# Comparative analysis of a novel geothermal and nanofluid-based solar-driven multigeneration system integrated with high-temperature Kalina cycle: Energy, Exergy, and exergo-economics (3E) analysis 

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#### Abstract

In this comparative study, the thermodynamic and economic investigation of an integrated system driven by a parabolic trough collector and geothermal water is carried out. The proposed multigeneration system is composed of a high-temperature modified Kalina cycle, an electrolyzer, a combined organic Rankine cycle-ejector refrigeration (ORC-EJR) cycle, a reverse osmosis (RO) desalination unit, and a domestic water heater. The absorption fluids applied in the solar collector are $\mathrm{AL}_{2} \mathrm{O}_{3}$ and CuO -based nanofluids, and Therminol VP1 as the base fluid. A comprehensive thermodynamic and exergoeconomic analysis is carried out for the proposed cycle. Engineering Equation Solver (EES) software is used in all conducted simulations. Nanoparticle percentage, solar irradiation, ambient temperature, and collector inlet temperature were the parameters studied to find their effects on the hydrogen production rate, total net power, useful energy achieved, energy and exergy efficiency, collector outlet temperature, and freshwater production rate. The highest outlet temperature of the solar collector was found to be 688.7 K , and the maximum energetic and exergetic efficiencies of the cycle were $34.45 \%$ and $17.25 \%$, respectively. The exergo-economic outputs show that the PEM with $1.99 \$ / h$ has the maximum exergy destruction cost rate. The results depicted that CuO-based nanofluid has better performance from the exergy and exergo-economic viewpoints compared to $\mathrm{AL}_{2} \mathrm{O}_{3}{ }^{-}$ nanofluid. Also, the results proved that the nanoparticle percentage and solar irradiation increases lead to the increase in hydrogen production rate and total net power produced by the system. Moreover, the freshwater production rate increases when the ambient temperature and the collector inlet temperature rise.


Keywords: Thermodynamic analysis, PTC, geothermal, multigeneration system, nanofluids.

## 1. Introduction

In recent years, one of the major worries of society is the rapid declining non-renewable fossil fuel reserves, world population growth,

[^0]environmental pollution, as well as growing energy demand [1]. This concern has led energy sector planners to use more environmentfriendly and more efficient systems and replace centralized power plants with new power generation methods [2]. One of the basic strategies to reach these goals is to use multigeneration systems. In multigeneration systems, in addition to providing electrical
power, waste heat can be used to meet various needs, including the production of cooling, heating, drinking water, and hot water [3]. Utilizing this technology, due to the reduction of many losses that occur when converting thermal energy into mechanical or electrical, can not only significantly increase efficiency but also reduce environmental impact and overall system costs [4].

Numerous investigations have been accomplished in the field of exergy-economic modeling and analysis of multigeneration cycles. Al-Ali and Dincer [5] proposed a novel geothermal-solar system producing power, cooling, heating, heat, and hot water. The results showed a $60 \%$ and $10 \%$ difference between energy and exergy efficiencies when considering sole generation and the proposed multigeneration system. Panchal et al. [6] studied the integrated solar-geothermal cycle, including the Rankine cycle, absorption chiller cycle, and drying process in residential applications. The effects of different parameters were investigated to show the system performance variations. The results cleared that the overall efficiency of the multigeneration cycle is much higher than the single production cycle. A thermo-economic analysis is carried out by Khalid et al. [7] for a renewable energybased multigeneration system to produce power, hot water, hydrogen, cooling, and heating. The proposed cycle was capable of producing 2.7 $\mathrm{kg} / \mathrm{h}$ of hydrogen. Moreover, the levelized cost of electricity was discovered to be $0.089 \$ / \mathrm{kWh}$. Waseem et al. [8] analyzed a solar-geothermal multigeneration system producing hydrogen, power, and cooling. The proposed cycle included an electrolyzer, Rankine cycle, and vapor absorption cycle. Two arrangements were comparatively investigated, and a $0.45 \%$ power efficiency difference was observed. Li et al. [9] developed a system including a geothermal cycle, solar cycle, and PEM system. The system generated electricity, hydrogen, heating, and hot water. To find the irreversibilities, energy, and exergy analyses were done. Compared to the sole geothermal system, the proposed system can produce $148.3 \%$ more power. A novel multigeneration solar-geothermal system producing electricity for a living area was introduced by Sen et al. [10]. The presented system could generate 2900 kW electricity and
$0.0185 \mathrm{~kg} / \mathrm{s}$ hydrogen. The thermodynamic analysis made it clear that the energy and exergy efficiencies of the cycle were $5.90 \%$ and $18.99 \%$, respectively. Atiz [11] investigated an integrated solar-geothermal system for electricity and hydrogen production purposes. The effects of three different solar collectors including parabolic trough solar collectors (PTSCs), evacuated tube solar collectors (ETSCs), and flat plate solar collectors (FPSCs) were studied and compared on the performance of the system. The results showed that the temperature of the geothermal resource is increased by solar collectors as well as the overall energy and exergy efficiencies. The integrated system could produce 414.93 MJ of electricity and 2758.69 g of hydrogen in a day. Sohani et al. [12] studied the effects of price inflation on the optimal performance of a solargeothermal system capable of combined production of hydrogen, power, freshwater, and heat. The results indicated that when inflation rises from 0.05 to 0.20 , annual freshwater production drops by $15 \%$ while annual freshwater increases by $12 \%$. By increasing inflation, the payback period increases from 6.11 to 7.39 years showing the economic viability of such a combined solar-geothermal system. A solar-driven multi-production system of power, cooling, and hydrogen generation was developed and exergetically assessed by Ghorbani et al. [13]. According to the results, the overall energy and exergy efficiencies were $90.77 \%$ and $92.19 \%$, respectively. The proposed hybrid system produced 4.36 MW power, 1.65 MW cooling, and $2026 \mathrm{~kg} / \mathrm{h}$ hydrogen. Mahmoudan et al. [14] investigated a novel multigeneration system driven by geothermal and solar energy. The integrated system was composed of parabolic trough collectors, a Kalina cycle, an ejector refrigeration system, a thermoelectric generator (TEG) unit, an organic Rankine cycle, and a PEM electrolyzer to produce electricity, domestic hot water, a cooling load, and hydrogen. Optimal exergy efficiency, total product cost, and hydrogen production rate were calculated to be $35.2 \%, 37.8 \$ / \mathrm{GJ}$, and $1.9 \mathrm{~kg} / \mathrm{h}$, respectively. A parametric study indicated that applying the TEG unit would enhance system performance. Hashemian and Noorpoor [15] proposed a novel geothermal-biomass-powered
multi-generation system through thermo-economic-environmental aspects. The system applied the Rankine cycle, a dual-effect absorption refrigeration unit, a proton exchange membrane water electrolyzer, and a biomass combustor to produce electricity, heating, cooling, as well as hydrogen and freshwater. They showed that the studied system provided 31.68 MW direct power, $23.3 \mathrm{~m} 3 / \mathrm{h}$ freshwaters, and $88.12 \mathrm{~kg} / \mathrm{h}$ hydrogen, and energetic and exergetic efficiencies were calculated to be $58.54 \%$ and $16.45 \%$. Moreover, the total product cost rate was attained at $1.629 \$ / \mathrm{s}$ and the exergo-environmental impact factor was found to be 0.9 .

The maximum temperature that can be produced by a parabolic trough solar collector (PTC) is $400^{\circ} \mathrm{C}$ with oil and $550^{\circ} \mathrm{C}$ with salt [16]. One of the ways to reach better and higher performance in PTC is by transferring more useful heat to the working fluid by using nanofluids [17]. Several studies considered the use of different nanofluids in PTCs. Subramani et al. [18] studied the thermal efficiency of a solar PTC collector applying Al2O3/DI-H2O nanofluids in different concentrations, mass flows, and turbulent regimes. The results showed that by using nanofluid compared with pure water, the efficiency of the collector increased up to $56 \%$. The thermodynamic analysis of a PTC receiver tube using oilAl 2 O 3 nanofluid with a concentration ratio of 86 was investigated by Mwesigye et al. [19] applying the entropy generation minimization method. The authors proved that by varying the nanofluid temperature from 350 to 600 K , the thermal efficiency of the PTC increased by up to $7.6 \%$. The volume fraction rising led to a reduction in the optimal Reynolds number. In a comparative study, Bellos et al. [20] studied the effect of using three working fluids including thermal oil, thermal oil with nanoparticles, and pressurized water as well as geometry on the thermal efficiency enhancement of the parabolic collector. According to the final results, nanofluid use increased the collector efficiency up to $4.25 \%$ while this amount is $4.55 \%$ for geometry. Wang et al. [21] investigated the application of Al2O3/synthetic oil nanofluid on the performance of the parabolic trough collector systems. The effects of different parameters on
the performance of the PTC were studied. It is discovered that by using the nanofluid, higher efficiencies could achieve and the temperature gradients and the maximum temperature in the absorber were decreased. Furthermore, the varying temperature and DNI caused a reduction in the inlet temperature and the inlet velocity. The comparison of applying $\mathrm{Al}_{2} \mathrm{O}_{3}$ and $\mathrm{Fe}_{2} \mathrm{O}_{3}$-based nanofluids in a parabolic trough solar collector and a parabolic dish solar collector was studied by Abid et al. [22]. The collectors were combined with a Rankine cycle and an electrolyzer to produce electricity and hydrogen. According to the results, the power obtained by the system using a parabolic dish collector is higher than the system using a parabolic trough collector. Compared to the base fluids, applying nanofluids leads to higher power production and better efficiency. Moreover, more hydrogen could be produced when nanofluids were utilized. Ratlamwala et al. [23] studied a solar-geothermal multigeneration system generating power, hydrogen, cooling, hot water, and freshwater. $\mathrm{Al}_{2} \mathrm{O}_{3}$ and $\mathrm{Fe}_{2} \mathrm{O}_{3}$ nanofluids were applied in the solar part of the proposed cycle. According to the results, both of the nanofluids showed excellent heat transfer properties, but $\mathrm{Al}_{2} \mathrm{O}_{3-}$ based nanofluid resulted in better performance compared with $\mathrm{Fe}_{2} \mathrm{O}_{3}$-based nanofluid. Furthermore, when the nanoparticle percentage increased, the amount of power and hydrogen rate production decreased. Bellos et al. [24] surveyed the thermal increment of parabolic trough collectors when the Syltherm $800 / \mathrm{Cu}$ nanofluid was applied. The investigation of three different collectors revealed that for cermet coating and $25 \mathrm{~L} /$ min flow rate, the amount of increment for the bare tube was $7.16 \%$, for the non-evacuated receiver was $4.87 \%$ and for the evacuated receiver was $4.06 \%$. Khan et al. [25] studied the effect of applying nanofluids in a solar-based cycle from energy, exergy, and exergo-environmental point of view. The influence of three nanofluids, including Fe2O3/Therminol VP1, SiO2/Therminol VP1, and $\mathrm{Cu} /$ Therminol VP1, were compared by considering different parameters. The results showed that among the studied nanofluids, $\mathrm{SiO} 2 / \mathrm{VP} 1$ presented better outcomes. Moreover, the amount of CO 2 was reduced for all the studied nanofluids.

Increasing nanoparticle concentration leads to an increment in the exergy efficiency and a decrease in the nanoparticle thermal conductivity. Ibrahim and Kayfeci [26] studied using two kinds of nanofluids in a parabolic trough solar collector of a trigeneration system. The results showed that by using ferrofluid and graphene as the nanoparticles, the overall performance of the proposed system improved. Graphene nanoparticles indicated better performance in comparison with ferrofluid nanoparticles. Allouhi et al. [27] studied the effect of using different nanoparticles for medium and high-temperature PTCs mathematically. They concluded that high operating temperatures are more appropriate for using nanofluids in PTCs as higher gains of energy are achieved. The maximum exergy efficiency of CuO-based nanofluid was calculated to be about $9.05 \%$. Kalbande et al. [28] studied the possibility of attaining a higher temperature range of above $200^{\circ} \mathrm{C}$ in oil-based thermal energy storage systems. They used aluminum oxide ( Al 2 O 3 ) and soybean oil nanofluid as the heat transfer fluid to store and transfer the sensible heat. After a mathematical analysis, they could achieve up to $220^{\circ} \mathrm{C}$ in a parabolic trough solar collector coupled with a thermal storage system. Tonekaboni et al. [29] studied the amount of enhancement of the solar collectors by adding porous media and nanofluid. They applied $90 \%$ porosity copper, CuO , and $\mathrm{Al2O} 3$ nanofluids to investigate the changing in the thermal properties of different types of solar collectors. The results showed that adding porous media and nano-fluids increased an average of $14.4 \%$ collector energy efficiency and $8.08 \%$ collector exergy efficiency. Moreover, the highest amount of energy and exergy efficiencies were $60.12 \%$ and $18.84 \%$, respectively which were obtained for parabolic solar collectors. Said et al. [30] surveyed a nanofluid-based parabolic trough collector from energy, exergy, economic, and environmental points of view. The results showed that by adding Ti3C2 to the silicon oil, the thermal conductivity of the nanofluid enhances from $70 \%$ to $89 \%$. It is found that, compared to pure oil, applying nanofluid reduces the system's cost by $0.021 \mathrm{M} \$$ and increases the gained energy by $1.51 \%$. For
energy analysis, the annual CO 2 reduction varies from 2.25 to 2.30 tons $\mathrm{CO} 2 /$ year.

According to the literature review, there is a great concern about applying the nanofluid in parabolic trough solar collectors. On the other hand, publications investigating the effects of nanofluids in high-temperature PTCs are rare. Moreover, to the best knowledge of the authors, there is no study of the combination of hightemperature Kalina cycle and nanofluid applied parabolic trough solar collectors. In this research a novel multigeneration system including solar and geothermal energy sources where $\mathrm{Al2O} 3-$ based and CuO -based nanofluids were used as working fluids in PTC. The proposed system contains a high-temperature modified Kalina cycle, an electrolyzer, a combined ORC-EJR cycle, an RO desalination unit, and a domestic water heater.

A summary of the main goals and novelties of this research is presented below:

- Introducing a new multigeneration energy system using solar and geothermal renewable sources.
- Applying a modified high-temperature Kalina cycle for retrieving the solar source energy.
- Presenting the overall performance of the proposed novel multigeneration system.
- Energy, exergy, and exergo-economic analysis of the proposed systems are applied to perform a comprehensive evaluation of the system.
- Comparing the use of three working fluids to analyze the performance of the system.
- Producing hydrogen and fresh water from the novel energy system.


## NOMENCLATURE

| $A$ | Area (m2) |
| :--- | :--- |
| $C$ | Cost per exergy unit (\$/GJ) |
| $C$ | Cost rate (\$/h) |
| $C_{p}$ | Specific heat capacity (J/kgK) |
| CRF | Capital recovery factor |
| cooling | Cooling |
| con | Condenser |
| $D$ | Destroyed |
| DWH | Domestic water heater |
| eje | Ejector |
| electric | Electricity |


| eva | Evaporator | $Z$ | Capital cost rate (\$/h) |
| :---: | :---: | :---: | :---: |
| exv | Expansion valve |  |  |
| $E$ | Energy input (kW) | Subscripts |  |
| $E x$ | Exergy rate (W) | 0 | Ambient |
| $F$ | Faraday constant (C/mol) | 1,2,.. | State points |
| $f_{k}$ | Exergoeconomic factor (\%) | $a$ | Anode |
| $F$ |  | $a c t$ | Activation |
| ${ }_{1}$ | Collector efficiency factor | $a p$ | Aperture |
| $F_{R}$ | Heat transfer factor | avg | Average |
| $f$ | Feed water | $b f$ | Base fluid |
| $\mathrm{H}_{2}$ | Hydrogen | $c$ | Cathode |
|  | Hydrogen | ch | Chemical |
| $\mathrm{H}_{2} \mathrm{O}$ | Water | $k$ | Kth component |
| $H X$ | Heat exchanger | KAL | Kalina |
| in | Inlet | mx | Mixer |
| $G_{b}$ |  | net | net |
| $h$ | Solar irradiation (W/m2) | $n f$ | nanofluid |
| $h$ | Heat transfer coefficient (W/m2K) | $n p$ | nanoparticle |
| $J$ | Current density (A/m2) | out | Outlet |
| $J_{0}$ | Exchange current density ( $\mathrm{A} / \mathrm{m} 2)$ | $p$ | Pump |
| $J_{a}^{r e f}$ | Pre - exponential factor ( $\mathrm{A} / \mathrm{m} 2$ ) | $\begin{aligned} & P E M \\ & \text { ph } \end{aligned}$ | Proton exchange membrane Physical |
| $K$ | The ratio of specific heats (CP/CV) |  | Physical |
| $k$ | Thermal conductivity (W/m K) | prh | Preheater |
| $L$ | Collector length (m) | $P T C$ | Parabolic trough collector |
| $M$ | Molecular weight ( $\mathrm{kg} / \mathrm{mol}$ ) | $r$ | Receiver tube |
| $m$ | Mass flow rate (kg/s) | re | Recuperator |
| $N_{x}$ | Outlet flow rate of fluid $\mathrm{x}(\mathrm{kg} / \mathrm{s})$ | reacted $r, i$ | Entering water of PEM electrolyzer Receiver inlet |
| $n_{c s}$ | Number of collectors in series | $r, o$ | Receiver outlet |
| $n_{c p}$ | Number of collectors in parallels | RO | Reverse osmosis |
| $P$ | Pressure (bar) | ORC- | Organic Rankine Cycle-Ejector |
| Q | Heat transfer rate (W) | EJR | Refrigeration cycle |
|  |  | sep | Separator |
| $Q_{u}$ | Useful energy gain | spl | Splitter |
| $R$ | Overall ohmic resistance | sun | sun |
| $R R$ | Recovery ratio | $t$ | Turbine |
| $r_{k}$ | Relative cost difference (\%) | th | Thermal |
| $S$ | Absorbed solar radiation | $v g$ | Vapor generator |
| $T$ | Temperature (K) | W | Work |
| TCF | Temperature correction factor | Greek Sym |  |
| $U_{L}$ | Solar collector's overall heat loss coefficient | $\alpha$ | Absorptivity of receiver |
| $V$ | Overpotential (V) | $\gamma$ | Correction factor for diffuse radiation |
| $V_{0}$ | Reversible potential (V) | $\sigma(x)$ | Ionic conductivity of the membrane |
| $V_{\text {ohm }}$ | Ohmic overpotential of the electrolyte (V) | $\eta$ | Efficiency Dynamic Viscosity |
| $W$ | Net output power (kW) | $\lambda(x)$ | Water content at location x |
| W | Collector width (m) | $\rho$ | Density (kg/m3) |
| $X$ $Z$ | Salt concentration (g/kg) The capital cost of a component (\$) | $\tau_{c}$ | Transmissivity of the cover glazing |


| $\tau_{p}$ | Effective transmissivity of the <br> parabolic trough collector |
| :--- | :--- |
| $\phi$ | Nanoparticle volume concentration |

## 2. System description

Figure 1 presents the schematic of the considered multigeneration system. As is shown, the principal components of the system are a low-temperature geothermal source, a nanofluid solar system-based modified Kalina system, an RO desalination unit, a combined ORC-EJR cycle, a PEM electrolyzer, and a domestic water heater. The outcomes of the proposed system are electricity, fresh water, hydrogen, hot water, heating, and cooling. The primary energy sources of the system are geothermal and solar cycles. The solar cycle employed in this system has two main functions: to act as the high-temperature energy source of the Kalina cycle and to increase the temperature of the flow extracted
from the ground. The parabolic trough collector absorbs energy by applying $\mathrm{Al}_{2} \mathrm{O}_{3}$, and CuO nanoparticles added to the Therminol_VP1 as the base fluid. The geothermal water, after gaining energy from the heat exchanger of the solar system, first enters the vapor generator of the combined ORC-EJR cycle to supply the cycle's energy needed to produce power. Then, it discharges into the reinjection well, passing the domestic water heater, which produces hot water. The generated power in the modified Kalina cycle is used as the electricity in the residential area and the source power of the RO desalination unit to purify the sea water and supply fresh water. The combined ORC-EJR cycle produces a cooling effect and power. The power generated in the combined ORC-EJR cycle is divided into two streams: employed in the residential area and applied in the PEM electrolyzer to produce hydrogen.


Fig. 1. Schematic diagram of the proposed multigeneration energy system

## 3. Mathematical modeling

The mathematical simulation evaluates the performance, energy, and exergy efficiency, and exergy destruction rate of the system. Engineering Equation Solver (EES) [31] was used to determine the properties of the working fluids and solve the relations of all components of the system. For simplifying the modeling process, some considerations are taken as follows [32,33]:

- The system functions under steady-state conditions.
- The changes in potential and kinetic energies, as well as pressure losses in the pipes and heat exchangers, are ignored.
- All turbines, pumps, condensers, and valves are assumed adiabatic systems.
- The outlet streams of the condensers and evaporators are considered to be saturated liquid and saturated vapor, respectively.
- Isentropic efficiency is assumed for pumps and turbines.
- Solar irradiation is presumed to be uniform and steady-state.
- Geothermal hot water is considered to be net water with no pollution.
To model the multigeneration cycle, the primary input and design values are exhibited in Table 1.

Table 1. The input parameters for the modeling of the present study

| Parameters | Unit | Value |
| :---: | :---: | :---: |
| GEOTHERMAL [34,35] Gn |  |  |
| Production well temperature, $T_{1}$ | ${ }^{\text {c }} \mathrm{C}$ ) | 120 |
| Production well Pressure, $P_{1}$ | (bar) | 7 |
| SOLAR [36-38] |  |  |
| Collector width, $\mathcal{W}$ | (m) | 5.76 |
| Collector length, $L$ | (m) | 1227 |
| Receiver outside diameter, $D_{o, r}$ | (m) | 0.07 |
| Receiver inside diameter, $D_{i, r}$ | (m) | 0.066 |
| Collector heat loss coefficient, $U_{L}$ | $\left(\frac{W}{m^{2} C}\right)$ | 3.82 |
| Receiver inlet temperature, $T_{n}$ | (c) | 180 |
| The heat transfer coefficient inside the receiver, $h_{f i}$ | $\left(\frac{W}{m^{2} C}\right)$ | 300 |
| The thermal conductivity of the receiver, $K$ | $\left(\frac{W}{m^{2} C}\right)$ | 16 |
| Solar radiation intensity, $G_{b}$ | $\left(\frac{W}{m^{2}}\right)$ | 850 |
| Cover glazing transmissivity, $\tau_{c}$ | - | 0.96 |
| PTC effective transmissivity, $\tau_{p}$ | - | 0.94 |
| Receiver absorptivity, $\boldsymbol{\alpha}_{r}$ | - | 0.96 |
| Correction factor for diffuse radiation, $\gamma$ | - | 0.95 |
| KALINA [39] |  |  |
| Turbine inlet ammonia concentration, $x_{9}$ |  | 0.49 |
| Turbine inlet temperature, $T_{9}$ | $\left({ }^{\circ} \mathrm{C}\right)$ | 330 |
| Turbine inlet pressure, $P_{9}$ | (bar) | 120 |
| RO[40,41] |  |  |
| Recovery ratio, $R R$ |  | 0.3 |
| Number of elements, $n_{e}$ |  | 7 |
| Number of pressure vessels, $n_{v}$ |  | 42 |
| Seawater salinity, $X_{f}$ |  | 43 |
| ORC-EJR [42-44] |  |  |
| Working fluid Turbine inlet pressure, $P_{36}$ |  |  |
| Evaporator temperature, $T_{44}$ |  | -5 |

### 3.1.Energy and exergy analysis

The main conservation equations, mass, and energy, employed in analyzing the proposed system are as follows [45]:

$$
\begin{align*}
& \sum m_{\mathrm{n}}=\sum \mathrm{m}_{\text {out }}  \tag{1}\\
& \sum(\mathrm{m} h)_{o u t}-\sum(\mathrm{m} h)_{\text {in }}=Q-W  \tag{2}\\
& E x=E x_{p h}+E x_{c h}  \tag{3}\\
& E x_{\text {ph }}=\sum m_{n t}\left[\left(h_{\text {in }}-h_{\text {out }}\right)-T_{o}\left(s_{i n}-s_{o u t}\right)\right] \tag{4}
\end{align*}
$$

$E x_{c h}=$
$m_{m}\left[\left(\frac{x}{M_{N H 3}}\right) e x_{c h, N H 3}^{o}+\left(\frac{1-x}{M_{H_{2} \mathrm{O}}}\right) e x_{c h, H_{2} \mathrm{O}}^{o}\right]$
In the above-mentioned equations, $x$ is the molar fraction of ammonia, and $c h$ and $p h$ represent the chemical, and physical exergy, respectively.

By employing the energy and exergy balance relations for the multigeneration system, the transferred heat and power, and exergy destruction rate of each component can be computed as listed in Table 2.

Table 2. Energy conservation and exergy destruction rate relations for the plant's elements

| Component | Energy balance equations | Exergy destruction rate equations |
| :---: | :---: | :---: |
| PTC field | $m_{8} h_{8}+Q_{u}=m_{5} h_{5}$ | $E x_{D, P T C}=E x_{\text {slu }}+E x_{8}-E x_{5}$ |
| Heat exchanger | $Q_{H X}=m_{1}\left(h_{2}-h_{1}\right)=m_{7}\left(h_{7}-h_{8}\right)$ | $E x_{D, H X}=E x_{1}+E x_{7}-E x_{2}-E x_{8}$ |
| Kalina evaporator | $Q_{\text {eva, }, \text { AAL }}=m_{5}\left(h_{5}-h_{6}\right)=m_{5}\left(h_{9}-h_{29}\right)$ | $E x_{D, e v a, K A L}=E x_{5}+E x_{29}-E x_{6}-E x_{9}$ |
| Kalina turbine | $W_{t, K A L}=m_{9}\left(h_{9}-h_{10}\right)$ | $E x_{D, t, K A L}=E x_{9}-W_{t, K A L}-E x_{10}$ |
| Kalina recuperator 1 | $Q_{r e 1, K A L}=m_{10}\left(h_{10}-h_{11}\right)=m_{28}\left(h_{29}-h_{28}\right)$ | $E x_{D, r e 1, K A L}=E x_{10}+E x_{28}-E x_{11}-E x_{29}$ |
| Kalina recuperator 2 | $Q_{\text {re2,KAL }}=m_{11}\left(h_{11}-h_{12}\right)=m_{18}\left(h_{19}-h_{18}\right)$ | $E x_{D, r e 2, K A L}=E x_{11}+E x_{18}-E x_{12}-E x_{19}$ |
| Kalina recuperator 3 | $Q_{\text {re } 3, K A L}=m_{20}\left(h_{20}-h_{21}\right)=m_{27}\left(h_{28}-h_{27}\right)$ | $E x_{D, r e 3, K A L}=E x_{20}+E x_{27}-E x_{21}-E x_{28}$ |
| Kalina recuperator 4 | $Q_{\text {re } 4, K A L}=m_{17}\left(h_{17}-h_{18}\right)=m_{22}\left(h_{23}-h_{22}\right)$ | $E x_{D, r e 4, K A L}=E x_{17}+E x_{22}-E x_{18}-E x_{23}$ |
| Kalina mixer 1 | $m_{12} h_{12}+m_{24} h_{24}=m_{13} h_{13}$ | $E x_{D, m x 1, K A L}=E x_{12}+E x_{24}-E x_{13}$ |
| Kalina mixer 2 | $m_{16} h_{16}+m_{21} h_{21}=m_{25} h_{25}$ | $E x_{D, m \times 2, K A L}=E x_{16}+E x_{21}-E x_{25}$ |
| Kalina condenser 1 | $Q_{\text {con } 1, K A L}=m_{13}\left(h_{13}-h_{14}\right)$ | $E x_{D, \text { cnn }, \text { KAL }}=E x_{13}-E x_{14}-Q_{\text {conn }, \text { KAL }}\left(1-\frac{T_{0}}{T_{14}}\right)$ |
| Kalina condenser 2 | $Q_{\text {con } 2, K A L}=m_{25}\left(h_{25}-h_{26}\right)$ | $E x_{D, \text { onn } 2, K A L}=E x_{25}-E x_{26}-Q_{\text {conn } 2, K A L}\left(1-\frac{T_{0}}{T_{26}}\right)$ |
| Kalina pump 2 | $W_{p 2, K A L}=m_{14}\left(h_{15}-h_{14}\right)$ | $E x_{D, p 2, K A L}=W_{p 2, K A L}+E x_{14}-E x_{15}$ |
| Kalina pump 3 | $W_{p 3, K A L}=m_{26}\left(h_{27}-h_{26}\right)$ | $E x_{D, p 3, K A L}=W_{p 3, K A L}+E x_{26}-E x_{27}$ |
| Kalina splitter | $m_{15} h_{15}=m_{16} h_{16}+m_{17} h_{17}$ | $E x_{D, s p l, K A L}=E x_{15}-E x_{16}-E x_{17}$ |
| Kalina separator | $m_{19} h_{19}=m_{20} h_{20}+m_{22} h_{22}$ | $E x_{D, \text { sep }, K A L}=E x_{19}-E x_{20}-E x_{22}$ |
| Kalina expansion valve 1 | $h_{23}=h_{24}$ | $E x_{D, e x v 1, K A L}=E x_{23}-E x_{24}$ |
| ORC generator | $Q_{v g, \text { ORC-ERC }}=m_{2}\left(h_{2}-h_{3}\right)=m_{35}\left(h_{36}-h_{35}\right)$ | $E x_{D, v g, O R C-E R C}=E x_{2}+E x_{35}-E x_{3}-E x_{36}$ |
| ORC turbine | $W_{t, O R C-E R C}=m_{36}\left(h_{36}-h_{37}\right)+m_{38}\left(h_{37}-h_{38}\right)$ | $E x_{\text {D, }, \text { ORC }- \text { RRC }}=E x_{36}-W_{t, O R C-E R C}-E x_{37}-E x_{38}$ |
| ORC ejector | $\mu_{\text {eje }}=\frac{m_{45}}{m_{37}}$ | $E x_{D, \text { eje,ORC-ERC }}=E x_{37}+E x_{45}-E x_{39}$ |
| ORC preheater | $Q_{p m, \text { ORC-ERC }}=m_{34}\left(h_{35}-h_{34}\right)=m_{40}\left(h_{40}-h_{41}\right)$ | $E x_{\text {D,ph }, \text { ORC-ERC }}=E x_{34}+E x_{40}-E x_{35}-E x_{41}$ |
| ORC pump 4 | $W_{p, O R C-E R C}=m_{33}\left(h_{34}-h_{33}\right)$ | $E x_{D, p 4, \text { ORC-ERC }}=W_{p 4, \text { ORC-ERC }}-E x_{33}+E x_{34}$ |
| ORC condenser 3 | $Q_{\text {con } 3, O R C-E R C}=m_{41}\left(h_{41}-h_{42}\right)=m_{48}\left(h_{49}-h_{48}\right)$ | $E x_{D, \text { Con } 3, O R C-E R C}=E x_{41}+E x_{48}-E x_{42}-E x_{49}$ |
| ORC expansion valve 2 | $h_{43}=h_{44}$ | $E x_{D, \operatorname{exv} 2, O R C-E R C}=E x_{43}-E x_{44}$ |
| ORC evaporator | $Q_{\text {eva,ORC-ERC }}=m_{44}\left(h_{45}-h_{44}\right)$ | $E x_{D, E V, O R C-E R C}=E x_{44}+E x_{46}-E x_{45}-E x_{47}$ |
| PEM | $W_{\text {PEM }}=\left(m_{52} h_{52}-m_{53} h_{53}-m_{54} h_{54}\right)$ | $E x_{D, P E M}=E x_{52}+W_{P E M}-E x_{53}-E x_{54}$ |
| DWH | $Q_{\text {DWH }}=m_{3}\left(h_{3}-h_{4}\right)=m_{50}\left(h_{51}-h_{50}\right)$ | $E x_{D W H}=E x_{3}+E x_{50}-E x_{4}-E x_{51}$ |
| RO | $W_{R O}=\left(m_{30} h_{30}-m_{31} h_{31}-m_{32} h_{32}\right)$ | $E x_{D, R O}=E x_{30}-E x_{31}-E x_{32}$ |

The equations used for the simulation of the RO unit and PEM electrolyzer are expressed in Tables 3 and 4, respectively.
3.2.Calculation of nanofluid heat transfer properties

In this study, $\mathrm{Al}_{2} \mathrm{O}_{3}$ and CuO nanoparticles have been applied inside Therminol_VP1 as the base fluid. The thermal properties of the nanoparticles are presented in Table 5.

Compared to the base fluids, the nanofluids offer different thermal properties, which can be calculated by the following formulas, applying the properties of the nanoparticles $(n p)$ and the base fluid (bf).

The density of the nanofluid $(\rho)$ is presented as [49]:

$$
\begin{equation*}
\rho_{n f}=\varphi \cdot \rho_{n p}+(1-\varphi) \cdot \rho_{b f} \tag{6}
\end{equation*}
$$

in which $\varphi$ is the volume concentration of the nanoparticle.

Table 3. RO unit required modeling relations [46]

| The recovery ratio | $R R=\frac{n \frac{n}{n}}{n+\frac{1}{6}}$ |
| :---: | :---: |
| Saline water flow rate |  |
| Osmotic pressure | $P_{\text {net }}=P_{\text {avg, } f}-75.85 \times X_{31}$ |
| Average Osmosis pressure | $P_{\text {cog }, f}=\frac{P_{30}+P_{32}}{2}=37.92 \times\left(X_{30}+X_{32}\right)$ |
| Temperature correction factor | $T C F=\exp \left\{2700 \times\left(\frac{1}{T+273}-\frac{1}{298}\right)\right\}$ |
| Membrane water permeability | $K_{w}=\frac{6.84 \times 10^{-8} \times\left(18.6865-0.177 \times X_{32}\right)}{(T+273)}$ |
| High-pressure pump power | $W_{p, R O}=\frac{m_{30} \times \Delta P}{\rho_{30} \times \eta_{p}}$ |

Table 4. PEM electrolyzer required modeling relations [47]

| Electrical energy consumption | $E_{\text {electric }}=J V$ |
| :---: | :---: |
| Electrolyzer voltage | $V=V_{0}+V_{\text {act }, c}+V_{\text {act }, a}+V_{\text {olm }}$ |
| Reversible equation | $V_{0}=1.229-0.00085\left(T_{\text {PEM }}-298\right)$ |
| Activation overpotential | $A_{u c t, i}=\frac{R T}{F} \sinh ^{-1}\left(\frac{J}{2 J_{0, i}}\right)=J_{a}^{r e f} \exp \left(\frac{-E_{a c t, i}}{R T}\right), i=a, c$ |
| Ohmic overpotential | $\begin{aligned} & V_{\text {olm }}=J R_{P E M}, R_{P E M}=\int_{0}^{L} \frac{d x}{\sigma[\lambda(x)]}, \lambda(x)=\frac{\lambda_{a}-\lambda_{t}}{D} x+\lambda_{t} \\ & \sigma[\lambda(x)]=[0.5139 \lambda(x)-0.326] \exp \left[1268\left(\frac{1}{303}-\frac{1}{T}\right)\right] \end{aligned}$ |
| Rate of produced $\mathrm{H}_{2}$ | $N_{H_{2}, \text { Out }}=\frac{J}{2 F}=N_{H_{2} \mathrm{O}, \text { reacted }}$ |

Table 5. Properties of the studied nanoparticles [48]

| Particle | $\boldsymbol{\rho}\left(\mathbf{k g} / \mathbf{m}^{3}\right)$ | $\boldsymbol{k}(\mathbf{W} / \mathbf{m K})$ | $\boldsymbol{c}_{\boldsymbol{\rho}}(\mathbf{k J} / \mathbf{k g K})$ |
| :---: | :---: | :---: | :---: |
| $\mathrm{Al}_{2} \mathrm{O}_{3}$ | 3970 | 40 | 0.765 |
| CuO | 6320 | 77 | 0.532 |

The specific heat capacity of the nanofluid $\left(C_{p}\right)$ can be expressed as [50]:
$c_{p, n f}=\frac{\rho_{n p} \cdot \varphi \cdot c_{p, n p}+\rho_{f f}(1-\varphi) \cdot c_{p, b f}}{\rho_{n f}}$
The nanofluid thermal conductivity $(k)$ is calculated as [51]:
$k_{n f}=k_{b f} \cdot \frac{k_{n p}+2 k_{b f}+2\left(k_{n p}-k_{b f}\right) \cdot(1+\beta)^{3} \cdot \varphi}{k_{n p}+2 k_{b f}-\left(k_{n p}-k_{b f}\right) \cdot(1+\beta)^{3} \cdot \varphi}$
In equation (8), $\beta$ is defined as the nanolayer thickness to the original particle radius and usually is taken as 0.1 [52].

The dynamic viscosity of the nanoparticle $(\mu)$ is estimated by the following correlation [53]:

$$
\begin{equation*}
\mu_{r f}=\mu_{b f} \cdot\left(1+2.5 \varphi+6.5 \varphi^{2}\right) \tag{9}
\end{equation*}
$$

The thermal properties of the nanofluids are displayed in Fig. 2. In the following figures, the properties, including the density, the specific heat capacity, and the thermal conductivity, are presented for different nanoparticle concentrations.


(c)

Fig. 2. Thermal properties of the studied nanofluids a) density, b) specific heat capacity, c) thermal conductivity

### 3.3 Parabolic trough collector

By passing through Parabolic trough collectors, the temperature of the geothermal fluid increases, which is calculated by the following equations [54,55]:
$Q_{u}=n_{c p} n_{c s} F_{R} A_{q p}\left[S-\frac{A_{t}}{A_{t p}} U_{L}\left(T_{r, i}-T_{0}\right)\right]$
where $n_{c p}$ and $n_{c s}$ are the numbers of collectors in series parallels, respectively. Also, $A$ and $F_{R}$ are the area and the heat removal factor. Moreover, $U_{L}$ expresses the total heat loss coefficient of the PTS collector. $S$ is absorbed solar radiation and is defined as:

$$
\begin{equation*}
S=G_{b} \eta_{r} \tag{11}
\end{equation*}
$$

$\eta_{r}=\gamma \tau_{c} \tau_{p} \alpha$
receiver, and $\tau_{c}$ and $\tau_{p}$ are the transmissivity of the cover glazing and effective transmissivity of the parabolic trough collector, respectively.

### 3.4 Exergo-economic analysis

To have a better insight into the overall system efficiency, the exergo-economic analysis should be determined, as the thermodynamic assessment alone is not enough to study the energy systems. The exergoeconomic analysis combines concepts of both exergy and economic evaluations and determines the cost per unit of products of the energy conversion system. The general equation for calculating the cost balance for each component can be defined as [56]:
$\eta_{\text {lh }, \text { ot }}=\frac{W_{K A L}+W_{\text {ORC-ERC }}+Q_{\text {cooding }}+Q_{\text {DWH }}+m_{53} H H V_{H 2}-W_{P E M}+m_{31} h_{31}-W_{R O}}{Q_{u}+m_{1} h_{1}}$
$\eta_{\text {ex,tot }}=\frac{W_{K A L}+W_{\text {ORC-ERC }}+E x_{x_{\text {odding }}}+E x_{53}+E x_{54}+E x_{51}-E x_{50}+E x_{31}}{E x_{\text {in, son }}+E x_{1}}$
in which $\gamma$ is the correction factor for diffuse radiation, $\alpha$ is the absorptivity of the

To find $F_{R}$ and $F_{l}$ the following equations can be used:
$F_{R}=\frac{m c_{p, c}}{A_{r} U_{L}}\left[1-\exp \left(-\frac{A_{r} U_{L} F_{l}}{m c_{p, c}}\right)\right]$
$F_{1}=\frac{\frac{1}{U_{L}}}{\frac{1}{U_{L}}+\frac{D_{r, 0}}{h_{f i}}+\left(\frac{D_{r, 0}}{2 k} \ln \frac{D_{r, 0}}{D_{r, i}}\right)}$
where $F_{l}$ is the collector efficiency factor, $h_{i}$ is the heat transfer coefficient of the inlet fluid and $D$ is the diameter.

The surface area of PTC is:

$$
\begin{equation*}
A_{q p}=(w-D) L \tag{15}
\end{equation*}
$$

in which $w$ is the width and $L$ is the length of the PTC.

Finally, the overall energy efficiency and second law efficiency of the proposed multigeneration system can be defined by the following equations:
$\sum_{\text {out }} C_{\text {out }, k}+C_{w, k}=\sum_{\text {in }} C_{i n, k}+C_{q, k}+Z_{k}$
$C_{\text {out }, k}$ :The outlet stream cost rate of components
$C_{w, k}:$ The work cost rate
$C_{i n, k}$ :The inlet stream cost rate of components
$C_{q, k}$ : The heat transfer cost rate
$Z_{k}$ indicated the purchase equipment cost rate of each component that can be expressed through the following equation [57]:

$$
\begin{equation*}
Z_{k}=\frac{Z_{k} \times \varphi \times C R F}{N} \tag{19}
\end{equation*}
$$

$\varphi$ :the operation and maintenance factor which is assumed as 1.06 .
$\tau$ :the time operation of the system which is assumed as 8000 .

CRF is the capital recovery factor that is defined as [58]:
$C R F=\frac{i_{r}\left(1+i_{r}\right)^{n}}{\left(1+i_{r}\right)^{n}-1}$
in which, $i$ and $n$ are interest and lifetime that are assumed as $12 \%$ and 20 years, respectively.

The cost and exergy are related by the specific exergy cost $\left(c_{j}\right)$ which is calculated as:

$$
\begin{equation*}
C_{j}=c_{j} E x_{j} \tag{21}
\end{equation*}
$$

In the exergo-economic analysis, there are some essential parameters such as the average cost per unit exergy of fuel $\left(c_{f, k}\right)$ and product $\left(c_{p, k}\right)$, and the exergy destruction cost rate $\left(C_{D, k}\right)$ which can be defined as follows [59]:
$c_{f, k}=\frac{C_{f, k}}{E x_{f, k}}$
$c_{p, k}=\frac{C_{p, k}}{E x_{p, k}}$
$C_{D, k}=c_{f, k} \times E x_{D, k}$
The exergo-economic factor $\left(f_{k}\right)$ indicates the ratio of the investment cost rate to the summation of investment cost and exergy destruction cost can be calculated as follows [60]:
$f_{k}=\frac{Z_{k}}{Z_{k}+C_{D, k}}$

The relative cost difference $\left(r_{k}\right)$ represents the relative increase in the average cost per exergy unit between fuel and product of the component [60]:

$$
\begin{equation*}
r_{k}=\frac{c_{p, k}-c_{f, k}}{c_{f, k}} \tag{26}
\end{equation*}
$$

Table 6 contains the cost functions of each component, which depend on some thermodynamic parameters. For performing the economic analysis, the cost balance is carried out for each element. Moreover, some auxiliary equations are needed to solve the equations. Table 7 includes the cost balance and auxiliary equations of the multigeneration system.

## 4. Validation

As the proposed multigeneration system is novel, therefore, to certify the accuracy of the modeling, some of the main parts of the system have been validated by previous studies. The results of the RO desalination unit have been compared with the results of Nafey and Sharaf [40] in Table 8. Moreover, Fig. 3 displays the comparison between the results of this study and the work of Zare and Moalemian [65]. According to the table and graphs, a reasonable agreement can be seen between the present modeling and the previous findings.

Table 6. Purchased equipment costs of components [61-64]

| Component | Purchase cost $(\$)$ |
| :---: | :---: |
| PTC field | $Z_{\mathrm{TIC}}=240 \mathrm{~A}_{\mathrm{ap}}$ |
| Heat exchangers | $Z_{H X}=130\left(\frac{A}{0.093}\right)^{0.78}$ |
| Evaporator | $Z_{e v a}=1397\left(A_{e v a}\right)^{0.89}$ |
| Turbine | $Z_{t}=4405\left(W_{t}\right)^{0.7}$ |
| Pump | $Z_{p}=1120\left(W_{p}\right)^{0.8}$ |
| Expansion valve | $Z_{\mathrm{exv}}=114.5 \mathrm{~m}_{\mathrm{xv}}$ |
| Vapor generator | $Z_{v g}=1397(A)^{0.89}$ |
| Ejector | $Z_{\text {eje }}=750 m_{40}\left(\frac{T_{45}}{P_{45}}\right)^{0.05}\left(P_{40}\right)^{-0.75}$ |
| PEM | $Z_{P E M}=1000\left(W_{p e m}\right)$ |
| RO | $Z_{R O}=n_{e} n_{p} c_{k}+n_{p} c_{p}+996\left(m_{31}\right)^{0.8}$ |

Table 7. The cost balance and auxiliary equations of each component of the proposed system

| Component | Cost balance equation | Auxiliary equation |
| :---: | :---: | :---: |
| PTC field | $C_{s s m}+C_{8}+Z_{\text {PTC }}=C_{5}$ | $c_{5}=c_{8}, c_{s s m}=0$ |
| Heat exchanger | $C_{1}+C_{7}+Z_{H X, \text { geo }}=C_{2}+C_{8}$ | $c_{1}=5\left\lfloor\frac{\$}{G J}\right\rfloor, c_{1}=c_{2}$ |
| Kalina evaporator | $C_{5}+C_{29}+Z_{\text {eva, } K \text { L }}=C_{6}+C_{9}$ | $c_{5}=c_{6}$ |
| Kalina turbine | $C_{9}+Z_{t, K A L}=C_{10}+C_{w, t, K A L}$ | $c_{9}=c_{10}$ |
| Kalina recuperator 1 | $C_{10}+C_{28}+Z_{\text {rel,KLL }}=C_{11}+C_{29}$ | $c_{10}=c_{11}, c_{28}=c_{29}$ |
| Kalina recuperator 2 | $C_{11}+C_{18}+Z_{\text {re2,KAL }}=C_{12}+C_{19}$ | $c_{11}=c_{12}, c_{18}=c_{19}$ |
| Kalina recuperator 3 | $C_{20}+C_{27}+Z_{\text {re3,KLL }}=C_{21}+C_{28}$ | $c_{20}=c_{21}, c_{27}=c_{28}$ |
| Kalina recuperator 4 | $C_{17}+C_{22}+Z_{\text {re4, KAL }}=C_{18}+C_{23}$ | $c_{17}=c_{18}, c_{22}=c_{23}$ |
| Kalina mixer 1 | $C_{12}+C_{24}+Z_{n n 1, K A L}=C_{13}$ | - |
| Kalina mixer 2 | $C_{16}+C_{21}+Z_{n 22, K A L}=C_{25}$ | - |
| Kalina condenser 1 | $C_{13}=C_{14}+Z_{\text {conl, }, \text { AL }}$ | $c_{13}=c_{14}$ |
| Kalina condenser 2 | $C_{25}=Z_{\text {con } 2, K A L}+C_{26}$ | $c_{25}=c_{26}$ |
| Kalina pump 2 | $C_{15}=Z_{p 2, K A L}+C_{14}+C_{w, p 2, K A L}$ | $c_{w, p 2}=c_{w, t, K A L}$ |
| Kalina pump 3 | $C_{27}=Z_{p 3, K A L}+C_{26}+C_{w, p 3, K A L}$ | $c_{w, p 3}=c_{w, t, K A L}$ |
| Kalina splitter | $C_{15}+Z_{\text {spl }, \text { ALL }}=C_{16}+C_{17}$ | $c_{16}=c_{17}$ |
| Kalina separator | $C_{19}+Z_{\text {sep,KAL }}=C_{20}+C_{22}$ | $c_{20}=c_{22}$ |
| Kalina expansion valve 1 | $C_{23}+Z_{e v, K A L}=C_{24}$ | - |
| ORC vapor generator | $C_{2}+C_{35}+Z_{\text {vg }, \text { ORC }}=C_{3}+C_{36}$ | $c_{2}=c_{3}$ |
| ORC turbine | $C_{36}+Z_{4, \text { ORC }}=C_{37}+C_{38}+C_{w, t, \text { ORC }}$ | $c_{36}=c_{37}, c_{36}=c_{38}$ |
| ORC ejector | $C_{37}+C_{45}+Z_{\text {eje, ORC }}=C_{39}$ | - |
| ORC preheater | $C_{34}+C_{40}+Z_{\text {prr,ORC }}=C_{35}+C_{41}$ | $c_{34}=c_{35}$ |
| ORC pump | $C_{34}=Z_{p, O R C}+C_{33}+C_{w, p, \text { ORC }}$ | $c_{w, p, \text { ORC }}=c_{w, t, \text { ORC }}$ |
| ORC condenser | $C_{41}+C_{48}+Z_{\text {con }, \text { ORC }}=C_{42}+C_{49}$ | $c_{41}=c_{42}, c_{48}=0$ |
| ORC expansion valve | $C_{43}+Z_{\text {evx }, \text { ORC }}=C_{44}$ | - |
| ORC evaporator | $C_{44}+C_{46}+Z_{\text {era,ORC }}=C_{45}+C_{47}$ | $c_{44}=c_{45}, c_{46}=0$ |
| PEM | $C_{v, P E M}+Z_{\text {PEM }}=C_{53}+C_{54}$ | $c_{w, P E M}=c_{w, \text {,ORC }}, c_{54}=0$ |
| DWH | $C_{30}+C_{50}+Z_{P E M}=C_{4}+C_{51}$ | $c_{3}=c_{4}, c_{50}=0$ |
| RO | $C_{30}+Z_{R O}+C_{w, R O}=C_{31}+C_{32}$ | $c_{w, R O}=c_{w, t, K A L}, c_{30}=c_{32}=0$ |

Table 8. Validation of the simulation results for the RO desalination unit.


Fig. 3. Comparison of the results of the present work with the results from Zare and Moalemian [65]

## 5. Results and discussion

After defining all required equations and constants for simulating the solar-geothermal multigeneration system, in the following section, the outputs and results of the studied system assessment are presented through different tables and diagrams and under varying different parameters. Firstly, the overall performance of the proposed system is expressed and then the exergo-economic analysis results are presented. In the end, by
parametric analysis, effects of the changing some critical parameters on the outputs of the studied system were investigated. For comparative purposes, the two most common and accessible nanofluids were examined, and then the most desired working fluid for the introduced system was discovered. Fig. 4 summarizes the main steps in conducting the present study. This flow diagram can be helpful to understand the procedure carried out to simulate this system.


Fig. 4. The flowchart of the pursued procedure in the present study.

### 5.1.Modeling results

In Table 9, the results of the operational characteristics of the proposed multigeneration system with different collector working fluids are listed and compared from the thermodynamic and cost of the component's point of view. The results show that applying nanofluid improves the performance of the system. Moreover, the comparison proves that CuO-based nanofluid results in the highest solar collector energy and exergy efficiencies, useful solar gained energy, net power output, and the amount of hydrogen produced.

Table 10 displays the quantity of fuel and product exergy, exergy destruction rate, and exergy efficiency of different components. One of the important scales in the exergy analysis
of a system is the exergy destruction rate of the components. The results show that the PTC field with 52002 kW has the maximum value of irreversibility. After that, the Kalina evaporator with 5156 kW possesses the second rank in the highest exergy destruction rate. The amount of fuel exergy transformed into product exergy is defined as the exergy efficiency. According to the results, the PTC field shows the lowest exergy efficiency with an amount of $5.92 \%$. The reason is that the PTC converts a large amount of the exergy of solar energy to low-grade energy in the form of a hot working fluid. Kalina condenser and ORC vapor generator have the lowest exergy efficiency after the PTC field which is because of the temperature difference between the fluids which flow inside these components.

Table 9. Thermodynamic performance and cost of components results for the proposed multigeneration system with different collector working fluids

| Parameters | Therminol_VP1 | $\mathrm{AL}_{2} \mathrm{O}_{3}$ nanofluid | CuO nanofluid |
| :---: | :---: | :---: | :---: |
| $\eta_{\text {en }}(\%)$ | 33.88 | 33.89 | 33.89 |
| $\eta_{e x}(\%)$ | 16.64 | 16.67 | 16.68 |
| $\eta_{\text {en, PTC }}(\%)$ | 67.62 | 67.82 | 67.88 |
| $\eta_{\text {ex }, P T C}(\%)$ | 66.44 | 66.62 | 66.67 |
| $Q_{u}(k W)$ | 6260 | 6279 | 6284 |
| $Q_{H X,, g e o}(k W)$ | 987.8 | 1007 | 1012 |
| $Q_{\text {eva, } K A L}(k W)$ | 5332 | 5333 | 5333 |
| $W_{\text {net }}(k W)$ | 1366 | 1369 | 1370 |
| $Q_{D W H}(k W)$ | 1362 | 1365 | 1365 |
| $m_{H \text { Hdrogen }}\left(\frac{g}{S}\right)$ | 0.00164 | 0.001174 | 0.001177 |
| $Z(\$)$ | 142183 | 142496 | 142582 |

Table 10. The quantity of fuel and product exergy, exergy destruction rate, and exergy efficiency of each component of the proposed system.

| Component | $E x_{f}(k W)$ | $E x_{p}(k W)$ | $E x_{D, k}(k W)$ | $\varepsilon(\%)$ |
| :---: | :---: | :---: | :---: | :---: |
| PTC field | 55277 | 3274 | 52002 | 5.924 |
| Heat exchanger | 367.2 | 265.9 | 101.2 | 72.43 |
| Kalina evaporator | 2931 | 2225 | 5156 | 75.94 |
| Kalina turbine | 1926 | 1686 | 239.5 | 87.56 |
| Kalina recuperator 1 | 415.6 | 212.4 | 203.2 | 51.1 |
| Kalina recuperator 2 | 547.8 | 236.8 | 311 | 43.22 |
| Kalina recuperator 3 | 20.08 | 7.29 | 12.8 | 36.27 |
| Kalina recuperator 4 | 64.19 | 38.13 | 26.06 | 59.4 |
| Kalina mixer 1 | 81887 | 81882 | 5.758 | 100 |
| Kalina mixer 2 | 40999 | 40994 | 5.789 | 100 |
| Kalina condenser 1 | 117.3 | 24.11 | 93.15 | 20.56 |
| Kalina condenser 2 | 44.37 | 5.847 | 50.22 | 13.18 |
| Kalina pump 2 | 4.56 | 3.2 | 1.35 | 70.28 |
| Kalina pump 3 | 53.56 | 37.46 | 16.1 | 69.95 |
| Kalina splitter | 81768 | 81768 | 0 | 100 |
| Kalina separator | 65423 | 65423 | 0 | 100 |
| Kalina expansion valve | 40959 | 40957 | 1.92 | 100 |
| ORC vapor generator | 617.2 | 83.76 | 92.21 | 13.57 |
| ORC turbine | 358.5 | 297.6 | 60.87 | 83.02 |
| ORC ejector | 88.99 | 57.75 | 31.24 | 64.9 |
| ORC preheater | 46.64 | 38.36 | 8.28 | 82.25 |
| ORC pump | 5.01 | 4.74 | 0.27 | 94.61 |
| ORC condenser | 104.5 | 40.44 | 64.02 | 38.71 |
| ORC expansion valve | 267.5 | 262.8 | 4.65 | 98.26 |
| ORC evaporator | 58.87 | 42.99 | 15.88 | 73.02 |
| PEM | 298.3 | 98.44 | 199.9 | 33 |
| DWH | 199.5 | 112.4 | 87.15 | 56.32 |
| RO | 14.61 | 5.34 | 9.27 | 36.57 |

Table 11. Exergo-economic parameters for different components of the proposed system

| Component | $c_{f, k}\left(\frac{\$}{G J}\right)$ | $c_{p, k}\left(\frac{\$}{G J}\right)$ | $Z_{k}\left(\frac{\$}{h}\right)$ | $C_{D, k}\left(\frac{\$}{h}\right)$ | $r_{k}(\%)$ | $f_{k}(\%)$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| PTC field | 0 | 0.351 | 0.131 | 0 | $\infty$ | 100 |
| Heat exchanger | 5 | 32.09 | 0.051 | 0.057 | 541.8 | 47.26 |
| Kalina evaporator | 0.351 | 1.76 | 0.329 | 0.206 | 400.9 | 61.44 |
| Kalina turbine | 9.6 | 43.49 | 6.262 | 0.262 | 353.2 | 95.98 |
| Kalina recuperator 1 | 9.6 | 10.02 | 0.381 | 0.384 | 4.42 | 49.79 |
| Kalina recuperator 2 | 9.6 | 3.57 | 0.083 | 0.256 | 62.81 | 25.25 |
| Kalina recuperator 3 | 3.57 | 7.08 | 0.05 | 0.005 | 98.34 | 90.67 |
| Kalina recuperator 4 | 7.41 | 11.38 | 0.151 | 0.022 | 53.47 | 87.25 |
| Kalina mixer 1 | 24.18 | 24.18 | 0 | 0 | 0 | 0 |
| Kalina mixer 2 | 11.93 | 38.32 | 0 | 0 | 0 | 0 |
| Kalina condenser 1 | 24.18 | 24.18 | 0.323 | 0.257 | 0 | 55.73 |
| Kalina condenser 2 | 38.32 | 38.32 | 0.194 | 0.219 | 0 | 46.91 |
| Kalina pump 2 | 43.49 | 142.6 | 0.029 | 0.0067 | 227.9 | 81.45 |
| Kalina pump 3 | 43.49 | 154.7 | 0.211 | 0.08 | 255.7 | 72.62 |
| Kalina splitter | 24.19 | 24.19 | 0 | 0 | 0 | 0 |
| Kalina separator | 3.57 | 3.57 | 0 | 0 | 0 | 0 |
| Kalina expansion valve | 38.75 | 38.76 | 0.005 | 0.008 | 0 | 36.72 |
| ORC vapor generator | 5 | 18.48 | 0.596 | 0.052 | 269.6 | 91.89 |
| ORC turbine | 27.01 | 87.27 | 1.856 | 0.187 | 223.1 | 90.83 |
| ORC ejector | 38.65 | 30.36 | 0.0152 | 0.137 | 21.44 | 9.98 |
| ORC preheater | 24.61 | 31.29 | 0.268 | 0.023 | 27.14 | 92.02 |
| ORC pump | 87.27 | 2.74 | 0.0009 | 0.047 | 96.86 | 1.85 |
| ORC condenser | 24.59 | 31.46 | 0.438 | 0.179 | 27.95 | 70.93 |
| ORC expansion valve | 31.46 | 32.07 | 0.0015 | 0.0167 | 1.932 | 8.23 |
| ORC evaporator | 32.07 | 87.41 | 0.2 | 0.058 | 164.1 | 77.49 |
| PEM | 87.27 | 263.9 | 0.00064 | 1.99 | 2.024 | 3.2 |
| DWH | 5 | 20.68 | 0.151 | 0.049 | 313.7 | 75.27 |
| RO | 43.49 | 72.75 | 5.095 | 0.046 | 67.3 | 99.11 |

The results of the exergo-economic analysis of the proposed system are presented in Table 11. Kalina turbine, Ro unit, and ORC turbine have the highest total costs which means the importance of these components from the exergo-economic point of view. PTC field and Kalina turbine have the highest amount of exergoeconomic factors which means that capital and maintenance costs have large shares of these high costs. Moreover, the ORC pump, PEM electrolyzer, and ORC expansion valve have the lowest amounts of exergoeconomic factor which show that the exergy destruction costs dominate the initial investment of these components. The economic investigation shows that the relative cost difference in the heat exchanger is the highest, which means that the average cost per exergy unit of product is superior to the average cost per exergy unit of fuel. Also, it can be inferred that the PEM electrolyzer has the highest exergy destruction cost rate. The reason is that the produced power enters the PEM electrolyzer as an exergy of fuel.

### 5.2.Parametric study

### 5.2.1.Effects of Nanoparticles on the performance of the system

By adding nanoparticles to the base fluid, the properties of the base fluid alter. The percentage of nanoparticles is one of the
factors that affect the properties of the base fluid considerably and is very decisive in preparing nanofluids. Fig. 5 demonstrates the hydrogen production rate while the nanoparticle percentage varies between $0 \%$ and $6 \%$. When the ratio is $0 \%$, both CuO and $\mathrm{Al}_{2} \mathrm{O}_{3}$-based Therminol_VP1 nanofluids generated $0.001164 \mathrm{~g} / \mathrm{s}$ amount of hydrogen. According to the graphs, at higher nanoparticle percentages, the hydrogen production rate for the studied nanofluids increased. The reason for the rise is that when the nanoparticle percentage increases, under the defined conditions, the overall production power by the ORC turbine increases, which leads to a rise in the $\mathrm{H}_{2}$ production rate. The rate of hydrogen production is higher for CuO than for $\mathrm{Al}_{2} \mathrm{O}_{3}$.

Figure 6 depicts the applying different percentages of CuO and $\mathrm{Al}_{2} \mathrm{O}_{3}$ nanoparticles to the total net power generated by the proposed system. For both nanofluids, the power production trend is increasing. According to the parametric results, for the range of $0 \%$ to $6 \%$ of nanoparticles, the power production of CuO-based nanofluid varied from 1366 to 1373, while for $\mathrm{Al}_{2} \mathrm{O}_{3}$ nanofluid, it increased from 1366 to 1372 . It can be seen that the graphs have a stepped behavior. It means that for some percentages of nanofluids, the generated power is steady, while for other amounts, it is rising.


Fig. 5. Effect of nanoparticle percentage on the total net power


Fig. 6. Effect of nanoparticle percentage on the hydrogen production rate

One of the factors that affect the useful energy gain by the solar collector is the nanoparticle percentage. Fig. 7 illustrates the changes in useful energy gained versus CuO and $\mathrm{Al}_{2} \mathrm{O}_{3}$ nanoparticles percentage. It is inferred that the nanoparticle percentage increase leads to increased useful energy gained from the PTC. According to equation 10 , the nanoparticle concentration change just alters the heat transfer factor. By increasing the nanoparticle concentration, the nanofluid mass
flow rate increases while the specific heat transfer reduces. The amount of increase in CuO -based nanofluid is superior to the amount of $\mathrm{Al}_{2} \mathrm{O}_{3}$-based nanofluid increase. By altering the ratio from 0 to 0.06 , the energy gain rises from 6260 to 6306 kW when CuO-based nanofluid was applied, while when $\mathrm{Al}_{2} \mathrm{O}_{3}-$ based nanofluid was used, it increased from 6260 to 6290 kW . CuO-based nanofluid compared to the $\mathrm{Al}_{2} \mathrm{O}_{3}$-based nanofluid shows better results.


Fig. 7. Effect of nanoparticle percentage on the useful energy achieved from the collector
5.2.2.Effects of solar irradiation on the Performance of the System

Another crucial factor that influences the efficiency of a solar collector is solar irradiation. The amount of energy that passes to the nanofluid in the PTC in the form of temperature is defined as solar irradiance.

Figure 8 demonstrates the trend of useful energy achieved by PTC and the hydrogen production rate when the solar irradiation changes from 400 to $1000 \mathrm{~W} / \mathrm{m}^{2}$ and the nanoparticle percentage is fixed at 0.03 . When solar irradiation rises, more energy can transfer to the Kalina cycle which increases the amount of the hydrogen production rate. Moreover, according to equation 10 , the absorbed solar radiation $(S)$ rises dramatically and the gained useful energy sharply increases. By investigating the graphs, it is clear that applying two introduced nanofluids has the same impact on the useful energy and hydrogen production rate, but for CuO -based nanofluids, the useful energy is slightly higher. By changing the solar irradiation from 400 to $1000 \mathrm{~W} / \mathrm{m}^{2}$, the CuO -based nanofluid alters from 2845 to 7430 kW , while for the $\mathrm{Al}_{2} \mathrm{O}_{3-}$ based nanofluid, this change is from 2842 to 7424 kW. Finally, for the studied solar irradiation range, both nanofluids increase from 0.00291 to $0.00538 \mathrm{~g} / \mathrm{s}$, although in some
parts, there is very little difference between the two nanofluids.

The energetic and exergetic efficiencies, as well as the total net power produced by the entire proposed system versus the effect of solar irradiation, are plotted in Fig. 9. There is a negligible difference between the nanofluids applied in this study. Energy and exergy efficiencies of the system decrease with increasing solar irradiation. The amount of reduction in the energetic efficiency is greater than the amount of decline in exergetic efficiency. By increasing solar irradiation, the energetic efficiency of the multigeneration system decreases from $36.77 \%$ to $33.26 \%$, while the exergetic efficiency reduces from $17.87 \%$ to $16.5 \%$. Moreover, it can be seen that the total net power production increases from 596 to 1638 kW when the solar irradiation changes from 400 to 1000 kW . The reason for the sharp rise in overall power production is that increasing solar irradiation leads to higher solar intensity bouncing back on the receiver, which increases the temperature of the nanofluids leaving the parabolic trough collector. The higher the nanofluid temperature leaving the receiver, the higher the efficiency of the PTC and more power will be produced in the turbines of Kalina and ORC cycles. Finally, more power production will happen, which leads to a higher production of hydrogen rate.


Fig. 8. Effect of solar irradiation on the useful energy achieved from the collector and the hydrogen production rate


Fig. 9. Effect of solar irradiation on the efficiency and the total net power

The intensity of solar radiation has a considerable impact on the outlet temperature leaving the PTC. The effect of solar irradiance on the outlet temperature of the collector and the heat rate produced by the Kalina cycle evaporator is shown in Fig. 10. The evaluation of the figure makes it clear that for both nanofluids, the outlet temperature rises when the solar radiation intensity increases. Although there is little difference between the studied nanofluids, both nanofluids have the same
trends. The outlet temperature of the collector is discovered to be enhanced from 543 K to 688 K by solar irradiance, changing from $400 \mathrm{~W} / \mathrm{m}^{2}$ to $1000 \mathrm{~W} / \mathrm{m}^{2}$. The higher the exit temperature of the solar collector, the higher the heat rate generated by the Kalina cycle evaporator. By increasing the amount of produced heat rate, the power generated by the Kalina cycle will increase too. By varying the solar irradiance, the heat rate generated by the evaporator changes from 1895 kW to 6480 kW .


Fig. 10. Effect of solar irradiation on the solar collector outlet temperature and the heat rate produced in the evaporator
5.2.3.Effects of ambient temperature on the performance of the system

To analyze the multigeneration system more comprehensively, the impacts of ambient temperature on the freshwater production rate, total net power generated, and outlet temperature of the collector have been studied.

Figure 11 shows the impact of ambient temperature on the freshwater production rate and total net power generated. For both studied nanofluids, the graphs display a rising trend for freshwater production rate and total net power. By changing ambient temperature from 275 to 325 K , the freshwater production rate varies from 4.867 to $4.898 \mathrm{~g} / \mathrm{s}$ while generated power alters from 1362 to 1378 kW . As it can be
inferred from the charts, it is clear that applying $\mathrm{AL}_{2} \mathrm{O}_{3}$ or CuO -based nanofluids does not show a remarkable difference, although using CuO as the nanofluid, the system produces a small amount of more net power.

It is evident in Fig. 12 that by enhancing ambient temperature, both collector efficiency and outlet temperature of solar collector go up. By increasing ambient temperature from 275 to 325 K , when AL2O3-based fluid is used, the outlet temperature rises from 651 to 653.5 , while for the case of CuO -based nanofluid, the change is from 650.5 to 652.6 . Moreover, the effect of ambient temperature change on the collector efficiency is very small, as the collector efficiency for both studied nanofluids varies between 67.5 to 68.25 .


Fig. 11. Effect of ambient temperature on the freshwater production rate and total net power


Fig 12. Effect of ambient temperature on the collector outlet temperature and collector efficiency
5.2.4.Effect of solar collector inlet temperature on the performance of the system

The solar collector's inlet temperature is another factor that impacts the efficiency of the multigeneration system. Fig. 13 shows the effect of solar collector inlet temperature on the energy and exergy efficiency of the studied system. The results of the three different cases were investigated, including base fluid (Therminol VP1), $\mathrm{AL}_{2} \mathrm{O}_{3}$-based nanofluid, and CuO -based nanofluid. As can be inferred from the diagrams, for both energy and exergy efficiencies, nanofluids show higher values compared with base fluid. Among the studied working fluids, CuO offers the best performance. For energetic efficiency, as the inlet temperature increases from 430 K to 490 K , the amount of net power produced by the system increases while the useful energy generated by the solar collector decreases. At first, the amount of energy produced is superior to the produced power and causes a decrease in the thermodynamic efficiency of the system, but then this superiority

disappears and causes an increase in the efficiency. For CuO-based nanofluid, the thermal efficiency first decreases from $34.91 \%$ to 33.81 $\%$ and then increases to $34.45 \%$. It can be seen that 470 K is an optimum point for the exergetic efficiency graph. By analyzing the exergetic efficiency graph, it can be concluded that increasing the PTC inlet temperature from 430 K to 490 K , increases the exergy efficiency for all working fluids.

Figure 14 displays the effect of the inlet temperature of the solar collector on the system's hydrogen production rate and freshwater production rate. According to the graphs, when the inlet temperature of the collector increases, the hydrogen production rate decreases while the freshwater production rate increases. By analyzing the results, it can be found that when the inlet temperature rises by about $14 \%$, the freshwater production rate increases by about $18.5 \%$, and the hydrogen production rate decreases by about $70 \%$. By comparing the graphs, CuO -based nanofluid showed a little better result than $\mathrm{AL}_{2} \mathrm{O}_{3}$-based nanofluid.


Fig. 13. Effect of collector inlet temperature on the energy and exergy efficiency


Fig. 14. Effect of collector inlet temperature on the hydrogen and freshwater production rates

## 6. Conclusions

In the present study, a multigeneration system was investigated in which solar and geothermal energies, as the two plentiful and clean sources of energy, are combined to satisfy the required demands of an area. To better understand the studied system, energy, and exergy research was performed. A comparative investigation among the base fluid and two introduced nanofluids was done to find the most suitable working fluid for the solar collector.

Some of the main conclusions are as follows:

- Although comparing $\mathrm{Al}_{2} \mathrm{O}_{3}$-based and CuO-based nanofluids show almost the same results, in general, it can be concluded that CuO -based nanofluid shows better results.
- Compared to the Therminol VP1 as the base fluids, applying nanofluids leads to higher power production and better efficiency.
- It was concluded that the hydrogen production rate for CuO nanofluid is higher than $\mathrm{Al}_{2} \mathrm{O}_{3}$ nanofluid when the nanoparticle concentration increases gradually.
- The results showed that by varying nanoparticle percentage, total net power produced and useful energy achieved from the collector has a rising trend, and

CuO-based nanofluid showed higher amounts.

- By changing the solar irradiation from 400 to $1000\left(\mathrm{~W} / \mathrm{m}^{2}\right)$, the total net power produced, hydrogen production rate, useful energy achieved from the collector, solar collector outlet temperature, and heat rate produced in the solar evaporator increased while the total energy and exergy efficiency of the system decreased.
- Ambient temperature is one of the factors that affect the performance of the studied multigeneration system. While the ambient temperature changes from 275 to 325 K , all the studied parameters, including freshwater production rate, total net power, collector efficiency, and collector outlet temperature, tend to increase.
- When the collector inlet temperature varies between 430 to 490 K , the energy efficiency of the proposed system first tends to decrease, but at 470 K , it starts increasing. For exergy efficiency, a rising trend can be seen. Moreover, by increasing inlet temperature, the hydrogen production rate decreases while the freshwater production rate increases.


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