MICROSTRUCTURE AND PROPERTIES OF COLLOIDAL SILICA BONDED MAGNESITE CASTABLE REFRACTORIES

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Abstract: Efforts have been carried out in order to use microsilica to develop a forsterite bond rather than other types of binders in the basic refractory castables. According to the higher drying rate and sinterability of colloidal silica, it has been proposed in the recent years. In the present work, effects of replacement of microsilica by colloidal silica evolution of forsterite bond have been studied in magnesia based refractory castables. In this way, physical properties of prepared samples with different amount of colloidal silica versus temperature were investigated. In addition, phase variation and microstructural evolution of sintered specimens at 1000, 1200 and 1400 °C were studied by X-ray diffraction (XRD) and scanning electron microscope (SEM) respectively. Results showed, due to Reaction of magnesia with microsilica and colloidal silica, magnesium hydrate and magnesium silicate hydrate formed in the dried samples strengthening the texture of the samples while forsterite formed from about 1000 °C and gradually increased with temperature rise. Also, better forsterite formation would be appeared by increasing the colloidal silica content. Further investigation carried out on the type of silica addition on properties of the castable refractory samples. It was found that the presence of micro silica and colloidal silica simultaneously (MS3C3 sample) at 1400 °C, caused modifying mechanical strength in compare with sample with only micro silica (MS sample).

Keywords: Magnesite refractory castables, Microsilica, Colloidal silica

1. INTRODUCTION

Refractory castables are combinations of refractory aggregates, matrix component, bonding agent, and admixtures [1]. Easy production of refractory castables along with cheap installation and similar performance to shaped products are the reasons for the growing market of monolithic refractories, at the expense of traditional bricks. Calcium aluminate cement is the most used hydraulic binder in refractory castables compositions. Nevertheless, CAC amounts higher than 2-3 wt% in Al2O3-CaO-SiO2 systems may induce liquid formation at lower temperatures [2]. In addition to the above disadvantageous property of high cement bonded conventional castable, it requires a high water demand on mixing, resulting low density and high porosity products. These drawbacks led to the development of low cement castable (LCC) and no cement castables (NCC) [3].

Micro silica, which is a by-product of ferrosilicon manufacture, when present in the bond system of refractory castable batches reduces the amount of mixing water required and facilitates flow and ease of placement. Magnesia–silica bond systems have been developed in magnesia-based, cement-free castables [4] forming forsterite in the bond on firing. The HDMOR reached a maximum at 12000C but deteriorated at higher temperatures due to the effect of raw material impurities [5].

Recently different kinds of sols which form coagulation bonding have been tried as binder in low, ultralow and no cement bonded refractory castable compositions. Colloidal silica sols are stable water-based suspensions, containing up to 50 wt% of nanometric spherical amorphous silica particles (8-15 nm diameter). When combined with other solid particles, colloidal silica can be linked together in branched chains, in a process known as gelation [3], which can be induced by water removal. During the drying step, the hydroxyl groups (Si-OH) on the surface of the particles generate siloxane bonds (Si-O-Si), which results in a three-dimensional network (figure 1) [5]. Gelation can also be induced by pH changes and a salt or water miscible organic solvent addition, named gelling agents. With the proper selection of these agents or the pH variation, the colloid is gelled around the solid particles providing mechanical strength to the
system after drying. The micro silica-magnesia-
colloidal silica system presents a pH value
around 4–6, which is closer to the pH where the
colloidal silica gelling rate is maximum (around 
pH 5). In this context, colloidal silica has been 
pointed out as an alternative for a calcium-free 
binder agent for refractory castables [6].

In the present research the influences of 
colloidal silica addition on casting process as 
well as properties are investigated.

2. EXPERIMENTAL PROCEDURE

2.1. Materials

Chemical analyses of the raw materials used in 
this study are listed in Table 1 and the recipes of 
the studied castables are provided in Table 2.

2.2. Castable Preparation and Testing

For each sample, a 3kg batch mixture using the 
formulation in Table 2 was prepared, by dry 
mixing for 2 minutes at slow speed, using a 
Hobart mixer with 5 liter capacity mixing bowl. 
Then wet mixing was prepared by addition of 
water, colloidal silica after 30 second and further 
3 minutes mixing at medium speed. Test samples 
were prepared by casting the mixture into the 
stainless steel moulds, having the dimensions of 
160×40×40 mm, vibrated at 50HZ for 1 minute. 
The castables were cured at 24°C and a relative 
humidity of 95% for 24 hours. After 24 hours 
curing time, the samples were allowed to be dried 
at the temperature of 110°C, and then fired at 
1000°C, 1200°C and 1400°C for 3 hours. The 
cold strength (CCS, CMOR) was measured 
according to the ASTM C109. Permanent linear 
changes (PLC) of the samples fired at 1000°C, 
1200°C and 1400°C for 3 hours were measured. 
Bulk density and apparent porosity were all

![Schematic representation of colloidal silica consolidation through the Gelling mechanism](image_url)

**Table 1. Chemical analysis of the raw materials**

<table>
<thead>
<tr>
<th></th>
<th>Magnesia (wt %)</th>
<th>Micro silica (wt %)</th>
<th>Colloidal silica (wt %)</th>
</tr>
</thead>
<tbody>
<tr>
<td>MgO</td>
<td>97 min</td>
<td>0.1</td>
<td>0</td>
</tr>
<tr>
<td>SiO₂</td>
<td>0.6 max</td>
<td>97.5</td>
<td>40.2</td>
</tr>
<tr>
<td>CaO</td>
<td>1.5 max</td>
<td>0.2</td>
<td>0</td>
</tr>
<tr>
<td>Fe₂O₃</td>
<td>1 max</td>
<td>0.4</td>
<td>0</td>
</tr>
<tr>
<td>Al₂O₃</td>
<td>1 max</td>
<td>0.4</td>
<td>0</td>
</tr>
<tr>
<td>K₂O</td>
<td>0</td>
<td>0.2</td>
<td>0</td>
</tr>
<tr>
<td>Na₂O</td>
<td>0</td>
<td>0.1</td>
<td>0</td>
</tr>
<tr>
<td>L.O.I</td>
<td>0.2 max</td>
<td>0.6</td>
<td>59.8</td>
</tr>
</tbody>
</table>

**Table 2. Composition of castables**

<table>
<thead>
<tr>
<th>Castable Composition (wt %)</th>
<th>Grade</th>
<th>MC</th>
<th>MS</th>
<th>C₃</th>
</tr>
</thead>
<tbody>
<tr>
<td>Magnesia</td>
<td>Iran</td>
<td>94</td>
<td>94</td>
<td>94</td>
</tr>
<tr>
<td>Micro silica</td>
<td>Ferroalloy of Iran</td>
<td>0</td>
<td>3</td>
<td>6</td>
</tr>
<tr>
<td>Colloidal silica</td>
<td>Beechems</td>
<td>6</td>
<td>3</td>
<td>0</td>
</tr>
<tr>
<td>Additive</td>
<td>Castament</td>
<td>0.3</td>
<td>0.3</td>
<td>0.2</td>
</tr>
<tr>
<td>Water</td>
<td></td>
<td>8.5</td>
<td>6.65</td>
<td>5.25</td>
</tr>
</tbody>
</table>
determined by standard methods. Phase analyses were performed using a JEOL JDX 8030 model X-ray diffractometer. For microstructural observation, scanning electron microscopy (SEM) was used.

3. RESULTS AND DISCUSSION

3.1. Apparent Porosity and Bulk Density

Apparent porosity and bulk density of the samples as a function of temperature are shown in figures 2, 3.

According to the evaporation of bonding water with increasing temperature till 1000 °C, the apparent porosity of all samples increased; moreover, the bulk density of them decreased. After increasing temperature till 1200 °C, in all the samples except MS3C3, the apparent porosity and bulk density were stable. Between 1200 °C to 1400 °C, as a result of sintering, the apparent porosity of all the samples decreased; in contrast, the bulk density increased. Figures 2 and 3 show that MS sample which contained 6 wt.% micro silica, had minimum apparent porosity and maximum bulk density in all temperatures. Apparent porosity and bulk density are directly related to the water content [2]. For the sintered magnesia–colloidal silica system the reaction was so fast that it was not possible to prepare a homogenous suspension. During mixing, silica particles immediately gelled around the magnesia ones, forming an impermeable external layer, which hindered the powder dispersion. Therefore we forced to add extra water in order to achieve suitable rheological condition. In this way, the increase in apparent porosity and the decrease in bulk density observed in which samples contain colloidal silica in compare with samples contain micro silica. Furthermore powder density of micro silica is more than the colloidal silica [3] which is another reason for increasing the bulk density of MS sample in compare with MC one.

3.2. Cold Mechanical Strength

In the next step variations of bending and cold crushing strength with firing temperature for different formulations are shown in figure 4 and 5.

Water content, particle size distribution and sintering condition of the castable samples are the most important factor which influences mechanical strength [7]. As figure 4 and 5 show, mechanical strength of MS sample is higher than other samples in 110 °C. Formation of MgO,SiO$_2$-H$_2$O phase in this temperature can be the reason of this matter [5]. According to figures 4 and 5 increasing temperature till 1200 °C, the
absence of bonding water, caused decreasing mechanical strength in all samples. After that, by increasing temperature till 1400 °C formation of forsterite bonds during sintering process resulted increasing mechanical strength in all samples. Higher mechanical strength for MC sample was expected than MS but because of fast gelation of colloidal silica and addition of extra water, mechanical strength in MS became higher than MC. The other point is that in 14000C, the mechanical strenght of MS$_3$C$_3$ sample was higher than MS sample. In fact using micro and colloidal silica simultaneously caused improving mechanical strength of the fired samples because micro silica improved particle size distribution consequently, flow of the castables and higher reactivity of colloidal silica caused to form higher amount of forsterite upon sintering.

3. 3. Permanent Linear Change (PLC)

Permanent linear changes of the samples in 1000,1200,1400 °C are shown in figure 6.

Permanent linear change (PLC) depends on sintering temperature, reaction between aggregates and bonding agents. Principally shrinkage is caused by sintering and also glass phase formation; on the other hand; expansion is caused by phase transition in the sample’s matrix.

As figure 6 shows, MS$_3$C$_3$ sample had lower Permanent linear change (PLC) than the other ones. Also by increasing forormation of forsterite during sintering process, more shrinkage resulted increasing the PLC.

3. 4. Phase Analyses and Microstructural Observations

XRD patterns showed that the presence of micro silica could inhibit the hydration of magnesia; furthermore it caused formation of magnesium silicate hydrate which bonded the aggregates together [8].

Forsterite is a crystalline magnesium silicate with chemical formula Mg$_2$SiO$_4$, which shows good refractoriness due to high melting point (1890 °C), relatively low thermal expansion (6×10$^{-7}$K$^{-1}$ between 20 and 500 °C), good chemical stability and excellent insulation properties (K=1.4 J/m.sec.0K at 1300 °C) even at high temperatures [8].

Increasing the silica content in magnesite refractories can change the CaO/SiO$_2$ molar ratio and phase formation. But in this study, in order to forsterite formation, specific content of silica was added. According to a) low content of impurity and b) short accessibility of impurities to SiO$_2$ in grains, formation of low melt phases would be limited. Furthermore this refractory has been considered for application in less than 16000C range. So the contents of low melt phases are negligible. On the other hand this small amount of low melting phases makes a little decrease in the service temperature of the refractories.

XRD pattern of MS$_3$C$_3$ sample after firing at 1000, 1200, 1400 °C are shown in figure 7. In addition to periclase as the main phase, forsterite and quartz phases formed at 1000, 1200 °C. Quartz phase eliminated at 1400 °C because of its reaction with magnesia in this temperature. Furthermore it can be seen, by increasing the temperature, forsterite phase formation was
developed and the intensity of forsterite peak was increased.

XRD pattern of MS$_3$C$_3$ and MC samples after firing at 1400°C are shown in figure 8. Result showed the bonding formation of forsterite started to form in the sample fired at 1200 °C which increased by further firing at 1400 °C. The intensity of forsterite’s peeks in MC sample is higher than MS$_3$C$_3$ in the sample. It may be needed to explain that many researchers introduce an optimum amount for colloidal silica addition as higher amount of colloidal silica deteriorate flow behavior and workability of the castable samples leading to strength decreased [5, 6, 8].

SEM micrographs of MS$_3$C$_3$ sample after firing at 1400°C with different magnifications are shown in figure 9. According to EDS analyses, high grey matrix indicated magnesia phase (point A), and light grey matrix indicated the forsterite phase (point B) which resulted by reaction between particles.

SEM micrographs of MC sample after firing at 1400°C with different magnifications are shown in figure 10. According to EDS analyses, high grey matrix indicated magnesia phase (point A), and light grey matrix indicated the forsterite phase (point B) which resulted by reaction between particles.

Fig. 7. XRD pattern of MS$_3$C$_3$ sample after firing at 1000, 1200, 1400°C

Fig. 8. XRD pattern of MS$_3$C$_3$ and MC samples after firing at 1400 °C
CONCLUSIONS

- The addition of colloidal silica individually caused to use extra water; therefore; mechanical strength decreased in all samples at 110°C, 1000°C, and 1200°C.
- The presence of micro silica and colloidal silica simultaneously in MS$_3$C$_3$ sample at 1400°C, caused better formation of forsterite phase and modifying mechanical strength in compare with MS sample. So there is an optimum amount of colloidal silica (3% wt) providing better workability of the castable as well as forsterite
formation upon sintering.
• In all the samples, the formation forsterite phase started approximately at 1000°C and gradually increased by increasing the temperature.
• X-ray analyses results showed that the intensity of forsterite’s peaks in MC sample are higher than those in MS$_3$C$_3$ sample at 1400°C.

REFERENCES