# AIR CONDITIONING BOOSTER 

Ehab Nader Tuffaha<br>Department of Mechanical Engineering, Hashemite Univesity-Jordan


#### Abstract

The study trying to utilize and allows to harness the natural and free solar energy gotten from the sun by converting the additional free enthalpy gotten as thermal and kinetic energy likewise into useful power; on other words the main concern is increasing the COP.This type of AC will use as much solar energy as is available and convert the solar energy directly to replace the equivalent amount of AC power from the mains provider. Under optimum conditions, this can save up to $25 \%-45 \%$ of your mains power usage during the summer. This will be most recent innovation in standalone power saving which is partially free of charge by the Sun.The study aims to renovate the traditional refrigeration system which is normally known as compression refrigeration cycle everywhere in our buildings, so that it can be developed to take advantage of the free solar energy without complicated changes neither a completely demolishing.


KEYWORD: HVAC, Solar radiation, R 134a, thermodynamics, momentum

## INTRODUCTION

Jordan is among of the most enthusastic developing country to promote utilization of renewable energy, as very well known the solar radiation in jordan average with in $4500 \mathrm{~Wh} / \mathrm{m} 2$ to $8000 \mathrm{~Wh} / \mathrm{m} 2$.

More than 300 sunny days in Jordan driving us to think more seriously in renewable energy resources and somehow reduce dependence on oil consumption.

Basic thermodynamics statement that as a greater temperature difference between substances the faster heat flow, so once the refrigerant gas temperature increased then a heat rejection is higher; same what happened when the ambient temperature increases, less heat can be rejected from the air-cooled condenser to the hotter ambient.


Figure - 1

The study here is utilizing thermal energy by using solar collectors see figure 1.

Adding an extra heat to the cycle in order to increase $\boldsymbol{\Delta T}$ between condenser and surrounding environment.
Not quite, but The future will be for renewable energy, however the first step should be renovating the transition phase of the hybridized types by paving the way for the renewable sources, as there is no doubt that renewable energy does not match in any way the traditional energy that we are accustomed to in term of cost neither the required efficiency.

## LITRATURE REVIEW

The main point is to drive the point number 2' of the below figure ( $\mathrm{P}-\mathrm{H}$ diagram) to point number 2 , in otherwords add a kinetek energy to the system in order to increase the enthlby.


Figure - 2
A heat exchanger will be used in this study to reach for a hybrid air conditioning system and achieve to targeted energy savings. By utilizing The First Law of Thermodynamics which is states that heat is a form of energy.

The First Law of Thermodynamics states that energy cannot be created or destroyed; so the sun radiation here or the heat itself will be the exerted energy to the closed refrigeration cycle.

The desired thermal output and rate of heat transfer help determine the optimal type and design of heat exchanger as some heat exchanger designs offer greater heater transfer rates and can handle higher temperatures than other designs.

## METHODOLGY

Heat transferee calculations:
a) $\mathrm{Qh}=\mathrm{mh} \times \mathrm{Cph}(\mathrm{T}$ hou $\mathrm{t}-\mathrm{Th}$ in) $\quad$ Versus
b) $\mathrm{Qc}=\mathrm{mc} \times \mathrm{Cpc}$ (Tc out- Tcin)

Equation a here represent water system versus equation $b$ for refrigeration cycle .

Where $\mathrm{Q}=$ heat energy (Joules, J ) $\mathrm{m}=$ mass of a substance $(\mathrm{kg}) \mathrm{c}=$ specific heat (unit's $\mathrm{J} / \mathrm{kg} \cdot \mathrm{K}$ ) $\Delta \mathrm{t}$ is a symbol meaning "the change in" temperatures

T cout $=$ Outlet refrigeration temperature from heat exchanger.
T cin $=$ Inlet refrigeration temperature from heat exchanger.
T hin $=$ Water Inlet temperature from the solar collector to heat exchanger.
T hout $=$ Water outlet temperature from heat exchanger returning to solar collector.

Cph $=$ The specific heat capacity of water is $4.18 \mathrm{~kJ} / \mathrm{g} \mathrm{C}$.
$\mathrm{Cpc}=$ Specific isobar heat capacity of R 134 a is $1.7 \mathrm{Kj} / \mathrm{Kg} . \mathrm{K}$
$\mathrm{mc} \& \mathrm{mh}$ are the mass flow rate of each fluid.
Capacity rate of hot liquid, $\mathrm{Ch}=\mathrm{mh} \times \mathrm{Cph}$
$\mathrm{mh}=$ mass flow rate of hot water $\mathrm{kg} / \mathrm{sec}$; here will be $1 \mathrm{~kg} / \mathrm{s}$ (Please see note 1 )
$\mathrm{Ch}=\mathrm{mh} \times \mathrm{Cph}=1 \times 4.18=4.18 \mathrm{~kW} / \mathrm{K}$
$\mathrm{Cc}=\mathrm{mc} \times \mathrm{Cpc}=0.2 \times 1.7==0.34 \mathrm{~kW} / \mathrm{K}$
It's clear now $\mathrm{Cmin}=\mathrm{Cc}=0.34 \mathrm{kw} / \mathrm{k}$
$\mathrm{Qmax}=\mathrm{Cmin} \mathrm{x}(\mathrm{T}$ hin $-\mathrm{T} \operatorname{cin})=0.34 \times(85-40)=15 \mathrm{KW}$;
$\mathrm{Qmax}=\mathrm{Ccx}(\mathrm{T}$ cout- $\mathrm{T} \operatorname{cin})=0.34 \mathrm{X}(\mathrm{T}$ cout -40$)$
$15 \mathrm{kw}=0.34 \mathrm{X}(\mathrm{T}$ cout -40$)$; so T cout $=84 \mathrm{C} \ldots$.
$\mathrm{Qmax}==\mathrm{Ch} x(\mathrm{~T}$ hin- T hout $)=4.18 \mathrm{X}(85-\mathrm{T}$ hout $)$, so T hout $=81 \mathrm{C} \ldots .$.
$\mathrm{Ch}=\mathrm{mh} \times \mathrm{Cph}=1 \times 4.18=4.18 \mathrm{~kW} / \mathrm{K} \ldots \ldots$. (1) Please see note 1
$\mathrm{Cc}=\mathrm{mc} \times \mathrm{Cpc}=0.2 \times 1.7==0.34 \mathrm{~kW} / \mathrm{K} \ldots \ldots$. (2) Please see note 2
In order to get mass flow rate of Water (inside heat exchanger) \& Refrigerant (R134 a):

Note 1, For water:
Assuming water inlet 85 c and outlet is 81 c :
$\mathrm{Q}=500$ GPM Delta T (F)
$15 \mathrm{KW} / 3.5=4 \times 12000=51429 \mathrm{BTU} \mathrm{h}$
$51429 /(500 \times(185-177)=10$ GPM Which is $0.7 \mathrm{Kg} / \mathrm{s}$ let say $=1 \mathrm{Kg} / \mathrm{s}$ to be more reasonable.

Note 2, For R 134 a :
As discharge temperature almost 40 C so enthalpy is $\mathrm{h} 2=289 \mathrm{kj} / \mathrm{kg}$, the same for suction line assume the temperature -10 C so enthalpy is $\mathrm{h} 1=245 \mathrm{kj} / \mathrm{kg}$.

Now the refrigerant effect is h2 @ T=-40 C -h1 @ T -10 C =h2 -h1= 244.5-$289=-44.5 \mathrm{kj} / \mathrm{kg}$ So for 1 KW we need $1(\mathrm{Kj} / \mathrm{s}) / 44.5(\mathrm{Kj} / \mathrm{Kg})=0.02 \mathrm{Kg} / \mathrm{s}$; so for 2 refrigeration Ton $=7 \mathrm{Kw}$ we need :
$7 \times 0.02=0.2 \mathrm{~kg} / \mathrm{s}$

## Heat exchanger design:




Figure - $3 \quad U^{*} A^{*} \Delta T m$

Heat transfer, $\mathrm{Q}=$
Wall heat transfer coefficient, $\mathrm{U}=1 /(1 / \mathrm{ho}+\mathrm{L} / \mathrm{K}+1 / \mathrm{h} 1$
Cylindrical wall heat transfer coefficient, $\mathrm{U}=1 /(1 / \mathrm{hi}+[\operatorname{RoLn}(\mathrm{Ro} / \mathrm{Ri})] / \mathrm{K}+\mathrm{Ri} / \mathrm{Ro})$
$i$ and o refer to inside and outside tube
surfaces.

$$
\begin{array}{rlcl}
\text { Large temperature difference, } \Delta \mathrm{Tbc}= & \mathrm{Ta}-\mathrm{Td} & \\
\Delta \mathrm{Tbc}= & 19 & \text { From Input Data below. } \\
\text { Small temperature difference, } \Delta \mathrm{Tad}= & \mathrm{Tb}-\mathrm{Tc} & \\
\Delta \mathrm{Tad} & = & 41 & \text { From Input Data below. } \\
\text { Logarithmic mean temp. difference, } \Delta \mathrm{Tm}= & (\Delta \mathrm{Tbc}-\Delta \mathrm{Tad}) / \ln (\mathrm{Tbc} / \Delta \mathrm{Tad}) \\
\text { Answer: } \Delta \mathrm{Tm}= & 28.6 & \operatorname{deg~C}
\end{array}
$$

The added resistance to heat transfer caused by corrosion is called fouling.
Fouling factor, R ranges between 0.0005 and 0.002 . See manufactures data.
Fouling factor, $\mathrm{R}=$ (1/Udirty) $-(1 /$ Uclean $)$
Forced Convection - in Coiled Tube Heat Exchangers

## Input Data

Turbulent flow in coiled tube.

| Temp. of water flowing in, $\mathrm{Ta}=$ | 85 | $\operatorname{deg} \mathrm{C}$ |
| ---: | :--- | :---: |
| Temp. of water flowing out, $\mathrm{Tb}=$ | 81 | $\left.\begin{array}{rl}\operatorname{deg} \mathrm{C} \\ \mathrm{Tc}= & 40 \\ \operatorname{deg~C} \\ \mathrm{Td}= & 104 \\ \operatorname{deg~C} \\ \text { Tube inside diameter, } \mathrm{Di}= & 13 \\ \mathrm{~mm} \\ \text { Tube outside diameter, } \mathrm{Do}= & 15 \\ \text { Velocity of water in tube, } \mathrm{V}= & 3.5 \\ \mathrm{~mm} \\ \mathrm{~m} / \mathrm{s}\end{array}\right]$ |

Water properties at temperature as per below table:

| Water density, $\rho=$ | 994.7 | $\mathrm{kg} / \mathrm{m}^{\wedge} 2$ |
| :---: | :---: | :---: |
| $\mathrm{Cp}=$ | 4.183 | $\mathrm{k} * / \mathrm{kg} * \mathrm{~K}$ |
| Water dynamic viscosity, $\mu=$ | $6.82 \mathrm{E}-04$ | $\mathrm{kg} / \mathrm{m}$ * s |
| Water conductivity, $\mathrm{k}=$ | 0.6283 | $\mathrm{W} / \mathrm{m} * \mathrm{~K}$ |
| Prandtl number, $\operatorname{Pr}=$ | 4.51 | - |
| Factor, $\mathrm{n}=$ | 0.40 |  |
| Velocity of water in tube, $\mathrm{V}=$ | 3 | $\mathrm{m} / \mathrm{s}$ |
|  | Calculations |  |
| Water bulk temperature, $\mathrm{Tb}=$ | $($ Tin + Tout) / 2 | deg C |
| Answer: $\mathrm{Tb}=$ | 83 | deg C |
| Reynolds Number, $\mathrm{Re}=$ | V*D / v |  |
| or, $\mathrm{Re}=$ | V*D* $/ \mu$ |  |



## Efficiency of the system:

$$
\eta=\frac{q_{\text {evap }}}{w_{\text {comp }}}=\frac{h_{1}-h_{A}}{h_{2 s}-h_{1}}
$$



Figure -4

W comp $=\mathrm{h} 2 \mathrm{~s}-\mathrm{h} 1=288-242=-46 \mathrm{kj} / \mathrm{kg}$
Q evap= $\mathrm{h} 4-\mathrm{h} 1$; as noted @ $\mathrm{T}=90 \mathrm{C} \& \mathrm{P}=10$ bar ,for isenthalpic system h3= $\mathrm{h} 4=38.5 \mathrm{kj} / \mathrm{kg}$
$=38.5-242=-204 \mathrm{kj} / \mathrm{kg}$; so Efficiency is $=204 / 47=4.3$
Now as we used solar energy injected to refrigeration cycle by direct heat exchanging so we have to add the additional gotten enthalpy so,

Qs $(2 S$ to 2$)=\mathrm{h} 2-\mathrm{h} 2 \mathrm{~s}=333-288=45 \mathrm{kj} / \mathrm{kg}$
So Qs +Q evap $=204+45=249 \mathrm{kj} / \mathrm{kg}$
Or
As it's a closed system; The sub cooling add will be the same h2-h2s $=\mathrm{h} 3 \mathrm{~s}-$ h3; by substituting the same parameters then the same values will be gotten.

New COP $=249 / 46=5.5$ which is mean we increase our COP by $25 \%$ from free solar radiation.

However, COP actually is increased more assuming constant pressure during thermal increasing whilst as below I will proof that new pressure has been increased and all values of the above should be taken @ 12 bar instead of 10 bar which is mean COP will be increased up to $35 \%$ more.

## SYSTEM DESIGN

The system will be as the below figure:


Figure - 5

Evacuated heat pipes will be the source of heating, as calculated previously:
Now as previously we got from below equation:
$\mathrm{Q}=500$ GPM Delta T (F)
$15 \mathrm{KW} / 3.5=4 \times 12000=51429$ BTU h 51429/ $(500 \times(185-177)=10$ GPM
Which is $0.7 \mathrm{Kg} / \mathrm{s}$ let say $=1 \mathrm{Kg} / \mathrm{s}$ to be more reasonable.
It seems we need 1 liter in one second so in one minute 60 liters, for factor of safety the tank will be 100 Liter.

Or:
$\mathrm{Pt}=(4.2 \times \mathrm{L} \times \mathrm{T}) \div 3600$. Pt is the power used to heat the water, in $\mathrm{kWh}, \mathrm{L}$ is the number of litters of water that is being heated and T is the difference in temperature from what you started with, listed in degrees Celsius.

We need 4.18 $\mathrm{KWh}=15048 \mathrm{Kj}=\mathrm{m} \times 4.18 \times(82-20)=55$ liter; let say 100 liters.

To get evacuated pipe size:
The capacity of each pipe of 58 mm by 1.8 m length almost a 10 liter so we need a 10 heat pipes. Usually each house of 4 persons are requiring from 200 to 300 -liter pipe collector so a 100 liter for such process could be connected easily without any new required collectors if we consider the diversity as well to guarantee the water outlet requirements and avoid the critical temperature of R134a which is 252 degree Fahrenheit or 122 degrees Celsius.

## Mpemba Effect on the Refrigeration Cycle

The basic advantage is a quicker phase change from gas to liquid leading to a better sub cooling and that to less flash gas then more cooling capacity at the evaporator and with the control of the unit then to the original cooling capacity at the evaporator but with less compressor work. All the known formulas in thermodynamics can't explain, because there is really none for the phase change except the amount of heat transferred at phase change.

We gain here 15 KW using our selected heat exchanger); Now a gas coming from a higher temperature has a much higher kinetic energy of the molecules and they kick much more into them as a result they can develop much quicker the interconnecting van der Waals forces and as a result they liquefy quicker. Since the gas gets quicker liquid, the remaining way in the condenser is used to cool down the fully liquid further.

In order to get new pressure @ $90 \mathrm{C}=363 \mathrm{k}$ of refrigerant; we have to use Van der Waals force consider the below:

$$
\left[P+a\left(\frac{n}{V}\right)^{2}\right]\left(\frac{V}{n}-b\right)=R T
$$

Where a and b are dimensions to be experimentally found. $\mathrm{p}=$ pressure, $\mathrm{n}=$ number of moles, $\mathrm{T}=$ temperature, $\mathrm{V}=$ volume and $\mathrm{R}=$ universal gas constant. Hence Its value will be nearby to general gas equation the given gas is at low pressure and high temperature.
$\mathrm{PV}=\mathrm{nRT} ; \mathrm{V}=\mathrm{As} x \mathrm{~L}$, as the selected exchanger is 0.311 m height,
P @ $90 \mathrm{C}=10 \mathrm{bar}=9.9 \mathrm{~atm} ; \mathrm{V}=0.5 \mathrm{~m} 3$
$\mathrm{R}=0.082$ lt. atm / deg.k.mole
$10 \times 0.5=n \times 0.082 \times 363$
So $\mathrm{n}=0.2$
However as very known from
$\mathrm{P} 1 \mathrm{~V}=\mathrm{n} 1 \mathrm{RT} \ldots ; \quad \mathrm{P} 2 \mathrm{~V}=\mathrm{n} 2 \mathrm{RT} \ldots$ Considering the volume is constant (Isochoric) as the heat exchanger is a small closed tank so: P2 X $0.5=0.2 \mathrm{x}$ $0.082 \times 363 \mathrm{k}$; So P2 = 12 bar .

## DISCUSSIONS AND IDEAS

Based on the below equations and numbers, a rapidly increasing in momentum during of heating the refrigerant gas R 134a:

As an estimation each pressure drop in heat exchanger $4 \mathrm{~m} / 100 \mathrm{~m}$ and we have a 10 meters pipes so we have a $0.4 \mathrm{bar}=4079 \mathrm{kgf} / \mathrm{m} 2(\mathrm{~kg}$ per sq. meter); let us multiply the factors as below:
$0.2 \mathrm{~kg} / \mathrm{s}$ (refrigerant flow rate) $\times 4079 \mathrm{~kg} / \mathrm{m} 2 \times 0.02 \mathrm{~m} 3 / \mathrm{kg}$ (specific volume @ $40 \mathrm{c} \& 2 \mathrm{bar})=16 \mathrm{~kg} . \mathrm{m} / \mathrm{s}$

This unit is a momentum, so let us try to get the momentum @ $90 \mathrm{c} \& 10$ bar:
$0.2 \mathrm{~kg} / \mathrm{s}$ (refrigerant flow rate) $\times 4079 \mathrm{~kg} / \mathrm{m} 2 \times 0.3 \mathrm{~m} 3 / \mathrm{kg}$ (specific volume @ 90 c \& 10 bar$)=245 \mathrm{~kg} . \mathrm{m} / \mathrm{s}$.

This is mean each change in momentum is a new specific impulse on other words this is producing a thrust force which is also a gain for the cycle.

## CONCLUSION

- The system clearly shows that COP is increased from 4.3 to 5.5 , by conveting solar thermal energy to subcooling additional for the refrigertion cycle.
- The system adds more 2 bars to the refrigeration cycle consequently increasing in the entropy as mean a quicker phase change due to quicker set-up of intermolecular bindings because of vV an der Waals forces yielding additional sub-cooling.
- The most important is We don't have to increase condenser surface. It is very well capable to remove the additional heat, the Formula $\mathrm{Q}=$ alpha X surface x delta $T$. Alpha is heat transfer coefficient and does not change, If Delta $T$ increases due to higher temperature of the refrigerant entering the condenser, the condenser just transfers more heat. The environment as the heat absorbing media is capable of taking up that additional heat. For more explanation, let say what happened with radiators: if you flush water with a temperature of $45^{\circ} \mathrm{C}$ through the radiator, it does not provide that much heat as if you flush water through with a temperature of $90^{\circ} \mathrm{C}$ so the radiator is capable of doing the job.

Basically $\mathrm{W}(\mathrm{KJ} / \mathrm{Kg})=\int \mathrm{p}^{*} \mathrm{dv}$, however the dv in heat exchanger considered zero by other words it's an Isochoric Work When V is held constant and P changes $\mathrm{W}=0$ Because gas is not doing any work but as surrounding adds energy to internal energy of gas That's why Cp is always greater than Cv Specific heat capacity at constant pressure is always greater than Specific heat capacity at constant volume, now once the gas goes through the condenser a quicker phase change occurred there cause When gas expands it has to do work to push
environment to create space for its expansion by spending energy from its internal energy that's why suddenly Gas cools down.

## Suggestions

The below figure shows as condensing temperature increased the efficiency of system will be reduced for all refrigerant gasses except R 717 which is rapidly increased, am suggesting to apply this methodology for such refrigeration cycles with R 717.


Figure 6

## REFERENCES:

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- Ron Zevenhoven Åbo Akademi University Thermal and Flow Engineering Laboratory / Värme- och strömningsteknik tel. (02 215)3223; ron.zevenhoven@abo.fi.


## ADDENDUM

## DATA ANALYSIS

The below study shows if reduce refrigerant up to certain limit could be useful for COP, however should we not reach to the maximum temperature of 120 c which is the critical temperature, however power consumption is reduced accordingly.

| Water side |  |  | R 134 a Side |  |  |
| :---: | ---: | ---: | ---: | ---: | ---: |
| $\mathrm{Kw} / \mathrm{k}$ | $\mathrm{kg} / \mathrm{s}$ | $\mathrm{kj} / \mathrm{kg} \cdot \mathrm{k}$ | $\mathrm{Kw} / \mathrm{k}$ | $\mathrm{kg} / \mathrm{s}$ | $\mathrm{kj} / \mathrm{kg} \cdot \mathrm{k}$ |
| Ch | mh | cph | Cc |  | mc |
| cpc |  |  |  |  |  |
| 4.18 | 1 | 4.18 | 0.34 | 0.2 | 1.7 |
| 4.18 | 1 | 4.18 | 0.238 | 0.14 | 1.7 |


| R 134 a Side <br> C degree |  | Water side <br> C degree |  |  |
| ---: | ---: | ---: | ---: | ---: | :---: |
| Delat Tc | Tc in | Tc out | Delta Th |  |
| Th in |  |  |  |  | Th out | 45 | 40 | 85 | 3.660287 |
| ---: | ---: | ---: | ---: |
| 64.28571 | 40 | 104.2857 | 3.660287 |

Table 1
The below are a figure out and illustrations using Mollier diagrams which is a very common for systems with fixed compressor speed = fixed mass flow. DC Inverter units with variable mass flows and many changes in circuit offer wide range of mollier diagrams will be a high improvement for the proposed system.


Figure - 7

Green =without solar collector


Figure -8

Green $=$ without solar collector:


Figure -9

## APPENDIX

Figure -1: Illustration of proposed system (By Author).
Figure -2: P-h diagram (Fundamentals and Applications, Fourth Editionby Yunus A. Çengel and Afshin JGhajar).

Figure -3: Heat exchanger design; (Fundamentals and Applications, Fourth Editionby Yunus A. Çengel and Afshin JGhajar).

Figure -4: T-S diagram (Fundamentals and Applications, Fourth Editionby Yunus A. Çengel and Afshin JGhajar).

Figure -5: Illustration of proposed system (By Author).
Figure -6: Refrigerant selection and COP
Figure -7;8 \& 9: Mollier diagrams illustrations of proposed system.

Table 1: Spread sheet of applying the same equation in excel sheet. (By author).

