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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. Its main components are air dehumidifier, solar liquid desiccant cooler, this cooling coil, desiccant absorbs water from air in the dehumidifier in the solar regenerator. This makes regeneration of a liquid desiccant more effective than conventional methods that require thermal energy. The heat can be obtained from sun with the help of a solar collector cum regenerator that enables storage of energy in the form of regenerator during non-shine hours. Detailed analysis of the performance characteristics of the system with a liquid desiccant air conditioning system and experimental performance of the system under actual weather conditions. A solar collector cum regenerator effective solar collector was made of corrugated stainless steel thermocol insulation board supporting a metal mesh. The solar collector cum regenerator was mounted on a metal stand so as to minimize corrosion.

This paper presents transient performance characteristics of increase in desiccant concentration, mass of water evaporated and mean daily usage of desiccant cum regenerator. LiCl and CaCl<sub>2</sub> were found to be 0.073313 & 17 kg and 36 & 43%, respectively. Typical results were obtained for different days above averaged over the month of May. The mass of water evaporated was derived by applying conservation of mass in the experimental procedure and it is very 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 system consists of air dehumidifier, solar liquid desiccant evaporative cooling system, a concentrator and a desorber. An air stream in an air dehumidifier flows from the solar regenerator. In the regenerator, diluted desiccant solution is regenerated by evaporating the moisture absorbed

dehumidifier using solar energy the possible solar liquid desiccant regeneration system was introduced in Chapter 4 and produced strong desiccant solution by evaporating water from dilute desiccant solution. The energy converter and simultaneous mass & heat transfer device. An open type solar liquid desiccant system that determines the cooling performance per kilogram of water absorbed in the dehumidifier, the same amount must be determined. The common desiccant in the system because of its excellent desiccation properties  $\text{CaO}_2\text{H}_2\text{O}$  solution is also used in the experiments since the nature of solar energy is intermittent, determining the transient performance of the experiments is crucial.

Several theoretical studies have been carried out on solar regeneration of liquid open/closed solar C/R. These papers describe methods of measuring the performance of collectors and identifying parameters that are used to indicate the performance of the system. The mass of water evaporated and energy rate ( $\text{W/m}^2\text{K}$ ) of wastewater evaporated (Peng & H. Kaushik et al., 1985; Kumar & Devidas, 1988); efficiency of water vaporized (Matta & 1988); temperature, evaporation rate, and concentration and film thickness after 0.25 temperature, concentration and evaporation rate,  $r_{\text{evap}} = 0.3(\text{H}_{\text{av}})^{0.75}(T_{\text{in}} - T_{\text{out}})^{0.5}$ ; mass transfer coefficient, evaporation rate, regeneration rate,  $k_{\text{m}} = 0.0005$ ; separation rate (Alizadeh & Samadi, 2002); separation rate  $r_{\text{separation}} = 0.0005k_{\text{m}}(T_{\text{in}} - T_{\text{out}})$ ; mass transfer rate, and friction factor  $f = 0.01993$  (Utong & Yang, 2010). One of the most common performance measures is the mass of water evaporated per unit area ( $\text{m}^2\text{h}^{-1}$ ) based on the mass of water evaporated. Edström & Åberg (2010) used mass balance of the ventilation inlet and the regenerator to estimate rate of evaporation of water for a case. However, even distillation scavenging may be (Utong & Yang, 2010).

In theoretical studies of operation of a desiccant system, conservation of solution mass with conservation of energy or species streams by partial differential equations were combined to develop analytical models (Kutman, 1988; Gandomi & Kaudinya, 1989; Kutman, 1989; Gandomi & Kaudinya, 1989; Alizadeh & Samadi, 2002; Katejane & Kumar, 2008), simplified (Gandomi & Kaudinya, 1983; Fagbenle & Karayannidis, 1988; Alizadeh & Samadi, 2002; Kaushik & Kaudinya, 1989). A combination of conservation of mass and conservation of energy equations were also used (Johansen & Grossqvist et al., 1992 and Utong & Yang, 2010). Various assumptions and give only approximate results (1992) for solar C/R. The most complicated components of desiccant system include adjustable performance procedures and methods of protection against research effort towards improving the performance of the system.

Estimating evaporation rate of moist air passing through over the solar C/R is particularly difficult because it is difficult to measure the flow of liquid desiccant and humidity simultaneously using pycnometer digital densitometer since quantity of flowing liquid cannot apply this conceptus flow and density measuring instruments. The variation of the liquid desiccant with temperature and throughout the regeneration chamber is measured in terms of solution stored by circulating the liquid desiccant over the solution apparatus throughout sunstroke. Vaporization of water occurs in the solar C/R throughout the pressure difference between ventilation air and vapor on. The short duration of a

solutions means a number of the solutions taken in time as it is the experimental procedure used to estimate the transient performance of the solar collector.

In the present work, evaporation rate increase of desiccant in the solution efficiency selected to indicate the transient performance of the solar C/R cycle liquid desiccant air cooling system the role of a liquid desiccant is the dehumidifier and rejection of the same in the water desiccant and also cyclically stored inside during storage. Moreover, moisture removal rate from the solution directly dependent on the inlet concentration of which desiccant is the rate of the desiccant from the evaporator. The water from the solar C/R is clearly regenerated transient concentration is much higher than in the work of Mousavi and Kabeel (2005) on regeneration of a solar C/R system performance the solar was reported collector efficiency, figure of merit, effectiveness, and reality, the function of the device is to collect the heat energy by the absorber plate and the solution, and utilize some fraction of energy by the absorber plate and the solution, and utilize some fraction of make the solar C/R energy converter. Hence, the proper emphasis is the combined efficiency of solar collection and evaporation: solar C/R helpful in estimating the net amount of solar energy (area of the solar C/R should be considered as one among the critical performance indicators of the

## 2. EXPERIMENTAL SETUP AND INSTRUMENTATION

The experimental system consists of a solar C/R and a centrifugal pump (P) solar was fabricated using gauge sheet of 2200 mm (length) x 12 cm (width) in a 2600 mm x 1200 mm polypropylene covering height of 85 mm from the absorber plate, solution is mixed in a PVC discharge pipe of length 2800 at the end of the sheet. The absorber is painted with anti-corrosion paint to prevent the desiccant solution from the plate continuously through the solution tank used to store the desiccant volume even though the C/R surface temperatures, pump rates, and other data essential to completely understand the performance of the system. Data of initial mass desiccant in the solution, solution temperature and the solar C/R is sufficient to determine its experimental parameters. The data were collected using RTD and Python help of a data logger every ten seconds regeneration. In addition, a densitometer was used to measure the solution in the tank at every 30 minutes. The RTD was calibrated at ice and hot water calibrating water according to the standard initial height of solution tank and absorber area of the plate were measured using a ruler and a tape rule, pyranometer (Kipp & Zonen) and a densitometer, respectively. The error in estimating solution volume was 0.625 mm error in height measurement tank using uncertainty of the instrument parameters the solar C/R was analyzed using uncertainty analysis (Dunn, 2005).

$$U_R^2 = \frac{N_{\text{dil}}}{Q_{\text{in}}^2} U_x^2 + U_{x,i}^2 \quad (1)$$

## 3. DATA REDUCTION

The initial solution was weighed and the weight was measured and solution taken in a glass of desiccant in the solution and then, the solution was circulated absorber plate for some time until thermal equilibrium was attained on the absorber plate. The absorber plate temperature three solution temperature at the beginning of experimentation, due to early morning radiation.

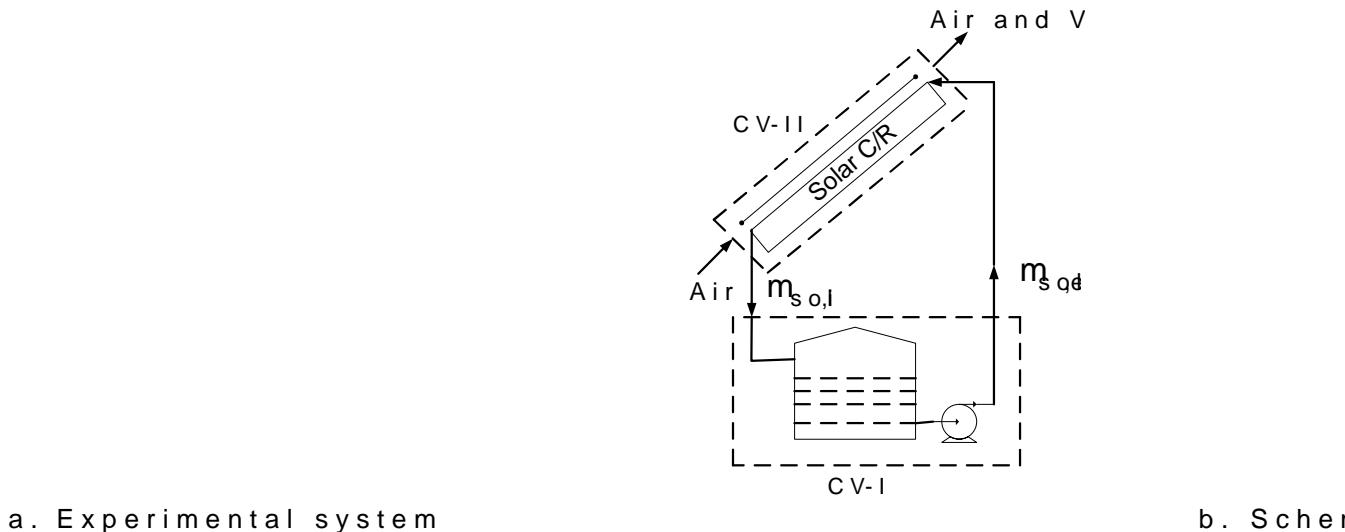


Figure Experimental system and its schematic diagram

### 3.1 Mass Evaporated

During experimentation, solar regeneration of aqueous solution of lithium chloride is carried out by continuously circulating the liquid desiccant over the absorber surface of the pump. Evaporation of few percent occurs in the solar regenerator as the solution passes through the absorber in contact with the atmospheric air trapped between the solution and the glazing glass. The outlet liquid desiccant from the solar regenerator is allowed to mix completely with the inlet liquid desiccant stored in the solution tank as shown in Figure 4(b). The difference in the initial and final mass of solution in the tank at the start of experimentation and the mass of solution remaining on the corrugated plate after 30 minute time interval; about 10% loss in the solution due to evaporation in the solution pipelines. Applying conservation of mass principle to the two containers gives the mass of water evaporated as

$$m_v \delta = m_{\text{solid}} \delta - m_{\text{solid}} \quad (2)$$

Measuring the final mass of solution contained in the pipelines, absorber plate, found from mass of desiccant in the solution bath, the initial and final masses are maintained throughout the solar regeneration processes, i.e., the final and the initial masses after regeneration are equal.

$$m_{d,i} \delta = m_{d,f} \delta @ m_{s,o,i} * \delta_1 \delta = m_{s,o,f} * \delta_f \delta = m_d \quad (3)$$

Thus, the final mass of sand in the tank can be expressed in terms of the initial & final concentrations and the initial sand size in the tank as

$$m_{s,0,f} = m_{s,0,i} \frac{\partial \chi}{\partial x} \quad (4)$$

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

$$m_v \ddot{\alpha} = m_{s,0,i} * \ddot{\alpha}_x * \frac{\ddot{\alpha}_y}{\ddot{\alpha}_x} - \frac{1}{\ddot{\alpha}_x} \quad (5)$$

From the solar regeneration experiment conducted, appreciable change in observed only after heating 6° R the ssoorhutit meir 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 used from time intervals of higher preexisting and present time as the initial and the final thermodynamic states of the liquid from the solution between this time intervals, "t", would be

$$m_{v,D} = m_{s,o,t_0} * \frac{\partial x}{\partial x} \frac{1}{\partial x} \quad (6)$$

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

$$m_{v,D} = m_{s,o,t_0} * \frac{\partial x}{\partial x} \frac{1}{\partial x} \frac{\partial x}{\partial x} \frac{1}{\partial x} \frac{\partial x}{\partial x} \frac{1}{\partial x} \quad (7)$$

The expression to the left of each time step is the mass of desiccant in the absorber at the corresponding times, and is always constant throughout the solar receiver. Expressions by the end of the iteration are used to determine the transient mass of water after solution. (7) thru the assumption of the transient mass of water to be zero.

$$m_{v,D} = m_d * \frac{\partial x}{\partial x} \frac{1}{\partial x} \frac{\partial x}{\partial x} \quad \text{where } d=1, 2, \dots \quad (8)$$

Equation (8) gives the mass of water evaporated during the known initial mass of the solution at beginning of each concentration of the solution and after some time interval. Mass of water evaporated per unit area of the absorber plate

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

### 3.2 Desiccant Concentration

Conde (2004) has reported empirical correlations of concentration of temperature and density of water, for lithium chloride and calcium chloride. 0.05 to 0.5 and 0.05 to 0.6, respectively. These expressions are rearranged to calculate the concentration of lithium chloride and calcium chloride in the solution measured solution density and temperature:

$$0.10078 \frac{\partial x}{\partial x} \frac{3}{\partial x} - 0.303 \frac{\partial x}{\partial x} \frac{2}{\partial x} + 0.540 \frac{\partial x}{\partial x} \frac{1}{\partial x} + 0.066 \frac{\partial r_{sol}}{\partial T_{sol}} \frac{\partial x}{\partial x} \frac{T_{sol}}{\partial r_{H_2O}} = 0 \quad (10)$$

$$0.10564 \frac{\partial x}{\partial x} \frac{3}{\partial x} - 0.430 \frac{\partial x}{\partial x} \frac{2}{\partial x} + 0.836 \frac{\partial x}{\partial x} \frac{1}{\partial x} + 0.044 \frac{\partial r_{sol}}{\partial T_{sol}} \frac{\partial x}{\partial x} \frac{T_{sol}}{\partial r_{H_2O}} = 0 \quad (11)$$

$$\frac{\partial r_{H_2O}}{\partial T_{sol}} = 0.322 \frac{\partial x}{\partial x} + 1.99373 \frac{1}{\partial x}^{1/4} + 1.09852 \frac{1}{\partial x}^{1/3} - 0.50944 \frac{1}{\partial x}^{5/3} \quad (12)$$

$$\frac{\partial r_{sol}}{\partial T_{sol}} = -1.7619 \frac{1}{\partial x}^{1/2} + 4.90024 \frac{1}{\partial x}^{1/3} - 7236928 \frac{1}{\partial x}^{10/3}$$

### 3.3 Solar Efficiency

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

$$\eta_{CR} = \frac{Q_{eva}}{I \cdot A_{ap}} \quad (13)$$

When the pressure of water in the solution approaches the air interface solution surface, water migrates to the air. This migration of water takes place by evaporation. The rate of evaporation is approximated by equation (14).

$$Q_{ev} = m_{vap} h_{fgav} = m_{vap} \Delta H_d$$

(14)

The average latent heat is an average value of the latent heat of vaporization at the previous mass and current time. The latent heat of vaporization of water is constant, so that corresponding to the vaporized pure water, by an average differential enthalpy of (C) (2004) the differential enthalpy of dilution is given by

$$\Delta H_d = \frac{6.90545.350086}{0.8405.0000} \frac{\partial x}{\partial u}^{0.196} \frac{\partial u}{\partial u}^{0.226} \quad (15)$$

$$\Delta H_d = \frac{-9.550 + 3.019700086}{0.8505.0000} \frac{\partial x}{\partial u}^{0.196} \frac{\partial u}{\partial u}^{0.226} \quad (16)$$

#### 4. RESULTS AND DISCUSSION

##### 4.1 Transient Performance of C/R Regenerator Solution

Initial mass of solution and desiccant in the tank can be obtained from the initial density, and initial temperature. The initial values are 1.4 kg/m³ and 20°C.

For this instantaneous irradiance remains constant at 400 W/m² during a time interval of 70% variation of dam and 10 seconds from 900 W/m². The initial solution temperature starts decreasing by the same after that. The irradiance solution temperature graphically depicted in Figure 2 with the hour mean data superimposed on it. In the mean solution temperature of evaporation rate over the period of 1 hour on solution temperature instead of a non-dimensional solar C/R together with the measured solution temperature is shown. This is due to the transient nature of the solution temperature. The transient variation of the mean irradiance data between two times the instantaneous irradiance is shown. The variation of the mean irradiance at the times intervals represent the variation of the mean irradiance at 900 W/m² and 740 W/m². The mean irradiance and standard deviation are 675.9 and 40.4 respectively.

Figure 2 Transient variation of irradiance & solution temperature w

4.1.1 Desiccant concentration of the measured solution during dehumidification is shown the density of solution was increased by time from 1.193 at 9 am to 1.2189 at 4 pm and that of 7.5% in aqueous) is a key indicator of evaporation of water from the solution increases due to sensible heat gain from 10 am to 4 pm (Figure 2). The concentration of LiCl was also obtained at 10 am to 4 pm (Figure 2). The concentration of LiCl at 9 am and 4 pm were 0.46 g/kg respectively. The total increase in water content was 0.016 g/kg. The uncertainty in centrifugal time was calculated to be  $\pm 0.001 \text{ kg/kg}$ . The concentration of LiCl for the remaining time intervals was transiently indicated in Figure 2. The transient trend of concentration increase is the same as the increase in increasing with time. This trend is different from the trend reported by Kaur et al. which were inadesimilarity to rate of evaporation of water with time. Hourly concentration of LiCl is also shown in Figure 3; with an peak decrease around 2.3% in changes of 1.386 to 1.400. The trend of concentration change of the desiccant with time was same as the rate of dehumidification of air. Ullal et al. (2011) reported that for dehumidification of air at the same condition, LiCl 0.4 g/kg respectively. However, Katrejanekarn & Kumar (2009) has used LiCl concentration which will result in lesser dehumidification of air.

4.1.2 Mass of water added in initial LiCl solution ( $m_{\text{sol}}$ ) and the initial mass of desiccant solution were 5.8 and 1.2 kg, respectively. The initial concentration was 0.53 mol/m<sup>3</sup>. Hence, the ratio between these two quantities were found to be 4.20 g/m<sup>3</sup> respectively. Likewise, the evaporation rates at different time intervals were calculated and the results are given in Table 1. The mass of water evaporated were 0.26 g/kg and 0.20 g/kg for initial volume solution density desiccant concentration and  $\pm 0.001 \text{ kg/kg}$  and  $\pm 0.03 \text{ kg/kg}$ , respectively; the maximum and minimum mass of water were 4.0% and 0.0% respectively. The mass of water evaporated over a day was 13.0 kg from the atmosphere and mass of water evaporated per unit area was 0.26 kg/m<sup>2</sup> day. The peak evaporation rate were 0.24 g/kg/m<sup>2</sup> day and 0.20 g/kg/m<sup>2</sup> day respectively. Kaur et al. (2011) reported peak evaporation rate of 0.26 kg/m<sup>2</sup> day under weather condition of Delhi.

4.1.3 Solar C/R efficiency heat of evaporation of water from aqueous solution and air were 270 and 32723.7 kJ/kg, respectively. The average ( $\dot{m}_{\text{air}}$ ) rate of evaporation was 2712.5 kJ/kg. Thus, the energy of water solution simultaneously with the evaporation was 4678.3 kJ with a  $\pm 0.001 \text{ kJ/kg}$  total energy intercepted by the solar collector in this time period was 12651.0  $\pm 0.001 \text{ kJ}$ . The overall C/R efficiency in this time period was 0.37 and the overall daily efficiency was about 3.9%. Similarly, the solar C/R efficiency was calculated at 0.36 from the experimental data shown graphically. During 8 MJ of energy falling on the aperture area of the solar collector was utilized for evaporating water and other losses due to convection, conduction and radiation.

#### 4.2 Transient performance of C/R regeneration of a solution

Similar solar regeneration performance analysis was done at three different concentrations shown in Figure 3. The initial concentrations were 38.4 mol/m<sup>3</sup>, 28.9 mol/m<sup>3</sup> respectively. From 9 am up to 4 pm, the concentration of LiCl decreased to 0.46 g/kg due to evaporation of 17 kg water. The peak water evaporation area was 0.26 g/kg/m<sup>2</sup> at an irradiation rate of 1360.0 W/m<sup>2</sup> and 1.2 and 1.12 m<sup>2</sup>. A peak value of regeneration was reported by Kabeel (2005) of 4.6% efficiency at 1360 W/m<sup>2</sup> and 4 pm with a peak value of 46% for this typical day. Daily efficiency of the solar C/R was 4.6%. The amount of energy intercepted by the solution for evaporating water were respectively. The regeneration performance was better than LiCl solution because water in solution is higher than LiCl solution for the same concentration and

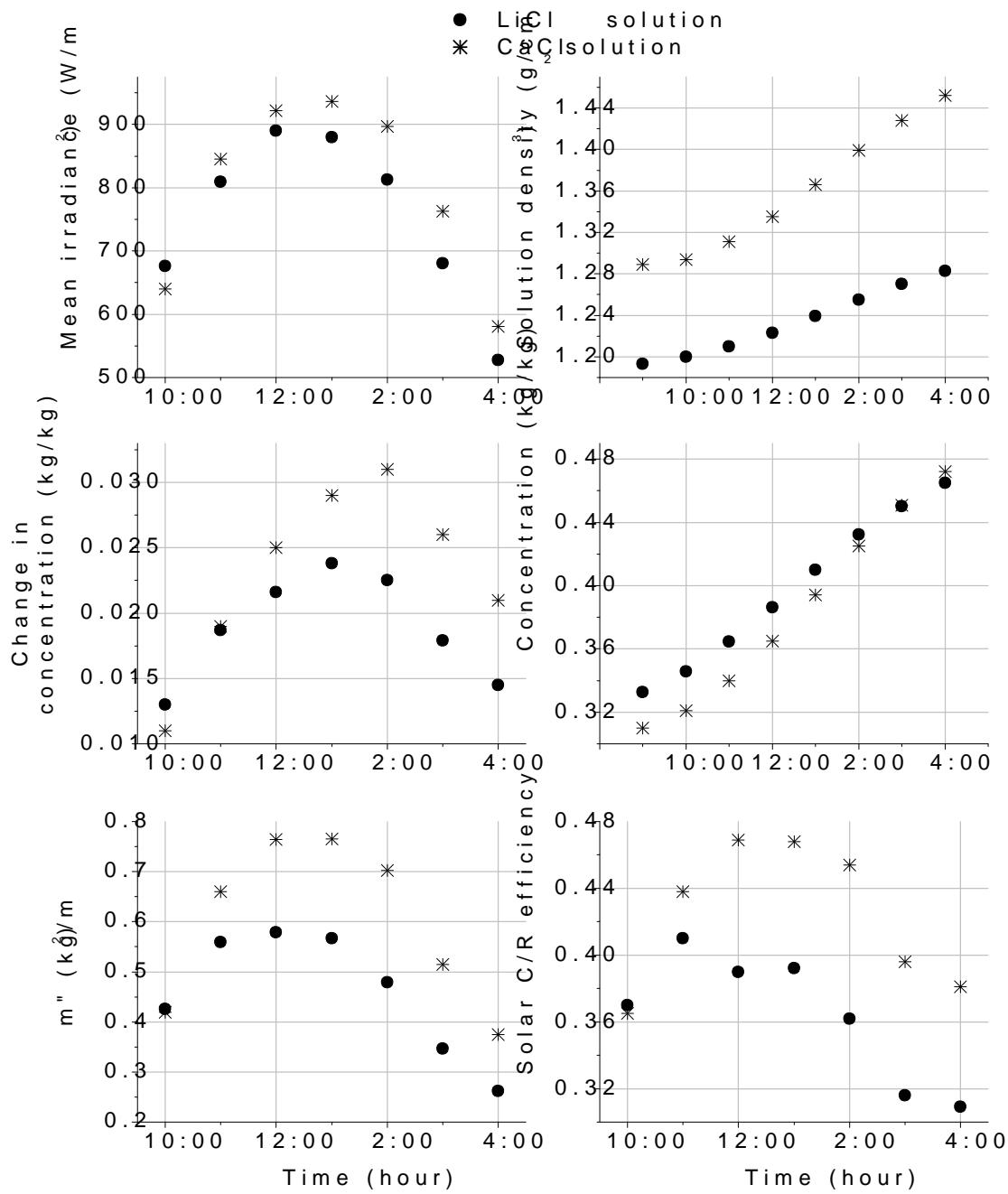


Figure 8 transient performance of a solar C/R on regeneration of LiCl and CaCl<sub>2</sub>

#### 4.3 Overall Transient Performance of C/R

The transient performance of glazed and unglazed solar C/R were evaluated regenerating LiCl and CaCl<sub>2</sub> solutions over hot & humid, hot & dry, and moderate temperatures in Delhi. And it was found that the mass of water evaporated from the liquid solution and inversely proportional to the desiccant in the solution. The trend of evaporation was the same as the irradiance. The solar C/R efficiency is inversely proportional to the evaporation off water and its concentration is increasing and it is same as the trend of solution density. Hence, the same

performance of the solar C/R in May (hot & dry) was better than in October. Evaporation rate of water and mean daily solar C/R efficiency ranges from respectively for LiCl solution of initial concentration about 0.32 and mean CaCl<sub>2</sub> solution these experimental values were 13 to 16.6 and 36.2 to 48.4%, re 0.31 and mean daily irradiance was found to be 7.0 W/m<sup>2</sup> for glazed and unglazed solar to each other. Contamination with dust, foreign bodies and clogging of di for the unglazed regenerator than the glazed one.

## 5. CONCLUSIONS

This paper presents an experimental design of a desiccant type solar C/R. The mass of water was kept constant to observe the performance of the solar C/R evaluated solar regeneration of LiCl and CaCl<sub>2</sub> and solar evaporation in terms of increase in concentration, mass of water vaporized per unit solar energy consumed during solar C/R. Between 9 am and 4 pm for LiCl solution initial concentration 0.3 kg/kg the average evaporation was 18.9 m<sup>2</sup>/s at mean daily solar C/R efficiency of about 39.9.

## NOMENCLATURE

A	area	(m <sup>2</sup> )
D <sub>d</sub>	difference in dilution	(kJ/kg)
I	mean irradiance	(W/m <sup>2</sup> )
m	mass	(kg)
m'	mass of water evaporated per unit time	(kg/m <sup>2</sup> )
Q	total energy over a period	(kJ)
R	arbitrary parameter	(variables)
T	mean temperature	(°C)
U	uncertainty of a measured result	(variables)
Ø	desiccant concentration	(kg/kg)
Ø <sub>r</sub>	mass density	(g/cm <sup>3</sup> )
Ø <sub>t</sub>	reduced temperature at water temperature	(°C)
Ø <sub>Ø=Ø Ø</sub>	reduced temperature at water temperature	(°C)
Ø <sub>hF</sub>	solar collecting efficiency	(%)

### Subscript

a <sub>t</sub>	absorber
a <sub>p</sub>	aperture
C/R;CF	collector cum regenerator
d	desiccant
e <sub>v</sub>	evaporation
s <sub>c</sub>	solution
v	evaporated water

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