

Working Principle of VA Refrigeration

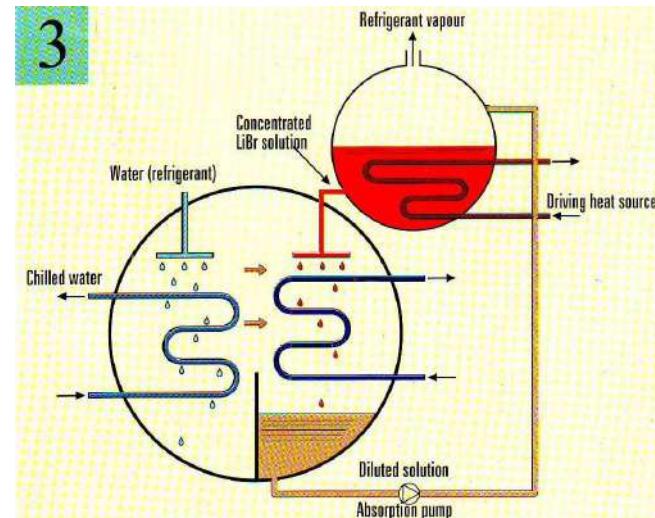
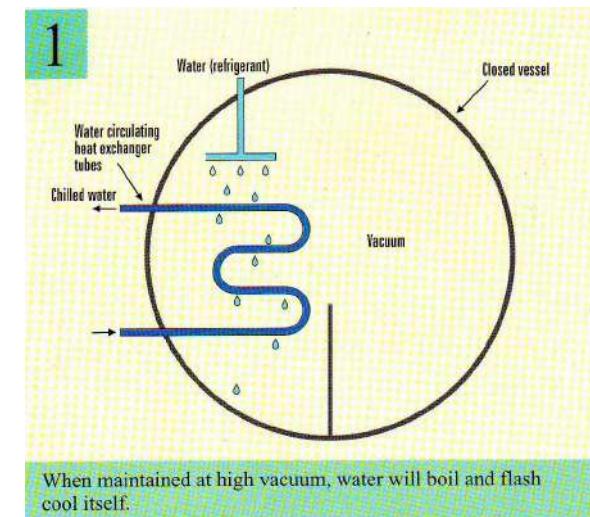
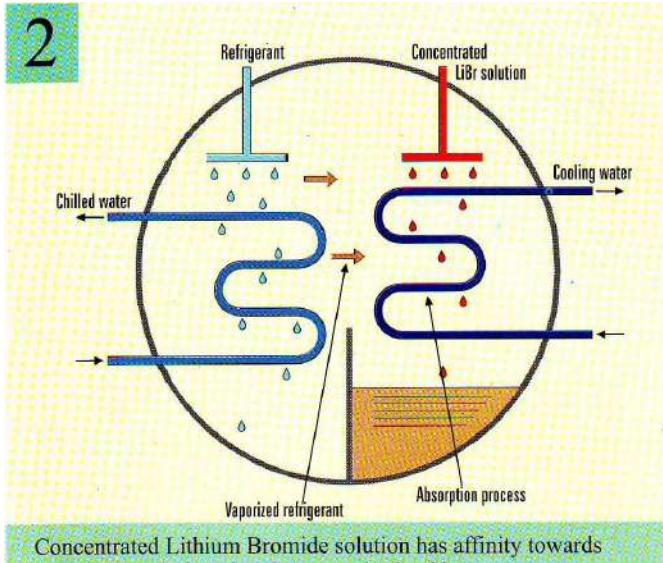
Absorption Refrigeration

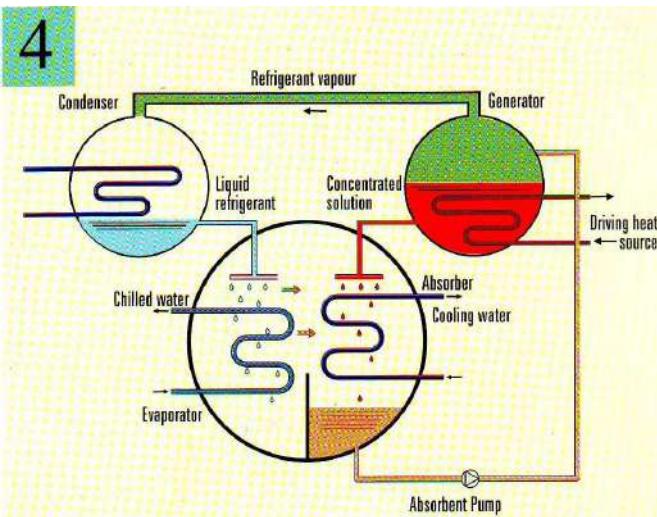
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ME 415: Refrigeration & Building Mechanical Systems





This heat causes the solution to release the absorbed refrigerant in the form of vapour. This vapour is cooled in a separate chamber to become liquid refrigerant.

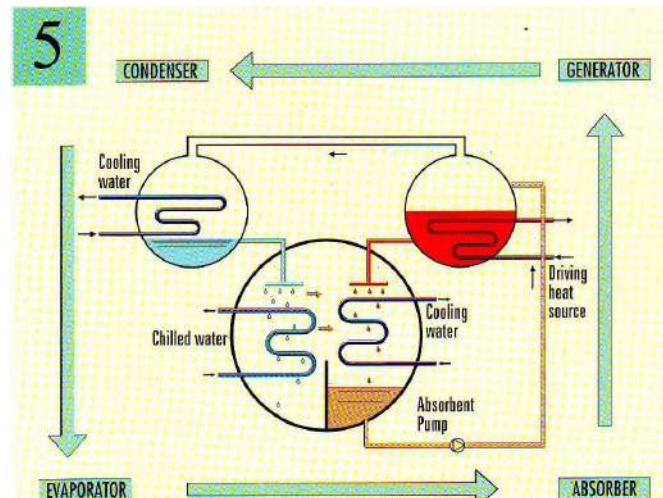
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Absorption Refrigeration

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The basic operation cycle of the single effect vapour absorption chiller.

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Absorption Refrigeration

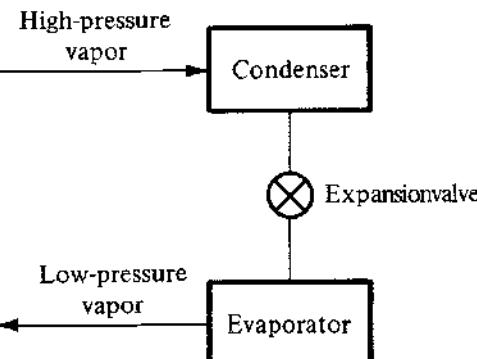
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VA & VC Comparison

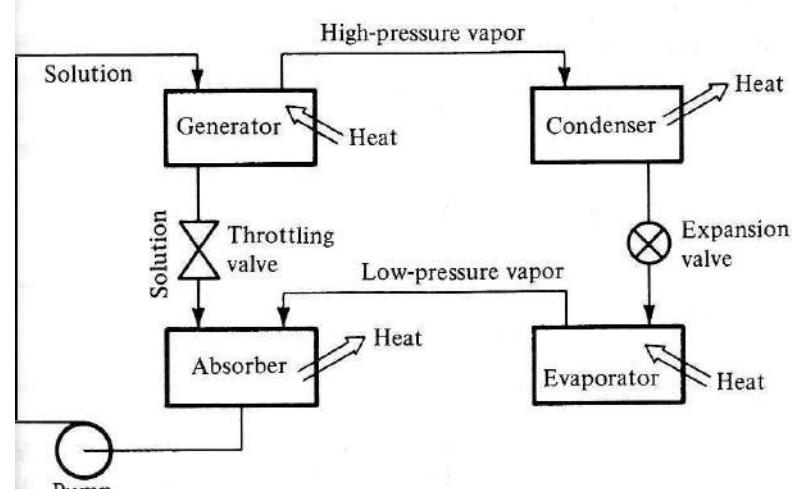
Vapor compression:

Absorption:

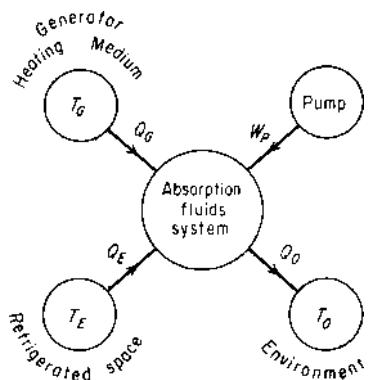
1. Absorb vapor in liquid while removing heat
2. Elevate pressure of liquid with pump
3. Release vapor by applying heat



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$$COP|_{max} = \frac{T_E}{T_G} \left[\frac{T_G - T_o}{T_o - T_E} \right]$$

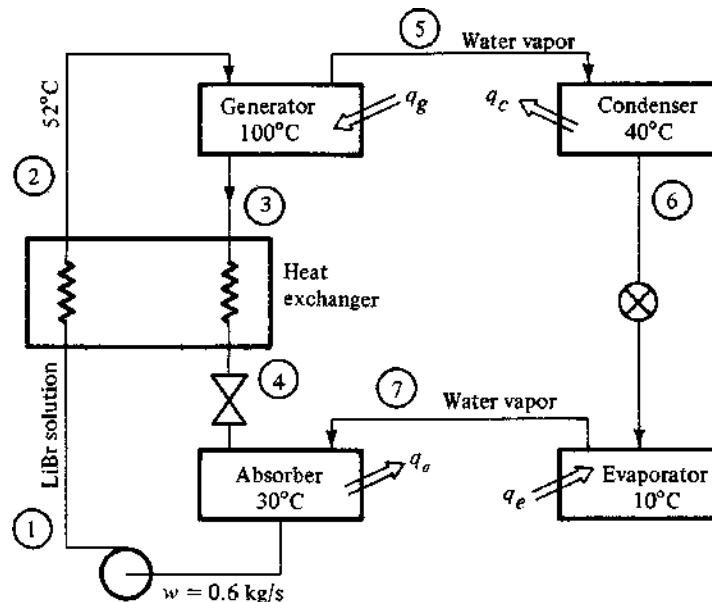
$T_G \uparrow \Rightarrow COP \uparrow, T_E \uparrow \Rightarrow COP \uparrow, T_o \uparrow \Rightarrow COP \downarrow$



Example

In an absorption system using LiBr, $T_G = 100^\circ\text{C}$, $T_E = 10^\circ\text{C}$, $T_a = 30^\circ\text{C}$, $T_c = 40^\circ\text{C}$. Estimate the values of COP for the following conditions:

- ① ideal/Carnot cycle
- ② a real cycle if pump delivers 0.6 kg/s solution
- ③ if a heat exchanger is inserted after the pump and water enters the generator at 52°C.
- ④ if condensing temperature is reduced to 34°C, is there any chance of crystallization?



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Without heat exchanger, (1) & (2) and (3) & (4) are the same.

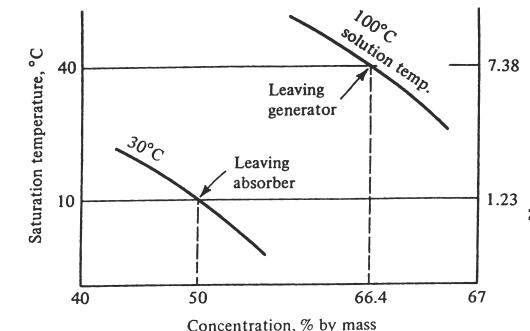
Analysis of VA System Example

1. Carnot cycle: Let, $T_o = (T_a + T_c)/2 = 35^\circ\text{C}$

$$COP = \frac{T_R}{T_G} \left[\frac{T_G - T_o}{T_o - T_R} \right] = \frac{10+273}{100+273} \frac{100-35}{35-10} = 1.97$$

Real Cycles using LiBr: Two pressures exist in the system:

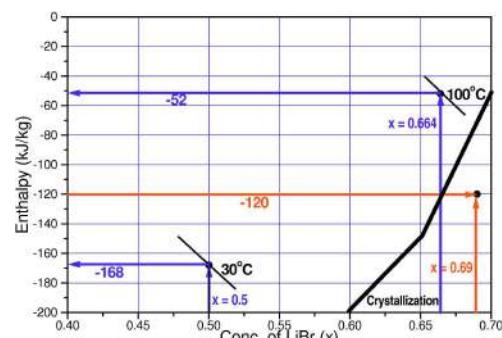
- ① High pressure (generator & condenser): $T_c = 40^\circ\text{C} \rightarrow P_H = 7.38 \text{ kPa}$
- ② Low pressure (absorber & evaporator): $T_E = 10^\circ\text{C} \rightarrow P_L = 1.23 \text{ kPa}$



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$$x_1 = 0.50 \text{ & } x_3 = 0.667$$



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Using LiBr solution Chart: $h_1 = -168 \text{ kJ/kg}$ & $h_3 = -52 \text{ kJ/kg}$

Using Steam Table:

- $h_5 = 2676.0 \text{ kJ/kg}$, saturated vapour at 100°C
- $h_6 = 167.0 \text{ kJ/kg}$, saturated liquid at 40°C
- $h_7 = 2520.0 \text{ kJ/kg}$, saturated vapour at 10°C



4. T_{cond} is reduced to 34°C :

- if $T_{cond} = 34^\circ\text{C} \Rightarrow P_{HP} = P_{sat} = 5.32 \text{ kPa}$.
- $\rightsquigarrow x_3 = 0.69 \Rightarrow m_3 = 0.435 \text{ & } m_5 = 0.165 \text{ kg/s}$
- $h_1 = -168, h_2 = -120 \text{ & } h_3(x = 0.69 \text{ & } 100^\circ) = -57 \text{ kJ/kg}$
- Energy balance in heat exchanger:
 $m_1(h_2 - h_1) = m_3(h_3 - h_4) \rightsquigarrow h_4 = -120 \text{ kJ/kg}$
- From chart, $x = 0.69 \text{ & } h_4 = -120$ \Rightarrow crystallized state.
- Crystallization is most likely to occur where the solution from the generator leaves the heat exchanger.
- An operating condition conducive to crystallization is low condensing pressure/temperature. Modern systems maintains higher condensing pressure even when low-temperature condensing water is available to avoid crystallization.



2. Without heat exchanger:

- LiBr balance: $m_1x_1 = m_3x_3 \Rightarrow m_3 = \frac{0.5}{0.667}0.6 = 0.452 \text{ kg/s}$
- $m_1 = m_3 + m_5 \Rightarrow m_5 = 0.6 - m_3 = 0.148 \text{ kg/s}$
- $q_g = m_3h_3 + m_5h_5 - m_2h_2 = 473.3 \text{ kW}$
- $q_e = m_7h_7 - m_6h_6 = 348.2 \text{ kW}$
- $COP = \frac{q_e}{q_g} = 348.2/476.6 = 0.735$

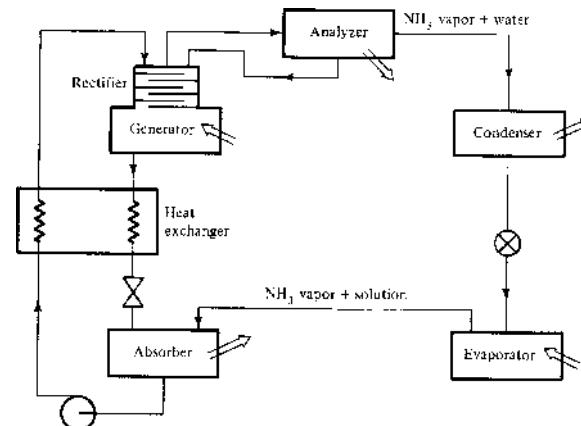
3. 50% Solution leaves heat exchanger at 52°C

- $h_2(x = 0.50 \text{ & } 52^\circ) = -120 \text{ kJ/kg}$
- $q_g = m_3h_3 + m_5h_5 - m_2h_2 = 444.5 \text{ kW}$
- $COP = \frac{q_e}{q_g} = 348.2/444.5 = 0.783$

Note that, maximum possible COP is only 1.97, with heat exchanger COP improves from 0.736 to 0.783.



Aqua-ammonia Absorption System



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Rectifier & analyser are used to minimize the presence of water vapour going to condenser and evaporator.



Aqua-ammonia vs. LiBr System

- two systems have comparable COPs.
- aqua-ammonia system can provide temperatures below 0°C, but commercial LiBr systems are limited to temperatures higher than 3°C.
- aqua-ammonia system requires extra components such as rectifier & analyser.
- aqua-ammonia system operates at pressures higher than atmospheric, LiBr systems operate at very low pressures.
- LiBr is very corrosive, hence require special inhibitors.

Problems with Vapour Absorption System

- high initial cost
- shorter life (15 yrs for VA systems, 25 yrs for VC systems.)
- requires more space
- requires more condenser water (4.4 gpm/ton for VA systems, 3.0 gpm/ton for VC systems), high capacity cooling tower and water treatment plant.
- requires chimney.
- require proper maintenance and longer downtime for overhauling.

