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Transient Performance of a Liquid Desiccant Solar Regenerator

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ABSTRACT

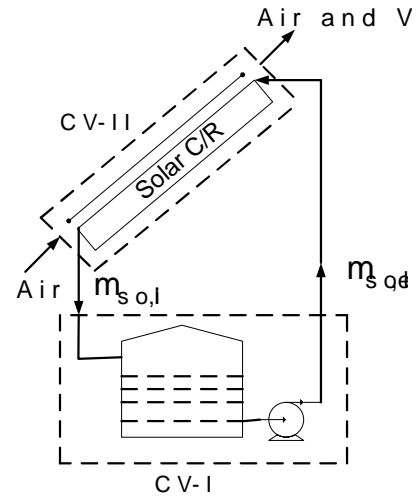
A solar liquid desiccant cooling system uses renewable energy and natural refrigerant, which is attractive. Its main components are air dehumidifier, solar liquid desiccant cooling system, and solar regenerator. In this cooling system, desiccant absorbs water from air in the dehumidifier and is regenerated in the solar regenerator. This makes regeneration of a liquid desiccant system more energy efficient. The regeneration energy can be obtained from sun with the type solar collector cum regenerator. A solar regenerator cum regenerator during sunshine hours. Detailed analysis of the performance of a liquid desiccant cooling system and experimental performance of a liquid desiccant cooling system were carried out. A solar collector cum regenerator was fabricated using corrugated sheet metal, thermal insulation, and supporting frame. The solar collector cum regenerator was mounted on a metal support structure. The corrugated absorber was coated with iron oxide to minimize corrosion.

This paper presents the results of an experimental study on the performance of a solar liquid desiccant cooling system. The effect of desiccant concentration, mass of water evaporated, and mean daily solar radiation on the performance of the system was studied. The results show that the mass of water evaporated and the mean daily solar radiation increase with the increase in desiccant concentration. The typical results were obtained during the month of May. The mass of water evaporated was 17 kg and 36% of the total mass of water evaporated. The experimental procedure and analysis are useful in designing solar components of open cycle liquid desiccant cooling systems.

Keywords: Solar energy, liquid desiccant, solar collector cum regenerator, solar

1. INTRODUCTION

Solar liquid desiccant cooling system mainly consists of air dehumidifier, solar liquid desiccant cooling system, and solar regenerator. In this cooling system, a concentrated solar radiation is used to regenerate the desiccant in an air dehumidifier. The desiccant is then used to dehumidify the airstream in an air dehumidifier. The desiccant is regenerated by evaporating the moisture ab-



a. Experimental system

b. Schematic diagram

Figure Experimental system and its schematic diagram

3.1 Mass of Water Evaporated

During experimentation, solar regeneration of aqueous solution of lithium chloride is carried out by continuously circulating the liquid desiccant over the absorber surface. Evaporation of water from the solution occurs in the solar regenerator as the solution flows over the absorber in contact with the atmospheric air trapped between the solution and the glazing gap by natural convection. The outlet liquid desiccant from the solar regenerator is allowed to mix with the solution in the solution tank as shown in Figure 1. The mass of water evaporated from the solution in the tank at the start of experimentation and the mass of solution in the tank after 30 minute time interval; about 10% of the solution and solution pipelines. Applying conservation of mass principle to the two control volumes gives the mass of water evaporated as

$$m_v \dot{\delta} = m_{s, o, i} \dot{\delta} - m_{s, o, f} \dot{\delta} \quad (2)$$

Measuring the final mass of solution contained in the pipelines, absorber plate and solution tank, it is found from mass of desiccant in the solution tank at the start and at the end of the solar regeneration processes, i.e., the final and the initial mass of solution in the tank after regeneration are equal.

$$m_{d, i} \dot{\delta} = m_{d, f} \dot{\delta} \quad (3)$$

Thus, the final mass of solution in the pipelines, absorber plate and solution tank can be expressed in terms of the initial mass of solution in the tank as

$$m_{s, o, f} \dot{\delta} = m_{s, o, i} \dot{\delta} \quad (4)$$

Substituting equation (4) into equation (2) and rearranging gives the mass of water evaporated as

$$m_v \dot{\delta} = m_{s, o, i} \dot{\delta} \left(\frac{\partial x}{\partial x} - \frac{1}{\partial x} \right) \quad (5)$$

From the solar regeneration experiment conducted, appreciable change in concentration of the solution is observed only after heating of the solution. The change in concentration of the

between inlet and outlet of the solar C/R is negligible. This is due to small solar collection area of the experimental setup was found to be insignificant at other phase transition and present time as the initial and the final thermodynamic states of the liquid from the solution between this time intervals, "t, would be

$$m_{v, \delta D} \delta = m_{s, o, t_0} \frac{\partial x_{o, \delta}}{\partial x_{o, \delta}} \delta - \frac{1}{\delta} \frac{\partial \rho}{\partial x_{o, \delta}} \delta \quad (6)$$

Between the time step, the mass of water evaporated will be

$$m_{v, \delta D} \delta = m_{s, o, t_0} \frac{\partial x_{o, \delta}}{\partial x_{o, \delta}} \delta - \frac{1}{\delta} \frac{\partial \rho}{\partial x_{o, \delta}} \delta \quad (7)$$

The expression to the left of the equals sign in (7) represents the mass of desiccant in the tank at the corresponding times, and is always constant throughout the solar radiation. The expressions by the right side of the equals sign in (7) through (8) are used to determine the transient mass of water evaporated from the solution. (7) through (8) can be simplified to

$$m_{v, \delta D} \delta = m_d \frac{\partial x_{o, \delta}}{\partial x_{o, \delta}} \delta - \frac{1}{\delta} \frac{\partial \rho}{\partial x_{o, \delta}} \delta \quad \text{where } \delta = 1, 2, \dots \quad (8)$$

Equation (8) gives the mass of water evaporated from the tank at the beginning of the solar radiation and after some time interval, the mass of water evaporated per unit area of the absorber plate

$$m' \delta = \frac{m_{v, \delta D}}{A_{a, b}} \quad (9)$$

3.2 Desiccant Concentration

Conde (2004) has reported empirical concentration as a function of concentration of temperature and density of water, for lithium chloride and calcium chloride. The expressions are rearranged to give the concentration of lithium chloride and calcium chloride in the solution measured solution density and temperature:

$$0.100 \frac{\partial x_{o, \delta}}{\partial x_{o, \delta}} \delta - 0.303 \frac{\partial x_{o, \delta}}{\partial x_{o, \delta}} \delta^2 + 0.540 \frac{\partial x_{o, \delta}}{\partial x_{o, \delta}} \delta^3 - \frac{\partial r_{s, o, l}}{\partial r_{H_2O}} \frac{\partial (x_{s, o, l}, \delta)}{\partial (r_{s, o, l}, \delta)} \delta = 0 \quad (10)$$

$$0.105 \frac{\partial x_{o, \delta}}{\partial x_{o, \delta}} \delta^3 - 0.43 \frac{\partial x_{o, \delta}}{\partial x_{o, \delta}} \delta^2 + 0.836 \frac{\partial x_{o, \delta}}{\partial x_{o, \delta}} \delta - \frac{\partial r_{s, o, l}}{\partial r_{H_2O}} \frac{\partial (x_{s, o, l}, \delta)}{\partial (r_{s, o, l}, \delta)} \delta = 0 \quad (11)$$

$$\frac{\partial r_{H_2O}}{\partial r_{s, o, l}} \frac{\partial (x_{s, o, l}, \delta)}{\partial (r_{s, o, l}, \delta)} = 0.392 \frac{\partial x_{o, \delta}}{\partial x_{o, \delta}} \delta + 1.9937 \frac{\partial x_{o, \delta}}{\partial x_{o, \delta}} \delta^{1/2} + 1.0985 \frac{\partial x_{o, \delta}}{\partial x_{o, \delta}} \delta^{2/3} - 0.5094 \frac{\partial x_{o, \delta}}{\partial x_{o, \delta}} \delta^{5/3} - 1.7619 \frac{\partial x_{o, \delta}}{\partial x_{o, \delta}} \delta^{1/2} + 4.4244 \frac{\partial x_{o, \delta}}{\partial x_{o, \delta}} \delta^{1/3} - 7.2369 \frac{\partial x_{o, \delta}}{\partial x_{o, \delta}} \delta^{1/3} + 2.602 \frac{\partial x_{o, \delta}}{\partial x_{o, \delta}} \delta^{10/3} \quad (12)$$

3.3 Solar Efficiency

The solar C/R efficiency can be defined as the ratio of the input energy. For a desired effect is the collection of solar energy to evaporate the water. It is given by the total insolation falling on the aperture area of the C/R. It is given by

$$\eta_{C/R} = \frac{Q_{e, va}}{I \cdot A_p \delta} \quad (13)$$

When the pressure of water in the solution approaches the air interface at the solution surface, water migrates to the air. This migration of water molecules to evaporate it. The rate of evaporation is given by equation (14)

4.1.1 Desiccant concentration of the measured solutions during the experiment. As shown in Figure 2, the density of solution was continuously high from 1.193 up to 1.283 g/cm³ and a total increase of 7.5% in weight is a key indicator of evaporation of water from the solution. The concentration of LiCl also obtained is equal to 10% at every hour; for instance the concentration at 9 am was 0.233 and 0.446/kg respectively by total increase in concentration of 10%. The uncertainty in concentration of water calculated is ± 0.01 kg/kg. The concentration of LiCl for the remaining time interval is also indicated in Figure 2. The transient trend of concentration increase is the same as the increase in density with time. This trend is different from the trend reported by Kabeel et al. (2015) which were indeed similar to rate of evaporation of water with time. Hourly concentration of LiCl is also shown in Figure 3; with the peak decrease of 2.13% in concentration of 0.386 to 0.409 in 1 hr and the trend of concentration change of the desiccant with time was same as the rate of evaporation. In (Wang & Yang et al., (2011) reported that for dehumidification of air from the initial humidity ratio of 0.4 to 0.3, respectively. However, Kabeel et al. (2009) has used LiCl concentration which will result in lesser dehumidification of air.

4.1.2 Mass of water and amount of LiCl solution ($m_{s,0}$) and the initial mass of desiccant solution ($m_{d,0}$) were 45.8 and 5.12 kg, respectively. The LiCl concentration (C_{LiCl}) was 0.53 ($m_{LiCl}/m_{s,0}$) and ($m_{d,0}$) between these two quantities ($m_{s,0}$ and $m_{d,0}$) were found to be 4.20 kg and 20 g/m² respectively. Likewise, the evaporation rate of water in regeneration intervals were calculated and the results are shown in Figure 3. The mass of LiCl in the solution and mass of water evaporated were (1.45 kg and 0.25 kg) respectively, volume of solution density desiccant concentration were 1.193 ± 0.01 kg/kg and 1.283 ± 0.01 kg/kg, respectively; the maximum amount of water evaporated was 4%, approximately. The mass of water evaporated over a 24 hr day was 13.06 kg from the atmospheric mass of water evaporated per unit area, 2.16 kg/m² and 0.26 kg/m² respectively. Kabeel et al. (2009) reported peak evaporation rate of LiCl solution evaporated under weather condition of Delhi.

4.1.3 Solar C/R The efficiency of evaporation of water from aqueous solution and air were 27 and 32723.7 kJ/kg, respectively. The average latent heat of evaporation was 2712.5 kJ/kg. Thus, the energy of evaporation simultaneously with the evaporation equation was 4678.3 kJ with a ± 0.01 error. The total energy intercepted by the solar panel (2.6 m x 1.2 m) in this time period was 12651.04 kJ. The coefficient of C/R efficiency in this time period was 0.37 and the average energy efficiency was about 3.9 % time interval. Similarly, the solar C/R efficiency was calculated for the evaporation shown graphically in Figure 3. The energy falling on the aperture area of the solar panel was utilized for evaporating water and over the average efficiency of evaporation,

4.2 Transient Solar C/R Regeneration Performance and Justification. Similar solar regeneration performance and justification was also shown in Figure 3. The initial concentration were 38.4 and 7.7% respectively. From 9 am up to 4 pm, the concentration of LiCl increased to 0.7 kg/kg due to evaporation of 17 kg water. The peak water evaporation area was 0.765 kg/m² and an irradiance of 86.0 W/m² and a peak value of evaporation was reported by Kabeel et al. (2015) for regeneration solution of concentration of 0.135 and C/R efficiency was 36.5% and 9.6% and 4 pm with peak value of 0.69 for this typical time. The daily efficiency of the solar C/R was 4 amount of energy intercepted by the energy utilized for evaporating water were respectively. Regeneration performance of LiCl solution was better than LiCl solution because water in solution is higher than LiCl solution for the same concentration and

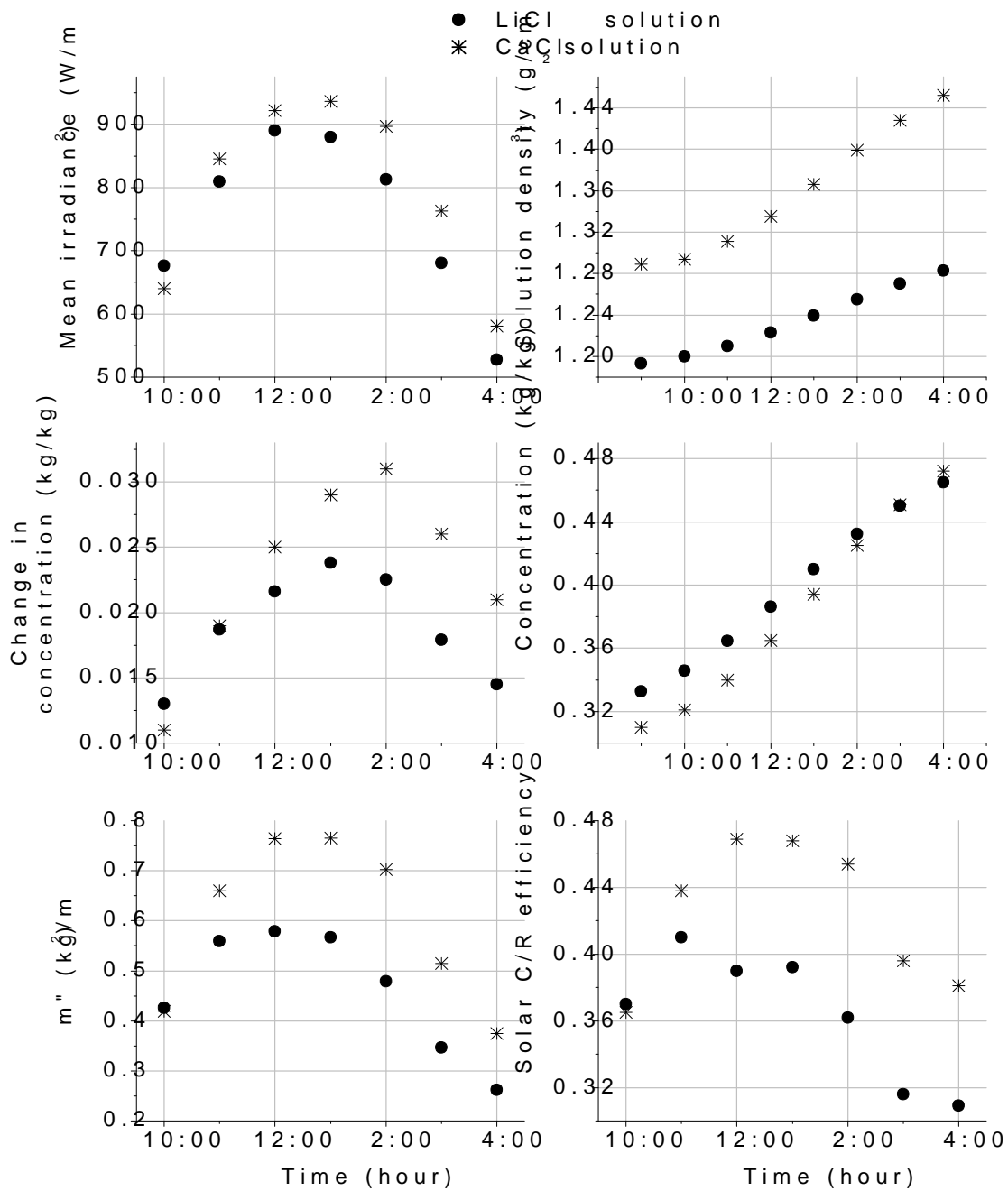


Figure 10. Transient performance of a solar C/R on regeneration of LiCl and CaCl2 solutions

4.3 Overall Transient Performance of Solar C/R

The transient performance of glazed and unglazed solar C/R were evaluated for regeneration of LiCl and CaCl2 solutions over hot & humid, hot & dry, and moderate temperature conditions in Delhi. And it was found that the mass of water evaporated from the liquid solution is proportional to the mean solar irradiance and inversely proportional to the concentration of the solution. The trend of the mass of water evaporated was the same as the irradiance. The solar C/R efficiency is inversely proportional to the concentration of the solution and its trend is related to the evaporation of water as the concentration is increasing and it is same as the trend of solution density. Therefore, the same

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