Preparation and Hydration Mechanism of Mine Cemented Paste Backfill Material for Secondary Smelting Water-granulated Nickel Slag

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ABSTRACT

Using secondary smelting water-granulated nickel slag (SWNS) after iron extraction as the main raw material, made composite activator (flu gas desulphurization gypsum (FGDG) and carbide slag (CS) are the main activators, Na₂SO₄ and cement clinker (CC) are the auxiliary activators) with cementing agents (CA), and then made into mine cemented paste backfill (CPB) material with iron ore tailings (IOT). The factors affecting the mechanical properties such as the composition of raw materials and the amount of activator were studied by means of mechanical property test, X-ray diffraction (XRD) and scanning electron microscope (SEM), the ratio of CPB material was optimized, and the hydration mechanism of CPB material was analyzed. The results show that when the mass fraction of SWNS, FGDG, CS, Na₂SO₄ and CC in CA is 85%, 5%, 5%, 3% and 2%. The 28d flexural strength and compressive strength of CPB material of SWNS reach 1.99 MPa and 3.38 MPa, which meets the strength requirements of mine CPB material. The results of X-ray diffraction (XRD), scanning electron microscope (SEM) and hydration mechanism analysis showed that both vitreous and crystalline substances in, CA of SWNS could be hydrated, and the main hydration products were ettringite (Aft) and Ca₃(PO₄)₂, Mg₅(OH)₆ containing hydrated aluminosilicate (C-M-S-H) gel.

Keywords: Secondary Smelting Water-Granulated Nickel Slag, Cemented Paste Backfill Material, Cementing Agents, Composite Activator, Ettringite

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1. INTRODUCTION

Cemented paste backfill (CPB) method refers to the backfilling mining method in which cementing agents (CA), aggregate and water are mixed in a certain proportion and stirred evenly, and then the filling slurry or paste is transported to the mined out area by gravity self-flow or pumping through the conveying pipeline [1]. The main function of CPB body is to meet the operational requirements of mining equipment and to provide artificial bottom pillars or false roofs, so the strength requirements vary according to the backfilling technology and the function of the backfilling body, which is generally 1~5 MPa [2]. High quality CPB mining can increase resource recovery rate by more than 30% compared with non-CPB mining. CPB is increasingly widely used as an effective mining technology for deep well, complex and special conditions ore deposits [3, 4]. CPB is a new backfilling technique for mines, which allows the tailings harmful to the environment to return to the underground roadway or stope, so as to maximize the safety, efficiency and productivity of the process/operation, it is of great significance to solve the safety hazards and environmental problems caused by a large number of tailings stockpiling [5-8].

CPB costs generally account for 25%~40% of the total mining costs, and CA costs account for 70%~80% of the backfilling costs [9, 10]. CPB materials are mainly composed of aggregate and CA, ordinary Portland cement (OPC) is used as cementing agent in most mines [11-14]. Cement, as the CA that has been used for the longest time and has been widely used in cementing backfilling, has the characteristics of stable strength development and reliable property, and has been widely used in mine filling. The most commonly used type of cement is OPC, but cement as a backfilling CA also has the following problems: (1) In the process of CC production, a large amount of coal, electricity, limestone, clay and other resources are consumed, and 0.95 t CO₂ will be emitted for 1 t of CC production, resulting in resource waste and environmental pollution. (2) When sulfide is present in tailings, the oxidized acid and sulfate of the sulfide will reduce the stability of the cement CPB material [15]. (3) The cost of cement CA accounts for 60%~80% of the backfilling cost. Therefore, many mining enterprises and researchers began to look for industrial byproducts with low cost, excellent performance, environmental protection and cementitious properties, such as fly ash [16-19], nonferrous metallurgical slag [20], steel slag [21], GBFS [22-26] and other additives [27-29], the solid wastes with potential cementitious activity are partly or completely substituted for cement to prepare CA, it can not only improve the comprehensive utilization rate of industrial solid waste, but also protect the environment and reduce the cost of CA, which has broad development prospect.

Non-ferrous metallurgical slag is the solid waste after the smelting of non-ferrous metals, mainly including the slag formed in the process of fire smelting and the residue
discharged in the process of wet smelting. After mechanical grinding and chemical excitation, it has a good potential activity, and can be used to prepare CA for CPB material, and improve the utilization rate [20, 30]. Nickel slag contains 30%~40% of iron in the form of silicates, and further reduction smelting of this nickel slag can extract most of the valuable metal at one time. However, a large amount of SWNS is produced in the process of iron extraction by secondary smelting, after water-granulated, its appearance is close to that GBFS, but its chemical composition and material structure are quite different from GBFS, and SWNS has certain potential hydraulic properties [31-34]. If enough strength can be generated by the activator, it can be used in backfilling. In this study, FGDG, CS, Na₂SO₄ and CC were used as activators. Through the factors affecting the mechanical properties, such as the composition of raw materials and the amount of activator, optimized the ratio of CPB material, and on this basis, the hydration process and mechanism of CPB material were analyzed by means of XRD, SEM and other testing methods. It provides a basis for the application of SWNS in mine CPB material.

2. MATERIALS AND METHODS

2.1 Experimental materials

The main materials used in the test are SWNS, CC, CS, FGDG, Na₂SO₄, IOT, its chemical composition is shown in Table 1, and the mineral composition of raw materials is shown in Figure 1.

<table>
<thead>
<tr>
<th>Materials</th>
<th>SiO₂</th>
<th>Al₂O₃</th>
<th>Fe₂O₃</th>
<th>FeO</th>
<th>MgO</th>
<th>CaO</th>
<th>Na₂O</th>
<th>K₂O</th>
<th>SO₂</th>
<th>LOI</th>
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<tr>
<td>SWNS</td>
<td>36.72</td>
<td>6.97</td>
<td>0.27</td>
<td>3.01</td>
<td>28.78</td>
<td>22.69</td>
<td>0.16</td>
<td>0.08</td>
<td>0.87</td>
<td>0.13</td>
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<td>CC</td>
<td>22.50</td>
<td>4.86</td>
<td>3.43</td>
<td>—</td>
<td>0.83</td>
<td>66.30</td>
<td>—</td>
<td>0.08</td>
<td>0.24</td>
<td>0.96</td>
</tr>
<tr>
<td>CS</td>
<td>3.45</td>
<td>1.94</td>
<td>0.078</td>
<td>0.29</td>
<td>—</td>
<td>59.54</td>
<td>0.28</td>
<td>0.001</td>
<td>3.55</td>
<td>30.11</td>
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<tr>
<td>FGDG</td>
<td>3.16</td>
<td>1.35</td>
<td>0.47</td>
<td>0.09</td>
<td>7.49</td>
<td>33.38</td>
<td>0.13</td>
<td>0.18</td>
<td>36.56</td>
<td>8.28</td>
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<tr>
<td>Na₂SO₄</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td>—</td>
<td>43.23</td>
<td>—</td>
<td>0.008</td>
<td>44.62</td>
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<tr>
<td>IOT</td>
<td>68.96</td>
<td>7.68</td>
<td>2.32</td>
<td>4.47</td>
<td>3.64</td>
<td>4.35</td>
<td>1.41</td>
<td>1.85</td>
<td>0.024</td>
<td>2.49</td>
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Table 2. Particle size distribution of IOT

<table>
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<tr>
<th>Sieve Size/ mm</th>
<th>+1.25</th>
<th>-1.25</th>
<th>+0.63</th>
<th>-0.63</th>
<th>+0.32</th>
<th>-0.32</th>
<th>+0.16</th>
<th>-0.16</th>
<th>+0.08</th>
<th>-0.08</th>
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<tbody>
<tr>
<td>Meter sieve/%</td>
<td>1.5</td>
<td>5.8</td>
<td>6.9</td>
<td>36.8</td>
<td>41.2</td>
<td>7.8</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Cumulative sieve/%</td>
<td>1.5</td>
<td>7.3</td>
<td>14.2</td>
<td>51.0</td>
<td>92.2</td>
<td>100</td>
<td></td>
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</table>

Figure 1. XRD spectrum of raw materials (a)-SWNS, (b)-CS, (c)-FGDG, (d)-IOT
SWNS. The SWNS produced by molten reduction of iron extraction from nickel smelting slag was selected. The chemical composition is shown in Table 1, and the XRD analysis results are shown in Figure 1(a). It can be seen from Figure 1(a) that there are more crystals in SWNS, including monticellite, kamaishilitie, grossularite, augite and so on. Fig. 2 shows the SEM microstructure of SWNS corroded, cleaned, dried and sprayed with 1% HF solution. As the hydrofluoric acid solution corrodes the glass phase on the surface, a large amount of crystalline materials is exposed on the glass phase. SWNS contains a large number of crystalline materials. EDS semi-quantitative analysis of crystalline materials and XRD analysis show that the crystalline materials of SWNS were monticellite, kamaishilitie, grossularite and augite (Fig. 2(a)). In Figure 2(b), the labeled region A is a large amount of glassy materials, and the labeled region A is monticellite. This is because the generation of SWNS after the extraction of iron from nickel slag is similar to the generation of GBFS. By water-granulated and rapid cooling of molten nickel slag, glassy substances and crystalline substances with low hydration potential are generated, which have certain potential hydraulic activity. Due to the different content and composition of glassy materials, the strength of SWNS as a cementitious material is very different. Similar to GBFS, SWNS is also composed of Al2O3 and SiO2 to form a threedimensional network space, and alkali metals and alkali earth metals are embedded in the gaps of the grid. Under the action of the activator, the network structure of [AlO4]4- tetrahedron and [SiO4]4- tetrahedron was destroyed. Mg2+, Ca2+, 6-coordination Al3+ and alkali metals were involved in the reaction of SWNS, forming hydrates and showing gelation [31-34]. However, there are great differences between SWNS and GBFS in this study, as shown in the following aspects: the chemical components of SWNS and GBFS differ greatly, and the formation of the final phase will follow the phase law of certain components, SWNS have more crystals. The formation of the hydration products of SWNS CA not only needs to stimulate the hydration reactivity of the glass phase, but also needs to stimulate the hydration reactivity of the crystalline phase to a certain extent.

(2) CC. CC is the OPC clinker produced by cement plant. The chemical composition is shown in Table 1, the main mineral components are C2S (24.6%), C3S (55.7%), C3A (8.8%) and C2AF (10.9%).

(3) CS. Its chemical constituents are listed in Table 1. The CaO content of CS is 60.16%, and its mineral composition is mainly Ca(OH)2 and CaCO3 (see Figure 1(b)), CS, and the residue of 0.08 mm square-hole sieve is 12.8%.

(4) FGDG. FGDG is the wet FGDG from the thermal power plant, and its main component is CaSO4·2H2O. Fig. 1(c). According to Table 1, the burning loss of FGDG was 8.28%, the contents of CaO and SO3 in FGDG were 33.38% and 45.70%, respectively. According to the calculation that all SO3 came from CaSO4, the mass fraction of CaSO4 in FGDG was 78.26%, and the SSA was 210 m2·kg-1.

(5) IOT. Its chemical constituents are listed in Table 1. The particle size of tailings with a mass fraction of about 78% is between 0.08 mm and 0.32 mm; those with a particle size greater than 0.32 mm account for 14.2%; and those with a particle size of about 7.8% are less than 0.08 mm (see Table 2). The content of SiO2 in the tailings reaches 68.96% (mass fraction, the same below), which belongs to high-silicon type IOT. The major minerals of IOT included: quartz, hornblende, biotite, plagioclase and a few amount of chlorite, calcite and magnetite (see Figure 1(d)).

2.2 Methods

The experimental process of SWNS preparing CPB material is shown in Figure 3. Firstly, SWNS, CS and FGDG were dried in oven at 105 °C for 24 h to make moisture content less than 1%, respectively. Then, crushed them using SM 500 mm × 500 mm laboratory ball mill at the speed of 48 r·min-1 to yield SSA of 550 m2·kg-1, 300 m2·kg-1 and 330 m2·kg-1. The grinding media consists of several steel balls and a steel forging. The loading mass of the grinding media should be 100 kg. The steel balls add up to 40 kg, in which the Φ 70 mm ball takes up 19.7%, the Φ 60 mm ball takes up 33.1%, the Φ 50 mm ball takes up 29.6%, and Φ 40 mm ball takes up 17.6%; the Φ 25 mm ×30 mm steel forging weighs 40 kg. CC was first crushed to 1~3 mm with a jaw crusher, and then the ball-milling to SSA 400 m2·kg-1. The prepared raw materials are mixed evenly in proportion to form CA, then, in accordance with binder sand ratio (CA/IOT) of 1:4, IOT was added as aggregate and mixed evenly with water in the binder sand mixer to make slurry concentration of 80% mortar. After that, 4 cm × 4 cm × 16 cm cement binder sand triple test model was injected, after 24 h of shaking molding on the cement and sand vibrating table, the mold was taken off for curing, and the mold was put into a standard curing box with a temperature of (20 ± 1) °C and a humidity of more than 90% for curing, because the early strength is not high, after 7 d, the mold is damaged and then put into the curing box for curing. Finally, the mechanical properties of the test block at different ages are determined. An optimized ratio was obtained by measuring the compressive strength of the samples, and the optimized ratio of CA was analyzed by XRD and SEM.

The SSA is measured using dynamic specific surface area analyzer (SSA-3200). The compressive strength test of samples was based on the Chinese National Standard GB/T 17671-1999 Method of testing Cements-Determination of strength. The hydraulic pressure testing machine (YES-300) with a maximum load of 300 KN and a loading rate of 2.0 ± 0.5 kN/s was used to measure compressive strength. The X-
ray diffraction (XRD) spectrum of the samples were performed using a D/Max-RC diffractometer (Japan) with Cu Kα radiation, voltage of 40 kV, current of 150 mA and 2θ scanning ranging between 5° and 90°. FE-SEM observation was performed to analyse the hydration products of the CA using a Zeiss SUPRA™55 scanning electron microscope coupled with a Be4-U92 energy spectrum.

![Figure 3. Process flow of CPB material prepared by SWNS](image)

**3. RESULTS AND DISCUSSION**

**3.1 Properties of CPB material**

**3.1.1 Effect of the amount of FGDG and CS on strength of CPB material**

![Figure 4. Effect of FGDG content on compressive strength of CPB material](image)

![Figure 5. Effect of Na2SO4 content on flexural strength of CPB material](image)

![Figure 6. Effect of Na2SO4 content on compressive strength of CPB material](image)

Similar to GBFS, SWNS itself cannot generate strength by their own hydration, but can be generated strength by activator excitation. In this study, two kinds of industrial solid waste FGDG and CS were selected as main activators, Na2SO4 and CC as auxiliary activators, and the appropriate content of these activators was determined through experiments. FGDG belongs to sulfate activator and CS belongs to alkali activator. Only when the two activators reach a reasonable ratio can SWNS produce the best excitation effect. First fixed SWNS content in CA, the main activator and the auxiliary activator quality percentage were 85%, 10% and 5%, and fixed in the auxiliary activator, the mass ratio of Na2SO4 to CC is 3:2 and FGDG accounted for the proportion of the main activator can be divided into 0, 10%, 20%, 30%, 40%, 50%, 60%, 70%, 80%, 90%, 100%, and then according to the test method specified in section 2.2, determine the proportion of FGDG in activator. Figure 4 shows the effect of FGDG content on the compressive strength of CPB material of SWNS. It can be seen from Figure 4 that when the mass ratio of FGDG and CS is 5:5 (namely 1:1) and the sum of them accounts for 10% of CA, the compressive strength of the CPB material is the highest. The compressive strength of CPB material samples of SWNS on 7d and 28d is 2.12 MPa and 3.14 MPa, respectively, meeting the strength requirements of mine CPB material.

**3.1.2 Effect of the amount of Na2SO4 and CC on strength of CPB material**

Fixed SWNS content in CA, the main activator and the auxiliary activator quality percentage were 85%, 10% and 5%, and fixed the main activator of FGDG and CS quality ratio is 1:1, change the auxiliary activator Na2SO4 quality percentage is 0%, 10%, 20%, 30%, 40%, 50%, 60%, 70%, 80%, 90%, 100%, the change of the auxiliary activator content on CPB material samples of SWNS compressive strength and flexural strength, the test results are shown in Figure 5 and Figure 6.

As can be seen from Figure 5 and Figure 6, when Na2SO4 accounted for 60% of the auxiliary activator content (Na2SO4: CC=3:2), CPB material samples of SWNS showed the highest flexural strength and compressive strength on 7d and 28d. Therefore, the mass ratio of Na2SO4 to CC is set as 3:2, that is, the mass fractions of SWNS, FGDG, CS, Na2SO4 and CC in CA should be 85%, 5%, 5%, 3% and 2%, at this point, the compressive strength and bending strength of CPB material samples of SWNS on 28 d are 3.38 MPa and 1.99 MPa, meet the requirements of CPB material for 28 d compressive strength ≥3.1 MPa and 28 d flexural strength ≥1.9 MPa.
3.2 Phase composition and morphology of CA

Figure 7. XRD spectrum of paste block of CA and SWNS

Figure 7 shows the XRD patterns of SWNS, Na₂SO₄, CC, FGDG and CS with corresponding mass fractions of 85%, 3%, 2%, 5% and 5% of CA after hydration for 7 d and 28 d. The main mineral phases are ettringite (AFt), monticellite, grossularite, kamaishilite, augite and pyrope. From the comparison of the three diffraction curves in Fig. 7, it can be seen that, compared with the diffraction curves of SWNS, only 9.645 Å, 5.576 Å and 4.679 Å, which represent AFt, can be clearly identified as the new crystalline phase after 7 d and 28 d hydration of CA prepared with SWNS as the main raw materials, as the diffraction peaks of AFt and other crystalline phases in SWNS overlap superimposed, they cannot be clearly identified in the XRD pattern. The diffraction peak of AFt at 7 d after CA hydration is sharper than that of AFt at 28 d. It is said that the crystallinity of the hydration products at 28 d is lower than that of the hydration products at 7 d, that is, a large number of amorphous substances are produced at 28 d.

Figure 8 shows the SEM photos of CA paste block on curing 7 d and 28 d. According to Figure 8(a), it can be seen that the micro morphology of the paste block at the time of CA hydration 7 d, after curing, the main hydration products of the paste block were a large amount of low-crystallinity or amorphous C-S-H gels and a small amount of fibrous hydration product AFt, the accumulation of hydration products is not close enough, indicating that the sample of CA has undergone a certain degree of hydration reaction. Figure 8(b) shows the enlargement of the labeled area in Figure 8(a), and a large number of AFt formation of acicular or fibrous hydration products can be seen.

Figure 8. SEM images and EDS spectrum of CA samples at different ages. (a) and (b)-7 d, (c), (d) and (e)-28 d
As can be seen from Figure 8(c), a large number of flocculent gels grew around the particles (see D point in Figure 8(c)) in the paste block (see Figure 8(c)), this indicates that the crystalline material and glass phase in SWNS are fully stimulated to participate in the hydration reaction. It can be seen in the Figure 8(c) that the particles are embedded in the hydration products, and the boundary of the particles is blurred, the hydration products glue a large number of particles together and form a piece, the gap between the particles is greatly reduced, which increases the compactness of the slurry. Figure 8(d) is a local enlarged view of Figure 8(c). It can be seen from the figure that a large number of long-rod-shaped AFt are observed, and most of them grow in radial bundles. Compared with hydration products at 7 d of hydration, hydration products at 28 d of CA become thicker and shorter, and the long rod-shaped AFt and crystalline C-S-H gels grow better, making the structure become very dense, and no obvious large holes can be observed, which makes the paste block have good mechanical strength on the macro level. The energy spectrum semi-quantitative analysis of these long rod-shaped products (Figure 8(e)) was conducted, which was speculated to be AFt, which was also consistent with the analysis results of XRD (Figure 7).

3.3 Hydration mechanism of CPB material

The strength source of CPB material of SWNS is a combination of alkali activator and sulfate activator to joint excitation. When the four activators of FGDG, CS, Na$_2$SO$_4$ and CC are matched according to the appropriate proportion, the CPB material of SWNS is of high strength.

In the early stage of OH$^-$ hydration, CaA and Ca$_2$AF in CC system react with CaSO$_4$·2H$_2$O to form Af$_3$ and Ca(OH)$_2$. Since the amount of CC is low, this reaction can only generate a small amount of Af$_3$, while the resulting CC hydration generates Ca(OH)$_2$ as alkali activator that will continue to react with SWNS. Meanwhile, FGDG and Na$_2$SO$_4$ in CA reacted with SWNS as sulfate activator and CS as alkali activator. CS provides sufficient Ca(OH)$_2$ in the early hydration reaction, so that the system contains higher concentration of OH$^-$, this part of OH$^-$ and the OH$^-$ released from the CC are the main sources of "alkali" in the alkali excitation of CPB material of SWNS. As can be seen from Fig. 4, when the content of FGDG is 100% (that is, CS is not added), the compressive strength of the samples prepared on 7 d and 28 d is very low, indicating that CS is particularly important in stimulating the bonding strength of SWNS.

The incorporation of Na$_2$SO$_4$ will greatly affect the dissolution of CS. This is because Na$_2$SO$_4$ reacts with Ca$^{2+}$ in the system to form CaSO$_4$·2H$_2$O with very low solubility:

$$\text{Ca(OH)}_2 + 2\text{Na}^+ + \text{SO}_4^{2-} + 2\text{H}_2\text{O} \rightarrow \text{CaSO}_4 + 2\text{H}_2\text{O} \cdot 2\text{Na}^+ + 2\text{OH}^-$$

(1)

This reaction indicated that the presence of Na$_2$SO$_4$ would accelerate the dissolution rate of CS, resulting in a higher pH value of the solution than that of saturated Ca(OH)$_2$ solution, thus continuously dissolving silicate and aluminosilicate vitreous under the erosion of high concentration OH$^-$, accelerating the alkali excited reaction of the system. At the same time, because the pH value of the system is much higher than that of OPC slurry, it can also stimulate the volcanic activity of some crystalline materials in SWNS so that they also participate in the hydration reaction. In addition, the introduction of Na$_2$SO$_4$ increased the concentration of SO$_4^{2-}$, so that more AFt could be generated in the early stage. In summary, the early strength of sulphate-induced SWNS is attributed to two factors: the acceleration of the early volcanic ash reaction (active silicon and aluminum from the dissolution of SWNS) and the formation of AFt. As an activator, FGDG provides necessary substances for the formation of AFt.

The Ca$^{2+}$ and Mg$^{2+}$ on the vitreous surface of SWNS can adsorb OH$^-$ to form hydroxides, thus damaging the surface structure of SWNS and promoting the hydration of SWNS. The bond strength of Ca-O and Mg-O bonds in the external body of SWNS network are much stronger than those of Si-O and Al-O bonds, so the network structure of SWNS glass body dominated by [AlO$_4$]$^-$ tetrahedron and [SiO$_4$]$^4-$ tetrahedron can be destroyed. Si-O-Al and Al-O-Al in the vitreous of SWNS were eroded by OH$_{-}$, and Al$^{3+}$ was gradually released. The concentration of AlO$_2^-$ in the solution increases and reacts with Ca$^{2+}$ and SO$_4^{2-}$ to form AFt (3CaO·Al$_2$O$_3$·3CaSO$_4$·32H$_2$O). A certain amount of alkali activator can destroy the vitreous network structure of SWNS and promote hydration. An appropriate amount of sulfate activator can provide Ca$^{2+}$, SO$_4^{2-}$ to produce the hydration product AFt.

4. CONCLUSIONS

(1) The SWNS after molten iron extraction from nickel slag contains both glass phase and crystalline material, and the excitation effect of FGDG and CS alone is weak. The special CA for CPB mining can be prepared by adding composite activator, which can meet the requirements of mine CPB.

(2) With FGDG and CS as the main activator, Na$_2$SO$_4$ and CC as the auxiliary activator, CA was prepared from SWNS, FGDG, CS, Na$_2$SO$_4$ and CC by mass ratios of 85%, 5%, 5%, 3% and 2%, then the CPB materials with a slurry concentration of 80% is made by the ratio CA/IOl of 1:4. The 28 d flexural strength and compressive strength can reach 1.99 MPa and 3.38 MPa, meet the requirements of CPB material (28 d flexural strength ≥ 1.9 MPa, compressive strength ≥ 3.1 MPa).

(3) The results of XRD, SEM analysis and hydration mechanism show that the hydration products of the prepared CA are C-S-H gel and AFt. During the hydration of CA, the network structure of [AlO$_4$]$^-$ tetrahedron and [SiO$_4$]$^4-$ tetrahedron in SWNS glass phase is destroyed. Ca$^{2+}$ and Mg$^{2+}$ participate in the reaction to form hydrated aluminosilicate (C-M-S-H) gel containing Ca$^{2+}$ and Mg$^{2+}$, which plays an important role in the continuous increase of the strength of CPB material in the later stage.

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REFERENCES


