energy, and the advection mechanism is already active in the heat transfer. For $t=7780$ s, the boundary condition is altered so that solidification occurs. One can note a thin layer of solidified material in the external region. The liquid is then confined between the two phase change boundaries. The sensible energy associated with the liquid is released to the two boundaries. This initially reduces the advance of the solidification in the external boundary. In the inner boundary, the melting continues until the thermal equilibrium is reached. Initially, the thickness of the solidified external layer is not uniform, as for when $t=7800$ s. This occurs due to the fact that it is in the upper region where the liquid with the lowest density and enthalpy level concentrates. This difference is small, being the more intense the larger is the temperature $T_w$ set to cause the melting. In the solidification, the vortices structure changes drastically, as can be observed for the instants $t=7780$ s, $t=7800$ s, and $t=8000$ s. As the solidification advances, the inner liquid and the solid reach thermal equilibrium, as for when $t=8750$ s. After this, the buoyancy effects become negligible, the velocities associated with the liquid are of such a magnitude that convection stops acting, so that diffusion predominates again, and the liquid density becomes homogeneous. The inner solid becomes “insulated”, without altering its energetic state until $t=10750$ s, when the two boundaries meet. From this point ahead, there is heat transfer from the inner solid by diffusion. This can be visualized for time instants later than $t=12500$ s.

In the solidification, the time and intensity of the natural convection actuation are larger for situation where the initial sensible energy, or equivalently the initial liquid temperature ($T_{int,L}$), is large and when the $T_w$ temperature is low. To evaluate the influence of the natural convection in the solidification process, it was plotted the behavior of the $T_{int}$, defined in “Eq. 42” with $m_L$ being the liquid mass.
Figure 10. On the left: behaviors of the $\frac{Nu_{\text{average}}}{Rac}$ ratio as a function of $Vf/V0$. On the right: the $Vf/V0$ ratio as a function of the dimensionless time $Fo$, for various wall temperatures $T_w$.

Figure 11. Comparison between the relations $\log \frac{Nu_{\text{average}}}{Rac}$ and $Vf/V0$ as functions of $Fo$ obtained from the numerical simulation and from the correlations.
In both situations, it was considered the same physical model, and the same boundary and initial conditions, the difference being solely on the solution method. The first case was dealt considering the solidification process to be purely diffusive; in the other case, the convection mechanism was taken into account.

\[
T_{\text{solidization}} = \left( \frac{1}{m} \int_{v_{\text{liquid}}} h \rho \text{dy} \right) - \lambda
\]

(42)

The observations show a faster decrease on \( T_{\text{mist}} \) in the simulation that took the natural convection into account, and this trend remained the same for the other simulations having different boundary and initial conditions. The difference between the solutions narrowed with the decrease of the liquid sensible energy in the beginning of the process. The time needed for the temperature homogenization in the interior of the liquid domain also decreased for the same reason; the decrease on \( T_{\text{ini,L}} \) leads to the same effects.

The influences of the natural convection, and of the sensible energy existing in the beginning of the process on the PCM solidification time, can be seen in “Fig. 14” (right side). The results for the two different solidification situations are exposed. For the first and second cases, it was set \( T_{\text{ini,L}}=8 \) °C and 16 °C. For both cases, \( T_{\text{ini}}=5.5 \) °C and the solutions were obtained for processes with pure diffusion and with natural convection.

Observing the left side of “Fig. 14”, one realizes the proximity between the purely diffusive and natural convection solutions, since the steeper decrease on \( T_{\text{mist}} \) when convection is taken into account is not sufficient to cause noticeable differences between them (solidification time). This behavior can be justified by the sensible energy being much smaller than the latent energy (that is, low Stefan numbers). The differences in the values of \( T_{\text{solidization}}/T_{\text{ini,L}} \) between the two observed cases in the left side are consequences of the fluid movement in the convective process, causing a faster homogenization of the temperature in the liquid. The amount of sensible energy in the liquid has small influence on the solidification speed, for temperatures around \( T_{\text{ini,L}}=8 \) °C, so that there is a coincidence between the solutions with natural convection and pure diffusion.

For higher temperatures, as for \( T_{\text{ini,L}}=16 \) °C, the solidification speed is smaller for the initial instants in the convective process, \( V_f/V_0>0.6 \). After the temperature homogenization is reached, for \( V_f/V_0>0.4 \), there is an inversion in the solidification speed, and the convective process becomes faster. In the end of the process, the solidified volume is identical for the two cases, and the time needed for this to occur, \( t_{\text{solidization}} \), was similar for all the simulations.
Figure 13. Results for the partial melting in the periods $t=8300$ s, $t=8750$ s, $t=10750$ s, $t=12500$ s and $t=13000$ s. On the left and right sides, respectively, the behaviors of the specific enthalpy and of the density together with the streamlines are presented.

Figure 14. Comparisons between numerical solutions for the solidification, considering the pure diffusive process and the presence of natural convection. On the left: the $T_{\text{max}}/T_{\text{ini},L}$ ratio as a function of $V_{\text{sol}}/V_0$ for temperatures $T_{\text{ini},L}=16^\circ$C and $T_\text{w}=-5.5^\circ$C, in the solidification process. On the right: the $V/V_0$ ratio as a function of the $V_{\text{max}}$ for temperatures $T_{\text{ini},L}=8^\circ$C and $T_{\text{ini},L}=16^\circ$C, with $T_\text{w}=-5.5^\circ$C.
Conclusions

Melting initiates by purely diffusive heat transfer. In the evolution of the process, the space occupied by the liquid phase increases. This causes the increase in the Rayleigh number, intensifying the natural convection, which in many cases can become the dominant heat transfer mechanism. The solidification process initiates in the presence of natural convection, since the larger portion of the domain is in the liquid phase. As time progresses, the thermal gradients and the spaces occupied by the liquid diminish, decreasing the Rayleigh number and causing, from a certain point in time, the diffusion to become the dominant heat transfer mechanism, as seen in “Fig. 13” for $\text{Ra}_\text{c}=63$. As expected, the dominant mechanism of heat transfer in the interior of the duct changes with time and the boundary conditions.

The vortices structure, which varies with time and the boundary conditions, drastically interferes in the change of the heat transfer mechanism. The degree of complexity of the heat transfer phenomenon, in the considered problems, is associated with the “singular” behavior of the water density. This behavior is responsible for the variation in the vortices structure and in the isothermals in the interior of the tube. In the melting process, this structure “absorbs” the energy on the wall and then “releases” it to the solid, intensifying the exchange mechanism in specific positions, where the fluid with low enthalpy concentrates. The vortices structures have influence on the phase change boundary. As long as the diffusive process is dominant, the solid maintains a cylindrical configuration; however, as the vortices structure intensifies, the geometry of the phase change boundary “deforms”. In the initial instants of the resolidification, the liquid sensible energy is released to the two phase change boundaries. The external phase change boundary maintains a homogeneous geometry, for the liquid sensible energy is rapidly consumed, establishing a homogeneous density and minimizing the effect of natural convection.

In the solidification, the assumption of pure diffusion leads to results that are close to the solutions that consider natural convection, for the engineering oriented cases that were considered in this work. In the melting, a larger discrepancy between the two solutions was observed.

Correlations for the Nusselt number, and for the melted volume as functions of time were also obtained for practical use in engineering.

References


