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EXERGETIC MODELLING OF OIL-FIRED STEAM BOILERS

I. O. Ohijeagbon^{1,*}, M. A. Waheed², S. O. Jekayinfa³ and O. A. Lasode⁴

^{1,4}MECHANICAL ENGINEERING DEPARTMENT, UNIVERSITY OF ILORIN, ILORIN, NIGERIA
 ²MECHANICAL ENGINEERING DEPARTMENT, UNIVERSITY OF AGRICULTURE, ABEOKUTA, NIGERIA
 ³AGRICULTURAL ENGINEERING DEPARTMENT, LADOKE AKINTOLA UNIVERSITY OF TECHNOLOGY, OGBOMOSO, NIGERIA
 E-mail addresses: 1 idehaiohi@yahoo.com, 2 akindoye@yahoo.com, 3 jekaysol@yahoo.com, 4 oalasode@yahoo.com

ABSTRACT

The performance variables and potential for energy savings in oil-fired industrial steam boilers were studied. Operational parameters of steam boilers using low pour fuel oil (LPFO) and diesel were used to determine thermodynamic properties of material streams and exergetic parameters. Analysis of thermodynamic properties on exergetic performance of boilers was conducted. Exergetic models developed are also presented. The ratio of the energy to exergy efficiencies of the heat exchanging unit and of the entire steam boiler was determined as 1.4 and 1.8614 respectively. The average overall boiler exergy destruction decreased by 14.62%, as the evaporation ratio increased from 10 to 14 though heat loss in boiler was found to increase with increases in evaporation ratio. Developed models in this study could be employed to analyse and estimate exergetic performance indices of oilfired steam boilers in order to enhance their efficiencies and reduce energy wastage.

Keywords: exergetic efficiency; steam boilers; oil-fired; operational parameters

LIST OF SYMBOLS

| actual and stoichiometric air-fuel ratio (<i>kg of</i> |
|--|
| air/kg of fuel) |
| molar chemical exergy of carbon dioxide (<i>CO</i> ₂), <i>kJ/kmol</i> |
| molar chemical exergy of fuel, <i>kJ/kmol</i> |
| molar chemical exergy of water vapour (<i>H₂O</i>), <i>kJ/kmol</i> |
| molar chemical exergy of oxygen (<i>O</i> ₂), <i>kI/kmol</i> |
| the energy input in the combustion unit, <i>kJ/s</i> |
| standard Gibbs function, <i>kJ/kmol</i> |
| <pre>standard Gibbs function of carbon dioxide (CO2), kJ/kmol</pre> |
| standard Gibbs function of water vapour (<i>H</i> ₂ <i>O</i>), <i>kJ/kmol</i> |
| standard Gibbs function of oxygen (<i>O</i> ₂), <i>kJ/kmol</i> |
| specific enthalpy, <i>kJ/kg</i> |
| reference state enthalpy, <i>kJ/kg</i> |
| high or gross heating value of fuel, <i>kJ/kg</i> |
| exergy destruction rate, <i>kJ/s</i> |
| low or net heating value of fuel, <i>kJ/kg</i> |
| low pour fuel oil |
| exit mass flow rate, <i>kg/s</i> |
| |

| \dot{m}_i | inlet mass flow rate, <i>kg/s</i> | | | |
|--------------------------|---|--|--|--|
| M P | molar mass of chemical substance, <i>kg/kmol</i> pressure, <i>N/m²</i> | | | |
| \dot{Q} | heat transfer rate to the system, <i>kJ/s</i> | | | |
| $Q_{H(loss)}$ | hHeat loss in heat exchanging unit, <i>kJ/s</i> | | | |
| \overline{R} | molar or universal gas constant, <i>kJ/kmol K</i> | | | |
| S | specific entropy, <i>kJ/kg K</i> | | | |
| S 0 | reference state entropy, <i>kJ/kg K</i> | | | |
| Т | temperature, ^o C, K | | | |
| T_{0} | reference state temperature, ^o C, K | | | |
| • • • • | | | | |
| | CHEMICAL SYMBOLS | | | |
| Ar | argon | | | |
| С | carbon | | | |
| CO_2 | carbon dioxide | | | |
| $C_x H_y$ | hydrocarbon (fossil fuel) molecules | | | |
| Н | hydrogen | | | |
| H_2O | water | | | |
| N_2 | nitrogen | | | |
| <i>0, 0</i> ₂ | oxygen | | | |
| SUBSCRIPTS | | | | |
| 0 | reference state | | | |
| 0a | reference state of air | | | |
| A | air | | | |
| | | | | |

Atm atmospheric

- B boiler
- *C* combustion unit
- F Fuel
- *G* exhaust flue gas
- *H* heat exchanging unit
- *in* in
- out out
- *P* hot products
- S Steam
- W feed water

SUPERSCRIPTS

- *ch* Chemical
- *ph* Physical

GREEK LETTERS

| the change in the standard Gibbs function, <i>kI/kmol</i> |
|--|
| specific exergy (<i>kJ/kg</i>) |
| exit specific flow exergy of material streams, <i>kJ/s</i> |
| specific chemical flow exergy, <i>kJ/kg</i> |
| specific physical flow exergy, <i>kJ/kg</i> |
| energy efficiency (%) exergy efficiency (%) |
| |

1. INTRODUCTION

Steam boilers are widely applied for process heating, sterilization, drying, material conditioning and saponification in bottling, pharmaceutical, tobacco and soap industries; and also for production of mechanical and electrical energy. Energy is consumed, conserved and dissipated in steam boilers when engaged in producing steam for different end uses by combustion of fuel.

For any given environment, energy which may be converted into other forms is called useful energy or exergy. And energy which cannot be converted into other energy forms is called useless energy or anergy [1]. When all processes of the system and the environment are reversible, exergy is fully conserved. Exergy is destroyed whenever an irreversible process takes place. Exergy destruction is a measure of the thermodynamic imperfections determined when an exergy analysis is performed on a plant such as an entire power station, a chemical processing plant or a refrigeration plant, and this refers to wasted work or wasted potential for the production of work [2].

Exergy is a useful concept for determining improvement, since it is a connection between the physical and engineering world and the surrounding environment, and expresses the true efficiency of engineering systems. Exergy analysis has been applied

in the design of engineering systems as well as sectoral energy and exergy consumption [3]. There has been an increasing interest in applying energy and exergy analysis modelling techniques for energyutilization assessments in order to obtain energy Exergy analysis is a savings in recent years [4]. methodology for the assessment of the performance of devices and processes, and involves analysing the exergy at different points in a series of energyconversion steps [6]. Consequently, efficiencies can be evaluated, and the process steps having the largest losses (i.e., the greatest margin for improvement) can be identified. Exergy analysis has been found to be a highly versatile method by many scientists, engineers, managers, decision and policy makers. Exergy analysis has been employed in many spectra of socioeconomic and industrial infrastructure such as, analysing and evaluating the energy utilization efficiency of countries, steam and thermal power plants, combustion and energy conversion processes, iron smelting processes, etc. It has been established to be a very useful tool for determining operational improvements and energy savings of industrial processes and products [7-10].

Suresh et. al [11] conducted a study to predict the possible improvement in efficiency obtained with thermal power plants. The study dealt with the comparison of energy and exergy analysis of thermal power plants based on advanced steam parameters in Indian climatic conditions. The study involved coalbased thermal power plants using subcritical, supercritical, and ultrasupercritical steam conditions. The design configurations of 500 MWe unit size were considered. The study encompassed the effect of condenser pressure on plant and exergy efficiency. The effect of high grade coal on performance parameters as compared to typical Indian low grade coal was also studied. The major exergy loss took place in coal combustion followed by the steam generator. Due to condenser pressure limitation, the maximum possible plant efficiency was found to be about 41% for supercritical steam power plant and 44.4% for the ultrasupercritical steam power plant. It was therefore concluded that installing coal-based thermal power plants based on advanced steam parameters in India will be a prospective option aiding energy self-sufficiency.

The persistent increase in energy demand for economic development and the need for the preservation of the environment are indications that energy systems ought to become more efficient [12]. Attempts have been made by other researchers to combine mathematical or economical methods and advanced thermodynamic processes to investigate and improve plant systems, and also to study and minimize the effects of irreversibility. Ataei and Yoo [13] performed the simulation of a 325 MW steam power plant in which the operational parameters of the Rankine cycle were optimized using the exergy concept combined with a pinch-based approach in the study of combined pinch and exergy analysis for energy efficiency optimization in a steam power plant. Srinivas et. al [14] conducted a thermodynamic analysis of the effect of 'n' feedwater heaters on performance of a steam power cycle with a generalized mathematical formulation. They carried out performance calculations formulated separately to a single feedwater heater and extended to 'n' feedwater heaters for parametric study. The work of Srinivas et. al [14] examined the improvements in efficiency with increases in boiler pressure, turbine inlet temperature and furnace temperature, and they were able to determine the maximum gain in the efficiency of the cycle. Gutierrez and Mendez [15] developed a theoretical analysis to minimize the entropy generation of a thermo-electric cooler. They applied an analysis of first and second law of Thermodynamics to thermo-electric cooler, and considered the entropy generation equation as the objective function and the first law as a restriction, in accordance with the variational calculus theory. Their numerical estimations indicated that minimum entropy generation and the better performance coefficients (COPs) were achieved with the largest figure of merit, which represented a relationship between the Peltier effect (cooling) and the Joule effect (heating).

It is imperative to adopt reliable methods to assess the performance of industrial facilities, such as steam boiler, for sustainable use. The performance of steam boilers could be assessed based on operational and output parameters which are of primary interest to a researcher or other stakeholders. In a study of an oil-fired boiler system, Orosun and Adamu [16] modelled steam boiler as a multivariable plant with two inputs, namely, feed water rate and oil-fired flow rate; and two outputs, which were steam temperature and pressure. The authors modelled the plant parameters using artificial neural network. Their result showed a correlation of 0.97 in the simulated and actual outputs for temperature and pressure respectively. They however did not consider the exergetic analysis of the

steam boiler. Igboanugo et. al [17] had also studied the performance characteristics of a boiler unit using failure mode effect and fault tree analyses, the energy source for the boiler was low grade fuel from palm oil bunch and fibre. The study revealed that in a single shift operating mode of eight hours, a time of 72 minutes was saved arising from the effectiveness of the plant overhaul carried out. They obtained an energy efficiency of 54.2% at boiler operating pressure of 700 kN/m². The effect of irreversibility, exergy destruction and efficiency were not analysed in the study.

This study is aimed at investigating the exergetic performances, developing of exergetic models and potential of energy savings through investigation of the effects of various thermodynamic and operational variables in oil-fired industrial steam boilers.

2. METHOD

2.1 Material streams in steam boiler

The basic influx of material streams in a boiler as exemplified by Figure 1 which is an exploded view of the model of Saidur et. al [18], takes place in the combustion unit and heat exchanging unit. The intake air (from ambient air supply) mixes with the combustion fuel, usually hydrocarbon in a chemical oxidation process to release heat energy in the furnace. Combustion of LPFO and diesel fuels was used in this study. The hot product of combustion exiting the furnace enters the heat exchanging unit in which the supplied heat energy is utilised to transform the feed water entering into the heat exchanging unit into saturated steam. The saturated steam may further be heated in the superheater by hot flue gases from the furnace to obtain superheated steam. The flue gases and heat loss exit the boiler to complete the material and energy flow process in the boiler.

2.2 Operational parameters

The general specifications of the boiler investigated are presented in Table 1. This shows that the boiler is a fire tube type and horizontal in orientation. It is internally fired, forced circulation, low pressure and stationary. It is also a multi-tube and operates on either LPFO or diesel fuels. The operating steam temperature, maximum steam pressure and operating steam pressure are 180 °C, 1600 kN/m² and 750kN/m² respectively. The steam generated from boiler is for the purpose of boiling, steam regeneration of carbon filter, and sterilization.

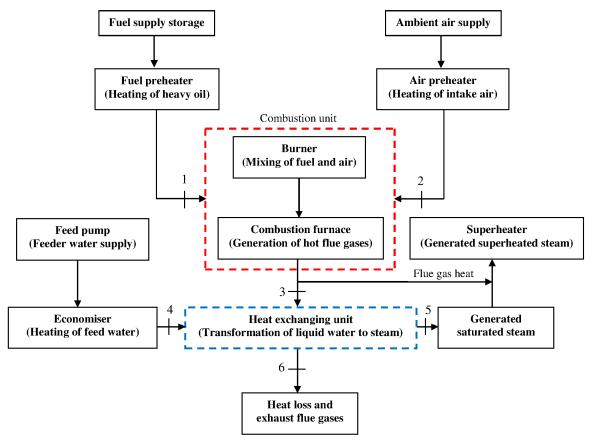


Figure 1: Material streams flow diagram in a steam boiler

The operational parameters used in this study are enumerated in Table 2 for both LPFO and diesel operation. Similar values had been adopted for the air temperature, steam pressure, mass flow rate of fuel, evaporation ratio, and excess air. The fuels have their peculiar specific heat capacities, stoichiometric and actual fuel ratios respectively as presented in Table 2. The specific heat capacity, stoichiometric and actual fuel ratios of LPFO operated boiler are 1.866 kJ/kgK, 13.428 and 16.114 kg of air/kg of fuel, while that of diesel operated boiler are 1.9612 kJ/kgK, 14.128 and 17.00 kg of air/kg of fuel respectively. The distinction in the peculiar thermodynamic properties is expected to influence the nature of exergetic performances of LPFO and diesel operated boilers.

Table 1: Specification of industrial steam boiler investigated

| investig | ated | (kg/s) | | |
|---------------------------|------------------|--|--------|--------|
| Details | Specification | Evaporation ratio, E _r | 12 | 12 |
| 1. Orientation | Horizontal | Specific heat capacity, c _p | 1.866 | 1.9612 |
| 2. Type of tube | Fire tube | (kJ/kgK) | | |
| 3. Type of firing | Internally fired | Stoichiometric air-fuel | 13.428 | 14.128 |
| 4. Type of circulation | Forced | ratio, AFR _{st} | | |
| 5. Type of pressure | Low | (kg of air/kg of fuel) | | |
| 6. Stationary or portable | Stationary | Actual air-fuel ratio, AAF | 16.114 | 17.00 |
| 7. Single or multi-tube | Multi-tube | (kg of air/kg of fuel) | | |
| 8. Type of fuel used | LPFO and Diesel | Excess air, EA (%) | 20 | 20 |

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| Details | Specification |
|---|--|
| 9. Operating steam temperature (°C) | 180 |
| 10. Maximum steam pressure (kN/m ²) | 1600 |
| 11. Operating steam pressure (kN/m ²) | 750 |
| 12. Purpose of steam generated | Boiling, steam regeneration of carbon filter, sterilization |
| | |

| Table 2: Parameters for the LPFO and diesel operation |
|---|
| of boiler [19-20] |

Parameter

Air temperature (°C)

Steam pressure (kN/m²) Rate of fuel consumption LPFO

operated

77.00

0.18

800.00

Diesel

operated

77.00

0.18

800.00

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2.3 Mass balance of material streams

Conservation of mass and energy balance was applied to the boiler system in addition to the evaporation ratio and air fuel ratio and were used to determine mass flow rates of substances of all material streams in the boiler, such as; air, \dot{m}_a , fuel,

 \dot{m}_{f} , hot products, \dot{m}_{p} , feed water, \dot{m}_{w} , steam, \dot{m}_{s} and flue gas, \dot{m}_{g} [19].

Hence,

.

 $\dot{m}_a + \dot{m}_f = \dot{m}_p \tag{1}$

$$\dot{m}_p = \dot{m}_g \tag{2}$$

$$\dot{m}_w = \dot{m}_s \tag{3}$$

The evaporation ratio of a boiler can be expressed by [19-21];

$$E_r = \frac{\dot{m}_s}{\dot{m}_f} \tag{4}$$

2.4 Energy and exergy analysis of boiler

The physical and chemical exergy associated with material streams for a flow process makes up the total flow exergy transfer which is the theoretically obtainable work when the substance is brought into total equilibrium with the local environment [22-25]. Ertesvag [26] presents the total flow exergy transfer by equation (5), while the specific physical flow exergy is given by equation (6).

$$\mathcal{E} = \mathcal{E}^{ph} + \mathcal{E}^{ch} \tag{5}$$

$$\mathcal{E}^{ph} = h - h_0 - T_0(s - s_0) \tag{6}$$

The method of determining the chemical exergy of fuels is similar to that used in methodology for the physical and chemical exergetic analysis of steam boilers [19] and cumulative exergetic assessment of LPFO utilized steam boilers [20], and are as follows [25, 27-28];

For complete combustion of hydrocarbon with atmospheric air,

$$C_x H_y + (x + \frac{y}{4})O_2 \to xCO_2 + \frac{y}{2}H_2O$$
 (7)

The molar chemical exergy of reactants and products is given by,

$$e_{xfuel}^{ch} + (x + \frac{y}{4})e_{xO_2}^{ch} \rightarrow -\Delta\overline{g} + xe_{xCO_2}^{ch} + \frac{y}{2}e_{xH_2O}^{ch}$$
(8)

$$e_{x\,fuel}^{\ ch} = -\Delta \overline{g} + \overline{R}T_0 \ln \left[\frac{n_{O_2}^{(x+y/4)}}{n_{CO_2}^x n_{H_2O}^{y/2}}\right] \tag{9}$$

The mole fraction of oxygen (n_{O_2}), carbon dioxide

 (n_{CO_2}) and water (n_{H_2O}) in a standard environment are presented in Table 3, and the values of *x* and *y* which are constant coefficients that characterises the hydrocarbon and combustion process are presented in Table 4. Where the molar or universal gas constant $\overline{R} = 8.3144 \text{ kJ/kmol K}$.

The standard Gibbs functions of formation of other components in equation (9) are presented in Table 3. The change in standard Gibbs function is expressed by,

$$\Delta \overline{g} = x \overline{g}_{CO_2} + \frac{y}{2} \overline{g}_{H_2O(v)} - (x + \frac{y}{4}) \overline{g}_{O_2}$$
(10)

The specific chemical exergy was determined by

$$\varepsilon^{ch} = \frac{e_x^{ch}}{M} \tag{11}$$

The chemical exergy of atmospheric air was determined by equation (12), where the mole fractions of the elements in a standard environment with respective chemical exergies are presented in Table 3.

$$e_{x(air)}^{ch} = \sum \left(n_i e_{x_i}^{ch} \right)_{air} \tag{12}$$

The specific chemical exergy of steam, water and combustion fuels presented in Ohijeagbon et. al [19-20] are shown in Table 5. Complete combustion analysis was assumed in determining the chemical exergises of fuels undergoing combustion.

Table 3: Chemical exergies, standard Gibbs functions of formation and mole fractions of components of atmospheric air at reference conditions of 25°*C and 1 atm [19, 26-27]*

| Components | Chemical exergies | Standard Gibbs | Mole fraction composition |
|------------------------------------|-------------------------|--------------------------|----------------------------|
| | (kJ/kmol) | function, \overline{g} | of a reference environment |
| | (70% relative humidity) | (kJ/kmol) | model |
| Nitrogen, N ₂ | 668.10 | 0 | 0.7565 |
| Oxygen, <i>O</i> ₂ | 3,930.50 | 0 | 0.2035 |
| Carbon dioxide, <i>CO</i> 2 | 19,610.00 | -394,390 | 0.0003 |
| Argon, <i>Ar</i> | 11,640.00 | 0 | 0.0091 |
| Water, $H_2O(g)$ | 9,474.00 | -228,590 | 0.0303 |
| Water, <i>H₂O</i> (liq) | 884.20 | -237,150 | - |
| Hydrogen, <i>H</i> ₂ | 236,098.00 | 0 | 0.0001 |
| Carbon monoxide, CO | 274,870.00 | -137,160 | - |

Table 4: Fuel specifications for LPFO and diesel

| Specifications | LPFO | Diesel |
|----------------------------------|-------------|-------------|
| | (No. 6 Oil) | (No. 2 Oil) |
| %Carbon (<i>C</i>) | 87.49 | 85.84 |
| %Hydrogen (<i>H</i>) | 9.92 | 12.46 |
| Gross heating value (HHV), kJ/kg | 42,657.34 | 45,482.52 |
| Net heating value (LHV), kJ/kg | 40,515.15 | 42,790.21 |
| CO ₂ max | 16.50 | 15.60 |
| %Sulfur (<i>S</i>) | 1.40 | 1.60 |
| $\%O_2$ | 1.190 | 0.100 |

Source: Ohijeagbon et. al [20], TSI Incorporated [29]

| in the boiler [19, 30] | | | |
|---|-----------------|--|--|
| Substance | Chemical exergy | | |
| | (kJ/kg) | | |
| Air | 58.79 | | |
| Steam | 526.33 | | |
| Water | 49.12 | | |
| LPFO (Before combustion) = HHV _{LPFO} | 42,657.34 | | |
| LPFO (During combustion) | 44,566.66 | | |
| Diesel (Before combustion) = HHV _{diesel} | 45,482.52 | | |
| Diesel (During combustion) | 44,569.69 | | |
| Exhaust flue gases (LPFO operated) | 147.97 | | |
| Exhaust flue gases (Diesel operated) | 148.77 | | |

Table 5: Specific chemical exergy of material streams

The energy and exergy values and efficiencies of the flow processes in the LPFO and diesel operated boiler which include the physical and chemical exergies were evaluated using equations (1) to (12), the exergetic formulae summarised in Table 6 and the thermodynamic properties of materials streams data presented in Table 7 which was computed according to the method used by Ohijeagbon et. al [19] to determine mass flow rate, temperature, enthalpy and entropy of material streams in a steam boiler at specified node points shown in the material streams flow diagram of Figure 1. The exergetic formulae summarised in Table 6 were determined through energy and exergy balance amalysis in the combustion and heat exchanging units of a boiler system [19-20]. The exergetic values and efficiencies which includes energy input, energy efficiency, exergy destruction and efficiency of the combustion unit; heat loss, energy efficiency, exergy destruction and efficiency of the heat exchanging unit; energy efficiency, overall exergy destruction and efficiency of the entire boiler accordingly. Consequently, a parametric analysis of reference state ambient conditions, entropy effects, and mass flow rates of fuel, feed water and steam temperatures and evaporation ratio on exergetic performance of boilers was carried out, and various exergetic empirical relations was also determined.

| | <i>c</i> 1 | 1 60 1 | | 1 11 540 007 |
|-----------------------------|----------------------|-------------------|--------------------|---------------------|
| Table 6 [,] Summar | rv of exergetic valu | es and efficienci | ies tormulae of st | eam boilers [19-20] |
| rubie of building | y or energette valu | es ana entrerener | co ioi manac oi ou | |

| <u>Combustion unit</u> | | | |
|---------------------------------|--|--|--|
| Energy input (kJ/s) | $\dot{E}_{in} = \dot{m}_f h_f + \dot{m}_a h_a$ | | |
| Adiabatic energy efficiency (%) | $\eta_C = \frac{\dot{m}_p h_p}{\dot{m}_f \times HHV}$ | | |
| Exergy destruction (kJ/s) | $ \dot{I}_{C} = \dot{m}_{a}[(h_{a} - T_{0}s_{a}) + \varepsilon_{a_{1}}^{ch}] - \dot{m}_{p}(h_{p} - T_{0}s_{p}) + \dot{m}_{f}[(h_{f} - T_{0}s_{f}) + \varepsilon_{f_{2}}^{ch} - \varepsilon_{f_{3}}^{ch}] $ | | |
| Exergy efficiency (%) | $\psi_{C} = \frac{\dot{m}_{p}(h_{p} - T_{0}s_{p})}{\dot{m}_{f}[(h_{f} - T_{0}s_{f}) + \varepsilon_{f_{2}}^{ch} - \varepsilon_{f_{3}}^{ch}]}$ | | |
| <u>Heat exchanging unit</u> | | | |
| Heat loss (kJ/s) | $\dot{Q}_{H(loss)} = \dot{m}_p (h_p - h_g) - \dot{m}_w (h_s - h_w)$ | | |
| Energy efficiency (%) | $\eta_H = \frac{\dot{m}_w (h_s - h_w)}{\dot{m}_p (h_p - h_g)}$ | | |

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| Exergy destruction (kJ/s) | $\dot{I}_{H} = \dot{m}_{p}(h_{p} - T_{0}s_{p}) + \dot{m}_{w}[(h_{w} - T_{0}s_{w}) + \mathcal{E}_{w_{4}}^{ch}] - \dot{m}_{s}[(h_{s} - T_{0}s_{s}) + \mathcal{E}_{s_{5}}^{ch}] - \dot{m}_{g}[(h_{g} - T_{0}s_{g}) + \mathcal{E}_{g_{6}}^{ch}]$ | | | | | | | |
|-----------------------------------|--|--|--|--|--|--|--|--|
| Exergy efficiency (%) | $\psi_{H} = \frac{\dot{m}_{s}[(h_{s} - T_{0}s_{s}) + \varepsilon_{s_{5}}^{ch}] - \dot{m}_{w}[(h_{w} - T_{0}s_{w}) + \varepsilon_{w_{4}}^{ch}]}{\dot{m}_{p}(h_{p} - T_{0}s_{p}) - \dot{m}_{g}[(h_{g} - T_{0}s_{g}) + \varepsilon_{g_{6}}^{ch})]}$ | | | | | | | |
| Entire boiler | | | | | | | | |
| Energy efficiency (%) | $\eta_B = \frac{\dot{m}_s (h_s - h_w)}{\dot{m}_f h_f}$ | | | | | | | |
| Overall exergy destruction (kJ/s) | $\dot{I}_B = \dot{I}_C + \dot{I}_H$ | | | | | | | |
| Overall exergy efficiency (%) | $\psi_B = \frac{\dot{m}_s[(h_s - T_0 s_s) - (h_w - T_0 s_w) + \varepsilon_{s_5}^{ch} - \varepsilon_{w_4}^{ch}]}{\dot{m}_f[(h_f - T_0 s_f) + \varepsilon_{f_2}^{ch} - \varepsilon_{f_3}^{ch}]}$ | | | | | | | |
| | | | | | | | | |

Table 7: Thermodynamic analysis of material streams for the diesel and LPFO operated boiler

| | | Mass flow rate (kg/s) | | Temperature (⁰ C) | | Enthalpy (<i>kJ/kg</i>) | | Entropy (<i>kJ/kgK</i>) | |
|-------|-------------------------------|-----------------------|----------|-------------------------------|----------|---------------------------|----------|---------------------------|----------|
| Point | Stream | LPFO | Diesel | LPFO | Diesel | LPFO | Diesel | LPFO | Diesel |
| | | operated | operated | operated | operated | operated | operated | operated | operated |
| 1 | Air, \dot{m}_a , | 2.90 | 3.06 | 77.00 | 77.00 | 350.49 | 350.49 | 1.8571 | 1.8571 |
| 2 | Fuel, \dot{m}_f | 0.18 | 0.18 | 1345.69 | 1289.13 | 40515.15 | 42790.21 | 1.6664 | 1.7337 |
| 3 | Hot products, \dot{m}_p | 3.08 | 3.24 | 342.03 | 327.66 | 2697.38 | 2708.25 | 4.3857 | 4.5088 |
| 4 | Feed water, \dot{m}_w | 2.16 | 2.16 | 100.00 | 100.00 | 419.10 | 419.10 | 1.3070 | 1.3070 |
| 5 | Steam, \dot{m}_s | 2.16 | 2.16 | 170.40 | 170.40 | 2768.30 | 2768.30 | 6.6616 | 6.6616 |
| 6 | Exhaust flue gas, \dot{m}_g | 3.08 | 3.24 | 148.84 | 146.32 | 223.46 | 218.85 | 2.6882 | 2.7475 |

3. RESULTS AND DISCUSSION

3.1 Thermodynamics of material streams flow

The same values of combustion air temperature 77.00 ^oC, steam pressure 800kN/m², mass flow rate of fuel 0.18 kg/s, evaporation ratio 12, and excess air 20% respectively used in the analysis of the boiler operation resulted in higher fuel combustion temperature of 1345.69 ^oC in the LPFO operated boiler compared with 1289.13 ^oC in the diesel operated boiler as presented in Table 7. This was attributed to lower values of the specific heat capacity and air fuel ratio of LPFO as presented in Table 2 and used for computing the combustion fuel temperatures in the energy balance equation of the combustion unit. Consequently, higher combustion fuel temperature resulted in higher temperatures of products of hot flue and exhaust gases as indicated in Table 7.

3.2 Exergetic modelling

Previously derived thermodynamic formulations to determine exergetic values and efficiencies in a steam boiler operation given in Table 6 were used to develop empirical models for estimations and applications of oil fired steam boilers. Averaged values of thermodynamic variables, standard reference state temperature, specific chemical exergy of material streams and heating values given in Tables 4, 5 and 7, the exergetic expressions contained in Table 6 and expression relating *AAF* to *AFR*_{st} [19] were used to obtain the exergetic models given in equations (13) to (28).

 $E_{in} = m_f [350.49(1 + EA)AFR_{st} + 41652.68]$ (13)

$$\eta_{C(LPFO)} = 5.8433[(1 + EA)AFR_{st} + 1]$$
(14)

$$\eta_{C(Diesel)} = 5.57[(1 + EA)AFR_{st} + 1]$$
 (15)

$$I_C = m_f [39270.29 - 1521.66(1 + EA)AFR_{st}]$$
 (16)

$$\psi_C = 3.39[(1 + EA)AFR_{st} + 1] \tag{17}$$

$$\dot{Q}_{H(loss)} = m_f [2481.66\{(1+EA)AFR_{st}+1\}-2349.20E_r]$$
(18)

$$\eta_H = \frac{94.66E_r}{[(1+EA)AFR_{st}+1]}$$
(19)

$$\dot{I}_H = m_f [1817.92\{(1 + EA)AFR_{st} + 1\} - 1230.74E_r]$$
(20)

$$\psi_H = \frac{67.7E_r}{[(1+EA)AFR_{st}+1]}$$
(21)

$$\eta_B = 5.64E_r \tag{22}$$

$$\dot{I}_B = m_f [296.26(1 + EA)AFR_{st} - 1230.74E_r + 41088.21]$$
(23)

$$\psi_B = 3.03E_r \tag{24}$$

From equations (14), (15) and (17), the relation between the energy and exergy efficiencies of an adiabatic combustion unit of a steam boiler can be expressed respectively for LPFO and diesel operated boilers as

 $\eta_{C(LPFO)} = 1.724\psi_C \tag{25}$

 $\eta_{C(Diesel)} = 1.643 \psi_C \tag{26}$

From (19) and (21), the energy and exergy efficiencies relation of a heat exchanging unit of a steam boiler may be expressed as

$$\eta_H = 1.4\psi_H \tag{27}$$

From (22) and (24), the energy and exergy efficiencies relation of an entire steam boiler may be expressed as:

 $\eta_B = 1.8614 \psi_B \tag{28}$

The models make it possible to estimate vital exergetic information on the performance evaluation of oil-fired steam boilers with minimum input operational parameters, such as mass flow rate of fuel, air-fuel ratio and evaporation ratio. The models developed assumed linearity, as the respective coefficient of correlation and standard error of estimate obtained was approximately 1.00 and 0.00 respectively. The ratio of energy to exergy efficiencies of steam boiler was obtained as 1.8614. Previous study by Mehdi and Amir [31], obtained the energy to exergy efficiencies relation of boiler as 1.962, suggesting relatively higher energy or lower exergy efficiencies compared to that represented by the developed models in this study; they obtained the energy and exergy efficiencies of boiler to be 89.21% and 45.48%, respectively. The higher energy obtained

in their study could be attributed to higher evaporation ratio of 15.26. Applying the evaporation ratio used by Mehdi and Amir [31] into the exergetic efficiency models developed and given by equations (22) and (24), results in energy and exergy efficiency of 88.21% and 46.24% respectively which compare favourably with those obtained by Mehdi and Amir

3.3 Exergetic losses with changes in evaporation ratio and mass flow rate of fuel

Thermodynamic variables are critical to the exergetic performance and energy savings of steam boilers; hence, it is highly imperative to understand the mode of performance outputs with operational variables, such as, evaporation ratio and mass flow rates of fuel. Figure 2 shows that exergetic losses increased in boiler with changes in mass flow rate of fuel at constant evaporation ratio, while Figure 3 shows reducing values in exergetic losses with changes in evaporation ratio at constant mass flow rate of fuel. This clearly indicates that reduction in exergetic losses and consequently energy savings can be achieved in steam boilers by maintaining the input fuel value and increasing the evaporation ratio of steam boiler operation. Table 7 shows that the same steam pressure of 800 kN/m² resulted in the same saturated steam temperature of 170.00 °C. This implies that higher energy loss and exergy destruction are experienced in the heat exchanging unit of the diesel operated boiler as against that of the LPFO operated boiler as shown in Figures 2 and 3.

The overall exergy destruction of LPFO and diesel operated boilers were nearly the same irrespective of changes in evaporation ratio and mass flow rate of fuel. This is attributed to the same values of evaporation ratio, excess air and mass flow rate of fuel assumed for both boiler operations. The average overall boiler exergy destruction in the LPFO and diesel operated boiler operations increased from 3121.76 to 9365.27 kJ/s as the mass flow rate increases from 0.1 to 0.3 kg/s, while the average overall boiler exergy destruction decreased from 6062.23 to 5176.10 kJ/s (14.62%) as the evaporation ratio increases from 10 to 14 respectively

[31].

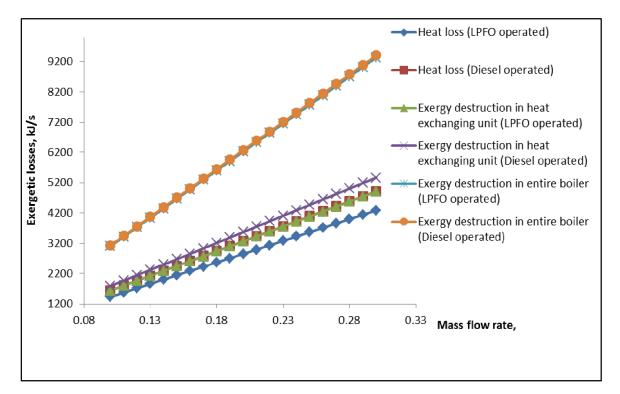


Figure 2: Exergetic losses in boiler with changes in mass flow rate of fuel at constant evaporation ratio

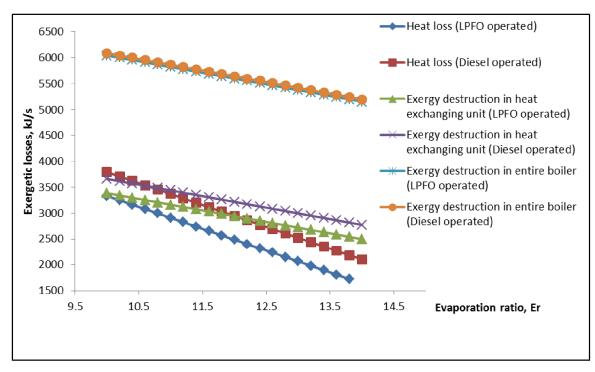


Figure 3: Exergetic losses in boiler with changes in evaporation ratio at constant mass flow rate of fuel

3.4 Effects of reference state temperature, entropy, feed water and steam temperatures

Figure 4 clearly shows reducing values in exergy efficiency with corresponding increases in reference state temperature, consequently, increases in exergy destruction in all units in a boiler. This is an indication that it is more advantageous to keep ambient temperature of a boiler system as low as possible. The combustion unit and entire boiler exergy efficiencies experienced slight decreases in exergy destruction and slight increases in exergy efficiencies as fuel entropy was increasing as shown in Figure 5, while Figure 8 shows that the heat exchanging unit and entire boiler experienced slight increases in exergy destruction and slight decreases in exergy efficiencies as feed water temperature and entropy

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was increasing. The average factor of exergy efficiency to entropy changes due to fuel entropy and feed water entropy was determined as 0.015 and 0.049 accounting as the lowest. The average factor of exergy efficiency to entropy changes due to exhaust flue gas entropy, combustion product entropy in heat exchanging and combustion units and superheated steam entropy were determined as 0.51, 0.76, 1.29 and 3.85 respectively from Figures 6, 7 and 9. This indicates that the largest amount of entropy changes in a boiler system is due to superheated steam entropy. As shown in Figure 9, increasing superheated steam temperature and entropy would result in decreases in exergy destruction and possible energy savings since superheated steam can be achieved by recycling of waste heat in boilers.

Figure 6 shows that combustion product entropy would result in increase in exergy destruction in the combustion unit while it leads to decreases in exergy destruction in the heat exchanging unit. Increases in flue gas entropy results in increases in exergy destruction in the heat exchanging unit as shown by Figure 7. The Figures 5 to 9 show that the key area to control irreversibility resulting from entropy effects is that due to combustion product entropy which leads to increased exergy destruction in the combustion unit.

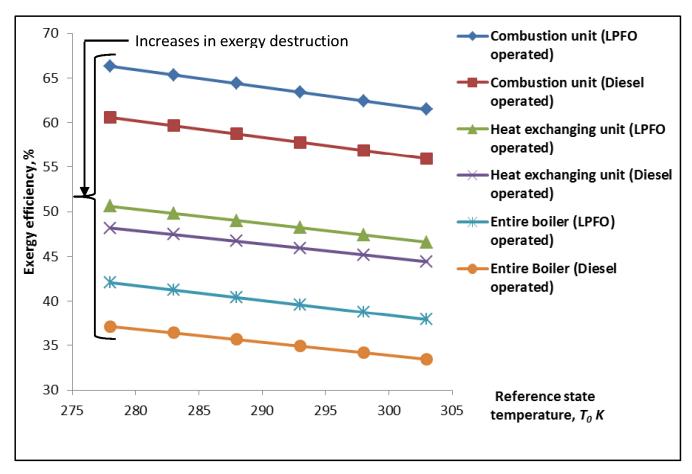


Figure 4: Exergy efficiency in boiler at different reference state

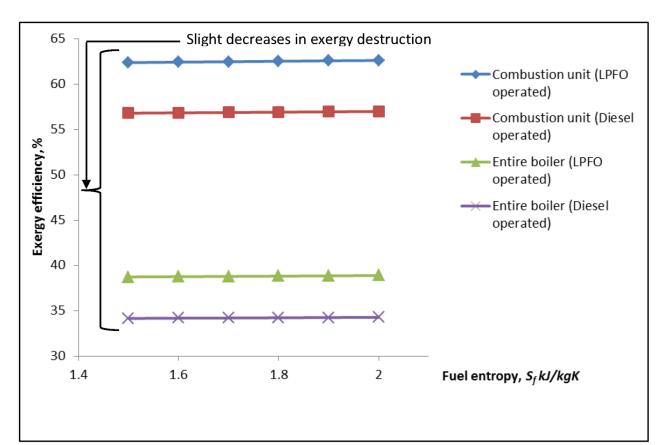


Figure 5: Exergy efficiency in boiler with respect to combustion fuel entropy

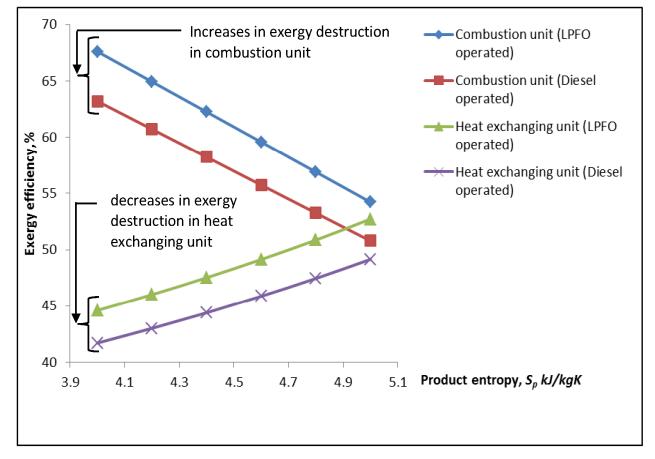


Figure 6: Exergy efficiency in boiler with respect to combustion product entropy

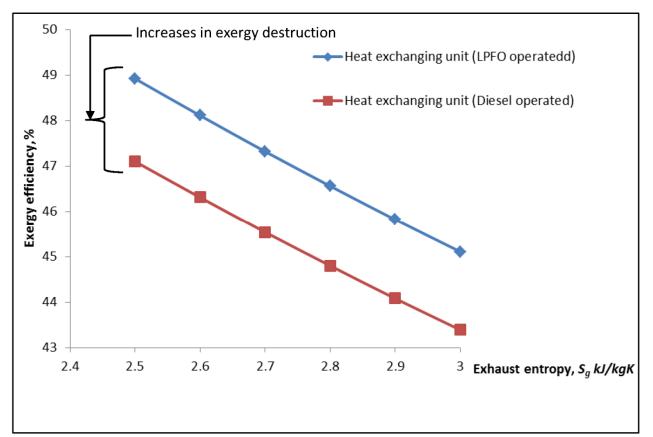


Figure 7: Exergy efficiency in boiler with respect to exhaust flue gas entropy

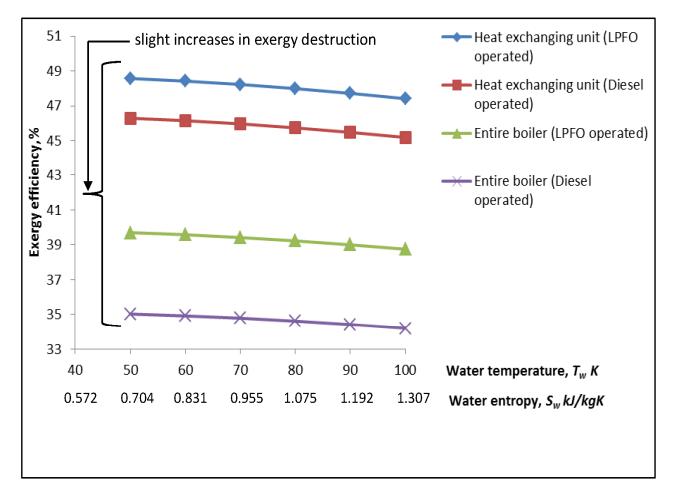


Figure 8: Exergy efficiency in boiler with respect to feed water temperature/entropy

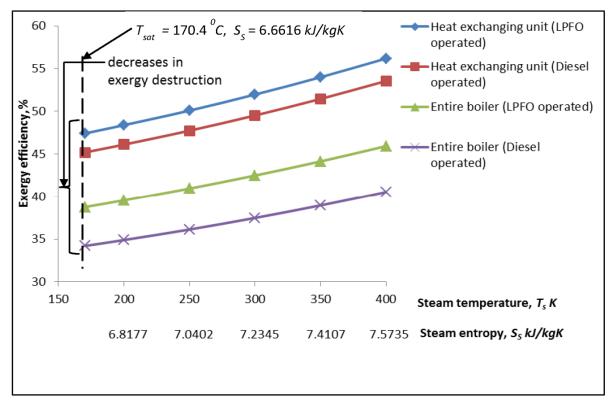


Figure 9: Exergy efficiency in boiler with respect to super heated steam temperature/entropy

4. CONCLUSION

Exergetic models to estimate exergetic quantities and efficiencies for performance evaluation of oil-fired steam boilers with minimum input operational parameters as mass flow rate of fuel, air-fuel ratio and evaporation ratio have been developed. The relation between the energy and exergy efficiencies of an adiabatic combustion unit for LPFO and diesel operated boilers, heat exchanging unit and of an entire steam boiler have been determined as

$$\eta_{C(LPFO)} = 1.724 \psi_C$$
, $\eta_{C(Diesel)} = 1.643 \psi_C$,

 $\eta_H = 1.4 \psi_H$ and $\eta_B = 1.8614 \psi_B$ respectively. Reduction in exergetic losses and subsequently gain in energy savings can be achieved in steam boilers by maintaining the input fuel value and increasing the evaporation ratio of steam boiler operation. The average overall boiler exergy destruction decreased from 6062.23 to 5176.10 kJ/s, that is, 14.62% as the evaporation ratio increased from 10 to 14. Increase in superheated steam temperature results in decreases in exergy destruction and possible energy savings as superheated steam can be achieved by recycling of waste heat in boilers. The major area to control irreversibility resulting from entropy effects is the combustion product entropy which leads to increased exergy destruction in the combustion unit. Increasing feed water temperature resulted in slight increases in exergy destruction of heat exchanging unit and entire boiler.

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