

# Analysis of Vapor Absorption Refrigeration Machine Performance with Different Configuration of Water Supply

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**ABSTRACT**

In this article we analyzed the performance of the heat based single effect vapor absorption refrigeration machine, with the different configuration of cooling water supply. The comparison in these configuration is done on the basis of concentration of libr in the weak as well as in the strong aqueous solution, and requirement of heat load into generator .also observed the percent reduction of heat load with composition of libr (XG) .all the configuration has a specific significance but we try to find out that appropriate configuration which is more economically feasible without reach the crystallization limit in the strong solution side.

**KEYWORDS:** vapor absorption refrigeration machine, strong solution, weak solution, generator

**INTRODUCTION**

To deal with the changes in global climate, rapid growth of energy demand and rapid development of renewable energy and energy utilization, much attention has been focused on the research and development (R&D) of absorption cycles, such as absorption heat pump and absorption refrigeration

cycle. In this work we try to increase the performance of vapor absorption refrigeration machine by changing the cooling water supply in three different configuration. These are as given below

**Nomenclatures**

<b>XG</b> -composition of libr in at exit of generator	<b>TC</b> -condenser temperature
<b>QG</b> -heat load at generator	<b>PC</b> -condenser pressure
<b>XA</b> -composition of libr in at exit of absorber	<b>TG</b> -Generator temperature
<b>COP</b> -coefficient of performance	<b>TGI</b> -Inlet temp to generator
<b>CR</b> -circulation ratio	<b>TGO</b> -Outlet temp from generator

**CASE 1: Cooling Water Enters First Absorber and Outlet of It Goes To Condenser**

In the first configuration cooling water enters first in the absorber than outlet of cooling water from absorber goes to condenser. We can see it from the fig: 1

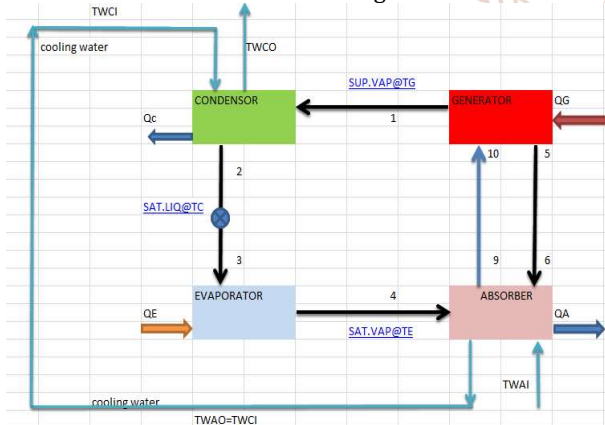


Fig: 1 cooling water enters first absorber and outlet of it goes to condenser

**CASE2: Cooling Water First Enters Into Condenser and Outlet Of It Goes To Absorber**

In the second configuration cooling water enters first in the condenser than outlet of it goes to the absorber. We can see it from the fig: 2 in this case TC reduced by aprox.5 degC but

the concentration increased by approx. 3% which tends to reach near the crystallization limits. That is not desirable.

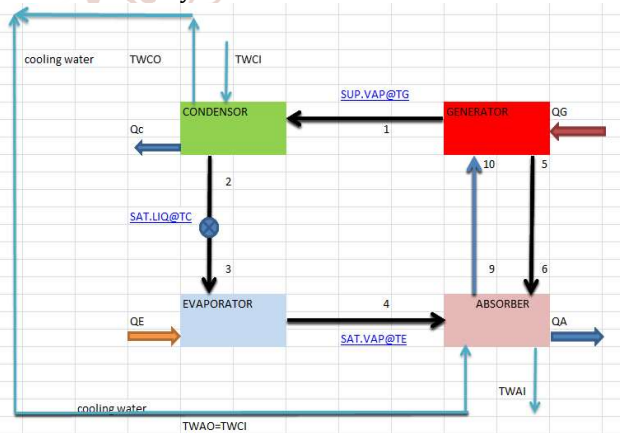


Fig:2 cooling water first enters into condenser and outlet of it goes to absorber

**Case 3: Cooling Water Enters Parallel Into Absorber AND Condenser**

In the third configuration cooling water enters parallelly into absorber and condenser. We can see it from the fig: 3 this case shows the maximum cop among all the configuration and show less amount of heat load because the TC also go down in this case. But requirement of cooling water is almost double as compare to first and second case.so this case economically not feasible.

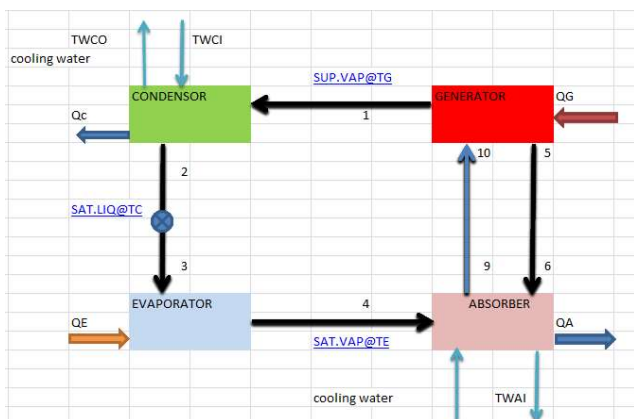


Fig:3 cooling water enters parallel into absorber and condenser

The effect of all these configurations analyzed on the basis of generator temperature, condenser pressure, absorber temperature, concentration of libr at the outlet of absorber side and generator side and also analyzed the coefficient of performance. While keeping the cooling load constant.

Third configuration shows high cop but on the verge of large supply of cooling water that is not feasible economically. While in the second configuration water enters into condenser first , this is also show the good cop but in this case we have to face large problem of crystallization means libr exceeds its solubility limit, when strong solution from generator, exchange its heat with the week solution in the solution heat exchanger. so it required rapid inspection of the system that is difficult task.

In the first configuration cop does not so high but in this configuration system can work beyond the crystallization limit and also at the low cost with respect to water supply.

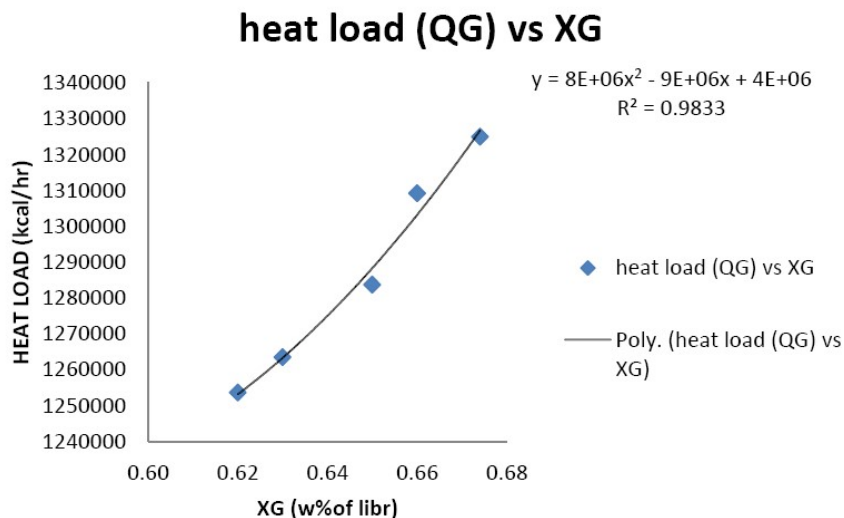
This study also shows that as the temperature of condenser decrease the requirement of heat source temperature reduced ,so ultimately cop increased but problem of crystallization also increased. So we have to decide the optimum condition among these entire configurations. That is shown in first configuration, in which cooling water first enters absorber than condenser. The tabulated form of the out comes as given below in the table1.

Table1: observations

Parameters	Units	case1cw in abs.	case2 cw in cond.	case3 cw in para
TC	C	45.9358	40.59363	40.78813
PC	mmhg	75.27203	57.30486	57.89823
TA	C	43.89835	47.74024	43.91214
XA=XGI	%	60.3157	62.3967	60.3463
XGO	%	64.6932	67.3967	64.9508
TGI	C	95.20067	95.38219	92.93262
TGO	C	103.9942	97.0734	94.43737
TG	C	125.446	118.408	115.8632
QG	kcal/hr	1306351	1323973	1301840
QE	kcal/hr	1043280	1043280	1043280
COP		0.79862	0.787992	0.801389
CR		12.93865	13.47933	14.10608

**VARIATION OF HEAT LOAD**

we also analyzed that amount of heat require to separate the water from the aqueous solution of libr not only depends on condenser temperature but also depends upon composition of libr because among all the above configuration if we compare first and second configuration we observed that temperature of condenser reduce when cooling water first enters into the condenser but as per the correlation[1]. it leads enhancement of concentration of libr because of this we require large heat supply in the second configuration(cooling water first enters into condenser).as we see in the graph 1 the heat load increase with increasing the composition of libr at constant TC.

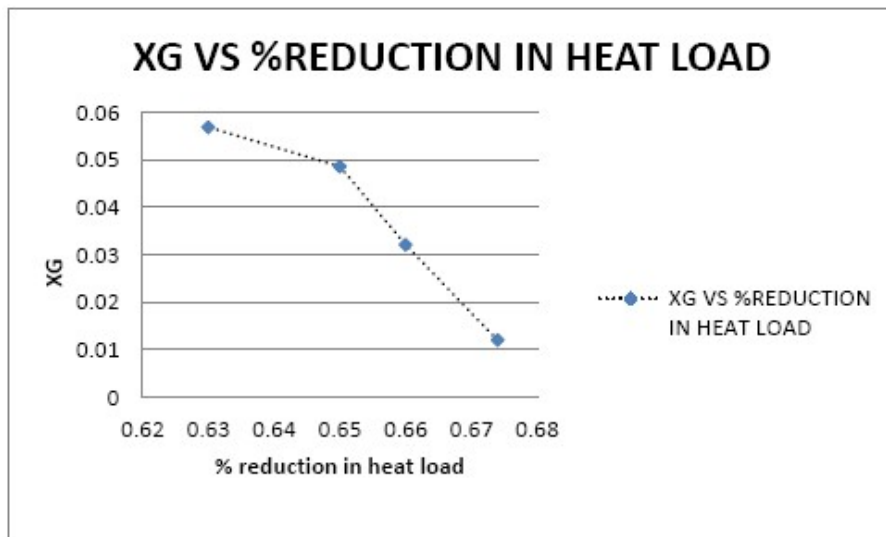


Graph:1 -variation of heat load with XG

To see the variation of heat supply with concentration of libr at constant condenser temperature and also at constant cooling load.(the calculation part of heat load has shown in given annexure) we can observe from table 1 that as the concentration decrease requirement of heat supply also decrease . percent reduction is approximately 6% for 57% concentration of libr in generator while for 66% libr we obtained only 1.2% heat load reduction. This variation can be easily observe by the graph2. But we are unable see this variation into first and second configuration for the fix out let temperature of cooling water( from the condenser in the first case and from the absorber in the second case). if we allow to increase the concentration due to this temperature of the absorber and the condenser goes below the outlet temperature of the cooling water that is not feasible .

Table: 2 variation of heat load with concentration

XG	H1 (Kcal/kg)	H5 (Kcal/kg)	H10 (Kcal/kg)	m5 (kg/hr)	m10 (kg/hr)	QG (kcal/hr)	%reduction
0.62	650.79	264.675	240.59	23263.1	25127.13	1324846	0
0.61	650.59	257.69	234.259	22741.18	24605.21	1309090	1.203
0.60	650.11	252.215	229.85	22368.37	24232.4	1283608	3.212
0.58	649.88	242.78	221.3022	21622.76	23486.79	1263402	4.863
0.57	649.78	238.205	217.16	21249.95	23113.98	1253528	5.689



Graph: 2-variation of XG with % reduction.

**CONCLUSION**

The effect of all these configurations analyzed on the basis of generator temperature, condenser pressure, absorber temperature, concentration of libr at the outlet of absorber side and generator side and also analyzed the coefficient of performance. While keeping the cooling load constant. This study also shows that as the temperature of condenser decrease the requirement of heat source temperature reduced ,so ultimately cop increased but problem of crystallization also increased. So we have to decide the optimum condition among these entire configurations. That is shown in first configuration, in which cooling water first enters absorber than condenser.

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## Sample Calculation

Objective → Cal. Amount of heat load require AT given  $T_c, Q_E$

Annexure  $Q_{c1} = m_4 H_1 + m_5 H_5 - m_{10} H_{10}$  — (i)

$m_r = m_1 = m_4 = m_2 = m_3 =$  flow rate of refrigerant

Given data →  $Q_E = 1043280 \text{ kcal/hr}, T_c = 40.78^\circ\text{C}, T_{c1} = 116.66^\circ\text{C}$

$$m_4 = \frac{Q_E}{H_4 - H_3} \text{ ----- (ii)} = 1864.03 \text{ kg/hr}$$

Enthalpy of sat. liq.

$$H_3 = (T_c - T_{ref}) \text{ , } H_4 = 600.4785 \frac{\text{kcal}}{\text{kg}} \text{ at } T_E \text{ \& } P_E$$

$$= 40.78 \text{ kcal/kg}$$

Enthalpy of sup. vapor

\*  $H_1 = (2501 + 1.88 \times T_{c1}) / 4.18 \text{ ----- (iii)}$   
 $= 650.79 \text{ kcal/kg}$

$$m_5 = m_4 \times \left( \frac{x_A}{x_{c1} - x_A} \right) \text{ , } m_{10} = m_4 \times \left( \frac{x_{c1}}{x_{c1} - x_A} \right) \text{ ----- (iv)}$$

$$= 23263.15 \text{ kg/hr} \text{ , } = 25127.13 \text{ kg/hr}$$

Enthalpy of solution @  $T_{c10}$

\*  $H_5 = 42.81 - 425.92 \times x_{c1} + 404.67 \times x_{c1}^2 + (1.01 - 1.23x_{c1} + 0.48x_{c1}^2) \times T_{c10}$

$$x_{c1} = 0.67, x_A = 0.62, T_{c10} = 95.3137, T_{c1} = 93.65$$

$$H_5 = \dots \text{ kcal/}$$

\* Enthalpy of solution @  $T_{c11}$

$$H_{10} = 42.81 - 425.92 \times x_A + 404.67 \times x_A^2 + (1.01 - 1.23x_A + 0.48x_A^2) \times T_{c11}$$

$$= 240593 \text{ kcal/kg} \text{ ----- (vi)}$$

after putting the all value in equation (i)

$$Q_{c1} = 1864.03 \times 650.79 + 23263.1 \times 264.6751 - 25127.13 \times 240593$$

$$= 1324846 \text{ kcal/hr.}$$

**REFERENCES**

[1] Prof. M Ramgopal, lesson on Vapour compression refrigeration systems, version 1 ME, IIT Kharagpur (<http://nptel.ac.in/courses/112105129/10>)