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Al-Dadah, Raya; Mahmoud, Saad; Youssef, Peter

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Performance Analysis of Four Bed Adsorption Water Desalination / Refrigeration System, Comparison of AQSOA-Z02 to Silica-gel

Peter G. Youssef*, Saad M. Mahmoud, Raya K. AL-Dadah

Abstract—Although desalination many water techniques have been introduced decades ago, there are still areas around the world suffering from fresh water shortages. The widespread of desalination technologies is limited due to their high energy consumption, cost and adverse environmental impact. Recently, adsorption technology for water desalination has been investigated showing potential of using low temperature waste heat (50-85°C) thus reducing energy consumption and CO₂ emissions. This work mathematically investigates the performance of 4 bed adsorption cycle using two different adsorbents, silica-gel and an advanced zeolite material AQSOA-Z02, produced by Mitsubishi-plastics for fresh water production and cooling. The work studied effects of evaporator and heat source temperatures on water production rate and cooling capacity. Results showed that at low chilled water temperatures below 20°C, AQSOA-Z02 outperform *silica-gel* with water production of 6.2 m³ of water/day and cooling of 53.7 Rton/ tonne of AQSOA-Z02 compared to 3.5 m³ of water/day and 15.0 Rton/tonne of silica-gel. While, at chilled water temperatures above 20°C, AQSOA-Z02 and silica-gel have comparable performance with around 7m³ of water/day and 60 Rton of cooling. Since cooling applications require chilled water temperature less than 20°C, therefore AQSOA-Z02 is more suitable for applications where cooling and fresh water are needed.

Keywords—Adsorption, AQSOA-Z02, Desalination, Refrigeration, Seawater, Silica-gel.

1. INTRODUCTION

Although about 70% of the earth is covered with water, 97% of this water is salty. Also, only 0.3% of the remaining 3% is usable by humans since the remaining fresh water is either underground or in the form of ice covering mountainous regions [1]. Therefore seawater

desalination is required to provide part of human needs for water. Different desalination technologies exist like thermal, membrane and chemical [2] but many of these suffer from consumption; excessive energy adverse environmental impact and high cost [3, 4]. Recently, adsorption technology was shown to provide water desalination with minimum energy consumption [5]. It is capable of producing fresh water with low salinity of 10 ppm, running cost of 0.2 m³ and CO₂ emissions of 0.6 kg/m³ [6]. In this technology, besides water production. cooling can be produced for air conditioning required in many areas around the world [7-11]. A major advantage of adsorption technology is its ability to utilize low grade waste heat sources (50 - 85° C) or solar energy and using environmentally friendly refrigerants leading to lower pollution and cost [8, 12]. A comparison between a lumped parameter model and experimental tests has been presented by Ng et al [13]. The comparison covered wide range of operating conditions for basic and hybrid *silica-gel*/water pair adsorption cycles that are able to produce cooling and desalination. In addition the energetic efficiency and life cycle cost (LCC) of adsorption plants have been compared to other conventional desalination technologies. It was found that adsorption desalination plants require electrical energy of 1.38 kWh/m³ and thermal energy of 38.8 kWh/m³. Comparing to other desalination methods, multi effect desalination (MED) and multi stage flash (MSF) needs 43.21 and 57.14 kWh/m³ of thermal energy respectively. Also reverse osmosis (RO) needs 3.5-5 kWh/m³ of electric energy. Moreover, studies showed that adsorption desalination and cooling cycle has the lowest cost of US \$2.7/ MWh compared to

^{*}P. G. Youssef is a PhD student at University of Birmingham, UK (phone: +44-0121-4143513; fax: +44-0121-4143598; e-mail: pgy348@ bham.ac.uk).

Nomenclature				
с	Uptake (kg.kg ⁻¹)	PR	Performance ratio (-)	
c^*	Equilibrium uptake (kg. kg ⁻¹)	Q_{st}	Isosteric heat of adsorption (kJ/kg)	
c_p	Specific heat at constant pressure (kg. kg ⁻¹ .K ⁻¹)	SCP	Specific cooling power (kW.kg ⁻¹)	
COP	Coefficient of performance (-)	SDWP	Specific daily water production $(m^3 t^{-1} day^{-1})$	
h	Enthalpy (kJ.kg ⁻¹)	Т	Temperature (K)	
Μ	Mass (kg)	X	Salt concentration (ppm)	
т	Mass flow rate (kg.s ⁻¹)	heta	Seawater charging flag (-)	
n	Adsorption/Desorption phase, flag (-)	γ	Brine discharge flag (-)	
OCR	Overall conversion ratio (-)	τ	No of cycles per day (-)	
Р	Pressure (kPa)			
Subscripts				
a	Adsorbent material	hw	Heating Water	
ads	Adsorption	HX	Heat exchanger	
b	Brine	in	inlet	
cond	Condenser	Mads	Master adsorber bed	
CW	Cooling Water	Mdes	Master desorber bed	
D	vapor	out	outlet	
d	Distillate water	S	Seawater	
des	Desorption	Sads	Slave adsorber bed	
evap	Evaporator	Sdes	Slave desorber bed	
f	Liquid	t	Time	

\$4.4/MWh for the combined RO and mechanical chiller or \$3.4/MWh for RO and absorption chiller.

The desalination/refrigeration adsorption system consists of four consecutive processes namely evaporation, adsorption, desorption and condensation. In the evaporator, seawater is evaporated due to the effect of adsorption by the dry adsorbent material while extracting heat from the chilled water passing through the evaporator coil thus producing cooling effect [14]. In the adsorption process, water vapour is adsorbed by the adsorbent material while in the desorption process the water vapour is regenerated by the waste heat. The desorbed water vapour is then condensed in the condenser producing fresh water [8, 15].

Various researchers have investigated the use of adsorption technology for water desalination and cooling using silica gel with various cycle configurations. Wang et al. [12], experimentally investigated the performance of a four bed *silica*- *gel* adsorption desalination system, studying the effects of cycle time, hot, cold and chilled water temperatures on water production rates and cycle coefficient of performance. They found that at chilled water temperature of 12.2° C, a maximum specific daily water production (SDWP) of 4.7 m³/day per tonne of silica gel was obtained using cycle time of 150 seconds, switching time of 40 seconds and low heat source temperature of 85°C. In addition, they have reported that this method of desalination produced potable water without any means of bio-contamination.

Thu et al. [16] studied experimentally the performance of a four bed silica gel adsorption desalination system that operates on either two or four bed configuration. In the two bed operation, each two beds are heated or cooled jointly while in the four bed operation mode, each two beds are heated or cooled sequentially in a master and slave arrangement. Water production and performance ratio were studied at different heat source temperatures with constant heat sink temperature using cycle time ranging from 950 to 2400 seconds for two bed operation or from 480

1920 seconds for four bed operation. 1 to 2 Experimental results showed that in four bed operation at low heat source temperature ($<65^{\circ}$ C), 3 a longer cycle time (1560 sec.) is required for the 4 production of maximum amount of fresh water of 5 6.28 m^3 of water per day per tonne of silica gel. 6 At the maximum heat source temperature of 7 85°C, the two bed configuration produced 8.79 8 m^{3}/day per tonne of silica gel while the four bed 9 master-slave configuration produced 10 m³ of 10 water per day per tonne of silica gel with 11 performance ratio of 0.61. 12

13 Mitra et al. [17], have analyzed a four bed single stage silica gel/water adsorption 14 desalination system that used solar energy for the 15 desorption process. Effects of condenser 16 temperature and cycle time were studied to find 17 the optimum operating conditions for maximum 18 water production and cooling outputs. Simulation 19 results showed that 600-900 seconds is the 20 optimum half cycle time for producing maximum 21 SDWP and specific cooling power (SCP) of 22 2.3m^3 /day and 18 Rton per tonne of silica gel 23 respectively at condenser temperature of 25°C. 24 They concluded that compromise is needed 25 between desalination and cooling capacities as 26 27 COP increases with increasing in cycle time while increasing condenser temperature degrades 28 the cycle performance. 29

Ng et al. [8], have developed a mathematical 30 model for a 4 bed adsorption system using silica 31 gel/water pair to produce both cooling and 32 desalinated fresh water. At different hot and 33 chilled water temperatures, cycle performance 34 35 was analyzed by calculating SCP, SDWP, and overall conversion ratio (OCR). It was found that 36 a silica gel adsorption cycle can produce $8 \text{ m}^3/\text{day}$ 37 and 51.6 Rton per tonne of silica gel when 38 optimized for water production at evaporator 39 temperature of 30°C or 3.8 m³/day and 22 Rton 40 41 per tonne of silica gel at evaporator temperature of 10°C. In addition, the cycle can reach a 42 43 maximum OCR of 1.4.

44 Ng et al. [10], have investigated
45 experimentally and mathematically a 4 bed
46 adsorption cycle that uses *silica-gel*/water pair.
47 The cycle is capable of producing two useful

48 effects which are desalination and cooling using 49 solar energy at low temperature namely (65 to 50 80° C). Measurements indicated that chilled water 51 at 7 to 10° C with a SCP of 25-35 Rton/tonne of 52 silica gel can be produced in addition to a SDWP 53 of 3-5 m³ per tonne of silica gel per day while the 54 OCR ranging from 0.8-1.1.

55 Youssef et al. [18], compared numerically the performance of an adsorption, two bed cycle 56 using AQSOA-Z02/water with the same two bed 57 cycle using Silica-gel/water. Different heating 58 source temperatures and evaporator water inlet 59 60 temperatures were applied to study their effect on cycle performance. It was found that at all heating 61 temperatures, and evaporator water 62 inlet temperatures above $25^{\circ}C$, Silica-gel 63 cycle produces more water and cooling. Silica-gel cycle 64 produces maximum SDWP of 8.4 m³ and 62.4 65 Rton of cooling at 30°C evaporator water inlet 66 temperature. However, as evaporator inlet water 67 temperature decreases, AQSOA-Z02 outperforms 68 *Silica-gel.* At 10°C evaporator inlet water 69 temperature, AOSOA-Z02 cycle produced 5.8 m^3 70 of fresh water per day and 50.1 Rton of cooling 71 while Silica-gel cycle generated only SDWP of 72 2.8 m^3 and SCP of 17.2 Rton. 73

74 AQSOA-Z02 was studied as an adsorbent but in other applications like cooling [19] and 75 dehumidification [20]. Verdi et al. [19] have 76 developed a numerical model for a truck air 77 conditioning system using AQSOA-Z02/ water 78 adsorption system that utilizes waste heat from 79 truck engine temperature $(80-90^{\circ}C).$ 80 at Experimental tests were also performed using a 81 82 lab-based adsorption chiller prototype. The specific cooling power produced at the laboratory 83 reached 600 W/kg of AQSOA-Z02 which is 84 180% higher than the amount of cooling 85 produced by Silica-gel system which is 334 W/kg 86 [21]. 87

Intini et al. [20] have analyzed numerically and experimentally the performance of an AQSOA-based desiccant wheel. Performance of the system was assessed by determining the effect of area ratio, process inlet temperature, humidity and air force velocity. It was found that maximum moisture removal capacity is achieved at equal

area split. In addition, inlet humidity ratio was 1 found to be important in determining moisture 2 removal efficiency while process inlet 3 temperature is not that much effective. 4

6 All reviewed work on water adsorption desalination, showed that *silica-gel* / water was 7 the only adsorption working pair investigated in a 8 4 bed cycle. This work, investigates the use of an 9 advanced zeolite adsorbent material, (AQSOA 10 FAM-Z02, produced by Mitsubishi plastics) in a 4 11 beds adsorption cycle for production of both 12 13 cooling and fresh water. The effect of operating conditions in terms of evaporator and hot water 14 temperatures were studied and compared to those 15 for a silica gel 4 beds adsorption desalination 16 system. 17

18

5

2. SYSTEM MODELLING 19

A full scale four bed adsorption machine is 20 21 simulated by Simulink to study its ability to 22 produce both cooling and fresh water. It comprises of four beds with the capacity of 890 23 kg of silica gel per bed in addition to an 24 evaporator and a condenser (Fig. 1). 25

hot water is passed through desorbing beds 32 (master then slave bed). Fresh water is obtained 33 34 from the condenser by condensing water vapour via external cooling water passing through 35 cooling coil while cooling is achieved at the 36 37 evaporator.

39 In order to study the cycle, energy equations are solved for evaporator, condenser, adsorber and 40 desorber beds in addition to mass and salt balance 41 equations for the evaporator [8] as shown in 42 43 equations 1-6:

45 Evaporator mass balance equation:

$$46 \quad \frac{dM_{s,evap}}{dt} = \theta m_{s,in}^{\cdot} - \gamma m_b^{\cdot} - \frac{dc_{Mads}}{dt} M_a$$

$$47 \qquad -n. \frac{dc_{Sads}}{dt} M_a \qquad (1)$$

48

53

38

44

4

49 *Evaporator salt balance equation:*

50
$$M_{s,evap} \frac{dX_{s,evap}}{dt} = \theta X_{s,in} m_{s,in} - \gamma X_{s,evap} m_{brine}$$

51 $- X_D \frac{dcM_{ads}}{dt} M_a - n X_D \frac{dc_{sads}}{dt} M_a$
52 (2)

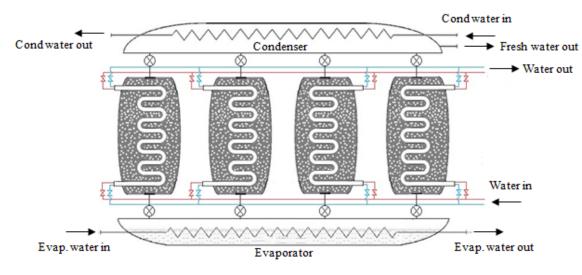


Fig. 1 Schematic diagram of the adsorption system

26

In this system, cooling water is passed through 27 28 first adsorbing bed (master bed) and the outlet stream continues the cooling process in the 29 second adsorber bed (slave bed). This cooling is 30 to absorb the heat of adsorption. For desorption, 31

54

Evaporator energy balance equation: 55

57
$$\left[M_{s,evap}c_{p,s}(T_{evap},X_{s,evap})+M_{HX,Evap}c_{p,HX}\right]\frac{dI_{evap}}{dt}=$$

$$1 \quad \theta \cdot h_f (T_{evap}, X_{s,evap}) \quad m_{s,in}^{\cdot} - h_{fg} (T_{evap}) \frac{dc_{Mads}}{dt} M_a$$

$$2 \quad -n \cdot h_{fg} (T_{evap}) \frac{dc_{Sads}}{dt} M_a - \gamma h_f (T_{evap}, X_{s,evap}) m_b^{\cdot}$$

$$3 \quad + m_{chilled} c_p (T_{evap}) (T_{chilled,in} - T_{chilled,out})$$

$$4 \qquad (3)$$

5

6 Master adsorption /desorption bed, energy
7 balance equation:

$$\begin{bmatrix} M_a c_{p,a} + M_{HX} c_{p,HX} + M_{abe} c_{p,abe} \end{bmatrix} \frac{d I_{Mads/Mdes}}{dt} = \\ 8 \qquad \pm m_{cw/hw} c_p (T_{cw/hw,in} - T''_{cw/hw,out}) \\ 9 \qquad \pm Q_{st} M_a \frac{d c_{Mads/Mdes}}{dt}$$
(4)

10 Slave adsorption /desorption bed, energy balance11 equation:

$$12 \left[M_{a}c_{p,a} + M_{HX}c_{p,HX} + M_{abe}c_{p,abe} \right] \frac{dT_{sads/Sdes}}{dt} =$$

$$13 \qquad \pm m_{cw/hw}c_{p} \left(T''_{cw/hw,in} - T_{cw/hw,out} \right)$$

$$14 \qquad \pm z \cdot Q_{st} M_{a} \frac{dc_{ads/des}}{dt} \qquad (5)$$

15 Where, z is a flag equals 0 in heat recovery phase16 and 1 in all other cases.

- 17
- 18 Condenser energy balance equation:

$$\begin{bmatrix}
 M_{cond}c_p(T_{cond}) + M_{HX,Cond}c_{p,HX}\end{bmatrix} \frac{dT_{cond}}{dt} = \\
 h_f \frac{dM_d}{dt} + h_{fg}(T_{cond}) M_a \left(\frac{dc_{Mdes}}{dt} + n.\frac{dc_{Sdes}}{dt}\right) \\
 20 + m_{cond}c_p(T_{cond}) \left(T_{cond,in} - T_{cond,out}\right)$$
(6)

21

22 For assessment of cycle performance, different parameters are calculated which are specific daily 23 water production (SDWP), performance ratio 24 (PR) which is the ratio between heat of 25 condensation to the heat of desorption, specific 26 power (SCP) and coefficient 27 cooling of performance (COP). For the determination of the 28 overall cycle performance where two useful 29 effects are obtained from the same heat source, 30 overall conversion ratio (OCR) is calculated. 31 OCR is the ratio between useful effects produced 32 (summation of heat of condensation and heat of 33 evaporation) over the input which is heat of 34

desorption [8]. These parameters are calculatedusing equations 7-14:

37

38
$$SDWP = \int_0^{t_{cycle}} \frac{Q_{cond}}{h_{fg}M_a} dt$$
 (7)

$$39 \quad PR = \frac{1}{t_{cycle}} \int_0^{t_{cycle}} \frac{m_d h_{fg}}{Q_{Mdes} + Q_{Sdes}} dt \tag{8}$$

40
$$SCP = \int_0^{t_{cycle}} \frac{Q_{evap}}{M_a} dt$$
 (9)

41
$$COP = \int_0^{t_{cycle}} \frac{Q_{evap}}{Q_{Mdes} + Q_{Sdes}} dt$$
 (10)

42
$$OCR = \int_0^{t_{cycle}} \frac{Q_{evap} + Q_{cond}}{Q_{Mdes} + Q_{Sdes}} dt$$
(11)

43

57

71

44 Where,

 $45 \quad Q_{cond} = m_{cond}c_p(T_{cond})(T_{cond,out} - T_{cond,in})$ $46 \tag{12}$

47
$$Q_{Mdes/Sdes} = m_{hw}c_p (T_{hw,in} - T_{hw,out})$$
(13)

48
$$Q_{evap} = m_{chilled} c_p (T_{evap}) (T_{chilled,in} - T_{chilled,out})$$
49 (14)

50 These set of energy and mass balance 51 equations are solved by Simulink with tolerance 52 value of 1×10^{-6} . A lumped simulation model was 53 used where the adsorbent, adsorbate and heat 54 exchangers are assumed to be momentarily at the 55 same temperature. Also perfect heat insulation is 56 assumed for all parts.

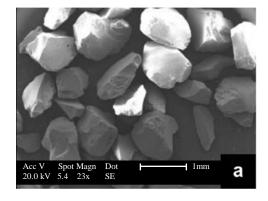
58 3. ADSORBENT MATERIAL59 CHARACTERISTICS

Two materials are compared in this work, *Silica-gel*-RD and *AQSOA-Z02*. Figure 2 (a –b),
shows SEM images for both materials and their
physical properties are listed in table I [19, 22,
23].

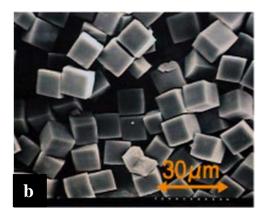
To investigate the performance of any adsorbent material, adsorption isotherms and adsorption kinetics are required.

Adsorption isotherms represent the maximum
amount of adsorbate that can be adsorbed per unit
mass of dry material at a specific vapor pressure.

5



1



2 3

4

5

TABLE I PHYSICAL PROPERTIES OF ADSORBENT MATERIALS SILICA GEL AQSOA-Z02 Property

Fig. 2 SEM images for (a) RD Silica-gel

(b) AQSOA-Z02

Granules size	0.18-1 mm	0.25-0.4 mm
BET surface area	840 m ² /g	650-770 m ² /g
Bulk Density	1 g/cm^3	0.5-0.7 g/cm ³

6 7

Different isotherm 8 models can predict performance adsorbent materials such 9 as Dubinin-Astakhov (D-A), Sips, Toth, Freundlich, 10 Modified Freundlich and Langmuir. Isotherms of 11 Silica-gel-RD, can be predicted by the Dubinin-12 Astakhov (D-A) model (equation 15) with the 13 constants given in table II [8]. 14

15

16
$$c^* = c_0 \exp\left[-\left(\frac{RT}{E}\ln\left(\frac{P}{P_o}\right)\right)^n\right]$$
 (15)

TABLE II **DUBININ-ASTAKHOV EQUATION CONSTANTS**

Symbol	Value	Unit ^a
c_0	0.592	kg/kg of adsorbent
E	3.105	kJ/mole
п	1.1	(-)
R	8314	J/mole.K

^aUnits are; kg = kilogram, K = Kelvin.

18 19

For AQSOA-Z02, water vapor uptake is 20 calculated via the model developed by Sun et al. 21 [24] as shown in equations (16-17). 22

23

24
$$\frac{c}{c_{max}} = \frac{K(P/P_s)^m}{1+(K-1)(P/P_s)^m}$$
 (16)
25

Where, 26

 $K = \alpha \exp[m(Q_{st} - h_{fa})/RT]$ 27 (17)

 $\alpha = 9 * 10^{-7}$, m= 3.18 and $Q_{st} = 3600 \ kJ/kg$ 28

29 30 c_{max} is maximum uptake, *m*, is Where, heterogeneity factor, h_{fg} , is the latent heat [kJ/kg], 31 *R*, is universal gas constant [J/mol.K]. 32

33

As adsorption process is time dependent, 34 adsorption kinetics are needed to determine the 35 rate of adsorption or desorption at the specified 36 operating conditions. Adsorption kinetics are 37 modeled by Linear Driving Force (LDF) model 38 39 for both materials, (equations 18-19) [8] with all constants given in table III. 40

41
42
$$\frac{dc}{dt} = k(c^* - c)$$
 (18)

43
$$k = (15 D_{so}/R_p^2) e^{\left(\frac{-Ea}{RT}\right)}$$
 (19)

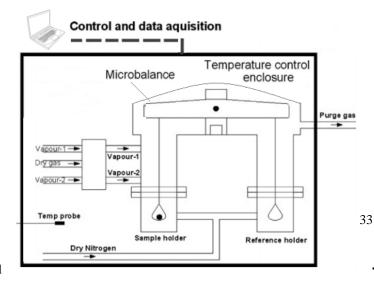
44

4

45 For Silica-gel, LDF model constants are obtained from Ng et al [8] while for AQSOA-46 Z02, tests by a dynamic vapor sorption (DVS) 47 48 gravimetric analyzer, fig. 3, were carried out at University of Birmingham to determine the 49 50 constants.

51 This DVS analyser consists of a temperature controlled chamber that contains a sensitive 52 53 recording microbalance (Cahn D200).

17



1

2 Fig. 3 Schematic diagram of the used DVS instrument3

A dry Nitrogen purge system is applied before 4 sample loading to ensure accurate uptake 5 measurements by preventing vapor condensation 6 on the balance head. The vapor pressure in the 7 chamber is regulated by mass flow controller 8 which determines the amount of dry and saturated 9 vapor gas. A PC microcomputer is connected to 10 the microbalance to record mass readings every 4 11 seconds while vapor pressure sensor and RTD 12 temperature sensor are used to measure testing 13 conditions. The accuracy of the DVS analyser 14 microbalance Cahn is verified by using 100 mg 15 standard calibration mass, where the expected 16 mass accuracy of the tested sample is ± 0.05 mg. 17

To measure adsorption kinetics, firstly a sample 18 of 10 mg (± 0.05 mg) is placed in the sample pan 19 and then dried at certain temperature until no 20 change in the mass is recorded. Secondly, 21 adsorption process is started at various partial 22 pressures starting from 0.1-0.95. During that 23 time, change of the mass is calculated every 4 24 25 seconds. This test is repeated at different temperatures starting from 20°C to 52°C. 26

27

Test results at temperature of 36° C and at different vapor partial pressures (0.1 to 0.95) are shown in Fig. 4. The kinetics of *AQSOA-Z02* were then fitted using equations 18, 19 and the constants are tabulated in table III.

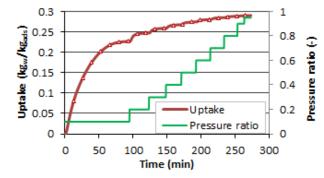


Fig. 4 DVS test results at 36°C for AQSOA-Z02

 TABLE III

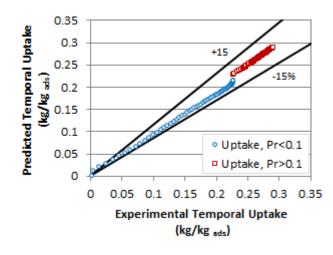
 LINEAR DRIVING FORCE, LDF EQUATION CONSTANTS

Symbol	SILICA GEL	AQSOA-Z02		Unit ^a
		Pr ^b >0.1	Pr <0.1	_
D_{so}	2.54 E-4	4.85 E-9	2.77 E-5	m ² /s
R_p	0.4 E-3	0.15 E-3	0.15 E-3	m
E_a	42000	17709.8	44423.5	J/mol

^aUnits are; m = meter, s = second, J = Joule, mol = mole.

^bPr is the pressure ratio between bed and heat exchanger

35 Figure 5, compares the LDF predicted temporal 36 uptake to the experimental ones showing 37 deviation of less than $\pm 15\%$.



40

41

34

38

39

Fig. 5 Comparison between the measured water uptake for AQSOA-Z02 and those predicted by the LDF model

1 4. METHODOLOGY

The developed numerical model of the 4 bed 2 adsorption system shown in fig 1 is validated 3 against the measured experimental results from a 4 5 silica-gel/water adsorption plant operating in a 4-Bed mode for desalination application and 6 7 cooling production [25]. Fig.6 shows the comparison between the simulation results and 8 experimental measurements 9 for the basic components of an adsorption desalination cycle. 10 11

TABLE IV ERROR RANGE FOR THE VALIDATION OF ADSORPTION DESALINATION CYCLE

	Maximum (%)	Minimum (%)
Bed 1	4.05	-8.34
Bed 2	4.6	-8.48
Bed 3	2.79	-9.89
Bed 4	3.67	-3.08
Condenser	5.59	-3.98
Evaporator	4.25	-1.37

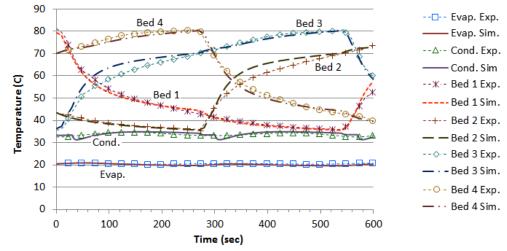


Fig. 6 Comparison of Basic components temperatures for numerical and experimental results for 4-Bed adsorption desalination cycle using silica-gel.

12

Comparison of cycle outputs i.e. SDWP and SCP are presented in fig.7, where results show that the current model can predict the performance of desalination and cooling cycles within $\pm 10\%$ error margin (table IV).



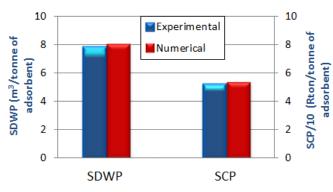


Fig. 7 Comparison of SDWP and SCP for numerical and experimental results for 4-Bed adsorption desalination cycle

19

20 To investigate the performance of the AQSOA-Z02, a parametric study was carried out to 21 compare the SDWP and SCP of AQSOA-Z02 22 23 against Silica-gel at different inlet hot water temperatures and evaporator water temperatures 24 ranging from $(65-85^{\circ}C)$ $(10-30^{\circ}C)$ 25 and respectively. 26

Bed cooling water and condenser cooling water temperatures are kept constant at 30°C with quarter cycle time of 150 sec and switching time of 30 sec.

31

32 5. RESULTS AND DISCUSSION

Figures 8 to 13 present SDWP and SCP for different evaporator and hot water temperatures. As shown in Figs 8 to 11, at low hot water temperatures, less than 75°C, *silica-gel* is better than *AQSOA-ZO2* at high evaporator temperatures

of more than 20°C. At evaporator water temperature of 30°C, and at hot water temperatures of 65°C, silica-gel can result in a SDWP and SCP of 5.4 m³ per day and 38.1 Rton per tonne of *silica-gel*, respectively while AOSOA-Z02 is capable only of producing 1.1 m^3 per day of fresh water and 6.2 Rton of cooling per tonne of AQSOA-Z02. However, as evaporator water temperature decreases below 18 °C and hot water temperature increases above 75°C, AQSOA-Z02 outperforms *silica-gel* as shown in fig. 10 to 10. At evaporator water temperature of 10°C, and at hot water temperatures of 85°C, AQSOA-Z02 can produce 6.2 m^3 per day and 53.7 Rton per tonne of AQSOA-Z02 while silica-gel is capable only of producing 3.39 m³ per day of fresh water and 15.7 Rton of cooling per tonne of silica-gel.

Another parameter which indicates the cycle performance is the overall conversion ratio, Figs 14 - 15. It is clear that OCR for *silica gel* is highly affected by varying chilled water temperature (Fig. 14), while it is not the case for *AQSOA-Z02* as shown in Fig. 15 which is a result of the "S" shaped isotherm of *AQSOA-Z02*. Moreover, Figs 15, proves that hot water temperature highly affects the performance of *AQSOA-Z02* cycle where OCR varies from 0.3 to 0.8 as hot source temperature increases from 65 to 85 °C. However, silica gel cycle performance is only better than *AQSOA-Z02* at high evaporator water temperature, above 20°C as it reaches 1.1 at 30° C.

According to these results, where heat source temperatures are available at temperatures above 75° C for the applications of space cooling, typically evaporator temperature (10-20°C), *AQSOA-Z02* is recommended. For applications where evaporator temperature above 20 °C is needed, *silica-gel* becomes more effective.

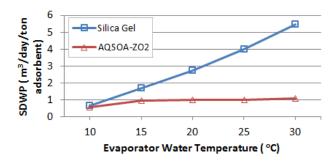


Fig. 8 SDWP at 65°C hot water temperature.

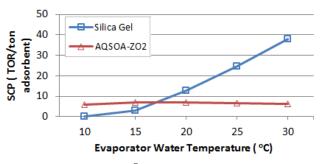


Fig. 9 SCP at 65°C hot water temperature.

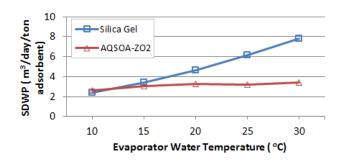


Fig. 10 SDWP at 75°C hot water temperature.

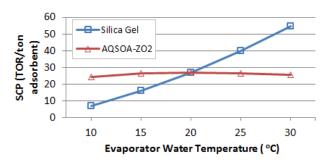


Fig. 11 SCP at 75°C hot water temperature.

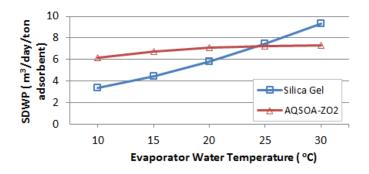


Fig. 12 SDWP at 85°C hot water temperature.

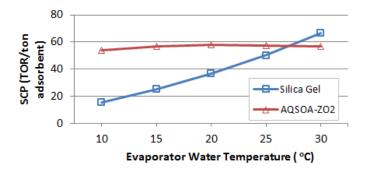


Fig. 13 SCP at 85°C hot water temperature.

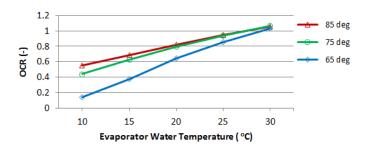


Fig. 14 Overall conversion ratio for *Silica-gel* at hot water temperatures (65-85°C).

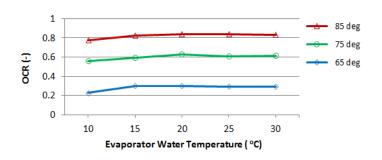


Fig. 15 Overall conversion ratio for *AQSOA-Z02* at hot water temperatures (65-85°C).

6. CONCLUSIONS

Adsorption based desalination technology outperform other desalination technologies in terms of its ability to use waste heat to produce potable water and simultaneously produce useful cooling effect. However, silica gel adsorbent material was the only material reported in the literature, this work investigates the use of advanced zeolite material *AQSOA-Z02* in a full scale four bed adsorption system for the purposes of fresh water production and cooling and compared the results to silica gel.

Results showed that for heat source $75^{\circ}C$ temperatures above and evaporator temperature below 20°C suitable for cooling applications, AQSOA-Z02 outperforms silica gel in terms of higher SDWP and SCP. While for application where evaporator temperature above 20°C is needed, *silica-gel* becomes more effective. Also, such results indicate that the use of combined system of silica gel and zeolites can cover wide range of evaporation temperature to achieve best combination of cooling and water desalination.

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