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Title: Boiling heat transfer and pressure drop of NH3/LiNO3 and NH3/(LiNO3+H2O) in a plate heat exchanger

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The mass flux greatly influenced the flow boiling heat transfer coefficient, whereas the addition of water only produced a slight increase. The measurements taken experimentally indicated that the parameters with a pronounced effect on the frictional pressure drop were vapour quality followed by mass flux. Finally, the correlations proposed in Táboas et al. [1] used to predict the flow boiling heat transfer coefficient and frictional pressure drop were well in agreement with the experimental results.

1	Boiling heat transfer and pressure drop of NH ₃ /LiNO ₃ and
2	$NH_3/(LiNO_3+H_2O)$ in a plate heat exchanger
3	
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11	
12 13 14	Keywords : flow boiling; plate heat exchanger; ammonia; lithium nitrate, absorption refrigeration
15	Highlights:
16 17	• Flow boiling heat transfer and frictional two-phase pressure drop for NH ₃ /LiNO ₃ and NH ₃ /(LiNO ₃ +H ₂ O) in a PHE are presented
18 19	• The effects of heat flux, mass flux and vapour quality on the flow boiling heat transfer and frictional pressure drop are analysed
20	• Mass flux has the most influence on the flow boiling heat transfer
21 22	• Those parameters with a pronounced effect on the frictional pressure drop are the vapour quality followed by the mass flux
23	

Abstract

This paper presents the experiments carried out to determine the flow boiling heat transfer coefficient and associated frictional pressure drop in a plate heat exchanger which uses the binary fluid mixture ammonia/lithium nitrate and the ternary fluid mixture ammonia/(lithium nitrate + water) with a water content in the absorbent of 20 % by weight. The effects on the flow boiling heat transfer coefficient and two phase frictional pressure drop are analysed with a heat flux range of 5 to 20 kW·m⁻², a mass flux of 50 to 100 kg·m⁻²·s⁻¹ and a mean vapour quality of 0 to 0.2.

The mass flux greatly influenced the flow boiling heat transfer coefficient, whereas the addition of water only produced a slight increase. The measurements taken experimentally indicated that the parameters with a pronounced effect on the frictional pressure drop were vapour quality followed by mass flux. Finally, the correlations proposed in Táboas et al. [1] used to predict the flow boiling heat transfer coefficient and frictional pressure drop were well in agreement with the experimental results.

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46 Nomenclature

47	Α	Heat transfer area (m ²)
48	b	Amplitude of the plate (m)
49	Во	Boiling number (-)
50	Ср	Heat capacity $(J \cdot kg^{-1} \cdot K^{-1})$
51	D_h	Hydraulic diameter (m)
52	е	Plate thickness (m)
53	8	Gravitational acceleration $(m \cdot s^{-2})$
54	G	Mass flux $(kg \cdot m^{-2} \cdot s^{-1})$
55	h	Heat transfer coefficient $(W \cdot m^{-2} \cdot K^{-1})$
56	Н	Height of the plate (m)
57	k	Thermal conductivity $(W \cdot m^{-1} \cdot K^{-1})$
58	Ke/V	Kinetic energy per unit of volume $(J \cdot m^{-3})$
59	L	Flow length of the plate (m)
60	LMTD	Log mean temperature difference
61	m	Mass flow rate $(kg \cdot s^{-1})$
62	Nu	Nusselt number
63	Р	Pressure (bar)
64	Pr	Prandtl number
65	Q	Heat power (W)
66	R	Thermal resistance $(K \cdot W^{-1})$
67	Re	Reynolds number
68	Т	Temperature (K)
69	U	Global heat transfer coefficient $(W \cdot m^{-1} \cdot K^{-1})$

70	и	Velocity $(\mathbf{m} \cdot \mathbf{s}^{-1})$
71	V	Volume (m ⁻³)
72	W	Width of the plate (m)
73	W	Liquid mass fraction
74	у	Vapour mass fraction
75	Z	Total mass fraction
76		
77	Greek syr	nbols
78	β	Corrugation angle
79	Λ	Pitch (m)
80	ρ	Density $(kg \cdot m^{-3})$
81	μ	Dynamic viscosity $(kg \cdot m^{-1} \cdot s^{-1})$
82	arphi	Chisholm two phase parameter
83	X_{LM}	Lockhart-Martinelli parameter
84	Subscript	S
85	cb	Convective boiling
86	exp	Experimental
87	f	Friction
88	h	Hydraulic
89	in	Inlet
90	out	Outlet
91	L	Liquid
92	LO	Liquid overall
02		Homogonous

94	man	Manifold
95	mon	Momentum
96	nb	Nucleate boiling
97	SS	Solution side
98	steel	Stainless steel
99	Т	Total
100	TP	Two phase
101	v	Vapour
102	W	Water
103	wall	Wall
104	WS	Water-side
105		

106 **1 Introduction**

107 Nowadays most absorption heat pumps and chillers use water/lithium bromide or 108 ammonia/water mixtures as working fluids. These working fluids have several 109 drawbacks, such as crystallization, corrosion and low-pressure operating conditions for 110 water/lithium bromide mixture. The refrigerant vapour leaving the generator needs to 111 be rectified and the temperature required for the heat source should be higher for 112 ammonia/water mixture. Other working fluids, such as ammonia/lithium nitrate (Gensch 113 [2], Aggarwal and Agarwal [3], Infante Ferreira [4], Antonopoulos and Rogdakis [5], 114 Kim and Machielsen [6]), have been studied for use as alternatives to overcome the 115 aforementioned drawbacks.

116 The absorption refrigeration cycle can be operated at lower generator temperatures 117 when using ammonia/lithium nitrate instead of ammonia/water and it is not necessary to rectify the refrigerant vapour leaving the generator. The advantages of this working pair, therefore, are the simplicity of the cycle and the greater potential for using solar cooling or low grade temperature heat sources. However, the drawback with this working fluid is its high viscosity, which penalizes heat and mass transfer processes in the absorber and generator.

123 Infante Ferreira [4] compiled thermodynamic and thermophysical properties of the 124 ammonia/lithium nitrate fluid mixture and proposed a series of correlations for their 125 determination. These correlations were later used by several authors to carry out 126 thermodynamic simulations of different absorption cycles such as single-effect 127 (Antonopoulos and Rogdakis [5], Kim and Machielsen [6], Niebergall [7], Sun [8]), and 128 double and half-effect ones (Bourouis et al. [9], Arzoz et al. [10] and Ayala et al. [11]).

129 Experimental data obtained with absorption cooling prototypes designed initially to 130 operate with the ammonia/water working pair (Ayala et al. [12], Infante Ferreira [13], 131 Heard [14]) and loaded with the ammonia/lithium nitrate working pair showed poor performance results. The authors of these experimental studies concluded that the main 132 133 reason for the poor performance lay in the absorber and was due to the high viscosity of 134 the fluid mixture compared with that of the ammonia/water mixture. High viscosity 135 reduces the cycle performance predicted by the thermodynamic models. This decrease 136 in the cycle performance is more pronounced at low temperatures of cooling-water due 137 to the fact that the viscosity of the working fluid in the absorber increases drastically.

Ehmke and Renz [15], and Bokelmann et al. [16] proposed the addition of water to the binary mixture ammonia/lithium nitrate used in absorption heat pumps. Later, Reiner and Zaltash [17] proposed the use of the ternary mixture for GAX systems as an alternative to using ammonia/water systems.

142 Ehmke and Renz [15] studied the effect of adding water on the solubility and viscosity 143 of the ternary mixture and suggested an optimal mass fraction of water between 0.20 144 and 0.25 in the absorbent mixture (lithium nitrate + water). These authors also 145 determined and correlated data for the density and vapour pressure of the mixture at 146 0.25 of water mass fraction in the absorbent. Bokelmann et al. [16] carried out an 147 experimental study dealing with the performance of an absorption heat pump working 148 with the ternary fluid mixture ammonia/(lithium nitrate + water). The data were 149 reported by Manago [18] in a study on new mixtures for absorption heat pumps for the 150 Heat Pump Program of The International Energy Agency. Reiner and Zaltash [17] 151 measured the densities and viscosities of the ternary mixture with an ammonia mass 152 fraction of 0.04 and a water mass fraction of 0.605, which are the typical values for 153 GAX systems.

In 1989, Bothe [19] published a comparative study carried out with ammonia/water and ammonia/(lithium nitrate + water) as working fluids for heat pumping applications. The author reported a significant improvement in COP using the ternary mixture compared to using the binary mixture.

158 Libotean et al. [20] obtained vapour pressure equilibrium data and Libotean et al. [21] 159 measured viscosities, densities and heat capacities for the ammonia/lithium nitrate and 160 ammonia/(lithium nitrate + water) mixtures. Linke [22], and Eysseltova and Orlova 161 [23], published experimental data regarding the solubility of the binary and ternary 162 mixtures. According to this data, the addition of a small amount of water improves the 163 solubility of the mixture. Recently, Cuenca et al. [29] presented new experimental data 164 on thermal conductivity for the ammonia/lithium nitrate and ammonia/(lithium nitrate + 165 water) mixtures.

Regarding the flow boiling heat transfer coefficient and pressure drop of these mixtures, Rivera and Best [24] published experimental data on flow boiling in a vertical tube with mass fluxes of 7.4 and 13.7 kg·m⁻²·s and a heat flux of between 11.8 and 16.4 kW·m⁻². The experimental local heat transfer coefficients were in the range of 1.3-4.0 kW·m⁻²·K⁻¹. The authors reported that both the forced convective and nucleate boiling mechanisms were significant and they proposed a heat transfer coefficient correlation based on their experimental data.

173Zacarías et al. [25] published data on the flow boiling heat transfer coefficient in plate174heat exchangers with the mixture ammonia/lithium nitrate. The data was obtained with a175maximum vapour quality generation of 0.03 and high subcooling of the solution176entering the heat exchanger. The solution flow rate was varied between 0.041 and 0.083177kg·s⁻¹. In the flow boiling region, the heat transfer coefficient ranged from 0.6 to 1.1178kW·m⁻² K⁻¹. The authors concluded from the experimental results, that nucleate boiling179appeared to be the dominant factor.

180 Regarding the boiling of the ammonia/(lithium nitrate + water) fluid mixture, 181 Sathyabhama and Ashok Babu [26] investigated the nucleate pool boiling heat transfer 182 coefficient applying an operating pressure of 4 to 8 bar, an ammonia mass fraction 183 ranging from 0 to 0.3 and different heat fluxes. The lithium nitrate concentration in the 184 solution was chosen from the range of 10-50% of mass ratio of lithium nitrate in pure 185 water. The effects of the concentration, heat flux, and pressure on the boiling heat 186 transfer coefficient were analysed. The authors concluded that the heat transfer 187 coefficient decreases with an increase in the ammonia mass fraction, increases with the 188 addition of lithium nitrate and increases with a rise in heat flux and pressure. To the 189 knowledge of the authors, there is currently no data available on flow boiling of the 190 ammonia/(lithium nitrate + water) fluid mixture.

The objective of the present work is to contribute to the technological development of brazed plate heat exchangers (BPHEs) as desorbers for ammonia based absorption refrigeration systems. For this purpose, the flow boiling heat transfer coefficient and associated frictional pressure drop in a plate heat exchanger were measured for the binary ammonia/lithium nitrate and ternary ammonia/(lithium nitrate + water) fluid mixtures. The experimental results were analysed whilst varying mass flux, heat flux, vapour quality, pressure and mixture concentration.

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9 **2 Experimental set-up**

Figure 1 shows a schematic diagram of the generator test bench, which consists of a solution circuit and three auxiliary circuits. The pre-evaporator circuit provides the desired vapour quality at the inlet of the test section. The heating circuit provides the heating for the test section and allows the heat flux to be fixed. The condensation circuit condenses the ammonia vapour which will be returned to the solution tank in order to regenerate the solution to the initial conditions.

The solution stored in the solution tank is pumped by the recirculation pump from the bottom through the pre-evaporator, where it is preheated to establish the required vapour quality at the generator inlet. Subsequently, the solution enters the plate heat exchanger (generator). The resulting two-phase flow at the outlet of the generator enters the solution tank and from the top of it the ammonia vapour is sent to the condenser where it condenses and is returned back to the solution tank to start the cycle again.





Figure 1. Schematic diagram of the experimental test bench

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The solution and water temperatures of the auxiliary circuits were measured with Pt-100 sensors located at the points indicated in Figure 1. The flow rate and concentration of the solution entering the test section were determined using a Coriolis flowmeter. The water flow rates in the auxiliary circuits were measured by magnetic flowmeters. The pressure was measured by two pressure sensors located at the inlet and outlet of the plate heat exchanger (test section). More details about the experimental set-up and its components can be found in Táboas et al. [1].

222 The section test consists of a plate heat exchanger manufactured by Alfa Laval, NB51,

223 with a chevron corrugation angle of 30 degrees to the horizontal and a plate thickness of

0.4 mm. The heat exchanger consists of four plates forming three channels. The solution
circulates through the central channel while the hot water circulates through the side
channels. A channel 100 mm wide was used to calculate the flow area. Figure 2 shows
the main parameters of the plate heat exchanger.





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230

Figure 2. Main parameters of the plate heat exchanger

231 **3 Methodology**

The design of the test bench allows for flow boiling experiments to be carried out varying the following operating parameters: solution mass flow rate, solution inlet temperature, heating water inlet temperature, solution inlet concentration, and system pressure. In each experiment, once a steady-state regime was reached, all operating conditions were registered and stored for about 25 minutes. The parameters characterizing the generator behaviour were calculated as reported in the following sections.

3.1 Properties of the working fluid

Vapour-liquid equilibrium of the ammonia/lithium nitrate and ammonia/(lithium nitrate
+ water) fluid mixtures was calculated using the correlations reported by Libotean et al.
[21]. The densities and viscosities of these fluid mixtures were calculated by the
correlations of Libotean et al. [20].

244 The liquid enthalpies of both ammonia/lithium nitrate and ammonia/(lithium nitrate + 245 water) fluid mixtures were obtained with the Haltenberger method [27] employed by 246 McNeely [28] for the conventional working pair water/lithium bromide. This method 247 was developed to obtain the liquid enthalpy of binary fluid mixtures in which only one 248 component is volatile. In the case of the ternary fluid mixture, the procedure was the 249 same, taking into account that the water content in the vapour phase was insignificant. 250 The enthalpy reference value for ammonia is 0 kJ/kg at 0°C, and the reference state for 251 the ammonia/lithium nitrate and ammonia/(lithium nitrate + water) fluid mixtures was 252 obtained with an ammonia mass fraction of 0.5 at 0°C. The thermal conductivity of both 253 fluid mixtures was determined from the data reported by Cuenca et al. [29].

3.2 Single-phase heat transfer coefficient and pressure

255 **d**

drop

The water-side heat transfer coefficient (h_{ws}) was determined by carrying out preliminary tests using water in both the hot and cold sides. The thermal load of the heat exchanger was obtained as the average value of the heat transfer rate calculated on the hot and cold sides (Eqs. (1-3)).

$$260 \qquad Q_{ws} = m_{ws} \cdot Cp_{ws} \cdot \left(T_{ws,out} - T_{ws,in}\right) \tag{1}$$

$$261 \qquad Q_{ss} = m_{ss} \cdot Cp_{ss} \cdot \left(T_{ss,out} - T_{ss,in}\right) \tag{2}$$

262
$$Q_{mean} = \frac{Q_{ss} + Q_{ws}}{2}$$
 (3)

263 Once the thermal load is obtained, the experimental global heat transfer coefficient can264 be calculated as follows:

265
$$U_{\exp} = \frac{Q_{mean}}{A \cdot LMTD}$$
(4)

266 Where LMTD is the logarithmic mean temperature difference involving inlet and outlet 267 temperatures in the generator (Eq. (5)):

$$268 \quad LMTD = \frac{(T_{ws,in} - T_{ss,out}) - (T_{ws,out} - T_{ss,int})}{Ln\left(\frac{(T_{ws,in} - T_{ss,out})}{T_{ws,out} - T_{ss,int}}\right)}$$
(5)

Eq. (4) is compared with the overall heat transfer coefficient expressed by Eq. (6) that combines the different thermal resistances between the solution and the coolant streams.

271
$$\frac{1}{U} = \frac{1}{h_{ws}} + \frac{1}{h_{ss}} + \frac{e}{k_{steel}}$$
 (6)

Where the single-phase heat transfer coefficient, expressed by the correlation of Eq. (7), was obtained by a least square adjustment between U_{exp} and U. The expression is valid for the turbulent regime (Re: 690 – 3100) with a coefficient of determination R²=0.998.

275
$$h = 0.577 \frac{k}{D_h} R e^{0.634} P r^{1/3}$$
 (7)

The single-phase friction pressure drop was calculated by subtracting the static and manifold pressure drops from the total pressure drop, as shown in Eq. (8):

278
$$\Delta P_f = \Delta P_{total} - \Delta P_{static} - \Delta P_{man}$$
(8)

279 The static pressure drop was calculated by Eq. (9):

$$280 \qquad \Delta P_{static} = g \cdot \rho_{mean} \cdot L \tag{9}$$

- 281 Where ρ_{mean} is the density calculated at the average temperature between the inlet and 282 outlet of the generator.
- The pressure drop at the manifolds was estimated using the correlation reported by Shahand Focke [30], Eq. (10):

$$285 \qquad \Delta P_{man} = 1.5 \cdot \left(\frac{\rho \cdot u^2}{2}\right) \tag{10}$$

Finally, the Fanning friction factor was calculated using equation Eq. (11):

$$287 f = \frac{\Delta P \cdot D_h \cdot \rho}{2 \cdot G^2 \cdot L} (11)$$

According to our experimental data, the correlation for the single-phase flow friction factor is given by Eq. (12) with $R^2=0.992$. This equation was obtained for the Reynolds number in the range 375-2500.

291
$$f = 4.778 \cdot \text{Re}^{-0.118}$$
 (12)

3.3 Flow boiling heat transfer coefficient

The approach used to calculate the flow boiling heat transfer coefficient on the solution side is described below. Firstly, the heat transfer rate was calculated from Eq. (13). The experimental overall heat transfer coefficient and the logarithmic mean temperature difference (*LMTD*) were calculated as in single-phase experiments by Eqs. (14) and (15):

$$298 \qquad Q_{ws} = m_{ws} \cdot Cp_{ws} \cdot \left(T_{ws,out} - T_{ws,in}\right) \tag{13}$$

$$299 \qquad U = \frac{Q_{ws}}{A \cdot LMTD} \tag{14}$$

$$300 \quad LMTD = \frac{\left(T_{ws,in} - T_{ss,out}\right) - \left(T_{ws,out} - T_{ss,in}\right)}{Ln\left(\frac{T_{ws,in} - T_{ss,out}}{T_{ws,out} - T_{ss,in}}\right)}$$
(15)

301 Finally, the flow boiling heat transfer coefficient was calculated from Eq. (16), while 302 h_{ws} was computed with Eq. (7):

$$303 \qquad \frac{1}{h_{TP}} = \frac{1}{U} - \left(\frac{1}{h_{ws}} + \frac{e}{k_{steel}}\right)$$
(16)

The global ammonia mass fraction of the solution was obtained using a Coriolis flowmeter to measure the density and temperature and by taking into consideration the density correlations from Libotean et al. [20]. The ammonia liquid mass fraction at the inlet and outlet of the heat exchanger was obtained by measuring the pressure and temperature and using the vapour-liquid equilibrium data of Libotean et al. [21]. Then the vapour quality of the mixture at the inlet and outlet of the heat exchanger was obtained by applying the liquid and global ammonia mass fraction, Eqs. (17)-(19):

$$311 \qquad m_L + m_V = m_T \tag{17}$$

$$312 \qquad m_L \cdot w + m_V = m_T \cdot z \tag{18}$$

$$313 \qquad x = \frac{m_V}{m_T} \tag{19}$$

314

The mean vapour quality was then assumed to be the average of the numerical values calculated at the inlet and outlet of the generator (Eq. (20)).

$$317 \quad x_{mean} = \frac{x_{in+}x_{out}}{2} \tag{20}$$

318

319 **3.4 Flow boiling two-phase pressure drop**

The total two-phase pressure drop in the plate heat exchanger is the sum of the frictional pressure drop, the static pressure drop, the momentum pressure drop, and the pressure drop produced by manifolds and ports. 323 The total pressure drop was determined experimentally using pressure transducers 324 located at the entrance and exit of the solution in the plate heat exchanger. The frictional 325 pressure drop, $\Delta P_{f,TP}$, associated with the two-phase solution flowing through the

327
$$\Delta P_{f,TP} = \Delta P_{total} - \Delta P_{static} - \Delta P_{mon} - \Delta P_{man}$$
(21)

328 The static pressure drop ΔP_{static} was determined as follows:

$$329 \quad \Delta P_{static} = g \cdot \rho_m \cdot L \tag{22}$$

330 where the average density ρ_m is obtained from the homogeneous model for two-phase

332
$$\rho_m = \left[\frac{x_{mean}}{\rho_V} + \frac{(1 - x_{mean})}{\rho_L}\right]^{-1}$$
(23)

333 The momentum pressure drop was estimated as follows:

334
$$\Delta P_{mon} = G^2 \cdot \left(\frac{1}{\rho_V} - \frac{1}{\rho_L}\right) \cdot \Delta x$$
(24)

The pressure drop resulting from manifolds and ports can be estimated using the correlation reported by Shah and Focke [30], Eq. (25):

$$337 \quad \Delta P_{man} \cong 1.5 \cdot \left(\frac{G^2}{2 \cdot \rho_m}\right) \tag{25}$$

338 Finally, after obtaining the value of $\Delta P_{f,TP}$, the Fanning friction factor for a two-phase

339 flow can be calculated as follows:

$$340 f_{TP} = \frac{\Delta P_{f,TP} \cdot D_h \cdot \rho_m}{2 \cdot G^2 \cdot L} (26)$$

341 3.5 Uncertainty of the measured and calculated

342 parameters

The approach used for determining the propagation of the uncertainty for the indirect variables was that proposed in the Technical Note 1297 of the National Institute of Standards and Technology (NIST) (Taylor and Kuyyat 1994). The EES (Engineering Equation Solver) software was used to perform the error analysis. Table 1 shows the accuracy of the variables measured and the combined uncertainty for the most relevant parameters calculated.

Measured variable	Accuracy
Temperature, T (°C)	± 0.1
Pressure, P (bar)	± 0.016
Water flow rate, $m_w (L/h)$	± 0.24
Solution flow rate, m_s (%)	± 0.1
Solution density, ρ (kg/m ³)	± 0.5
Calculated parameter	Combined uncertainty
Vapour Quality	± 0.0050-0.012
Flow boiling heat transfer coefficient, h (%)	±3.32-14.51
Friction factor, f (%)	±5.36-34.5

349	Table 1.Accuracy of	of measured	variables and	combined	uncertainty	of the	e calculated	parameters
						./		/

350

352 **4 Results and discussion**

Experimental results which were obtained on the generator test bench using the binary ammonia/lithium nitrate and ternary ammonia/(lithium nitrate + water) fluid mixtures are examined in this section. Operating conditions selected for temperature, pressure, solution mass flux, heat flux and concentration are summarized in Table 2. The effects of solution mass flux and heat flux on the experimental flow boiling heat transfer coefficient and associated frictional pressure drop are presented below and discussed with the variation of the mean vapour quality in the test section.

360

Table 2. Operating conditions

Parameters	Range
Heat flux, $(kW \cdot m^{-2})$	5.0, 10.0 and 20.0
Solution mass flux, (kg·s ⁻¹ .m ⁻²)	50 - 100
Ammonia mass fraction of the solution at the generator inlet (binary mixture)	0.49-0.54
Ammonia mass fraction of the solution at the generator inlet (ternary mixture)	0.42-0.46
Water content in the absorbent for the ternary mixture, (mass fraction)	0.20
Mean generator pressure, (bar)	12-15
Mean vapour quality, (kg vapour / (kg solution +kg vapour))	0-0.16

361

Moreover, the experimental values of the flow boiling heat transfer coefficient obtained with the binary mixture ammonia/lithium nitrate have been compared with those reported by Zacarías et al. [25] for the desorption process of the same fluid mixture in a plate heat exchanger which operated at the following experimental operating conditions: mass flux ranging from 10 to 20.2 kg·m⁻¹·s⁻², pressure from 9.78 to 16.06 bar, ammonia mass fraction from 0.452 to 0.462 and heat flux from 1.203 to 4.618 kW·m⁻².

369 **4.1 Two-phase Pressure drop**

The total pressure drop for the ammonia/ lithium nitrate and ammonia / (lithium nitrate
+ water) fluid mixtures was measured on the test bench, and the frictional pressure drop
was determined as described in section 3.4.

373 Figure 3 shows the experimental frictional pressure drop in the PHE as a function of the 374 mean vapour quality for the binary and ternary fluid mixtures. Heat fluxes were imposed at 5, 10 and 20 kW·m⁻², operating pressure at between 12 and 15 bar and 375 solution mass flux between 50 and 100 kg \cdot m⁻² \cdot s⁻¹. In the case of the ammonia/lithium 376 nitrate fluid mixture, the ammonia concentration ranged from 0.49 to 0.54, while in the 377 378 case of the ammonia/(lithium nitrate + water) fluid mixture the ammonia concentration 379 was between 0.42 and 0.46. The water content in the absorbent remained at 20%. The 380 results are in concordance with the results reported by other authors, Hsieh and Lin 381 [31], Yan and Lin [33], Claesson [33], Táboas et al. [1], who concluded that vapour 382 quality followed by mass flux have the most significant effect on frictional pressure 383 drop. For the selected experimental conditions, it is noteworthy that operating pressure, 384 ammonia mass fraction and heat flux have no significant effect on frictional pressure 385 drop.



Figure 3. Two-phase frictional pressure drop versus mean vapour quality at different mass fluxes
and heat fluxes for: (a) NH₃/LiNO₃ and (b) NH₃/(LiNO₃+H₂O)

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Figure 4 shows the friction factor versus mean vapour quality at solution mass flux of 50, 75 and 100 kg·s⁻¹·m⁻². The results indicate that the friction factor significantly decreases when the solution mass flux is increased. Furthermore, the friction factor decreases significantly with an increase in mean vapour quality for vapour qualities lower than 0.15.



395

Figure 4. Two-phase friction factor versus mean vapour quality at different mass fluxes and heat
fluxes for: (a) NH₃/LiNO₃ and (b) NH₃/(LiNO₃+H₂O)

398 **4.2 Flow boiling heat transfer**

399 4.2.1 Flow boiling of ammonia/lithium nitrate fluid mixture

Figure 5 shows the influence of mean vapour quality on the flow boiling heat transfer coefficient when using the ammonia/lithium nitrate fluid mixture at different mass fluxes and three heat fluxes, namely, (a) 5 kW·m⁻², (b) 10 kW·m⁻² and (c) 20 kW·m⁻². The experiments were carried out with an ammonia mass fraction of 0.50 and at an average operating pressure of 15 bar. The results show that the flow boiling heat transfer coefficient is greatly dependent on the solution mass flux at all selected operating conditions. This indicates the importance of the effects of convective boiling in the experimental data. The results are well in agreement with the data of Táboas et al. [1] obtained for ammonia/water and the same plate heat exchanger (PHE) and also with the data reported by Yan and Lin [32] for R-134a in a PHE.

411



Figure 5. Flow boiling heat transfer coefficient versus mean vapour quality for NH₃/LiNO₃ at
different mass fluxes and heat flux of: (a) 5 kW·m⁻², (b) 10 kW·m⁻² and (c) 20 kW·m⁻²

It is noteworthy that the experimental results presented in this work, like those of Táboas et al. [1] and Yan and Lin [32], were obtained with a mass flux considerably higher than that used by other authors, who suggested that flow boiling heat transfer in PHE was dominated by nucleation effects (Hsieh and Lin [31], Longo and Gasparella [34], Palm and Claesson [35], Zacarías et al. [25]).

422 Figure 6 shows the influence of heat flux on the flow boiling coefficient for NH₃/LiNO₃ at three values of mass flux, namely, (a) 100 kg·m⁻², (b) 75 kg·m⁻² and (c) 50 kg·m⁻²s. 423 424 As is observed, the heat flux has a limited effect compared to that of the mass flux. In 425 the case of a fluid mixture, the presence of the second component in the liquid phase 426 causes a resistance to the boiling process which results in the boiling heat transfer 427 coefficient being less dependent on heat flux. Nevertheless, the experimental works 428 available in the literature on flow boiling heat transfer coefficient in PHEs with pure 429 fluids, which report correlations fitted to their own data (Hsieh and Lin [31], Yan and 430 Lin [32], Han et al. [36], Djordjevic and Kabelac [37]), always show a lower heat flux 431 exponent than the typical value of 0.7 used for smooth tubes.



434

435 Figure 6. Flow boiling heat transfer coefficient versus mean vapour quality for $NH_3/LiNO_3$ at 436 different heat fluxes and mass flux of: (a) 100 kg·m⁻²s⁻¹, (b) 75 kg·m⁻²s⁻¹ and (c) 50 kg·m⁻²s⁻¹

437

Therefore, the flow boiling process of ammonia/lithium nitrate in the plate heat exchanger tested comprises both convective and nucleate effects. Convective effects are dominant for low heat fluxes and high solution mass fluxes, nucleate effects are present for low solution mass fluxes. It is also worthwhile mentioning that even in the zone of apparent nucleation (high heat flux and low mass flux) the mass flux clearly has an influence on the flow boiling heat transfer coefficient.

444 The heat transfer coefficients reported by Zacarías et al. [25] with $NH_3/LiNO_3$ and a 445 similar plate heat exchanger, are considerably lower than those in the data obtained in 446 the present work. However, it is not possible to make a direct quantitative comparison 447 because, although the authors present experimental results with a heat flux close to 5 448 kW/m^2 , the experiments were performed with a maximum mass flux of 19.8 kg·m⁻²s⁻¹ 449 and a maximum vapour quality of 3%.

450 4.2.2 Comparison of the ammonia/lithium nitrate with other451 ammonia based fluid mixtures

Figure 7 shows a comparison of the flow boiling heat transfer coefficient between ammonia/lithium nitrate and ammonia/(lithium nitrate + water) fluid mixtures at: (a) mass flux of 100 kg·m⁻²·s⁻¹ and heat flux of 20 kW·m⁻² and (b) mass flux of 50 kg·m⁻ 2 ·s⁻¹ and heat flux of 10 kW·m⁻². It is seen that the flow boiling heat transfer coefficient is somewhat higher in the case of the ternary fluid mixture. This is attributed to the lower viscosity and higher thermal conductivity of the ternary fluid mixture which improve the boiling coefficient.





Figure 7. Comparison between the flow boiling heat transfer coefficients of the binary and ternary
fluid mixtures at: (a) heat flux of 20 kW·m⁻² and mass flux of 100 kg·m⁻²·s⁻¹; (b) heat flux of 10
kW·m⁻² and mass flux of 50 kg·m⁻²·s⁻¹

465 Figure 8 compares the flow boiling heat transfer coefficients obtained in the present work at mass fluxes of 100 and 75 kg·m⁻² s⁻¹ and a heat flux of 20 kW·m⁻², with those 466 467 obtained by Táboas et al. [1] for ammonia/water fluid mixture at similar operating 468 conditions. The experimental data for ammonia/water was obtained with a mass fraction of 0.42, an operating pressure of 15 bar, mass fluxes of 100, 70 kg \cdot m⁻² \cdot s⁻¹ and a heat 469 flux of 20 kW \cdot m⁻². The flow boiling heat transfer coefficients obtained in the present 470 471 work are at least 35% lower than those presented by Táboas et al. [1]. Again, the higher viscosity and the lower thermal conductivity of the ammonia/lithium nitrate fluid 472 473 mixture are considered responsible for the lower values of the flow boiling heat transfer 474 coefficient obtained.





476 Figure 8. Comparison of the flow boiling heat transfer coefficient with NH₃/LiNO₃ fluid mixture
477 (present work) with data obtained by Táboas et al. [1] for NH₃/H₂O

478

479 Besides, Táboas et al. [1] who carried out their work in the same experimental set-up 480 using ammonia/water as a working pair, at mass fluxes higher than 70 kg·m⁻²·s⁻¹ 481 observed that the flow boiling heat transfer coefficient is not dependent on the heat flux 482 but is clearly dependent on the vapour quality. This region of operating conditions was 483 considered a purely convective region, since heat flux has no effect on the boiling heat 484 transfer coefficient while mass flux and vapour quality have. The experimental data of 485 the present work do not show the pure convective boiling region that Táboas et al. [1] 486 observed in their data at high mass fluxes. The higher viscosity of the binary fluid 487 mixture used in this study, reduces the turbulence in the heat exchanger at the same 488 mass flux and could delay the presence of pure convective boiling effects in the heat 489 exchanger.

4.3 Modelling of two-phase frictional pressure drop and 490 flow boiling heat transfer coefficient

4.3.1 Modelling of two-phase frictional pressure drop in the 492 493 plate heat exchanger

494 From the single-phase friction factor calculated with Eq. (26) and developed in section 495 3.4, the two-phase frictional pressure drop was correlated using the Lockhart-Martinelli 496 approach and the Chisholm parameter C (Eqs. (27)-(29)).

497
$$\varphi_L^2 = \frac{\left(\frac{\Delta P}{\Delta L}\right)_{TP}}{\left(\frac{\Delta P}{\Delta L}\right)_L}$$
(27)

498
$$\varphi_L^2 = 1 + \frac{c}{x_{LM}} + \frac{1}{x_{LM}^2}$$
 (28)

499

491

$$X_{LM}^{2} = \frac{\left(\Delta P / \Delta L\right)_{L}}{\left(\Delta P / \Delta L\right)_{V}}$$
(29)

501 According to Táboas et al. [38], a value equal to 3 was obtained for the parameter C in 502 experiments performed with ammonia/water fluid mixture and the plate heat exchanger 503 employed later in the present work. Figure 9 shows the results obtained using the same

value of parameter C and, as seen, this same value also correlates reasonably well with the two-phase frictional pressure drop. Figure 9a shows the comparison between the measured and the predicted two-phase frictional pressure drop for the two fluid mixtures studied in the present work, namely ammonia/lithium nitrate and ammonia/(lithium nitrate + water). Figure 9b incorporates the set of data obtained in the present work and that obtained by Táboas et al. [38] for ammonia/water.





511 Figure 9. Two-phase frictional pressure drop for: (a) binary and ternary fluid mixtures, (b) binary,
512 ternary and NH₃/H₂O fluid mixtures

513

The experimental two-phase frictional pressure drop values obtained for the ammonia/lithium nitrate fluid mixture have also been compared with the experimental data reported by Zacarías et al. [25] who presented the two-phase frictional pressure drop against the kinetic energy per unit volume computed by the homogeneous model, Eq. (30).

$$520 \qquad \frac{KE}{V} = \frac{G^2}{2 \cdot \rho_m} \tag{30}$$

521 Figure 10 shows that the slopes of both sets of data and operating conditions are522 different.



523

524 Figure 10. Frictional pressure drop per unit length versus the kinetic energy per unit volume.
525 Comparison between the present work and Zacarías et al. [25].

526 4.3.2 Modelling of the flow boiling heat transfer in plate heat527 exchangers.

528 Táboas et al. [38] showed that the flow boiling heat transfer coefficient for the 529 ammonia/water fluid mixture could be calculated by the following set of equations:

530 If
$$u_V < -111.88 \cdot u_L + 11.848$$
 $h_{TP} = h_{nb} = 5 \cdot Bo^{0.15} \cdot h_{LO}$

531 If
$$u_V > -111.88 \cdot u_L + 11.848$$
 h_{TP} higher of
$$\begin{cases} h_{nb} = 5 \cdot Bo^{0.15} \cdot h_{LO} \\ h_{cb} = \left(1 + \frac{3}{X_{LM}} + \frac{1}{X^2}\right)^{0.2} \cdot h_{LO} \end{cases}$$
(31)

These relationships depend on easy-to-calculate properties of binary fluid mixtures used in absorption refrigeration systems. In addition, the convective enhancement factor was characterized by the Chisholm equation, so the convective boiling term can be calculated from the experimental data of a two-phase frictional pressure drop, which makes use of the single-phase friction factor equation.

537 Figure 11 shows the predictions of the flow boiling heat transfer coefficient for the

ammonia/lithium nitrate fluid mixture using the Eq. (31) proposed by Táboas et al. [38]

539 for ammonia/water mixtures in plate heat exchangers. The predictions are presented at different mass fluxes and three values of heat flux, namely, 5, 10 and 20 kW·m⁻². As 540 541 observed, the equations predict the trend and data reasonably well.



543

544

545 Figure 11: Comparison between the values predicted by the correlation of Táboas et al. [38] and 546 experimental data of the flow boiling heat transfer coefficient for NH₃/LiNO₃ at different mass fluxes and heat flux of: (a) 5 kW·m⁻², (b) 10 kW·m⁻² and (c) 20 kW·m⁻² 547

548

549 Figure 12 shows a similar comparison to Figure 11 for the binary fluid mixture 550 NH₃/LiNO₃ but at different heat fluxes and three values of mass flux, namely, 50, 75 and 100 kg \cdot m⁻² \cdot s⁻¹. As shown in this figure, the exponent chosen for the boiling number 551 552 was able to predict the heat transfer coefficient trend reasonably well.



555

556

557 Figure 12: Comparison between the predicted values by the correlation of Táboas et al. [38] and 558 experimental data of the flow boiling heat transfer coefficient for $NH_3/LiNO_3$ at different heat 559 fluxes and mass flux of: (a) 100 kg·m⁻²·s⁻¹, (b) 75 kg·m⁻²·s⁻¹ and (c) 50 kg·m⁻²·s⁻¹

560

561 Figure 13 shows a comparison between the experimental flow boiling heat transfer 562 coefficient and the predicted one for both ammonia/lithium nitrate and 563 ammonia/(lithium nitrate + water) fluid mixtures using Eq. (27). Although the 564 experimental data presented for the ternary fluid mixture are scarce, the comparison 565 shows that most of the experimental values are within a maximum deviation from the 566 predicted values of 20%.



569Figure 13: Comparison between the experimental flow boiling heat transfer coefficient and the570predicted one using Eq. (27) for: (a) NH₃/LiNO₃ and (b) NH₃/(LiNO₃+H₂O)

571

572 **5 Conclusions**

573 Experimental results on flow boiling heat transfer and two-phase pressure drop of 574 ammonia/lithium nitrate and ammonia/(lithium nitrate + water) fluid mixtures in a plate 575 heat exchanger are presented. The experiments were carried out adhering to the 576 operating conditions of absorption equipment for HVAC applications. The effect of 577 mass flux and heat flux on the flow boiling heat transfer coefficient and in the two-578 phase frictional pressure drop were analysed at vapour qualities ranging from 0 to 0.20. The solution mass flux was varied between 50 and 100 kg \cdot s⁻¹·m⁻² and the heat flux 579 between 7.5 and 20 kW \cdot m⁻². 580

581 The major conclusions are summarized in the following paragraph:

The solution mass flux is the parameter that most influences the flow boiling
 coefficient at the applied operating conditions. This is characteristic of convective
 boiling. This trend, observed mainly for low heat fluxes and high solution mass

fluxes, showed that the boiling coefficient improves when the solution mass flux isincreased.

For the binary fluid mixture of ammonia/lithium nitrate the increase of the flow
boiling coefficient, is more significant at low values of heat flux, when the mass
flux is increased. However, with the ternary fluid mixture of ammonia/(lithium
nitrate + water) the improvement of the flow boiling coefficient, achieved by
increasing the solution mass flux, is similar for all the values considered for heat
flux.

For the ternary fluid mixture of ammonia/(lithium nitrate + water) the flow boiling
 coefficient was not influenced by the heat flux variation. It indicated a
 predominance of convective boiling at the operating conditions applied, and as in
 the case of the binary fluid mixture, the effect of vapour quality on the flow boiling
 coefficient was almost insignificant.

The frictional two-phase pressure drop in the plate heat exchanger (PHE) increased almost linearly with the mean vapour quality. The vapour quality followed by the mass flux had the most influence on the frictional two-phase pressure drop in the PHE. The influence of the heat flux on the frictional pressure drop was almost insignificant.

The Chisholm correlation with the constant C=3 obtained by Táboas et al. [38], can
 satisfactorily predict the two-phase pressure drop in a plate heat exchanger (PHE).

The correlation of Táboas et al. [38] initially proposed for ammonia/water in the
 PHE, can satisfactorily predict the flow boiling heat transfer coefficient for
 ammonia/lithium nitrate and ammonia/(lithium nitrate + water) fluid mixtures.

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611	
612	

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729 Figures caption

- 730 Figure 1: Schematic diagram of the experimental test bench
- Figure 2: Main parameters of the plate heat exchanger
- Figure 3: Two-phase pressure drop versus mean vapour quality at different mass fluxes
- and heat fluxes for: (a) $NH_3/LiNO_3$ and (b) $NH_3/(LiNO_3+H_2O)$
- Figure 4: Two-phase friction factor versus mean vapour quality at different mass fluxes
- and heat fluxes for: (a) $NH_3/LiNO_3$ and (b) $NH_3/(LiNO_3+H_2O)$
- 736 Figure 5: Flow boiling heat transfer coefficient versus mean vapour quality for
- 737 NH₃/LiNO₃ at different mass fluxes and heat flux of: (a) 5 kW·m⁻², (b) 10 kW·m⁻² and 738 (c) 20 kW·m⁻²
- Figure 6: Flow boiling heat transfer coefficient versus mean vapour quality for NH₃/LiNO₃ at different heat fluxes and mass flux of: (a) 100 kg·m⁻²s⁻¹, (b) 75 kg·m⁻²s⁻¹
- 741 and (c) 50 kg \cdot m⁻²s⁻¹
- Figure 7: Comparison between the flow boiling heat transfer coefficients of the binary
- and ternary fluid mixtures at: (a) heat flux of 20 kW \cdot m⁻² and mass flux of 100 kg \cdot m⁻² \cdot s⁻²

744 ¹; (b) heat flux of 10 kW \cdot m⁻² and mass flux of 50 kg \cdot m⁻² \cdot s⁻¹

- Figure 8: Comparison of the flow boiling heat transfer coefficient with $NH_3/LiNO_3$ fluid mixture (present work) with data obtained by Táboas et al. [1] for NH_3/H_2O
- 747 Figure 9: Two-phase frictional pressure drop for: (a) binary and ternary fluid mixtures,
- 748 (b) binary, ternary and NH_3/H_2O fluid mixtures
- Figure 10: Frictional pressure drop per unit length versus the kinetic energy per unit
- volume. Comparison between the present work and Zacarías et al. [25].
- Figure 11: Comparison between the predicted values by the correlation of Táboas et al.
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- at different mass fluxes and heat flux of: (a) $5 \text{ kW} \cdot \text{m}^{-2}$, (b) $10 \text{ kW} \cdot \text{m}^{-2}$ and (c) 20
- 754 $kW \cdot m^{-2}$
- Figure 12: Comparison between the predicted values by the correlation of Táboas et al.
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- at different heat fluxes and mass flux of: (a) $100 \text{ kg} \cdot \text{m}^{-2} \cdot \text{s}^{-1}$, (b) 75 kg $\cdot \text{m}^{-2} \cdot \text{s}^{-1}$ and (c) 50
- 758 $kg \cdot m^{-2} \cdot s^{-1}$
- Figure 13: Comparison between the experimental flow boiling heat transfer coefficient
- and the predicted one using Eq. (27) for: (a) $NH_3/LiNO_3$ and (b) $NH_3/(LiNO_3+H_2O)$
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- 762

763 **Tables caption**

- 764 Table 1: Accuracy of measured variables and combined uncertainty of the calculated765 parameters
- 766 Table 2: Operating conditions
- 767