

FUEL ECONOMY AT THE HEAT TREATMENT FURNACES OF STEEL CAST PIECES

Lilica IVANESCU, Alexandru IVANESCU,
Antoaneta ENE

"Dunarea de Jos" University of Galati
e-mail: ivanescu.lilica@ugal.ro

ABSTRACT

The paper presents a study made at the annealing furnaces for homogenization of cast pieces regarding the replacement of marsh gas with blast furnace gas, in a proportion of 100%. In this way an important economy of marsh gas is achieved, representing a special interest because its international deficit and its putting up the price.

KEYWORDS : blast furnace gas, oxygen, fuel economy

1. Introduction

At present, in Mixed Foundry of Mittal Steel Iron and Steel Works at Galati, at the heat treatment furnaces of steel moulded pieces, the marsh gas is used as fuel [1-3] which is, as we know, an expensive and defficitary fuel. On the other hand in many foundries, as is the case of the above mentioned one, there are fuel wastes such as coke gas or furnace gas.

The paper presents a study made at the annealing furnaces of the steel moulded pieces homogenization regarding the replacement of marsh gas with furnace gas in a 100% proportion. Thus an important economy of march gas is obtained. The provision that has to be complied with is that the same amount of heat as in the case of the replaced marsh gas should be obtained in the working space of

the furnace, in the different options of fuel utilization. In order to achieve that, in the case of the combustion of the blast furnace gas to a 100% degree, the using of oxygen-enriched air during combustion has to be done. In this paper the concentration of oxygen in the oxygen-enriched air is determined in the case of the hundred percent combustion of furnace gas.

2. Technical characteristics of the annealing furnaces of the steel moulded pieces homogenization

- The furnace capacity (pieces + supports), $C=75t$;
- The bedstone surface, $S_v=32\text{ m}^2$.
- The maxim piece temperature, $t_m=1100\text{ }^\circ\text{C}$;

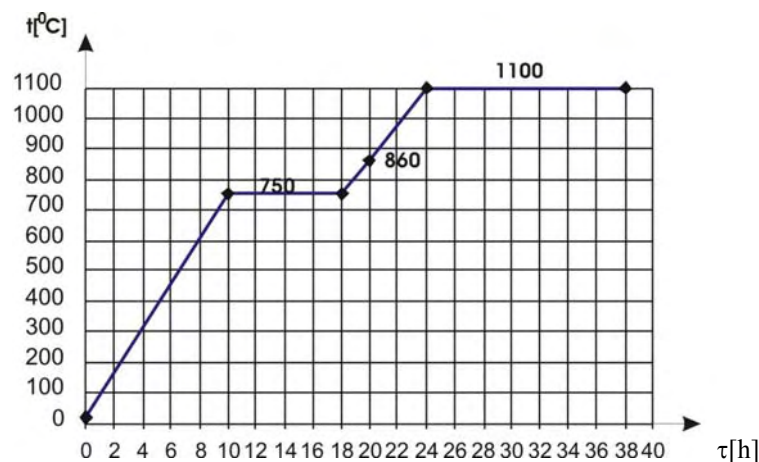


Fig.1. The diagram of the steel moulded pieces heating in the homogenizing annealing furnaces

- The furnace temperature (technological),
 $t_{\text{tech}} = t_m + (100-150)^{\circ}\text{C}$;
- The burning unit:
12 burners x 50 Nm³/h
12 burners x 20 Nm³/h
- Marsh gas consumption,
 $D_{\text{CH}_4} = 400 \text{ Nm}^3/\text{h}$;
- The cycle duration, $\tau_{\text{cycle}} = 38\text{h}$;
- The heating diagram (see fig.1).

3. The calculus of the fuel burning

The burning calculus is made in two varieties, namely marsh gas and furnace gas in a 100% proportion, respectively.

The calculus relations are the following [4]:

- the fuel chemical composition:

$$\begin{aligned} \text{CH}_4 + \text{CO} + \text{H}_2 + \text{CO}_2 + \text{O}_2 + \text{N}_2 + \text{H}_2\text{O} &= 100 \% \\ [\text{CH}_4] + [\text{CO}] + [\text{H}_2] + [\text{CO}_2] + \\ + [\text{O}_2] + [\text{N}_2] + [\text{H}_2\text{O}] &= 1 \text{ Nm}^3 \end{aligned} \quad (1)$$

- the inferior calorific power H_i , in [kJ/Nm³ fuel]:

$$\begin{aligned} H^i &= [\text{CH}_4] \cdot H_i^{\text{CH}_4} + [\text{CO}] \cdot H_i^{\text{CO}} + \\ &+ [\text{H}_2] \cdot H_i^{\text{H}_2} \end{aligned} \quad (2)$$

where:

$[\text{CH}_4], [\text{CO}], [\text{H}_2]$ - the quantities of the combustion elements, in [Nm³/Nm³ fuel];

$H_i^{\text{CH}_4}, H_i^{\text{CO}}, H_i^{\text{H}_2}$ = inferior calorific powers of the combustion elements, in [kJ/Nm³];

$$H_i^{\text{CH}_4} = 35500 \text{ kJ/Nm}^3$$

$$H_i^{\text{CO}} = 12740 \text{ kJ/Nm}^3$$

$$H_i^{\text{H}_2} = 12765 \text{ kJ/Nm}^3$$

- the minimum amount of oxygen necessary for the combustion, O_2^{min} , in [Nm³O₂/Nm³ fuel]

$$O_2^{\text{min}} = 2[\text{CH}_4] + \frac{1}{2}[\text{CO}] + \frac{1}{2}[\text{H}_2] - [\text{O}_2] \quad (3)$$

-the theoretic air necessary for the combustion, L_0 , in [Nm³air/Nm³ fuel]:

$$L_0 = \frac{100}{21} \cdot O_2^{\text{min}} \quad (4)$$

- the real air used for the combustion, L_r , in [Nm³O₂/Nm³ fuel]:

$$L_r = \lambda L_0$$

where:

λ – the coefficient of excess air; the adopted value:
 $\lambda = 1.05$

-the quantity of the burned gases, V_{ga} , in [Nm³ ga/Nm³ fuel]:

$$V_{\text{ga}} = V_{\text{CO}_2} + V_{\text{H}_2\text{O}_{\text{vap}}} + V_{\text{N}_2} + V_{\text{O}_2\text{excess}} \quad (5)$$

where:

$$V_{\text{CO}_2} = [\text{CO}_2] + [\text{CH}_4] + [\text{CO}]$$

$$V_{\text{H}_2\text{O}_{\text{vap}}} = 2[\text{CH}_4] + [\text{H}_2] + [\text{H}_2\text{O}]$$

$$V_{\text{N}_2} = 0.79L_r + [\text{N}_2]$$

$$V_{\text{O}_2\text{excess}} = 0.21(L_r - L_0)$$

-the participations of the component elements of the burned gases, r_i :

$$r_{\text{CO}_2} = \frac{V_{\text{CO}_2}}{V_{\text{ga}}} \quad (6)$$

$$r_{\text{H}_2\text{O}_{\text{vap}}} = \frac{V_{\text{H}_2\text{O}_{\text{vap}}}}{V_{\text{ga}}}$$

$$r_{\text{N}_2} = \frac{V_{\text{N}_2}}{V_{\text{ga}}}$$

$$r_{\text{O}_2\text{excess}} = \frac{V_{\text{O}_2\text{excess}}}{V_{\text{ga}}}$$

$$\sum_{i=1}^4 r_i = 1$$

-the real burning temperature [4], t_r , [^oC]:

$$t_r = \eta_{\text{pyr}} \frac{0.98 \cdot H_i}{V_{\text{ga}} \cdot c_{\text{ga}}} \quad (7)$$

where:

η_{pyr} - the pyrometric efficiency; the adopted value

$\eta_{\text{pyr}} = 0.7$;

H^i – the inferior calorific power, [kJ/Nm³ fuel];

V_{ga} – the volume of the burned gases, [Nm³/Nm³ fuel],

c_{ga} – the specific heat of the burned gases [kJ/Nm³grd].

-the specific heat of the burned gases c_{ga} , [kj/Nm³grd]:

$$c_{\text{ga}} = \sum_{i=1}^4 c_i \cdot r_i \quad (8)$$

where:

c_i – the specific heats of the components of the burned gases, which are labelled as a function of temperature. The adopted value for the temperature of

the burned gases is $t_r = 1000\text{ }^{\circ}\text{C}$ and $c_{ga}^{1000\text{ }^{\circ}\text{C}}$ can be calculated using the values:

$$c_{\text{CO}_2}^{1000\text{ }^{\circ}\text{C}} = 2.2035 \text{ kJ/Nm}^3\text{grd}$$

$$c_{\text{H}_2\text{O}_{\text{vap}}}^{1000\text{ }^{\circ}\text{C}} = 1.7229 \text{ kJ/Nm}^3\text{grd}$$

$$c_{\text{N}_2}^{1000\text{ }^{\circ}\text{C}} = 1.3917 \text{ kJ/Nm}^3\text{grd}$$

$$c_{\text{O}_2}^{1000\text{ }^{\circ}\text{C}} = 1.4775 \text{ kJ/Nm}^3\text{grd}$$

It results:

3.1. The burning of the marsh gas

$$c_{ga}^{1000\text{ }^{\circ}\text{C}} = 0.091 \cdot 2.2035 + 0.182 \cdot 1.7229 + 0.717 \cdot 1.3917 + 0.01 \cdot 1.4775$$

$$c_{ga}^{1000\text{ }^{\circ}\text{C}} = 1.529 \text{ kJ/Nm}^3\text{grd}$$

$$t_r = \frac{0.7 \cdot 0.98 \cdot 35500}{11 \cdot 1.529} = 1447.9\text{ }^{\circ}\text{C} \cong 1448\text{ }^{\circ}\text{C}$$

$$\text{error} = \frac{t_r' - t_r}{t_r'} 100 =$$

$$= \frac{1448 - 1000}{1448} 100 = 30.9\% \gg 5\%$$

With the admitted error of 5%, $c_{ga}^{1448\text{ }^{\circ}\text{C}}$ is calculated, by determining the specific heats of the components by interpolation, as a function of their values at the temperatures of $1400\text{ }^{\circ}\text{C}$ and $1500\text{ }^{\circ}\text{C}$.

$$c_{\text{CO}_2}^{1448\text{ }^{\circ}\text{C}} = 2.3136 + \frac{48 \cdot 0.0218}{100} = 2.324$$

$$c_{\text{H}_2\text{O}}^{1448\text{ }^{\circ}\text{C}} = 1.828 + \frac{48 \cdot 0.0247}{100} = 1.8398$$

$$c_{\text{N}_2}^{1448\text{ }^{\circ}\text{C}} = 1.4348 + \frac{48 \cdot 0.0092}{100} = 1.4392$$

$$c_{\text{O}_2}^{1448\text{ }^{\circ}\text{C}} = 1.5203 + \frac{48 \cdot 0.0091}{100} = 1.5246$$

$$c_{ga}^{1448\text{ }^{\circ}\text{C}} = 0.091 \cdot 2.324 + 0.182 \cdot 1.8398 + 0.717 \cdot 1.4392 + 0.01 \cdot 1.5246$$

$$c_{ga}^{1448\text{ }^{\circ}\text{C}} = 1.593 \text{ kJ/Nm}^3\text{grd}$$

$$t_r'' = \frac{0.7 \cdot 0.98 \cdot 35500}{11 \cdot 1.593} = 1389.7 \cong 1390\text{ }^{\circ}\text{C}$$

$$\text{error} = \frac{1390 - 1448}{1390} 100 = -4\% < -5\%$$

$$t_r^{\text{CH}_4} = 1390\text{ }^{\circ}\text{C}$$

3.2. The burning of the furnace gas

$$c_{ga}^{1000\text{ }^{\circ}\text{C}} = 0.2571 \cdot 2.2035 + 0.056 \cdot 1.7229 + 0.6824 \cdot 1.3917 + 0.0045 \cdot 1.4775 = 1.619$$

$$t_r' = \frac{0.7 \cdot 0.98 \cdot 3716}{1.581 \cdot 1.619} = 995.9 \cong 996\text{ }^{\circ}\text{C}$$

$$\text{error} = \frac{996 - 1000}{996} 100 = -0.4\%$$

$$t_r^{\text{GF}} = 996\text{ }^{\circ}\text{C}$$

The thermodynamic parameters at the burning with atmospheric air of the marsh gas or furnace gas respectively, and the chemical composition of the fuel are given in Tables 1 and 2.

The thermodynamic parameters at the burning with atmospheric air

Table 1

Fuel	The chemical composition of the fuel [%]								The calorific power, H_i [kJ/Nm ³]	O_2^{min} [Nm ³ /Nm ³ fuel]
	CH ₄	CO	H ₂	CO ₂	O ₂	N ₂	H ₂ O	Tot.		
Marsh gas	100	-	-	-	-	-	-	100	35500	2
Furnace gas	0.20	22.45	6.15	18.00	0.20	50.70	2.30	100	3716	0.145

Table 2

Fuel	L ₀ [Nm ³ / Nm ³ fuel]	L _r [Nm ³ / Nm ³ fuel]	V _{ga} [Nm ³ /Nm ³ fuel]					Participations in the burned gases, r _i					Burning temp.
			V _{CO₂}	V _{H₂O}	V _{N₂}	V _{O₂ex}	Tot.	r _{CO₂}	r _{H₂O}	r _{N₂}	r _{O₂ex}	Tot.	
Marsh gas	9.52	9.996	1	2	7.89	0.1	11	0.091	0.182	0.717	0.01	1	1390
Furnace gas	0.690	0.724	0.4065	0.0885	0.0789	0.0071	1.581	0.2571	0.0560	0.6824	0.0045	1	996

4. Study regarding the utilization of the furnace gas in a proportion of a hundred percent at burning

As it results from § 3.2. the burning temperature of the furnace gas with atmospheric air is 996 °C. The technological process of the homogenizing

annealing furnace necessitates an optimal temperature in the working space of 1200-1250 °C.

A method to increase the burning temperature is the utilization, at burning, of the oxygen enriched air.

The oxygen concentration in the enriched air (fig. 2) is calculated with the relation [5]:

$$x = \frac{\lambda \left\{ \frac{1}{2} [\text{CO}] + \frac{1}{2} [\text{H}_2] + 2[\text{CH}_4] - [\text{O}_2] \right\}}{\frac{\eta_{\text{pyr}} \cdot 0.98 \cdot H_i}{t_f \cdot c_{\text{ga}}} - \frac{1}{2} [\text{CO}] - [\text{CH}_4] - [\text{CO}_2] - [\text{H}_2\text{O}] - [\text{N}_2] - \frac{1}{2} [\text{H}_2] - [\text{O}_2]} \quad (9)$$

considering $t_f = t_m + 150 = 1250$ °C.

$c_{\text{ga}}^{1250^\circ\text{C}}$ is calculated as follows,

$$c_{\text{ga}}^{1250^\circ\text{C}} = \sum_{i=1}^4 r_i \cdot c_i$$

using the values:

$$c_{\text{CO}_2}^{1250^\circ\text{C}} = 2.2768$$

$$c_{\text{ga}}^{1250^\circ\text{C}} = 0.2571 \cdot 2.2768 + 0.056 \cdot 1.7898 + 0.6824 \cdot 1.4197 + 0.0045 \cdot 1.5055 = 1.661$$

$$x = \frac{1.05 \left(\frac{1}{2} \cdot 0.2245 + \frac{1}{2} \cdot 0.0615 + 2 \cdot 0.002 - 0.002 \right)}{\frac{0.7 \cdot 0.98 \cdot 3716}{1250 \cdot 1.661} - \frac{1}{2} \cdot 0.2245 - 0.002 - 0.18 - 0.023 - 0.507 - \frac{1}{2} \cdot 0.0615 - 0.002}$$

$$x = \frac{1.050 \cdot 0.145}{1.2277 - 0.857} = 0,41$$

For $t_f = t_m + 100 = 1200$ °C

$$c_{\text{CO}_2}^{1200^\circ\text{C}} = 2.2638$$

$$c_{\text{H}_2\text{Ovap}}^{1200^\circ\text{C}} = 1.7769$$

$$c_{\text{N}_2}^{1200^\circ\text{C}} = 1.4143$$

$$c_{\text{O}_2}^{1200^\circ\text{C}} = 1.5005$$

$$c_{\text{ga}}^{1200^\circ\text{C}} = 0.2571 \cdot 2.2638 + 0.056 \cdot 1.7769 + 0.6824 \cdot 1.4143 + 0.0045 \cdot 1.5005 = 1.652$$

and

$$x = \frac{1.05 \cdot 0.145}{1.285 - 0.857} = 0.355$$

For $t_m = 860$ °C, $t_f = 860 + 150 = 1010$ °C

$$x = \frac{1.05 \cdot 0.145}{1.558 - 0.857} = 0.217$$

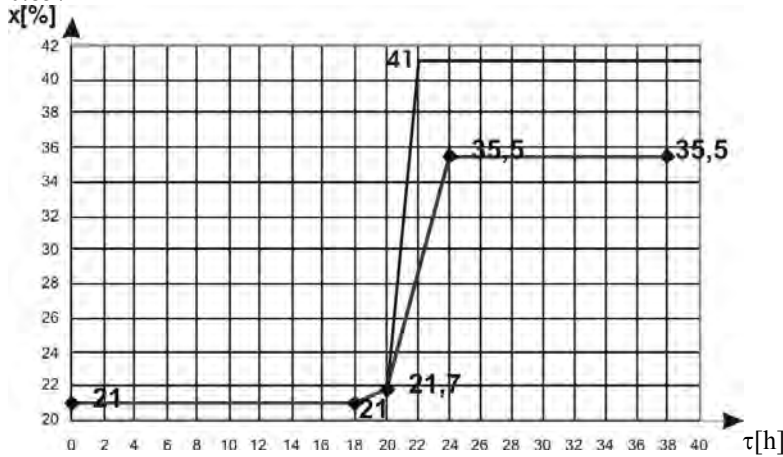


Fig.2. The oxygen concentration in the enriched air in the case of the furnace gas burning.

5. Economic efficiency

The costs are the following:

- CH₄ = 189.18 USD/1000 Nm³
- GF = 13.55 USD/1000 Nm³
- Oxygen = 86.15 USD/1000 Nm³
- Air = 7.68 USD/1000 Nm³

5.1. The case of the marsh gas burning

Using the values of the average flowrates of marsh gas taken from Fig. 3, the costs are calculated as following:

a) during the period $\tau_1=(0-10)$ h

$$D_{CH_4} = 135 \text{ Nm}^3/\text{h}$$

$$D_{air} = L_r D_{CH_4}$$

$$D_{air} = 9.996 \cdot 135 = 1349.5 \text{ Nm}^3/\text{h}$$

$$C_1 = 135 \cdot \frac{189.18}{1000} + 1349.5 \cdot \frac{7.68}{1000} = 35.9 \text{ USD/h}$$

b) during the period $\tau_2=(10-14)$ h

$$D_{CH_4} = 245 \text{ Nm}^3/\text{h}$$

$$D_{air} = 9.996 \cdot 245 = 2449 \text{ Nm}^3/\text{h}$$

$$C_2 = 245 \cdot \frac{189.18}{1000} + 2449 \cdot \frac{7.68}{1000} = 65.15 \text{ USD/h}$$

c) during the period $\tau_3=(14-24)$ h

$$D_{CH_4} = 330 \text{ Nm}^3/\text{h}$$

$$D_{air} = 9.996 \cdot 330 = 3298.7 \text{ Nm}^3/\text{h}$$

$$C_3 = 330 \cdot \frac{189.18}{1000} + 3298.7 \cdot \frac{7.68}{1000} = 87.76 \text{ USD/h}$$

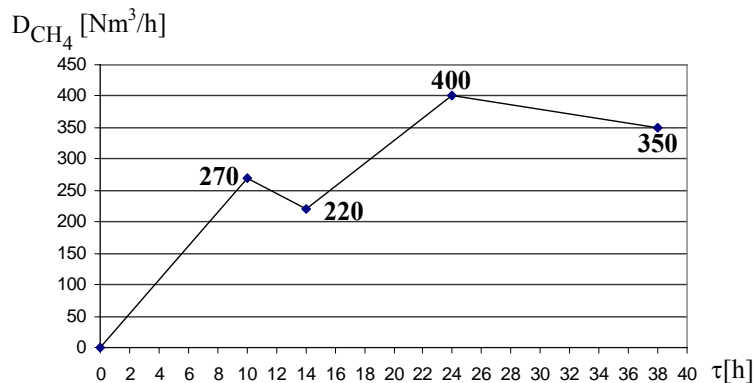


Fig.3. The variation of the marsh gas flowrate as a function of the treatment duration, in conformity with the exploitation data.

d) during the period $\tau_4=(24-38)$ h

$$D_{CH_4} = 375 \text{ Nm}^3/\text{h}$$

$$D_{air} = 9.996 \cdot 375 = 3748.5 \text{ Nm}^3/\text{h}$$

$$C_4 = 375 \cdot \frac{189.18}{1000} + 3748.5 \cdot \frac{7.68}{1000} = 99.72 \text{ USD/h}$$

The total cost is:

$$C_{\text{CH}_4} = \sum_{i=1}^4 C_i \tau_i$$

$$C_{\text{CH}_4} = 35.9 \cdot 10 + 65.15 \cdot 4 + 87.76 \cdot 10 + 99.72 \cdot 14 = 2893.28 \text{ USD/cycle}$$

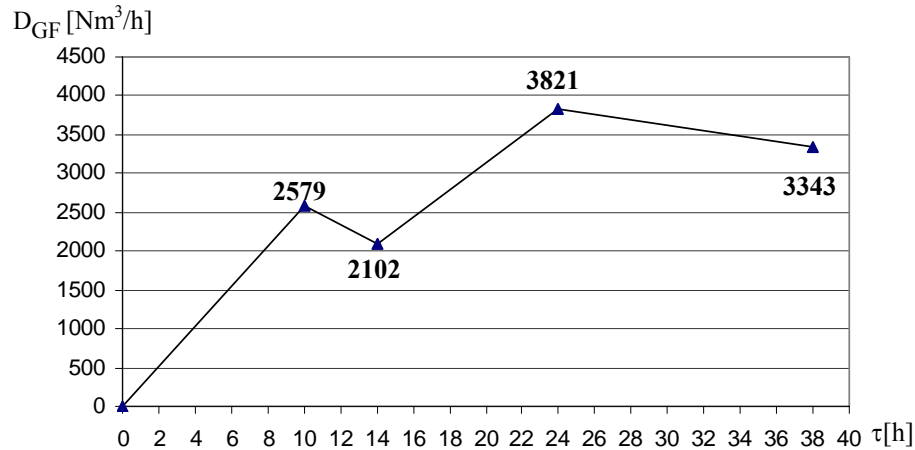


Fig.4. The variation of the furnace gas flowrate as a function of the treatment duration, determined from the calculations.

5.2. The case of the furnace gas burning

The values of the furnace gas flowrates in Fig. 4 have been calculated with the relation [6]:

$$D_{GF} = D_{\text{CH}_4} \frac{H_i^{\text{CH}_4}}{H_i^{\text{GF}}} \quad (10)$$

where:

D_{GF}, D_{CH_4} - the flowrates of the marsh gas and the furnace gas, respectively, in [Nm³/h];

$H_i^{\text{CH}_4}, H_i^{\text{GF}}$ - the calorific powers, [kJ/Nm³].

a) For $\tau_1=18$ h burning of the furnace gas with atmospheric air, the average flowrate of furnace gas, calculated in conformity with Fig. 4, is:

$$D_{GF} = 1780 \frac{\text{Nm}^3}{\text{h}}$$

and using the value

$$L_r = 0.724 \text{ Nm}^3 \text{ air/Nm}^3 \text{ GF}$$

the cost is:

$$C_1 = \left(1780 \frac{13.55}{1000} + 0.724 \cdot 1780 \frac{7.68}{1000} \right) 18 = 612.16 \text{ USD}$$

b) In the case of the furnace gas burning with oxygen enriched air:

- the oxygen enriched air L'_r , [Nm³/Nm³ fuel] is:

$$L'_r = \frac{100}{x} \text{ O}_2 \text{ min} \quad (11)$$

where x is oxygen concentration [%];

- the flowrate of the oxygen enriched air, D'_{air} , in [Nm³/Nm³ fuel] is [6]:

$$D'_{\text{air}} = D_{GF} L'_r \quad (12)$$

$$\begin{cases} D'_{\text{air}} = D_{\text{O}_2} + D_{\text{air}} \\ x = \frac{D_{\text{O}_2} + 0.21 \cdot D_{\text{air}}}{D_{\text{O}_2} + D_{\text{air}}} \end{cases} \quad (13)$$

It results:
$$\begin{cases} D_{\text{air}} = \frac{D'_{\text{air}} (1-x)}{0.79} \\ D_{\text{O}_2} = \left(\frac{x-0.21}{0.79} \right) D'_{\text{air}} \end{cases} \quad (14)$$

• For the range $\tau_2=(18-24)$ h, it is considering an average oxygen concentration in air $x=30\%$, and an average value for the furnace gas flowrate of

$$D_{GF} = 3305.3 \text{ Nm}^3 / \text{h}$$

taken from Fig. 4, resulting:

$$L'_r = \frac{100}{30} \cdot 0.145 = 0.483 \frac{\text{Nm}^3}{\text{Nm}^3 \text{ fuel}}$$

$$D'_{\text{air}} = 3305.3 \cdot 0.483 = 1596.5 \text{ Nm}^3 / \text{h}$$

$$D_{\text{air}} = \frac{1596.5(1-0.3)}{0.79} = 1414.6 \text{ Nm}^3 / \text{h}$$

$$D_{\text{O}_2} = \frac{0.3-0.21}{0.79} \cdot 1596.5 = 181.88 \frac{\text{Nm}^3}{\text{h}}$$

• For the range $\tau_3 = (24-38)\text{h}$, it is considering an average oxygen concentration in air $x = 37\%$, and an average value for the furnace gas flowrate of

$$D_{\text{GF}} = 3582 \text{ Nm}^3 / \text{h}$$

taken from Fig. 4, resulting:

$$L'_r = \frac{100}{37} 0.145 = 0.392 \text{ Nm}^3 / \text{Nm}^3 \text{ fuel}$$

$$D'_{\text{air}} = 3582 \cdot 0.392 = 1404.1 \text{ Nm}^3 / \text{h}$$

$$D_{\text{air}} = \frac{1404.1(1-0.37)}{0.79} = 1119.7 \text{ Nm}^3 / \text{h}$$

$$D_{\text{O}_2} = \frac{0.37-0.21}{0.79} 1119.7 = 226.7 \text{ Nm}^3 / \text{h}$$

$$C_3 = \left(3582 \frac{13.55}{1000} + 1119.7 \frac{7.68}{1000} + 226.7 \frac{86.15}{1000} \right) 14$$

$$C_3 = 1073.3 \text{ USD}$$

$$C_{\text{GF}} = C_1 + C_2 + C_3 = 2113.36 \text{ USD/cycle}$$

$\begin{aligned} \text{Economy} &= C_{\text{CH}_4} - C_{\text{GF}} = \\ &= 779.92 \text{ USD/cycle} \end{aligned}$
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6. Conclusion

In the case of burning of the furnace gas with atmospheric air, a burning temperature of 996°C has resulted. This burning temperature assures the heating of the pieces from the homogenizing annealing furnace in the first period of the 18 hours cycle. Forward, in order to increase the burning temperature

$$C_2 = \left(3305.3 \frac{13.55}{1000} + 1414.6 \frac{7.68}{1000} + 181.88 \frac{86.15}{1000} \right) 6$$

$$C_2 = 427.9 \text{ USD}$$

of the furnace gas, oxygen enriched air is used, reaching an average oxygen concentration in air of approx. 37%.

Calculating the costs for the two varieties, first burning of the furnace gas with atmospheric oxygen enriched air and second burning of the marsh gas with atmospheric air, respectively, a smaller cost has resulted in the case of burning the furnace gas than in the case of the marsh gas, the economy being of 779.92 USD/cycle.

The economy of marsh gas, achieved by the utilization of the furnace gas, presents a special interest because of the deficit of the marsh gas on the global market and its putting up the price.

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