Plasma-chemical modification of concrete

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Abstract – Plasma chemical modification (PCM) improves chemical resistance of a glazed decorative coating and expands opportunities for designing various types of the decoration of concrete. However, PCM exposes the material to a significant thermal shock that leads to dehydration of hydrosilicates in the cement matrix followed by loss of strength of the concrete coating. This results in a reduction in the service characteristics of the concrete protective and decorative layer such as adhesion strength and freeze-thaw resistance and shortens the coating durability. Therefore, the improvement of the PCM method is a hot topic of the study and it is one of the most effective and modern technologies, which will result in enhancement of serviceability of concrete with the glazed decorative coating.

This study investigated the effect of incorporation of calcium aluminate cement as a silicon-based intermediate layer between the glazed decorative layer and the concrete surface.

The study demonstrated that the calcium aluminate cement intermediate layer after plasma chemical modification was dehydrated. Based on mineral composition and structural analysis, the dehydrated layer was visually divided into three zones: the upper dense layer; an intermediate layer with micro cracks; and the bottom dense layer. The microstructure of the dehydrated zone is represented by small non-developed microcracks less than 50 μm. Magnesium spinel was the main crystal phase in a calcium aluminate cement matrix. The dense dehydrated zone had a shell-like fracture, which is typical for spinel with imperfect cleavage.

This study developed a glazed decorative coating and a protective calcium aluminate cement-based intermediate layer with the adhesion strength of 4.1 MPa and freeze-thaw resistance of more than 50 cycles that exceed service characteristics of concrete with the coatings using other materials.

Keywords: Plasma-chemical modification, protective-decorative coating, plasma jet, face layer.

I. INTRODUCTION

The implementation of the Presidential program “Affordable Housing for Russian Citizens” considers an increase in the production of wall building materials, particularly concrete products. Protective and decorative coatings on concrete significantly enhance the architectural and artistic merits of buildings and structures.

One of the most effective modern technologies is the plasma-chemical modification of protective-decorative coatings applied on concrete. To expand the color characteristics and minimize dehydration, various fillers made of ceramics, glass, expanded clay, chamotte, waste from the mining or metallurgical industry, etc. are used
before the plasma-chemical modification in the molding of concrete “face down” or “face up” [1-9]. Protective-decorative coatings on concrete have relatively low quality indicators [10, 11]. The adhesion strength of the coating with the substrate does not exceed 1 MPa and is 0.1-0.2 MPa, and the freeze-thaw resistance does not exceed 15-30 cycles.

It is proposed to use alumina cement as the basis of a protective and decorative coating, and as a filler - certain fractions of colored container glasses and the breakages of sanitary ceramics. According to a number of authors, when plasma reflows of concrete products are processed, the surface treated by the high-temperature jet is heated to a depth of several millimeters [10-14]. In this regard, the compositions of mixtures based on alumina cement have been developed for preliminary application to the front surface of concrete.

II. Results and discussions

To study the effect of PCM on the formation and accumulation of fusion, samples of alumina cement were made with a size of 20×20×20 mm. After hardening for 3 days, samples were reflorew at the speed of the plasma jet passing through the face at a rate of 2 mm / s. The high-temperature source was the plasma plant "Gorynych" with a plasma jet temperature of 6000 K. The plasma torch's operation parameters were 6 amperes, 140-160 volts (Fig. 1).

The average chemical composition of the glass determined by the X-ray fluorescent method is presented in Table I and differs from the chemical composition of the alumina cement. The change in the chemical composition of the amorphous phase is associated with the processes of evaporation and thermal diffusion.

TABLE I. CHEMICAL COMPOSITION OF THE AMORPHOUS PHASE AFTER PLASMA-CHEMICAL MODIFICATION

<table>
<thead>
<tr>
<th>Chemical composition, mass. %</th>
<th>Al₂O₃</th>
<th>CaO</th>
<th>MgO</th>
<th>SiO₂</th>
<th>V₂O₅</th>
<th>Fe₂O₃</th>
<th>MnO</th>
<th>TiO₂</th>
</tr>
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<tbody>
<tr>
<td>Chemical composition</td>
<td>64.81</td>
<td>17.02</td>
<td>16.01</td>
<td>2.44</td>
<td>0.41</td>
<td>0.17</td>
<td>0.08</td>
<td>0.06</td>
</tr>
</tbody>
</table>

The thickness of the fused zone is affected by the speed of passage of the plasma jet over the face of the concrete. With a decrease in the reflow rate of the sample from 20 to 10 mm / s, a continuous glassy coating with a thickness of 500 ± 25 μm was formed on their front surface, and 2500 ± 25 μm at a velocity of 2 mm/s (Fig. 2).

As can be seen from Fig. 2, with an increase in the rate of the plasma jet passing over the face of the sample at a rate of more than 15 mm/s, there was no formation of a continuous coating on the front surface of samples from alumina cement.

At a plasma flow rate of 2 mm/s, the temperature of the fusion heating and the kinetics of its cooling were determined (Fig. 3). For this purpose, an optical pyrometer and a platinum-platinum-rhodium thermocouple were used.

As can be seen from Fig. 3, heating to 2000 °C occurred almost in fractions of a second. Cooling of the facial zone to 300 °C occurred after 5 minutes.

The structure and distribution of elements of the fused layer of dehydrated alumina cement are presented in Figures 4, 5.
The melted zone was examined by X-ray phase analysis (Fig. 6).

As can be seen from Fig. 6, the fused layer is represented by an amorphous phase. The composition of the amorphous phase is represented by Mg-Ca-Al glass.

The microstructure of the glass phase is shown in Fig. 7; the fusion is not homogenized and had, in addition to gas inclusions, an area of microinhomogeneity.

The phase composition of the dehydrated zone (3) and its sections: a, b, c was studied by X-ray phase analysis. The phase composition of the upper dense layer (a) is represented by calcium monoaluminate and spinel (Fig. 9).
The composition of the spinel \( \text{Mg}_{0.4}\text{Al}_{2.4}\text{O}_4 \) was identified by the magnitude of the interplanar distances.

In this layer (a) there were no microfractures (Fig. 10).

The phase composition of layer (b) is represented by spinels with different substitution of the crystallographic positions: 

\[ \text{(Mg}_{0.68}\text{Al}_{0.32})(\text{Al}_{0.84}\text{Mg}_{0.16})_2\text{O}_4 \] and

\[ \text{(Mg}_{0.68}\text{Al}_{0.32})(\text{Al}_{0.68}\text{Mg}_{0.32})_2\text{O}_4 \] and calcium monoaluminate (Fig. 11).

Spinels with different substitution of crystallographic positions were identified by the magnitude of interplanar distances using the computer program "Crystallography" (Table II).

In the layer (b) (Fig. 12), mixed spinels were formed under the action of a plasma jet.

As can be seen from Fig. 12 (a, b), in this layer in addition to the crystalline phase and pores, there was an insignificant number of undeveloped microfractures, the size of which did not exceed 50 μm.

### Table II. Spinels identified in the protective-decorative coating after PCM

<table>
<thead>
<tr>
<th>Compositions</th>
<th>Intensity, %</th>
<th>Intensity, %</th>
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<th>Intensity, %</th>
<th>Intensity, %</th>
<th>Intensity, %</th>
</tr>
</thead>
<tbody>
<tr>
<td>MgAl_2O_4</td>
<td>10</td>
<td>2.4</td>
<td>6</td>
<td>2.0</td>
<td>4</td>
<td>1.5</td>
</tr>
<tr>
<td>Mg_0.4Al_2O_4</td>
<td>10</td>
<td>2.4</td>
<td>3</td>
<td>2.0</td>
<td>1</td>
<td>1.5</td>
</tr>
<tr>
<td>(Mg_{0.68}Al_{0.32})(Al_{0.32}Mg_{0.68})_2O_4</td>
<td>10</td>
<td>2.4</td>
<td>5</td>
<td>2.0</td>
<td>4</td>
<td>1.5</td>
</tr>
<tr>
<td>MgAl_{1.94}Fe_{0.1}O_4</td>
<td>10</td>
<td>2.4</td>
<td>5</td>
<td>2.0</td>
<td>4</td>
<td>1.5</td>
</tr>
</tbody>
</table>

Note: * – spinel with different substitution of crystallographic positions of Mg\(^{2+}\)Al\(^{3+}\)

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Fig. 9. Powder x-ray diffractogram of dehydration zone (a – upper dense layer): ○ – MgAl_2O_4; Δ – CaAl_2O_4

Fig. 10. Microstructure of the dehydration zone (a – upper dense layer): 1 – crystalline phase; 2 – pores

Fig. 11. A powder X-ray diffractogram of the dehydration zone in a sample of alumina cement (layer b): ○ – MgAl_2O_4; Δ – CaAl_2O_4

Fig. 12. A microstructure of the dehydrated layer (b): 1 – matrix of alumina cement; 2 – fractures; 3 – pores
As it was shown by the X-ray phase analysis of the lower layer (c), the main phases are calcium monoaluminate and spinel of composition \( \text{MgAl}_{1.9}\text{Fe}_{0.1}\text{O}_{4} \). The microstructure of the layer (c) is shown in Fig. 13 (a, b, c).

The crystalline phase has a conchoidal fracture, which is typical of spinel having imperfect cleavage.

The formation of spinels contributed to the sintering and consolidation of the fused layer and, as a consequence, to an increase in such operational parameters as adhesion strength and freeze-thaw resistance.

![Fig. 13. A microstructure of the dehydrated layer (c): 1 – crystals; 2 – pores; 3 – elements of a conchoidal fracture](image)

III. Conclusion

The conducted research made it possible to establish regularities in the formation of the structure of protective-decorative coatings on concrete using color metal salts. Using a plasma jet, it is possible to regulate the medium that will make it possible to expand the color scale of coatings.

High microhardness of coatings obtained after plasmochemical modification of fine-grained concrete will improve its resistance to scratching and possible shock loads during operation. This will increase the reliability and durability of fine-grained concrete with protective-decorative coatings.

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