# MANGANESE ALLOYS PRODUCTION: IMPACT OF CHEMICAL COMPOSITIONS OF RAW MATERIALS ON THE ENERGY AND MATERIALS BALANCE

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#### ABSTRACT

For the production of Iton of manganese alloys, the energy balance has been calculated for the charge mixtures. The theoretical energy consumption for some typical South African ores is mixted, and the individual contribution to the energy consumption is discussed. However, the with different chemical compositions, behave differently in the furnace and lead to products of the energy compositions. In this work, the MnO in the final slag has been assumed to 20%, the South African ores will give a high basicity slag. The mass and energy balances have shown the composition of three different ores will affect the metal analyses, the slag/metal ratio and the pasicity of the ore affect the slag/metal ratio. Of the ores investigated, the most than the parameter affecting the differences in energy consumption is the presence of carbonates ore. It has also been shown that there is a relationship between the CO/CO2 gas ratio, the carbon consumption and the electrical energy consumption.

KEYWORDS: Manganese ore, ferromanganese alloys, energy and materials balance.

#### INTRODUCTION

During the last two decades, a considerable improvement of thermodynamics data have prediction of equilibrium conditions as well as factors that affect the extend of reactions in the last last been studied. Hence, the effect of some components of the raw material namely alumina on have been investigated and have shown an influence on the viscosity which is ultimately pendent on the temperature [1], thus on the energy consumption and the slag composition.

wither, a comparison between oxidized ores, semi-oxidized ores and carbonated ore was a semi-oxidized and have shown that the reduction of the higher manganese oxides are all exothermic and to low power consumption [2]. This means that the ore containing more carbonates would more power than the one containing only oxides.

and the temperature [3]. The relationship between the degree of prereduction have shown energy consumption decreases when the degree of prereduction increases [4].

It is important to understand that the power consumption is dependent on the energy released in different zones in the furnace. Reactions below show how higher manganese oxides large are reduced to MnO and confirm that the higher the oxidation level, the lower the consumption [4 - 6]:

$$MnO_2 + \frac{1}{2} CO_{(g)} = \frac{1}{2} Mn_2O_3 + \frac{1}{2} CO_{2(g)}$$
  $\Delta H = -99 kJ$  (1)

$$\frac{1}{2} \text{ Mn}_2 \text{O}_3 + \frac{1}{6} \text{ CO}_{(g)} = \frac{1}{3} \text{ Mn}_3 \text{O}_4 + \frac{1}{6} \text{ CO}_{2 \text{ (g)}} \qquad \Delta H = -29 \text{ kJ}$$
 (2)

As it can easily be seen that reaction 1 to 3 are highly exothermic, thus releasing energy benefiting the process whilst reaction 5 is detrimental in terms of energy to the process. The power consumption is one key parameters to viably produce ferromanganese. From the above reference it can be seen that the chemical composition of the raw material and the extent of reaction 5, is a utmost importance when one would like to efficiently produce ferromanganese.

This paper investigates and compares theoretical mass and energy balances of different type of South African manganese ores. Manganese ores used were Nchwaneng 1 and Nchwaneng 2 and Gloria.

#### 2. ORE PROPERTIES

#### 2.1. Raw material

Manganese ore used in the calculation were Nchwaneng 1 and 2 and Gloria as presented in table 1 below with their chemical compositions. The basicity have been calculated according to the chemical composition based on the basicity formula  $Basicity = (CaO + MgO)/(SiO_2 + Al_2O_3)$ .

Table 1: Chemical analyses of manganese ores

	Element/ Compound	Unit	Ore type			
1			Ore 1	Ore 2	Ore 3	
	Mn	wt%	46	42.5	37	
	Fe	wt%	10	15.3	5.1	
ا ۾	SiO <sub>2</sub>	wt%	6.1	6.1	6.3	
ţį.	CaO	wt%	6.8	5.7	15.1	
composition	MgO	wt%	0.7	0.7	2.7	
	Al <sub>2</sub> O <sub>3</sub>	wt%	0.4	0.3	0.2	
	P	wt%	0.004	0.004	0.03	
Chemical	S	wt%	0.1	0.1	0.04	
	Na <sub>2</sub> O	wt%	0.1	0.1	-	
	K <sub>2</sub> O	wt%	0	0		
	B	ppb	628	628	380	
	Mn/Fe	wt%/wt%	4.6	2.8	7.3	
	Basicity		1.1	1.0	2.7	

In this paper, all the Mn has been considered as Mn<sub>2</sub>O<sub>3</sub> and all the Fe as Fe<sub>2</sub>O<sub>3</sub>. From the chemical analysis of the ores in table 1, it can be seen that Ore 3 has the lowest Mn content. The Mn

ontent has already been reported to have an impact on the power consumption through the change at slag/metal ratio [7]. This will be discussed in the theoretical results on the power consumption. It important to mention two things:

1) The CaO and MgO are basic oxides in Ore 1 and Ore 2, whilst in the Ore 3 they are present

a carbonates.

2) To obtain a basicity of 1.1 from the 2.7 as in table 1 above for Ore 3, theoretical amount of pure quartz was added to obtain a charge basicity of 1.1. To obtain this basicity 97.8 kg of SiO<sub>2</sub> per on of Ore 3 was added.

## Zone Limitations and assumptions in the furnace

The furnace has been subdivided into three zones with the following temperatures:

Zone 1: In the upper zone the raw materials enter the zone at 25°C and leave the zone at %C. The following reactions will occur:

$$Mn_2O_3 + CO = Mn_3O_4 + CO_2$$
 (I)

$$Fe_2O_3 + CO = 2FeO + CO_2$$
 (II)

Zone 2: In the middle Zone the materials will be heated from 700°C to 1100°C. In this zone it assumed that nothing melts and that the ores are still in solid state. The reactions are as follows:

$$Mn_3O_4 + CO = 3MnO + CO_2$$
 (IV)

$$FeO + CO = Fe + CO_2 (V)$$

$$C + CO_2 = 2CO (VI)$$

The degree of prereduction can be calculated according to Tangstad et al [4]. But in this reper, in the base case, it was assumed that 50% of CO<sub>2</sub> produced in this zone will react according reaction (VI).

Zone 3: In the Lower zone the material is heated from 1100°C to 1500°C. In the reduction the following reactions occur:

$$MnO_{(s)} + C = Mn_{(l)} + CO$$
  
 $SiO_2 + 2C = Si + 2CO$   
 $C = \underline{C}$ 

In addition in zone 3, the melting of the oxides and the metallic iron will occur. To calculate the enthalpy of the molten slag, the heat of solution would not be neglected given that the slag is a of oxides in which activities have been found to depart from ideality [8]. In this paper the and the assumption of Turkdogan (1983). According to Turkdogan, the heat of formation and heat capacity of many slag systems have very values and similar temperature dependence. For that reason, based on the similarity of the reactives and on similar heats of formation at 298 K, the enthalpy of molten slag at 1773K timated in J/g-atom. The value was obtained from Pyrometaalurgy course by Professor P.C When the state of of mol-atoms of the slag was calculated. The enthalpy of the molten slag was then the enthalpy of molten slag subtracted from the sum of enthalpy of each solid compound in the slag at 1773 K.

The metal and slag composition were calculated assuming that all iron from the ore was collected in the metal and that Si in metal was fixed to 0.06%. In addition, and maybe most important, as the basicity is in the area of one, the MnO content in the slag is assumed to be 20 %.

### 3. RESULTS AND DISCUSSIONS

#### 3.1. Mass Balance

The resulting slag and metal tapped is shown in table 2 and 3. The MnO in the slag was considered 20 % for the three ores, value taken from industrial data [9].

The Mn in the alloy was obtained by difference between the Mn form the ore and the one reported in the slag.

The carbon content in the alloy was calculated using the Li and Morris (1997) interactions parameters by assuming to be at saturation, therefore the activity of C would be 1 [10] whilst Fe was obtained by difference. The unreducible oxides reported all to the slag. SiO<sub>2</sub> was calculated knowing the amount in the Si content in the ore and assumed to be 0.06 % in the metal.

Table 2: Slag composition

Compounds	Ore 1	Ore 2	Ore 3
MnO	20	20	20
MgO	4	4.4	6.4
CaO	39	35.8	35.6
SiO <sub>2</sub>	34.6	37.9	37.6
$\frac{\text{SiO}_2}{\text{Al}_2\text{O}_3}$	2.3	1.9	0.5
Basicity (MgO+ CaO)/ (SiO <sub>2</sub> +Al <sub>2</sub> O <sub>3</sub> )	1.2	1	1.1

Table 3: Metal composition

Elements (Wt%)	Ore 1	Ore 2	Ore 3
Mn	75.1	67.03	79.1
Fe	17.3	25.6	13.3
	0.06	0.06	0.06
<u>SI</u>	7.5	7.3	7.6
Slag/Metal ratio	0.30	0.27	1.10
Mn/Fe ratio in alloy	4,3	2.6	5.9

From the data obtained in table 2 and 3, one can see the following results:

1) The Mn/Fe ratio calculated from the ore, is shown in table 1 Ore 2 had the lowest (2.8 ratio whilst the same ratio in the metal was of 2.6, still the lowest from the three alloys produced a indicated in table 3. The typical Mn/Fe ratio in the metal is 78/15, this means that ore (1,2) and Ore 3 should to be mixed with high Mn/Fe ore. This can explain the reason why the iron content is high and a low C content compared to Ore 1 and Ore 3. The more iron the lower the carbon content.

2) From table 3, it is shown that the slag/metal ratio with Gloria is the highest. This is mainly due to quartz that has been added to decrease the basicity, thus increasing the slag/metal ratio.

ides, which also will give a higher slag/metal ratio. If the Gloria ore was mixed with a high Mn, gh acid ore like Groote Eylandt or Comilog, this would give a much lower slag/metal ratio. The ics (1,2) slag/metal ratios were in the same low range around 0.3. Previously reported slag/metal for high basicity slag operation is higher, and this is due to several reasons. First, usually a mix various ores is used, and hence Ore (1,2) ore is mixed with Ore 3, a higher slag/metal ratio would expected. Also, if acid fluxes are used to lower the basicity a bit below 1, this would increase the cometal ratio. At last, metal loss in the slag would increase the apparent slag/metal ratio of 11.

#### Energy consumption

The energy consumptions per zone in the furnace for the different manganese ores used are cented in table 4 with a degree of prereduction of 50% whilst the change of energy consumption is the degree of prereduction is presented in table 5.

ble 4: Energy consumption per zone for different manganese ore

ZONE		Manganese ore	
(Energy consumption kwh/t metal)	Ore 1	Ore 2	Ore 3
Üpper	81	86	328
Middle	230	246	864
Lower	1111	992	986
Total Energy consumption/t metal	1422	1324	2177
otal Energy consumption (if 20% loss)/t metal	1778	1656	2723
Assumed Degree of prereduction	50	50	50

able 5: Energy consumption versus degree of prereduction for different manganese ore

Degree of prereduction	Energy	consumption kWh/ t me	tal metal
	Ore 1	Ore 2	Ore 3
20	1924	1819	3001
30	1875	1764	2908
50	1778	1656	2723

It is shown from table 4 that the energy consumption from upper zone to the lower zone traces for Ore 1 and 2, but for Ore 3 the increase in the second zone increased quite considerably middle zone. This can be attributed to the presence of carbonates. It is important to state the thowing:

The values in table 4 were obtained under conditions that the upper zone and middle zone not adiabatic, thus there was no thermal equilibrium. However, if one used the adiabatic modern in the first zone, the temperature of the gases leaving the top zone would be lower than sumed 700°C. In the mid zone there is a huge energy shortage. If one assumes that no electric is developed in this zone, this shortage could be covered by evaporation in the high relature zone and condensation in the mid zone of species like Mn and K gas. This could also the a narrow mid zone, and hence that the heat transfer would go through diffusion and lation from the high temperature zone. This has also been seen in previous calculations Tangstad 2007).

The energy consumption is low because the off-gas has been considered leaving the furnace at low temperature of 25°C whilst if the off-gas were considered leaving the furnace at about 500°C, the energy consumption would have been higher than the one obtained in table 4. In practice they leave at around 500°C and for each 100°C, 70kWh that must be added. This accounts for about 330 kWh extra. Also, if some metal is lost to the slag, the energy consumption will increase by the same percentage. Together this could be in the area of 500 kWh.

Table 5 shows the energy consumption per ton of metal produced against the degree of prereduction. It can be seen that the energy consumption decreased when the degree of prereduction increased. This can be attributed to the fact that the lower the degree of prereduction, the more CO<sub>3</sub>

available from the Boudouard reaction which is endothermic.

#### 4. CONCLUSIONS

The energy consumption per ton of metal produced was found to be dependent on the chemical composition of the ore. For three manganese ores, all Mn<sub>2</sub>O<sub>3</sub> ores, used in this paper and considered to be at the same degree of prereduction, the carbonates ore namely Ore 3 has a energy consumption higher than the Ores 1 and 2 which contain no carbonate. Also, the energy consumption was increasing with the decrease of the degree of prereduction for the three manganese ore. This is due to more CO<sub>2</sub> available for the Boudouard reaction when the degree of prereduction decreases. The Boudouard reaction is endothermic, this explains the increase in energy demand within the furnace. In addition to the above, the upper zone and the middle zone were not adiabatic as the temperatures were slightly higher than what they should be.

The overall energy consumption looked slightly lower than the values obtain practically in the plants. This could be due to the assumptions that the gases were leaving at around room temperature whilst in practice they leave at around 500°C and for each 100°C, there is 70kWh that must be added. If this is considered, the energy consumption can increase and might be close to industrial

values.

It was found that Ore 2 that had the lowest Mn/Fe ratio, had the lowest C content in the metal

produced compared to Ore 1 and Ore 3 which had a relatively high Mn/Fe ratio.

In addition, Ore 3 had the highest slag/metal ratio which explained the high amount of slap produced per ton of metal. This was attributed to the quartz added to modify the basicity to the same range as the Ores1 and 2 as well as a relatively low Mn content in the ore.

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