INCEPTION OF TRANSFORMATION OF HEMATITE TO MAGNETITE DURING MECHANICAL ACTIVATION: A THERMODYNAMICAL APPROACH*

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Abstract– In this research the thermodynamic of mechanochemical activation by high energy milling of hematite has been studied. It has been shown that transformation reaction due to ball milling of \( \alpha \)-Fe\(_2\)O\(_3\) has been started after 15 hours milling. As the sum of amorphization and dislocation energies is a suitable approximation of total stored energy, the contribution of these two energies has been identified. The contribution of amorphization and dislocation energies after 15 hours milling have been calculated at about 76 and 9 kJ/mol, respectively. Thus, the increase in molar enthalpy of milled hematite is about 85 kJ/mol. The total Gibbs free energy of mechanochemical transformation of hematite to magnetite after 15 hours milling has also been determined. The temperature for transformation of activated hematite to magnetite has been calculated at about 300 K. The bulk temperature of powder after 15 hours milling has been measured at about 325 K. Therefore, at this temperature hematite cannot be stable and will transform to magnetite during milling.

Keywords– Hematite, magnetite, stored energy, mechanical activation

1. INTRODUCTION

The iron oxides; hematite, maghemite and magnetite; are important electrical and magnetic materials. These oxides are conventionally prepared by oxidation–reduction process at elevated temperatures. Mechanochemical treatment is an alternating process to produce fine particles of these oxides.

During the last two decades, extensive studies have been made on the transformation reactions induced due to ball milling of hematite in various milling conditions. Kaczmarek and Ninham [1, 2] produced magnetite from hematite through ball milling in various milling atmospheres, i.e. vacuum, argon and air. The reverse transformation, magnetite to hematite, was explained to occur by ball milling in air [3]. Zdujic et al. [4, 5] explained the transformation process of \( \alpha \)-Fe\(_2\)O\(_3\) to Fe\(_3\)O\(_4\) and subsequently to FeO. They stated that transferring of sufficient energy to the particles was necessary to change the thermodynamical stability of \( \alpha \)-Fe\(_2\)O\(_3\) to Fe\(_3\)O\(_4\) and subsequently to FeO. Randrianantoandro et al. [6] showed that the \( \alpha \)-Fe\(_2\)O\(_3\) to \( \gamma \)-Fe\(_2\)O\(_3\) phase transformation could be induced by mechanical grinding within an ethanol medium. They showed that during high energy milling process, the oxide powders were exposed essentially to the compressive and shearing stresses. Mitov et al. [7] investigated the mechanochemical activation of magnetite and maghemite with close structures and different cation distributions. They demonstrated that the different rates of phase transformations could be explained by a phonon mechanism of energy dissipation for maghemite and a mixed phonon-electron mechanism for magnetite. Novikov [8], by means of X-ray diffraction and Mössbauer spectroscopy, demonstrated that the discrepancies in the available data on the cation distribution in cation-deficient compounds could be

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associated with the structural features of Fe$_3$-vO$_4$ compounds. Hofmann, Campbell and Kaczmarek [9-14] made comprehensive studies on the production of deficient magnetite from hematite by ball milling. Their comparisons by means of neutron diffraction patterns of as-milled samples at room temperature and 950 K combined with chemical analysis indicated that a significant fraction of the milled products had a disordered or amorphous form. They calculated the maximum fraction of γ-Fe$_2$O$_3$ at about 8%. The formation of γ-Fe$_2$O$_3$ was likely to be due to a shearing transformation involving movement of the oxygen included planes. They also concluded that the main transformation of α-Fe$_2$O$_3$ to Fe$_3$–vO$_4$ occurred by rupturing of the surface layers of α-Fe$_2$O$_3$. This rupture led to a release of oxygen and transformation of α-Fe$_2$O$_3$ to Fe$_3$–vO$_4$.

Although the mechanism of transformation of hematite to magnetite has been studied by various researchers, they have not showed the onset conditions of transformation and the way of thermodynamic functions changes. The present study attempts to gain a deeper insight into the thermodynamical features of this solid state reduction. In other words, the aim of this study is to clarify, for the first time, the reason of transformation of activated hematite to magnetite during ball milling at room temperature from a thermodynamic point of view.

2. EXPERIMENTAL PROCEDURE

Commercial grade α-Fe$_2$O$_3$ powder of 99% purity with particle size of 30-60 microns was used as the starting material. Mechanochemical treatment was performed in a planetary ball mill, Fritsch Pulverized 6. A hardened steel vial of 250 cm$^3$ volume filled with hardened steel balls of 20 mm diameter was used for milling. In all experiments, the ball to powder mass ratio was 25:1 with a powder mass of 20 g. All milling experiments were performed in 300 rpm under air atmosphere without any additives. It should be pointed out that the milling experiments were carried out under closed milling conditions, i.e. the vial was not opened during all the milling periods. Different milling times 2, 5, 10, 15, 20, 30, 45, 60, 75 and 90 h were considered in order to follow the solid state reactions. The morphology of the milled particles was studied by scanning electron microscopy (PHILIPS XL30) and their crystallographic structure was investigated by Philips PW-1730 X-ray diffractometer (XRD) using Cu Kα radiation.

3. RESULTS AND DISCUSSION

Figure 1 shows The XRD patterns of milled powder in various milling times. The patterns of the starting powder and the powders milled up to 10 hours show only the presence of α-Fe$_2$O$_3$ phase. It is seen that after 2 hours milling the peaks intensity decrease significantly due to the decrease of the size of hematite crystallites. The traces of magnetite can be observed after 15 hours milling. Although Fe$_3$O$_4$, is crystallographically isomorphous with γ-Fe$_2$O$_3$, having a lattice parameter (a $\approx$ 0.840 nm) slightly larger than that of γ-Fe$_2$O$_3$ (a $\approx$ 0.835 nm) [15], but according to the Hofmann et al. [9, 11] studies it can be concluded that a large amount of powder should be magnetite.

a) Energy contributions

Heegn [16] has reported that the following structural defects illustrate the energetic condition of mechanically activated solids:

(1) Dislocation concentration with its specific energy;
(2) Degree of amorphization with its energy;
(3) Surface area to the surroundings or to a second phase with its specific energy;

Pourghahramani [17, 18] has proved that the contribution of amorphization energy to the total energy changes is about 93–98.5%. Similar energy distributions are found for quartz, calcite, magnesite, kaolinite,
iron and periclase [17]. It is concluded that the sum of amorphization and dislocation energy is a suitable approximation of total energy.

Fig. 1. XRD patterns of α-Fe₂O₃ powder milled for various milling times.

1. Energy contribution of dislocations: The classical energy balance studies on the propagation of a small crack in an elastically isotropic solid establish a functional relationship between the critical tensile stress for crack propagation and flaw size [19]. Other studies propose that such ideal behavior does not apply to the majority of crack sensitive materials and should be modified [19]. Such modifications can be applied where crack tip plasticity is confined to a small plastic enclave in an elastic matrix, leading to a recast of Griffith’s ideas in terms of fracture mechanics concepts:

\[ K_{IC} = Y\sigma_c (a)^{1/2} \]  

where \( K_{IC} \) is the critical stress intensity factor for crack propagation, \( Y \) is a geometrical factor related to the shape of the crack, \( \sigma_c \) is critical tensile stress for crack propagation and \( a \) is the flaw size. This equation shows that by decreasing flaw size, \( a \), higher tensile stresses are needed to propagate a crack, i.e. \( \sigma_c \sim (a)^{1/2} \). Hence, initial fracture and size reduction of hematite occurs by propagation of the largest flaws at relatively low \( \sigma_c \). With increasing stored energy by milling, the particle and its flaw size decrease to micron or submicron in which \( \sigma_c \) becomes very large. This stored energy generates more dislocations rather than fracture of particles [20]. Figure 2 shows the SEM micrograph of 10 hours milled α-Fe₂O₃ in comparison with the initial powder. The effect of ball milling on the morphology and size distribution of powders has been previously investigated by Moshksar [21]. In this research, clusters of particles are formed during milling but decreasing the size of particles in Fig. 2b is in good agreement with the above discussions.
The increase in molar Gibbs free energy of milled hematite due to dislocations, $\Delta G_d$, is related to the changes in molar enthalpy $\Delta H_d$ and molar entropy $\Delta S_d$ associated with these dislocations:

$$\Delta G_d = \Delta H_d - T \Delta S_d$$  (2)

According to Cottrell hypothesis, the dislocations entropy is small. Thus, compared to $\Delta H_d$, $T \Delta S_d$ is negligible [22]. Hence, $\Delta G_d$ is almost equal to:

$$\Delta G_d \approx \Delta H_d = \rho_d m_v \left( \frac{b^2 \mu_s}{4\pi} \right) \ln \left( \frac{2 \rho_d^{1/2}}{b} \right)$$  (3)

where $b$ is Burger's vector of the dislocation, $\mu_s$ is the elastic shear modulus, $\rho_d$ is the dislocation density and $m_v$ is the molar volume of hematite [20]. This equation provides a quantitative estimation of the increase in molar free energy (stored energy) due to dislocations.

The direction of Burger's vector of hematite is similar to Corundum [17]. Therefore, the length of Burger's vector of hematite can be estimated as $5.03 \times 10^{-10}$ m [17]. $M_v = 3.047 \times 10^{-3}$ m$^3$mol$^{-1}$ is the molecular weight of hematite (0.15969 kg mol$^{-1}$) divided by its density 5240 (kg m$^{-3}$) [23]. The value of $\mu_s$ which is equal to 225 GPa, represents an average value for hematite polycrystalline aggregates [24]. The minimal dislocation density of polycrystalline hematite powders has been calculated using the following formula [2, 17, 25]

$$\rho_d = \frac{3}{d^2}$$  (4)

where $d$ is average crystallite size (m). Williamson–Hall approach can be used to estimate the crystallite size of the ball milled hematite powders [26]. The calculated mean crystallite size from XRD patterns which is corrected for instrumental broadening has been drawn versus milling time of Fe$_2$O$_3$ in Fig. 3.
The calculated dislocation density according to equation 4 up to 15 hours milling is shown in Fig. 4. The effect of dislocation density on the stored energy of hematite powders is shown in Fig. 5. It is obvious that the stored energy ($\Delta H_d$) increases with increasing the milling time.

![Fig. 4. Dislocation density of $\alpha$-Fe$_2$O$_3$ powder milled for various milling times.](image1)

![Fig. 5. Effect of dislocation density on stored energy, $\Delta H_d$, of hematite powder.](image2)

### 2. Energy contribution of amorphization:

The content of the amorphous phase has been determined by the equation proposed by Ohlberg and Stickler [27]. The degree of crystallinity is defined as:

$$X = \frac{U_o I_X}{I_o U_X} \times 100$$

where $U_o$ and $U_X$ refer to the background intensity of non-activated and activated hematite, respectively. $I_o$ and $I_X$ are also integral intensities of diffraction lines of non-activated and activated hematite.

In order to estimate the amount of amorphous phase after each milling time, the average relative intensity $\left(\frac{I_X}{I_o}$ and $\frac{U_X}{U_o}\right)$ of the four intensive peaks of milled hematite, ($20=24, 33, 41, 49.5$ degrees) has been calculated. The change of the intensities of the above mentioned peaks of the hematite after 15 hours milling has been given in Fig. 6.

Then, amorphization energy of milled hematite could be estimated as:

$$\Delta H_{am} = \left[1 - \left(\frac{U_o I_X}{I_o U_X}\right)\right] \Delta H_{am}$$
where $\Delta H_{am} = 90.99$ (kJ.mol$^{-1}$) is molar amorphization energy of hematite [28]. The energy contributions of dislocation density, $\Delta H_d$, amorphization, $\Delta H_{am}$, and overall change in enthalpy, $\Sigma \Delta H$, are shown in Fig. 7.

![Amorphization of 15 hours milled powder compared to initial powder.](image1)

*Fig. 6. Amorphization of 15 hours milled powder compared to initial powder.*

![Energy contributions of dislocation, amorphization, in total stored energy.](image2)

*Fig. 7. Energy contributions of dislocation, amorphization, in total stored energy.*

It is obvious that transferring sufficient energy to the hematite particles is necessary to start the transformation of hematite to magnetite. Figure 7 shows that the contribution of amorphization and dislocation energy up to 15 hours milling (starting of magnetite formation) are 76 and 9 kJ.mol$^{-1}$, respectively. Therefore, the increase in molar total enthalpy (required molar stored energy for transformation) of milled hematite prior to transformation to magnetite should be about 85 kJ/mol.

### b) Estimation of molar Gibbs free energy of milled hematite

The following reaction has been considered to estimate the molar Gibbs free energy of milled hematite:

$$\text{Fe}_2\text{O}_3 = 2/3\text{Fe}_3\text{O}_4 + 1/6\text{O}_2$$  \hspace{1cm} (7)

where the standard Gibbs free energy of this reaction is [4]:

$$\Delta G^o = G^o_2 - G^o_1 = 116596.67 - 67.8T \quad \text{J.mol}^{-1}$$  \hspace{1cm} (8)

and at the ambient pressure:
Inception of transformation of hematite to magnetite...

\[ \Delta G = \Delta G^0 + RT \ln P_{O_2}^{1/6} = 116596.67 - 67.8T + 8.314T \ln(0.2)^{1/6} = 116596.67 - 70.03T \text{ J.mol}^{-1} \] (9)

According to this equation, the equilibrium temperature is 1665 K. Therefore, magnetite cannot be stable below 1665 K.

During high energy ball milling up to 15 hours, the energy level of hematite, \( G^0 \), will increase to

\[ \Delta G = G^0 - (G^0 + \Delta G_d) \left(1 - \frac{U_o}{I_o} \frac{I}{U_X} \right) \left( \Delta G_{am} \right)_T + RT \ln P_{O_2}^{1/6} \] (10)

\[ \Delta G = \Delta G^0 + RT \ln P_{O_2}^{1/6} - \Delta G_d - \left[1 - \left( \frac{U_o}{I_o} \frac{I}{U_X} \right) \right] \left( \Delta G_{am} \right)_T \] (11)

while \( \Delta G_d \) and \( 1 - \left( \frac{U_o}{I_o} \frac{I}{U_X} \right) \left( \Delta G_{am} \right)_T \) are the increase in molar Gibbs free energy of milled particles due to dislocation and amorphization, respectively. \( \Delta G_d \) has been calculated before in Eq. (3). Due to the similarity of the amorphous phase and liquid phase, \( \Delta G_{am} \) could also be calculated by the following equation [20]:

\[ \left( \Delta G_{am} \right)_T = H^F \left( T_m - T \right) \] (12)

where \( H^F \) is the enthalpy of fusion at the melting point \( T_m \) and \( \left( \Delta G_{am} \right)_T \) is the estimated change in molar Gibbs free energy due to amorphization. By substituting 122.9 kJ.mol\(^{-1}\) and 1895 K as the enthalpy of fusion and melting point of hematite, \( \Delta G_{am} \) becomes [29]:

\[ \left( \Delta G_{am} \right)_T = 122900 - 64.85T \text{ J.mol}^{-1} \] (13)

According to equation (11), the total change in molar Gibbs free energy of milled hematite could be estimated as:

\[ \Delta G = 116596.67 - 67.8T + RT \ln P_{O_2}^{1/6} - \rho_d M V \left( b^2 \mu / 4\pi \right) \ln \left( 2 \rho_d^{1/2} / b \right) \] (14)

and finally:

\[ \Delta G = 4765.67 - 15.89T \text{ J.mol}^{-1} \] (15)

Figure 8 shows the \( \Delta G \) of non-activated and activated hematite versus temperature according to Eqs. (9) and (15). According to this figure, the equilibrium temperature for the 15 hours milled powder is 300K, above which the \( \alpha-\text{Fe}_2\text{O}_3 \) phase is not stable and the reaction proceeds toward the formation of \( \text{Fe}_3\text{O}_4 \).

The vial temperature (bulk temperature of the powder) after 15 hours milling has been measured by thermometer at about 325 K. It can be seen from Fig. 8 that at this temperature, \( \Delta G \) of reaction (7) in the activated condition is negative. Therefore it proceeds toward the formation of magnetite.
It should be mentioned that the calculation of $\Delta G$ according to equation 15 at temperatures above 600 K is meaningless and excluded from Fig. 8 because high temperature can recover the disorder structure by recrystallization of the amorphous phase and decrease dislocation density. The negative value of $\Delta G$ near 325 K shows that formation of magnetite has been started a few moments before the 15 hours milling.

**Fig. 8. Total Gibbs free energy of mechanochemical reaction before and after 15 hours milling**

### 4. CONCLUSION

The results of this investigation suggest that:

1) The increase in molar enthalpy (required molar stored energy for transformation) of milled hematite prior to transformation to magnetite is about 85 kJ.mol$^{-1}$.

2) The contribution of amorphization and dislocation energy in total stored energy up to 15 hours milling (starting of magnetite formation) is 76 and 9 kJ.mol$^{-1}$ respectively.

3) According to the calculated total Gibbs free energy function of mechanochemical reaction after 15 hours milling the transformation temperature is about 300 K.

4) At the bulk temperature of the powder, the total Gibbs free energy function of 15 hours milled powder is negative and reaction proceeds toward the formation of magnetite.

5) The negative value of $\Delta G$ near 325 K shows that formation of magnetite has started a few moments before the 15 hours milling.

### REFERENCES


