

## Exercise 2 (biomass)

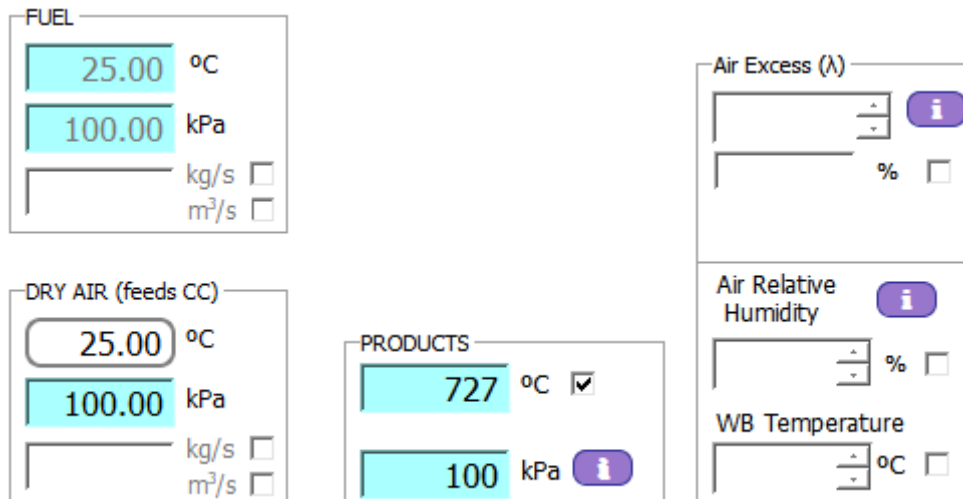
A solid fuel (rice husks) is burned in a non-adiabatic furnace. Knowing that dry air and fuel enter in standard conditions (25°C and 1 atm) and the flue-gas leaves at 727°C. The heat released to the environment is approximately 12502.7 kJ/kg of fuel.

Decide:

- Excess air coefficient assuming rich combustion
- Excess air coefficient assuming poor combustion
- Dry and wet combustion products
- Dew point temperature of combustion products
- Calorific powers
- Efficiency
- Irreversibility
- Ostwald diagram
- Kissel diagram

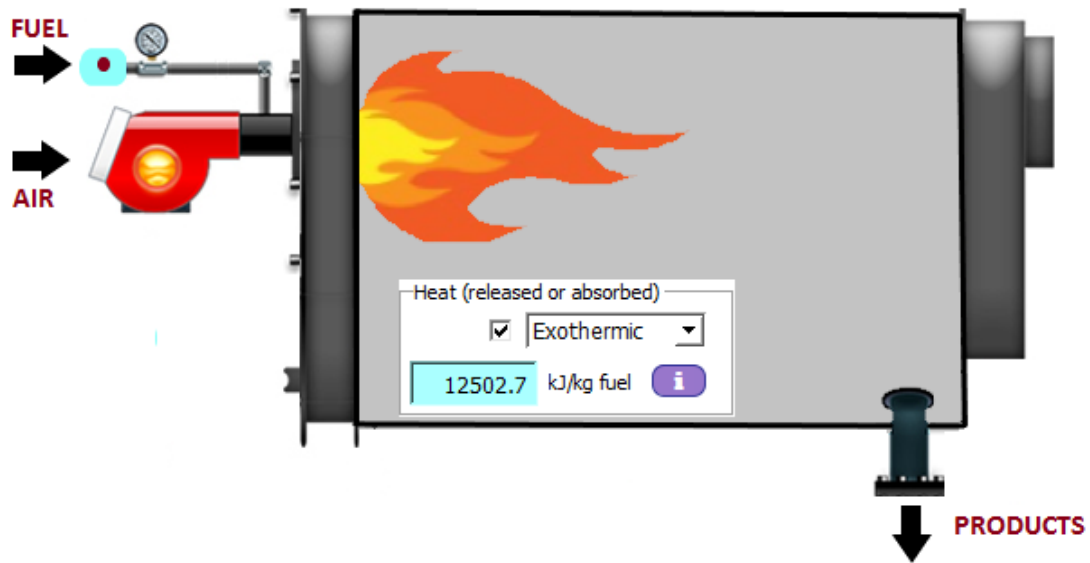


Solution by means THERMOCombustion:



The screenshot displays the software interface with the following settings:

- FUEL:** Temperature: 25.00 °C; Pressure: 100.00 kPa; Mass flow rate (kg/s) and Volume flow rate (m³/s) are set to 0.
- DRY AIR (feeds CC):** Temperature: 25.00 °C; Pressure: 100.00 kPa; Mass flow rate (kg/s) and Volume flow rate (m³/s) are set to 0.
- PRODUCTS:** Temperature: 727 °C (checked); Pressure: 100 kPa.
- Air Excess (λ):** Input field is empty; checkbox for percentage is unchecked.
- Air Relative Humidity:** Input field is empty; checkbox for percentage is unchecked.
- WB Temperature:** Input field is empty; checkbox for °C is unchecked.



FUEL selection

- Solid fossil fuel
- Coal sample
- Wood and woody biomass
- Herbaceous and agricultural biomass**
  - Grasses
  - Straws
  - Other residues
- Animal biomass
- Mixture biomass
- Waste fuel
- Biodiesel (FAME)
- Liquid fuel
- Empirical formula

Fuel

- Rice husks
- Palm fibres-husks
- Palm kernels
- Pepper plant
- Pepper residue
- Pistachio shells
- Plum pits
- Rice husks**
- Soya husks
- Sugar cane bagasse
- Sunflower husks



Biomass is used as a renewable solid fuel on power plants to provide CO<sub>2</sub>-neutral electricity and heat, where mainly wood and, in some cases, also straw have been applied in recent years. Biomass combustion has mainly been performed in grate boilers; however, wood and straw are used as fuels in suspension-fired power-plant boilers.

Molar Analysis		Ultimate Analysis		FUEL	
mol/kg fuel	Molar ratio %	Mass ratio %			
30.08650	43.99371	36.1369	C	CARBON	<input type="button" value="i"/>
22.18016	32.43274	4.4713	H <sub>2</sub>	HYDROGEN	
0.20933	0.30609	0.5864	N <sub>2</sub>	NITROGEN	
10.01003	14.63707	32.0321	O <sub>2</sub>	OXYGEN	<input type="button" value="i"/>
0.01827	0.02672	0.0586	S	SULPHUR	
5.88389	8.60366	10.6000	H <sub>2</sub> O	MOISTURE	
14.62241	g/mol (as AF)	16.1000	ASH	ASH	<input type="button" value="i"/>
15.99890	g/mol (as DAF)	0.0880	Cl	CHLORINE	
As received <input type="button" value="i"/>		100.0733	Total (%)		

Rice husk is the main byproduct of industrial processing of rice, and is a widely used byproduct in worldwide as biomass in energy production, reducing environmental impacts associated with improper disposal. Rice husk is the most prolific agricultural residue in rice producing countries around the world. It is one of the major by-products from the rice milling process and constitutes about 20% of paddy by weight. Rice husk, which consists mainly of lingo-cellulose and silica, is not utilized to any significant extent and has great potential as an energy source.

The ash generated in rice husk combustion has potential economic applications in a variety of sectors. However, the properties of rice husk ash are strongly dependent on the combustion method used. Rice is the second most consumed food item worldwide, with annual global demand of approximately 471 million tons and mean per capita consumption around 57 kg year. Accounting for roughly 20% of the bulk grain weight, rice husk is the most representative byproduct of industrial processing of rice.

**EMPIRICAL FORMULA**

C   H   O   N   S

3.009   4.436   2.002   0.042   0.002

Number of times the proportion is repeated:   Unknown  Rounding

Molecular formula   Normalized form   Formula known

Solid fuel

**PRODUCTS**

Incomplete Combustion i

Hydrogen in the products

There is hydrogen i

Water gas shift reaction

**CO + H<sub>2</sub>O ⇌ CO<sub>2</sub> + H<sub>2</sub>**

User Equilibrium temperature  °C

**K<sub>p</sub>** =

Molar ratio H<sub>2</sub>/CO

REFUSE

**Uncoked Coal**

**Uncoked Coal**

FC  %

ASH  %

VM  %

Total (%) -----

Through an iterative process applied to the energy balance, previously determined the enthalpy of biomass formation, the software determines that variable not defined in the initial menu (coefficient of excess air, heat exchanged, temperature of combustion products), in this case the coefficient of excess air. It is necessary to know a priori if the air-fuel mixture is rich or lean (both cases have been done in this problem).


Air/Fuel mixture

Lean,  $\lambda > 1$

Rich,  $\lambda < 1$

**CALCULATING..... wait please iiiii**

Excess Air Coefficient i



0.90000

**Equivalence Ratio**  
*(Relative Ratio, F<sub>R</sub>)*

1.11111

Flue gas (combustion products) composition is significantly affected by the combustion conditions. Where combustion takes place in complete, oxidizing gas species such as CO<sub>2</sub>, O<sub>2</sub>, SO<sub>2</sub>, and H<sub>2</sub>O. Neutral complete combustion is termed here when carbon and hydrogen in fuel is totally oxidized to carbon dioxide and water vapor, and no additional oxygen is present in the flue gases. In contrast, when combustion reactions proceed in air-deficient conditions, reducing gas species such as CO and H<sub>2</sub>. This last situation can also occur even with extra air in the combustion



**FLUE GAS (Combustion Products)**  On wet basis  On dry basis

	mol/kg fuel	<input type="checkbox"/> kg/kg fuel	Mole %	Mass %
CO <sub>2</sub>	23.84962	1.04962	14.5448	22.3542
CO	6.23688	0.17470	3.8036	3.7207
H <sub>2</sub> O	28.05165	0.50536	17.1074	10.7629
N <sub>2</sub>	105.79256	2.96362	64.5180	63.1175
O <sub>2</sub>				
SO <sub>2</sub>	0.01827	0.00117	0.0111	0.0249
H <sub>2</sub>				
Ar				
HCl	0.02468	0.00090	0.0151	0.0192
<b>TOTAL</b>	<b>163.9736</b>	<b>4.6954</b>	<b>100.00 %</b>	<b>100.00 %</b>

Water Dew Point T\* (and P<sub>sat</sub>)    H<sub>2</sub>O (liquid)     mol/mol fuel  
 %     kg/kg fuel

**56.73** °C     17.11 kPa

The dew point is the temperature where air is no longer capable of holding the water vapor that is contained within it. At the dew point temperature, water vapor condenses into liquid water. At all times, the dew point temperature is equal to or less than the air temperature. We can observe that this temperature is 56.73°C. A thorough knowledge about dew point temperature is very important in various industries because moisture can affect corrosion rates.



**FLUE GAS (Combustion Products)**  On wet basis  On dry basis

	mol/kg fuel	<input type="checkbox"/> kg/kg fuel	Mole %	Mass %
CO <sub>2</sub>	23.84962	1.04962	17.5466	25.0506
CO	6.23688	0.17470	4.5886	4.1695
H <sub>2</sub> O				
N <sub>2</sub>	105.79256	2.96362	77.8333	70.7308
O <sub>2</sub>				
SO <sub>2</sub>	0.01827	0.00117	0.0134	0.0279
H <sub>2</sub>				
Ar				
HCl	0.02468	0.00090	0.0182	0.0215
<b>TOTAL</b>	<b>135.9220</b>	<b>4.1900</b>	<b>100.00 %</b>	<b>100.00 %</b>

*FUEL without moisture+DRY AIR* Total O2 (REACTIVE) ▾

mol/kg fuel  kg/kg fuel

*FUEL with moisture+WET AIR*

mol/kg fuel  kg/kg fuel

*Reactants (fuel + dry air)*

kg/kg fuel

(Assume 1 kg fuel is being combusted)

Air/fuel ratio (dry air)

*Theoretical* *Actual* AIR ▾

Dry air Dry air

mol/kg fuel

kg/kg fuel

Experimental correlations (mass fractions)

Burnham: C,H,N,O,S

*(Based on the dry and mineral matter free elemental fuel composition)*

**Lower Heating, LHV**      **Higher Heating, HHV**  
**(DAF)**

18095.60      kJ/kg fuel      19435.16

We can check the calculation of the enthalpy of formation of a solid fuel:

For  $\lambda=1$  (neutral reaction condition):

**FLUE GAS (Combustion Products)**       On wet basis       On dry basis

	mol/kg fuel	<input type="checkbox"/> kg/kg fuel	Mole %	Mass %
CO <sub>2</sub>	30.08648	1.32410	17.1236	25.8427
H <sub>2</sub> O	28.05016	0.50533	15.9647	9.8626
N <sub>2</sub>	117.52191	3.29220	66.8872	64.2543
SO <sub>2</sub>	0.01840	0.00118	0.0105	0.0230

$$H_{\text{form}} \text{ (kJ/mol fuel)} = \text{PCS} + n_{\text{H}_2\text{O}} H_f(\text{liq})_{\text{H}_2\text{O}} + n_{\text{CO}_2} H_f(\text{gas})_{\text{CO}_2} + n_{\text{SO}_2} H_f(\text{gas})_{\text{SO}_2} =$$

$$19435.16 \text{ kJ/kg fuel} + 28.05016 \text{ moles H}_2\text{O/kg fuel} \cdot (-285.83) \text{ kJ/mol H}_2\text{O} + 30.08648 \text{ moles CO}_2/\text{kg fuel} \cdot (-393.52) \text{ kJ/mol CO}_2 + 0.0184 \text{ moles SO}_2/\text{kg fuel} \cdot (-296.991) \text{ kJ/mol SO}_2 =$$

$$= 19434.16 - 8017.57 - 11834.63 - 5.46 = -427.5 \text{ kJ/mol fuel (approximate value to that obtained by ThermoCombustion, the differences are due to the decimals used)}$$

**Properties of the FUEL** ( $T_0=25\text{ }^\circ\text{C}$ ,  $P_0=100\text{ kPa}$ )

**Enthalpy of formation** Experimental correlations   Energy balance

(DAF)  kJ/kg fuel  (As received)

**Absolute entropy** Eiserman et al. 1980 (coal, char, ash)  (Experimental expression)

(DAF)  kJ/kg fuel K

**Standard chemical exergy** Experimental correlations  (Experimental expression)  Exergy balance

(DAF)  kJ/kg fuel  (As received)

**Standard Gibbs Free Energy of Formation**

(DAF)  kJ/kg fuel

**H/C molar ratio**  mol H/mol C

**O/C molar ratio**  mol O/mol C

**Carbon content**  kg C/kg fuel

**Hydrogen content**  kg H/kg fuel

**C/H mass ratio**  kg C/kg H

**N/C molar ratio**  mol N/mol C

**Oxygen content**  kg O/kg fuel

We can check the heat transferred to the environment:





**Heat (Released or Absorbed)** i Exothermic  
Exo = release ENERGY

12502.689 kJ/kg fuel

kJ/kg fuel     kW

$$Q_f = \sum_{\text{Products}} n_{\text{out}} h_{\text{out}}(T_{\text{out}}) - \sum_{\text{Reactants}} n_{\text{in}} h_{\text{in}}(T_{\text{in}})$$

Reactants -12934.12 kJ/kg fuel    -431.44

Products

$Q_f = H_P(T_{\text{out}}, P) - H_R(T_{\text{in}}, P) = H_P(T_{\text{out}}, P) - H_P(T_{\text{adr}}, P)$

$H = H_{\text{form}} + H(T, P) - H(25^\circ\text{C}, 1 \text{ atm})$

Standard Enthalpy of Formation

**FUEL** -6.309 kJ/mol fuel    -431.436 kJ/kg fuel i

ENERGY BALANCE: REACTANTS    ?    X

	kJ/mol	kJ/kg fuel		
O <sub>2</sub>	0.000	0.000	A I R	
N <sub>2</sub>	0.000	0.000		
Ar				
CO <sub>2</sub>				
H <sub>2</sub> O	0.000	0.000		
TOTAL	0.000	0.000		
Fuel (DAF)	-6.309	-431.436	<input checked="" type="checkbox"/>	F U E L
Ash	0.000	0.000	<input checked="" type="checkbox"/>	
TOTAL	-6.309	-431.436		
TOTAL	-6.309	-431.436		(AIR+FUEL)

ENERGY BALANCE: FLUE GAS

	<i>kJ/mol</i>	<i>kJ/kg fuel</i>
CO <sub>2</sub>	-360.1150	-8588.6064
CO	-88.8440	-554.1094
H <sub>2</sub> O	-215.8420	-6054.7241
O <sub>2</sub>	0.0000	0.0000
N <sub>2</sub>	21.4600	2270.3081
H <sub>2</sub>	0.0000	0.0000
SO <sub>2</sub>	-262.7260	-4.8000
HCl	-88.8377	-2.1925
Ar	0.0000	0.0000
<b>TOTAL</b>	<b>-994.9047</b>	<b>-12934.1230</b>

**Indirect Method: Heat Loss Method** Boiler/Furnace Efficiency

*Dry flue gas loss (Sensible heat)*  %

*Presence of H<sub>2</sub> in fuel (Latent heat)*  %

*Unburned fuel, CO*  %

*Moisture in fuel, H<sub>2</sub>O*  %

*Moisture in air, RH*  %

*Unburned carbon\* in fly ash*  %

*Unburned carbon\* in bottom ash*  %

*Surface loss (radiation, convection) & unaccounted losses*

Industrial fire tube/package boiler  %

DAF

Referred to HHV

\* Only for solid fuel

**100 - Σ loss** **59.220** %

Dry Refuse LOST

%C

%VM

%HHV



Entropy Terms (kJ/kg fuel K) AVAILABILITY

**Generated Entropy ( $S_g$ )** 24.32 ●

$$S_g = -\frac{Q_f}{T_f} + \sum_{\text{Products}} n_{\text{out}} S_{\text{out}} - \sum_{\text{Reactants}} n_{\text{in}} S_{\text{in}}$$

Entropy balance ▾  Reactants  Products

i
12.50
11.82
i

⊗

ENTROPY BALANCE: FLUE GAS ? ×

	<i>J/mol K</i>	<i>kJ/kg fuel K</i>	
O <sub>2</sub>	218.009	6.119	A I R
N <sub>2</sub>	193.462	20.426	
Ar			
CO <sub>2</sub>			
H <sub>2</sub> O	0.000	0.000	
TOTAL	411.471	26.545	
Fuel	22.382	1.531	F U E L
(DAF)			
Ash	0.000	0.000	
TOTAL	22.382	1.531	
TOTAL	433.853	28.076	(AIR+FUEL)

ENTROPY BALANCE: FLUE GAS

	<i>J/mol K</i>	<i>kJ/kg fuel K</i>
CO <sub>2</sub>	285.2448	6.8030
CO	261.6030	1.6316
H <sub>2</sub> O	247.2776	6.9365
O <sub>2</sub>	0.0000	0.0000
N <sub>2</sub>	231.7007	24.5122
H <sub>2</sub>	0.0000	0.0000
SO <sub>2</sub>	358.2118	0.0065
HCl	307.6989	0.0076
Ar	0.0000	0.0000
<b>TOTAL</b>	<b>1691.7367</b>	<b>39.8975</b>

Considering a lean mixture, with excess air:

Air/Fuel mixture

Lean,  $\lambda > 1$

Rich,  $\lambda < 1$

FLUE GAS (Combustion Products)

On wet basis  On dry basis

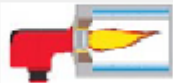
	<i>mol/kg fuel</i>	<input type="checkbox"/> <i>kg/kg fuel</i>	<i>Mole %</i>	<i>Mass %</i>
CO <sub>2</sub>	30.08648	1.32410	14.0763	20.2927
CO				
H <sub>2</sub> O				
N <sub>2</sub>	169.73137	4.75477	79.4107	72.8700
O <sub>2</sub>	13.87781	0.44409	6.4929	6.8060
SO <sub>2</sub>	0.01827	0.00117	0.0085	0.0179
H <sub>2</sub>				
Ar				
HCl	0.02468	0.00090	0.0115	0.0138
<b>TOTAL</b>	<b>213.7386</b>	<b>6.5250</b>	<b>100.00 %</b>	<b>100.00 %</b>

**FLUE GAS (Combustion Products)**  On wet basis  On dry basis

	mol/kg fuel	<input type="checkbox"/> kg/kg fuel	Mole %	Mass %
CO <sub>2</sub>	30.08648	1.32410	12.4432	18.8339
CO				
H <sub>2</sub> O	28.05165	0.50536	11.6016	7.1882
N <sub>2</sub>	169.73137	4.75477	70.1978	67.6316
O <sub>2</sub>	13.87781	0.44409	5.7396	6.3167
SO <sub>2</sub>	0.01827	0.00117	0.0076	0.0166
H <sub>2</sub>				
Ar				
HCl	0.02468	0.00090	0.0102	0.0128
<b>TOTAL</b>	<b>241.7903</b>	<b>7.0304</b>	<b>100.00 %</b>	<b>100.00 %</b>

Water Dew Point T<sup>\*</sup> (and P<sub>sat</sub>)    H<sub>2</sub>O (liquid)     mol/mol fuel  
 °C     kPa     %     kg/kg fuel

Excess Air Coefficient:



**Equivalence Ratio (Relative Ratio, F<sub>R</sub>)**

We verify the heat transferred to the environment:



**Heat (Released or Absorbed)** i **Exothermic** **Exo = release** **ENERGY**

12507.313 kJ/kg fuel

kJ/kg fuel     kW

Reactants

Products

$$Q_f = \sum_{\text{Products}} n_{\text{out}} h_{\text{out}}(T_{\text{out}}) - \sum_{\text{Reactants}} n_{\text{in}} h_{\text{in}}(T_{\text{in}})$$

-12938.75 kJ/kg fuel    -431.44

$Q_f = H_P(T_{\text{out}}, P) - H_R(T_{\text{in}}, P) = H_P(T_{\text{out}}, P) - H_P(T_{\text{adr}}, P)$

$H = H_{\text{form}} + H(T, P) - H(25^\circ\text{C}, 1 \text{ atm})$

Standard Enthalpy of Formation

FUEL

-6.309 kJ/mol fuel    -431.436 kJ/kg fuel    i

ENERGY BALANCE: FLUE GAS ?    X

	kJ/mol	kJ/kg fuel
CO <sub>2</sub>	-360.1150	-10834.5918
CO	0.0000	0.0000
H <sub>2</sub> O	-215.8420	-6054.7241
O <sub>2</sub>	22.7070	315.1234
N <sub>2</sub>	21.4600	3642.4353
H <sub>2</sub>	0.0000	0.0000
SO <sub>2</sub>	-262.7260	-4.8000
HCl	-88.8377	-2.1925
Ar	0.0000	0.0000
<b>TOTAL</b>	-883.3536	-12938.7500

**Indirect Method: Heat Loss Method** Boiler/Furnace Efficiency

Dry flue gas loss <i>(Sensible heat)</i>	30.905 %
Presence of H <sub>2</sub> in fuel <i>(Latent heat)</i>	7.929 %
Unburned fuel, CO	0.000 %
Moisture in fuel, H <sub>2</sub> O	2.104 %
Moisture in air, RH	0.000 %
Unburned carbon * in fly ash	0.000 %
Unburned carbon * in bottom ash	0.000 %

*\* Only for solid fuel*

Surface loss (radiation, convection) & unaccounted losses  
Industrial fire tube/package boiler  %

DAF  **57.062 %**

Referred to HHV

Dry Refuse LOST  
 %C   
 %VM   
 %HHV

There is a higher efficiency (indirect method), because there is no loss due to unburnt.

Entropy Terms (kJ/kg fuel K)

**Generated Entropy (S<sub>g</sub>)** 26.93 ●

Entropy balance  Reactants  Products

**AVAILABILITY**

$$S_g = -\frac{Q_f}{T_f} + \sum_{\text{Products}} n_{\text{out}} S_{\text{out}} - \sum_{\text{Reactants}} n_{\text{in}} S_{\text{in}}$$

In this case, the exergy destruction is greater than in the rich mixture case.



ENTROPY BALANCE: FLUE GAS ? X

	<i>J/mol K</i>	<i>kJ/kg fuel K</i>	
O <sub>2</sub>	218.009	9.824	A I R
N <sub>2</sub>	193.462	32.796	
Ar			
CO <sub>2</sub>			
H <sub>2</sub> O	0.000	0.000	
<b>TOTAL</b>	411.471	42.620	
<hr/>			
Fuel	22.382	1.531	F U E L
(DAF)			
Ash	0.000	0.000	
<b>TOTAL</b>	22.382	1.531	
<hr/>			
<b>TOTAL</b>	433.853	44.151 (AIR+FUEL)	

ENTROPY BALANCE: FLUE GAS ? X

	<i>J/mol K</i>	<i>kJ/kg fuel K</i>	
CO <sub>2</sub>	286.5424	8.6211	
CO	0.0000	0.0000	
H <sub>2</sub> O	250.5067	7.0271	
O <sub>2</sub>	267.2321	3.7086	
N <sub>2</sub>	230.9991	39.2078	
H <sub>2</sub>	0.0000	0.0000	
SO <sub>2</sub>	361.3613	0.0066	
HCl	310.9607	0.0077	
Ar	0.0000	0.0000	
<b>TOTAL</b>	1707.6023	58.5789	



Thermodynamic analysis of ashes:

**Ash constituents and Properties of the Ash** ? ×

Coal fly ash dependent of the region  Biomass  Lignitic ash  Bituminous ash

Reset

	Mass ratio %	Molar ratio %	Molecular weight (kg/kmol)	Standard enthalpy (kJ/kmol)	Standard entropy (kJ/kmol)	Standard chemical exergy (kJ/mol)
SiO <sub>2</sub>	94.4800	95.8610	60.0	-911300	41.9	1.6400
Al <sub>2</sub> O <sub>3</sub>	0.2100	0.1253	102.0	-1674400	51.1	4.4800
Fe <sub>2</sub> O <sub>3</sub>	0.2200	0.0837	160.0	-825900	87.0	17.6600
CaO	0.9700	1.0545	56.0	-634600	38.8	129.8800
MgO	0.1900	0.2892	40.0	-601500	26.8	62.4200
TiO <sub>2</sub>	0.0200	0.0152	80.0	-945200	50.2	21.2200
Na <sub>2</sub> O	0.1600	0.1571	62.0	-418200 *	75.0	296.3200
K <sub>2</sub> O	2.2900	1.4831	94.0	-418200 *	102.0	412.5400
SO <sub>3</sub>	0.9200	0.7001	80.0	-437900	257.0	242.0000
P <sub>2</sub> O <sub>5</sub>	0.5400	0.2315	142.0	-1505990	117.0	377.1200
Others	0.0000	0.0000	44.0	-393520	214.0	20.1400
<b>Total</b>	<b>100.000</b>	<b>100.001</b>	<b>60.877</b> <i>kg/kmol</i>	<b>-903552</b> <i>kJ/kmol ash</i>	<b>44.498</b> <i>kJ/kmol ash °C</i>	<b>11.696</b> <i>kJ/mol ash</i>

High-temperature ash analysis (815 °C)

**Enthalpy of formation**  
-14793.809  
*kJ/kg ash*

**Enthalpy of formation**  
-2381.803  
*kJ/kg fuel*

**Ratio of bottom ash to fly ash**  
 %  
 29.3761 TCO + 30.67  
**0.731** *kJ/kg ash °C*  
**0.118** *kJ/kg fuel °C*  
**192.122** *kJ/kg ash*  
**30.932** *kJ/kg fuel*

\* Na<sub>2</sub>O + K<sub>2</sub>O

One of the main problems of substituting fossil fuels by biomass in power-plant boilers is the biomass ash properties. Biomass fuels usually have a high content of alkali metals, which, together with other mineral components of the ash, give rise to severe ash deposition, thereby reducing the heat transfer and inducing increased boiler tube corrosion.

