Effects of *Bacillus subtilis* biocementation on the mechanical properties of mortars

*Efeitos da biocimentação promovida por Bacillus subtilis nas propriedades mecânicas de argamassas*

Abstract

This study aims to evaluate the influence of *B. subtilis* AP91 spores addition on the mechanical properties of mortars. *B. subtilis* strain AP91, isolated from rice leaves of the needle variety, which has an early cycle of production, was used at the concentration of 10^5 spores/mL in mortars with cement-to-sand ratio of 1:3 (by weight) and water-to-cement ratio (w/c) of 0.63. These spores were added in two different ways: in the mixing water and by immersion in a solution containing bacterial spores. Scanning Electron Microscope (SEM) analysis showed crystals with calcium peaks on the EDS, which possibly indicates the presence of bioprecipitated calcium carbonate. The results obtained in the mechanical analysis showed that the bioprecipitation of CaCO₃ by *B. subtilis* strain AP91 was satisfactory, particularly when the spores were added in the mixing water, increasing the compressive strength up to 31%. Thus, it was concluded that the addition of *B. subtilis* AP91 spores in the mixing water of cement mortars induced biocementation, which increased the compressive strength. This bioprecipitation of calcium carbonate may very well have other advantageous consequences, such as the closure of pores and cracks in cementitious materials that could improve durability properties, although more research is still needed on this matter.

Keywords: calcium carbonate, biocementation, *B. subtilis*, mechanical properties, SEM.

Resumo

Este estudo tem como objetivo avaliar a influência da adição de esporos de *B. subtilis* AP91 nas propriedades mecânicas de argamassas. *B. subtilis* 'AP91 isolada de folhas de arroz da variedade agulha precoce foi utilizada na concentração de 105 esporos/mL em argamassas com traço de 1:3 (em massa) e relação água/cimento (a/c) de 0,63. Os esporos bacterianos foram adicionados de duas diferentes formas: na água de amassamento e por imersão em solução contendo os esporos. A análise por Microscopia Eletrônica de Varredura (SEM) mostrou cristais com picos de cálcio, o que possivelmente indica a presença de carbonato de cálcio bioprecipitado. Os resultados obtidos a partir da análise mecânica mostraram que a bioprecipitação de CaCO₃ pela bactéria *B. subtilis* AP91 foi satisfatória, particularmente quando os esporos foram adicionados à água de amassamento, aumentando a resistência à compressão em até 31%. Portanto, conclui-se que a adição de esporos de *B. subtilis* AP91 na água de amassamento das argamassas induziu a biocimentação, que aumentou a resistência à compressão. A bioprecipitação de carbonato de cálcio pode ter outras consequências benéficas, como o fechamento de poros e fissuras em materiais cimentícios, que poderiam melhorar a durabilidade, embora mais pesquisas neste aspecto ainda sejam necessárias.

Palavras-chave: carbonato de cálcio, biocimentação, *B. subtilis*, propriedades mecânicas, MEV.

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1. Introduction

Except for water, the cementitious materials are the most consumed substances on Earth [1]. Because of that, they should be studied to present satisfactory performance and, for that matter, several works are being performed nowadays to improve the mechanical properties of such materials. In view of the growing concern about improving the characteristics and durability of cementitious materials, additions of bacteria, mainly of the genus *Bacillus*, are being studied for crack filling and increasing of the compressive strength through calcium carbonate (CaCO$_3$) precipitation [2]. Because of that, the bioprecipitation of CaCO$_3$ by bacteria in cementitious materials is also called biocementation [4]. It should be noted that bacteria already exist for billions of years, however many of their biotechnological applications are not yet widespread [5]. Although many researchers are interested in the bioprecipitation of CaCO$_3$, the whole process is not yet clear and defined. Once the bacteria can precipitate calcium carbonate through several mechanisms, the physiology and genetic involved are complex and difficult to understand [6], [7], [8].

Some bacteria that have the ability of producing minerals were used to repair limestone monuments [9], [10], [11], [12], [13], [14] and to fill pores and cracks in concrete and other cementitious materials [5], [8], [15], [16], [17], [18], which can improve their mechanical properties and durability. Among the bacteria used in researches about biocementation, the bacteria *B. subtilis* stand out. These bacteria can produce calcite, a crystalline form of CaCO$_3$, which is precipitated when they’re exposed to a medium with a calcium source [25], [26], [27].

The biocementation can be induced by two ways of addition of bacteria: in the mixing water of mortars [1] [3], [15], [19], [22], [28], [29] or by immersion of the specimens in a solution containing bacteria [18], [30]. However, the studies that confront these two ways of bacteria addition are rare, so that researches comparing both are important. The aim of this study was to evaluate the influence of biocementation, induced by two different ways of spores addition, on the mechanical properties of mortars. Thus, the spores of bacteria *B. subtilis* strain AP91 were used.

2. Materials and methods

2.1 Bacteria and growth conditions

*B. subtilis* AP91, isolated in Brazil from rice leaves of the early cycle needle variety, was used [31]. To achieve the required concentration, the bacteria was cultivated in a Luria-Bertani (LB) medium under agitation of 170 rpm at 37°C for 48 hours. The medium was poured into 15 mL polystyrene tubes, which were centrifuged at 4,000 rpm and 23°C for 3 minutes, in order to separate the bacterial cells from the growing medium. The supernatant was discarded and the pellet containing the bacterial cells was re-suspended in a phosphate buffer solution, homogenized and centrifuged again. This procedure was performed four times.

The obtained concentration was quantified in a spectrophotometer with a reading of 600 nm, from the Equation 1 described by [15]:

\[ X = 8.59 \cdot 10^{-3} \cdot Y^{1.3627} \] (1)

Where *X* is the concentration in cells/mL and *Y* is the reading of the spectrophotometer at 600 nm.

From the found concentration, the dilution calculations were made in order to achieve a concentration of 10$^5$ cells/mL. The dilution was made in a phosphate buffer solution, then stored at 8°C for 48 hours for the spores formation.

2.2 Mortars

The mortars were produced using:

- Portland cement (CP-II E-32, Brazilian denomination);
- Quartz fine sand: fineness modulus of 1.51, maximum size of 1.18mm and specific gravity of 2.604 g/cm$^3$;
- Potable water;
- Phosphate buffer solution: 1.05 g/L dibasic-anhydrous sodium phosphate (0.07% of the cement bulk), 0.36 g/L monobasic sodium phosphate (0.02% of the cement bulk) and 8.17 g/L sodium chloride (0.51% of the cement bulk);
- *B. subtilis* AP91.

The spores of bacteria, in a concentration of 10$^5$ spores/mL, were added to the mixture in two different ways: in the mixing water and by immersion in a phosphate buffer solution containing the spores. For that, four variations of mortars were produced in this study; the nomenclature and composition of those are presented in Table 1. The cement-to-sand ratio used was 1:3, by weight, and the bacterial culture or water-to-cement ratio (w/c) was kept at 0.63. The mixture proportions are shown in Table 2.

2.3 Experimental methods

The microstructure of the mortars was evaluated by Scanning Electron Microscopy (SEM) and Energy Dispersive System (EDS). In these analysis it was found crystals of calcium carbonate (CaCO$_3$) in the mortars with addition of spores. In order to evaluate the influence of CaCO$_3$ crystals on the mechanical properties of mortars, all of them were tested by compressive strength, indirect tensile strength (Brazilian test) and dynamic modulus of elasticity.

<table>
<thead>
<tr>
<th>Table 1 Nomenclature of the mortars</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Mortars</strong></td>
</tr>
<tr>
<td>REF_WT</td>
</tr>
<tr>
<td>SP_IM*</td>
</tr>
<tr>
<td>REF_PB</td>
</tr>
<tr>
<td>SP_MW</td>
</tr>
</tbody>
</table>

* Spores were added during the curing time, when this mortar was submerged in a tank with phosphate buffer with a concentration of 105 spores/mL.
2.3.1 SEM analysis

SEM was used to observe the mortar microstructure and the precipitation of calcium carbonate (CaCO₃). The microanalysis was performed using an FEI-Quanta 200 Scanning Electron Microscope with an accelerating voltage of 25 kV for determination of the chemical elements. The mortars specimens of 10x10x10mm had their surface covered with gold and were tested after 7 and 28 days of curing.

Table 2
Mixture proportions

<table>
<thead>
<tr>
<th>Mortars</th>
<th>Cement (g)</th>
<th>Sand (g)</th>
<th>Water (mL)</th>
<th>Phosphate buffer (mL)</th>
<th>Spores of B. subtilis/mL</th>
</tr>
</thead>
<tbody>
<tr>
<td>REF_WT</td>
<td>2362.5</td>
<td>7087.5</td>
<td>1488</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>SP_IM</td>
<td>2362.5</td>
<td>7087.5</td>
<td>1488</td>
<td>–</td>
<td>10⁵*</td>
</tr>
<tr>
<td>REF_PB</td>
<td>2362.5</td>
<td>7087.5</td>
<td>–</td>
<td>1488</td>
<td>–</td>
</tr>
<tr>
<td>SP_MW</td>
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<td>7087.5</td>
<td>–</td>
<td>1488</td>
<td>10⁵*</td>
</tr>
</tbody>
</table>

* The concentration of 10⁵ spores/mL was dispersed in the phosphate buffer solution.

Figure 1
SEM of mortars after 7 days of curing at magnification of 12.000x. (a) REF_WT. (b) SP_IM. (c) REF_PB and (d) SP_MW
2.3.2 Mechanical properties of the mortars

To study the compressive strength [32], indirect tensile strength [33] and dynamic modulus of elasticity, cylindrical specimens of 50mm diameter and 100mm height were molded according to [32]. All specimens were cured by immersion at room temperature until testing after 7 and 28 days of curing. The mortars REF_WT, REF_PB and SP_MW were cured in a tank of lime-saturated water and the mortar SP_IM was cured in a tank containing a phosphate buffer solution with $10^5$ spores/mL of *B. subtilis* AP91, also saturated with lime.

3. Results and discussions

3.1 Microscopic investigation

To search for evidence of microbial calcium carbonate precipitation, the mortar specimens were examined under a Scanning Electron Microscope (SEM). Fig. 1 shows a scanning electron micrograph of the mortars after 7 days of curing, in which Fig. 1(a) shows the reference mortar (REF_WT) highlighting the cement hydration products, such as calcium silicate hydrate (C-S-H) and ettringite (et). Fig. 1(b) shows the mortar with the addition of spores of *B. subtilis* strain AP91 by immersion (SP_IM), in which were highlighted the bioprecipitation of calcium carbonate (CaCO$_3$) can be observed. Finally, Fig. 1 (c) shows the mortar with phosphate buffer replacing water (REF_PB), where can be noted the presence of calcium hydroxide (CH), calcium silicate hydrate (C-S-H) and an intensive amount of ettringite. At last, Fig. 1(d) shows the addition of spores of *B. subtilis* in the mixing water, in which was highlighted the bioprecipitated calcium carbonate (CaCO$_3$).

Analyzing the SEM in Fig. 1 it can be observed that the mortars have shown the formation of cement hydration products, like...

Figure 2
SEM of mortars after 28 days of curing at magnification of 12.000x. (a) REF_WT. (b) SP_IM. (c) REF_PB and (d) SP_MW
C-S-H, ettringite (etr) and CH. The presence of CaCO$_3$ was observed only in mortars with addition of *B. subtilis* spores (Fig. 1 b and d), both by immersion and in the mixing water. After 28 days of curing, other SEM images were taken to verify if the calcium carbonate was still present in the samples with addition of *B. subtilis* spores. Fig. 2 shows all mortars after 28 days of curing. Considering the images in Fig. 2, it was found the precipitation of calcium carbonate crystals only in the mortars with addition of *B. subtilis* spores (Fig. 2 b and d). After 28 days of curing, about 95% of cement is hydrated, so in these images, no hydration products can be seen. Similar crystals of CaCO$_3$, with rounded structures, were found by Park et al. [2], Hammes et al. [6], Braissant et al. [35], Dupraz et al. [36] and Daskalakis et al. [37].

To complete this analysis, the specimens were submitted to an Energy Dispersive System (EDS), to verify the peaks of the chemical elements present in the sample. From the EDS shown in Fig. 3, it was observed that all mortars present peaks of calcium, but the peaks are higher when the *B. subtilis* spores have been added (Fig. 3 b and d). It is known that the cement has calcium, but in the mortars analyzed there was a greater presence of calcium when the spores were added, probably because the spores might have used the calcium present in cement hydration products and in the lime of the curing water to precipitate CaCO$_3$. The greater presence of calcium indicates that the calcium carbonate crystals deposit themselves in the cement matrix (Fig. 2 b and d). Besides that, it should be noticed that the gold peaks present in all the samples are due to the coating, so that high-resolution images could be obtained.

### 3.2 Compressive strength

The medium compressive strength of different mortars (n = 5) after 7 and 28 days of curing are summarized in Fig. 4. The compressive strength statistically increased in the mortar specimens that contained *B. subtilis* AP91 spores added to the mixing water (SP_MW), both after 7 and 28 days of curing. When the mortars were immerged (SP_IM) there were no significant differences.
differences on the compressive strength in both studied ages, compared to the control mortar (REF). That may be explained by the fact that the spores would have to penetrate the specimens to have access to the hydration products in order to react and precipitate CaCO$_3$; moreover, the spores could find it difficult to penetrate in the specimens because of the use of release agent to mold them. Perhaps, given the sufficient amount of time and the absence of the release agent, this procedure of spores’ addition could have been more efficient.

The same was found in the mortar with phosphate buffer (PB) at 7 days of curing, but at 28 days it was verified a significant decrease on the compressive strength presented by these samples. Ramachandran et al. [15] found that the addition of phosphate buffer in mortars caused better mechanical performance when compared to saline solution. However, these authors do not analyze the influence of phosphate buffer compared to a reference mortar.

It should be noticed that the results of addition of bacteria in the mixing water corroborate with what was presented in other works by Ghosh et al. [38], Reddy et al. [39], Chahal et al. [40] e Chahal and Siddique [41], who added different types of bacteria in the concentration of 10$^5$ cells/mL and found an increase on compressive strength.

The improvement in compressive strength by $B.\ subtilis$ strain AP91 is induced by the deposition of calcium carbonate on the cell surfaces, which may have the ability of closing pores and cracks of cementitious materials [15, 42, 43, 44).

3.3 Indirect tensile strength (Brazilian test)

Fig. 5 shows the effect of addition of $B.\ subtilis$ strain AP91 on the indirect tensile strength of mortars. In this test, the same tendency of the compressive strength test was verified, in which the highest indirect tensile strength, both at 7 and 28 days of curing, was found in the mortar with addition of $B.\ subtilis$ AP91 in the mixing water. It can be pointed out that the improvement on tensile strength is consequence of the improvement of the compressive strength, since the former is usually around 10% of the latter.

![Figure 5](image_url)

**Figure 5**
Effect of $B.\ subtilis$ strain AP91 on the indirect tensile strength of mortars after 7 and 28 days of curing. Values are mean ± Sd (n = 3)

3.4 Dynamic modulus of elasticity

The modulus of elasticity was calculated from the velocity of propagation of the ultrasonic waves, by direct transmission, through the Equation 2 described in the Brasilian standard 15630/2008 [34]. The results obtained are shown in Fig. 6.

$$Ec = \frac{V^2 (1 + \nu) x (1 - 2\nu)}{(1 - \nu)} \times 10^{-3}$$

(2)

It was verified that there was a small gradual increase in the modulus of elasticity of the mortars when were added the buffer solution and also the bacteria spores, both by immersion (SP_IM) and in the mixing water (SP_MW). It should be noticed that the mortars with the highest modulus of elasticity also corresponded to the higher strengths, for both tensile and compression. It was expected that the modulus of elasticity of the mortar containing phosphate buffer would present a lower value when compared to the reference mortar, once it also presented lower compressive strength. However, this test was performed with air-dried specimens that could still have water inside the pores, which can affect the propagation of ultrasonic waves. This could also be related to presence of salts in the buffer solution, which could have filled smaller pores in the matrix, also affecting the velocity of the ultrasonic waves.

Furthermore, the pore filling by biocementation could cause an increase in the velocity of propagation of the ultrasonic waves and, consequently, promote a higher modulus of elasticity of the specimens containing bacteria when compared to the reference.

4. Conclusion

The SEM and EDS analysis verified that the addition of...
B. subtilis AP91 spores in mortars induced biocementation through the bioprecipitation of calcium carbonate crystals;

- When the spores of B. subtilis AP91 were added in the mixing water, they presented better mechanical performance in all tests when compared to the addition by immersion;

- B. subtilis AP91 added into mixing water has shown to be a microorganism with great potential for application in cementitious materials, since the precipitation of CaCO₃ can increase both compressive and tensile strength.

- Because of the precipitation of CaCO₃, the application of this microorganism for the closure of pores and fissures in cementitious materials should continue to be studied, in order to prevent early deterioration.

5. Acknowledgments

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