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W.J. Quist, S.J.C. Granneman & R.P.J. van Hees (eds.)
MITIGATING SALT DAMAGE IN LIME-BASED MORTARS WITH MIXED-IN CRYSTALLIZATION MODIFIERS

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KEYWORDS

Crystallization Modifiers, Self-Healing, Lime Mortar

ABSTRACT

This paper describes some of the most important results of a four year PhD research on the use of crystallization modifiers mixed in lime mortar to mitigate salt crystallization damage. The research focused on two of the most damaging salts, sodium chloride and sodium sulfate, and suitable crystallization modifiers (sodium ferrocyanide and borax). We report the major findings related to the effectiveness of the modifiers when mixed in the mortar, and the results of characterization of the additivated mortars in comparison with reference mortars. Moreover, the durability of the developed mortars to salt decay is discussed, based on the results of an accelerated salt weathering test carried out in laboratory. No major effects of the modifiers on the fresh and hardened mortar properties were observed which might restrain the application of crystallization modifiers in restoration mortars. Additionally, the mortars with mixed-in modifiers showed a considerable improvement of the salt resistance when compared to reference mortars. Considering these results an outlook for future research pathways is given.

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INTRODUCTION

Salt crystallization damage in porous building materials is a ubiquitous threat to our built cultural heritage. Low mechanical strength makes lime-based mortars especially susceptible to salt damage. In restoration or renovation works, replacement of renders and plasters often constitutes a large part of the total costs of the project. Current solutions, such as using a stronger binder or changing the moisture transport properties of the mortar, usually have a limited resistance to salt decay and low compatibility with the existing materials [1, 2].

Alternatively, the use of crystallization modifiers has been proposed [3]. Crystallization modifiers do not aim to alter the material properties, but the damaging mechanism itself. Using crystallization modifiers in porous building materials has gained wide research interest in the last years (see e.g., [3-5]). However, the use of modifiers in a fresh mortar, thereby giving the mortar “self-healing properties”, is relatively new. By mixing modifiers in during mortar production, they can become active as soon as the damaging salts enter the porous material. Promising results have already been obtained in a pilot study [6].

To further study the feasibility of the use of crystallization modifiers to mitigate salt crystallization damage, a PhD research was started. This research had the following aims: i) Identify suitable modifiers for two of the most abundant and damaging salts: sodium chloride and sodium sulfate, ii) Study the modifier-salt interaction in bulk solution to elucidate the working mechanism and find a suitable concentration, iii) Test the effect of modifier addition on mortar properties (which might limit their application), and iv) Test the durability of additivated mortars in an accelerated salt weathering test.

From literature research, sodium ferrocyanide and borax (sodium tetraborate decahydrate) were identified as potential modifiers for sodium chloride and sodium sulfate, respectively [7]. Ferrocyanide is a well-known modifier for sodium chloride. Therefore, this research focused on the study of its effect on fresh and hardened mortar properties and on its capability of mitigating salt decay in mortar. Borax on the other hand is less well-known and was therefore first studied in bulk solution experiments. In this research, the effect of borax addition on solution properties and its effect on crystallization of sodium sulfate were studied. As has been reported in Ref. [8], depending on the starting concentration of the solution, two different phases of borax precipitate, each having a different effect on sodium sulfate crystallization.

In this paper the major findings relating to the effect of modifier addition on mortar properties and to the durability of additivated mortars are reported. First, an experiment to test whether borax’ effectiveness is affected by the carbonation process is described. Then, the effect of modifiers on mortar properties such as workability, water absorption and drying and flexural and compressive strength are reported. Finally, the durability of the additivated mortars with respect to salt crystallization damage is discussed based on the results of an accelerated salt weathering test.

MATERIALS AND METHODS

Mortar characterization

Two types of specimens were prepared: lime only specimens and mortar (lime + sand) specimens. The first, used to study the effect of borax on sodium sulfate crystallization, were prepared by mixing calcium hydroxide powder (Sigma-Aldrich, ≥ 96% purity) with distilled water only or with water additivated with the modifier. After carbonation, a blank specimen was treated with borax solution. Then, the blank and the two borax specimens
were contaminated with sodium sulfate solution. After drying, the specimens were broken and the cross section was studied using SEM. By comparing specimens additivated with borax prior to carbonation and those to which borax was added later on, the effect of carbonation on the effectiveness of borax as modifier of sodium sulfate crystallization was assessed. Full details of this experiment can be found in Ref. [9].

Mortar specimens were prepared (according to EN1015-2) to test fresh and hardened mortar properties and assess the durability towards salt decay of the additivated mortars. Commercial air lime (Supercalco90) and standard sand (EN 196-1) were used. The modifier was added to the water used to prepare the specimens: 0.94 wt% sodium ferrocyanide and 3.2 wt% borax were added with respect to the lime. Several fresh and hardened mortar properties were tested according to standard procedures or techniques: workability (EN1015-3), water absorption and drying (EN1015-18), porosity and pore size distribution (Mercury Intrusion Porosimetry) and flexural and compressive strength (EN1015-11). Full details on the preparation of the mortars and the testing methods can be found in Ref. [9].

![Figure 1: Temperature and RH cycles used in the accelerated salt weathering test. This entire cycle was repeated 5 times.](image)

**Accelerated salt weathering test**

The salt crystallization resistance of the reference and additivated mortar specimens was tested with a custom designed salt weathering test, shown in Figure 1, simulating circumstances found in practice. 80 RH% is above the equilibrium relative humidity of sodium chloride (RH\textsubscript{eq} = 75%), but below that of the sodium sulfate phases. Consequently, the sodium chloride crystals will deliquesce when the humidity goes up and recrystallize when the humidity goes down again. This ensures multiple crystallization cycles for sodium chloride, a requirement for this specific salt for damage development. Recrystallization of sodium sulfate was obtained only by rewetting with liquid water at the end of a cycle.

Before the test, the specimens were contaminated with salt solution via capillary absorption. A precise amount of solution was used to ensure contamination with 1 wt% sodium sulfate or 2 wt% sodium chloride with respect to the mortar. In total 3.46 (reference NaCl), 3.91 (ferrocyanide), 1.77 (reference Na\textsubscript{2}SO\textsubscript{4}) or 1.91 (borax) gram salt was added to the specimens during the complete test. For each mortar type, 3 replicas were tested.
After each complete cycle, all specimens were rewetted with water equal to the initial amount used to contaminate the specimens. After 3 cycles, salt solution (exact amount to obtain again 1 wt% sodium sulfate or 2 wt% sodium chloride) was used for rewetting, in order to replenish the brushed off salt. After rewetting, any loose material was brushed off and the specimens started a new cycle. The brushed off material was washed and dried in order to separate the salt efflorescences from the debris. The debris was weighed and the amount of salt calculated by the difference. In total the specimens were tested for 5 cycles. Full details of the experiment can be found in Ref. [10].

RESULTS

Mortar properties

The effect of borax on sodium sulfate crystallization can be observed in the SEM pictures in Figure 2. When Fig. 2A is compared to 2B/C, it is clear that the crystal habit of sodium sulfate without borax is different from the crystal habit in the presence of borax. If subsequently Figures 2B and 2C are compared, a similar crystal habit can be seen in both figures. This means that the carbonation process has no effect on the effectiveness of borax as modifier for sodium sulfate.

A selection of the measured fresh and hardened mortar properties is summarized in Table 1 (additional characterization results can be found in Ref. [9]). When the values for additivated and reference specimens are compared, no notable differences can be observed. It can therefore be concluded that there are no negative consequences to mixing these quantities of modifiers in the mortar during production, as the addition of these modifiers does not negatively affect the mortar properties.

<table>
<thead>
<tr>
<th>Property</th>
<th>Method</th>
<th>Reference</th>
<th>Ferrocyanide</th>
<th>Borax</th>
</tr>
</thead>
<tbody>
<tr>
<td>Workability</td>
<td>Flow table test</td>
<td>170 mm</td>
<td>161 mm</td>
<td>161 mm</td>
</tr>
<tr>
<td>Water content</td>
<td></td>
<td>15.95 wt%</td>
<td>15.14 wt%</td>
<td>14.59 wt%</td>
</tr>
<tr>
<td>WAC [kg/m²h⁰.⁵]</td>
<td>Capillary rise</td>
<td>8.05</td>
<td>7.62</td>
<td>7.84</td>
</tr>
<tr>
<td>Density [kg/m³]</td>
<td>Saturation at atm pressure</td>
<td>26.7 ± 0.19</td>
<td>25.9 ± 0.24</td>
<td>27.1 ± 0.06</td>
</tr>
<tr>
<td>Open porosity (%V/V)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Bulk density [g/ml]</td>
<td>Mercury Intrusion</td>
<td>1.977</td>
<td>1.971</td>
<td>1.961</td>
</tr>
<tr>
<td>Open porosity (%V/V)</td>
<td>Porosimetry</td>
<td>25.1 ± 0.11</td>
<td>25.4 ± 0.36</td>
<td>25.1 ± 0.55</td>
</tr>
<tr>
<td>Tensile strength (N/mm²)</td>
<td></td>
<td>0.79 ± 0.11</td>
<td>0.85 ± 0.03</td>
<td>0.92 ± 0.11</td>
</tr>
<tr>
<td>Compressive strength (N/mm²)</td>
<td></td>
<td>2.01 ± 0.33</td>
<td>2.08 ± 0.18</td>
<td>2.61 ± 0.22</td>
</tr>
</tbody>
</table>

Table 1: Fresh and hardened mortar properties of the different 1:3 lime:sand mortar mixtures. Results previously reported in [9].
Figure 2: SEM images of lime-only specimens contaminated with sodium sulfate. A: reference specimen; B: specimen with 3.2wt% borax mixed in during preparation (thus before carbonation); C: Specimen additivated with borax after full carbonation of the specimen. A clear difference in crystal habit can be observed between the reference specimen and both specimens with borax. Contrarily, no distinction can be observed between B and C, meaning that the carbonation process has no effect on the borax effectiveness as modifier of sodium sulfate crystallization.
Salt durability [10]

During the accelerated salt weathering test, the specimens were monitored both visually and gravimetrically. The material loss (with respect to the mortar) is plotted in Figure 3, and the salt loss is visualized in Figure 4. It is clear that for both salts, the reference specimens suffer considerable material loss after 5 cycles. Contrarily, the additivated mortars show no or only minor material loss. The ferrocyanide stimulates efflorescence of the salt, i.e. crystallization outside the material, as does borax but to a lesser extent. Figure 5 compares specimens contaminated with sodium chloride at the start and end of the test. The reference specimen shows considerable surface loss at the end of the test. Contrarily, the specimen with ferrocyanide shows no material loss, but extensive efflorescence, which developed very fast just after rewetting and brushing of the specimens. In Figure 6 the specimens contaminated with sodium sulfate are compared. At the end of the test, both specimens show damage at the surface, but this is in the case of the specimen additivated with borax considerably less than in the reference specimen. Both the material loss and the visual observations show that both sodium chloride and sodium sulfate have the potential to cause considerable damage in the reference specimen. However, when the mortars are additivated with modifiers, damage does not occur or is considerably less.

Figure 3: Cumulative material loss, comparison between reference and additivated specimens
Figure 4: Cumulative salt loss, comparison between reference and additivated specimens.
Figure 5: Comparison between reference (A/B) and specimens with mixed-in ferrocyanide (C/D), both contaminated with sodium chloride. A/C show the specimens at the start of the test, B/D show the specimens ~15 minutes after brushing after the 5th cycle. The reference specimen (B) shows sanding of the surface. The specimen with ferrocyanide shows no surface damage, only a large amount of efflorescence.

Figure 6: Comparison between reference (A/B) and specimens with mixed-in borax (C/D), both contaminated with sodium sulfate. A/C show the specimens at the start of the test, B/D show the specimens after brushing after the 5th cycle. The reference specimen (B) shows clear damage at the surface. The specimen with borax shows only minor surface damage at the lower left corner (D).
CONCLUSIONS

The additivation of mortars with crystallization modifiers during production has been proposed here to mitigate salt crystallization damage in porous building materials. Suitable crystallization modifiers for sodium chloride (sodium ferrocyanide) and sodium sulfate (borax) were identified to be mixed in a mortar during production. In this research, at first the effectiveness of borax as a modifier for sodium sulfate crystallization when mixed in lime was assessed and confirmed. In a next step, additivated mortars were characterized and compared to reference mortars in order to identify potential (negative) effects on fresh and hardened mortar properties. None of the tested properties was affected by the addition of the modifiers, meaning that there are no contra-indications to mixing them in the mortar in the used concentrations during production.

Finally, the salt crystallization resistance of the additivated mortars was assessed using a custom designed accelerated salt weathering test. The mortars with mixed-in modifiers showed a considerable improvement of the salt resistance when compared to reference mortars. Combining all these results it can be concluded that additivating mortars with crystallization modifiers during their production is a feasible method in order to mitigate salt crystallization damage in porous building materials.

OUTLOOK

The PhD research described in this paper shows the viability of using crystallization modifiers to mitigate salt weathering damage in porous building materials. Although the proof-of-principle has been shown on the laboratory scale, more research is needed to develop the material into a commercial product, suitable for renovation or restoration works. Interesting research paths to further develop the mortar designed in this project are:

- Studying the effect of modifier in mortars with different composition (e.g. cement-based).
- Studying the speed of modifier leaching and if necessary developing possible solutions, e.g. encapsulation.
- Assessing the effect of the identified modifiers on other salts and on salt mixtures and the possibility of combining different modifiers.
- Assessing the workability and effectiveness of the developed mortar (on test panels) in situ.

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REFERENCES


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