Incorporation of Optical Density into the Blending Design for a Biocement Solution

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Abstract: The engineering practices for applying the microbial precipitation of carbonates require a design of the blending biocement solution (BCS). The BCS is usually blended with concentrated strains NO-A10, reaction media, such as urea and CaCl₂, and a solvent, i.e., water or seawater. To characterize the BCS, the unknown microbial characteristics, such as the cell viability, are complex factors. Therefore, the optical density (OD) was redefined as Rcv OD*, in which OD* was the tentative OD of the BCS used and Rcv was the conversion rate concerning the cell viability. To determine Rcv values, a standard precipitation curve based on the precipitation rate at 24 h was determined. It was found that the curve was expressed by λ₁OD + λ₂OD², in which λ₁ and λ₂ were 8.46 M and –17.633 M, respectively. With this, the Rcv and OD values of unknown BCS were estimated from the results of precipitation tests using arbitrary OD* values. By extending the testing time, the second order term of OD or OD* was negligible. Accordingly, the precipitation amount is expressed as 8.46 OD, in which the OD can be estimated by precipitation tests using arbitrary OD* values of BCSs. Unless the Ca²⁺ value is dominant, the optimum blending of BCS can be determined by OD. Thus, it is concluded that the blending design of BCS is achieved using 8.46 OD, or 8.46 Rcv OD*, and the standard precipitation curve was defined in this study.

Keywords: urease; standard precipitation of carbonates; optical density; viability of cells; blending design

1. Introduction

The cementation process of natural sediments by carbonates is called carbonate diagenesis. The carbonates existing in the sediments are attributed to the buried effect of dead foraminifers and coccoliths [1–4] or biomineralization [5,6]. It was shown that natural deposited marine soils exhibited significant relationships between unconfined compressive strength and calcium carbonate contents [3]. The results showed that the silty surface sediments at depths of up to 6 m below the sea bottom in Tokyo Bay and Osaka Bay showed increasing rates of 7.5 kPa vane shear strength per a 1.0% increase in carbonate content. On the other hand, silty sediments at a depth of approximately 24 m below the sea bottom off Haneda Airport showed an increase in unconfined compressive strength of 64 kPa per 1.0% carbonate content. Since then, improvement in the carbonation technique in soil has been desired.

For example, Mobley and Hausinger [7] illustrated microbial ureases in terms of their significance, regulation, and molecular characterization, and contributed to expanding...
their medical knowledge on urease to a wide range of readers in other fields. On the other hand, subsequent chemical reactions in carbonate production after urease activity can be seen in agricultural papers [8]. Carbonate precipitation has been used to inhibit the loss of ammonia as fertilizer to the air and to retain the ammonium ions in the soil.

Carbonate diagenesis is distinguished from microbially precipitated carbonates because of natural processes. However, the results and evaluation in engineering may have similarities. The microbial precipitation of carbonate using ureolytic activity has been increasingly studied worldwide and the objectives of studies have been expanding [9–18]. These are based on microbially induced carbonate precipitation and are often called MICP techniques.

In the field of soil improvement, there may be no doubt that the most dominant factors are the carbonate content induced microbially and the soil characteristics. Other factors are environmental conditions, such as temperature, pH, etc. For example, it is known that the mechanical strength of soils has been shown in terms of carbonate content [19]. Van Paassen [20] investigated the microbially induced carbonate solidification of medium- to high-density sands. The study showed that the UCS, deviatoric stress, and tensile strength increased exponentially with the CaCO$_3$ content and initial density of the specimens. Non-destructive measurements were also applied to investigate the carbonate-cementing effects [21,22]. Fukue and Lechowicz [23] showed that, using the pocket penetrometer, the microbially induced carbonate content was the most dominant factor in controlling the strength development of sandy soils. They found that, for loose to medium density sands, the unconfined compressive strength (UCS) estimated by measurements was expressed as a linear function of the carbonate content. It was also shown that the proportional constants varied with the soil type, such as the grain size, distribution, and initial density. Similar results have been reported by many studies [24].

Thus, regardless of whether it is natural or artificial, the evaluation of microbial carbonate precipitation can be achieved with ultimate amounts of induced carbonates. However, it is required to predict or design the ultimate amounts of induced carbonates for engineering practice.

In nature, urease (enzyme) is produced by many organisms, such as animals and plants, including microorganisms. The enzyme activity, i.e., ureolysis due to the urease, is well known as:

\[
(\text{NH}_2)_2\text{CO} + [\text{e}] + \text{H}_2\text{O} \rightarrow 2\text{NH}_3 + \text{CO}_2
\]  

Reaction (1) is known as hydrolysis with the catalysis of urease [e]. A microbe’s system is shown in Figure 1, in which, the first products are NH$_3$ and CO$_2$ [7]. The 2NH$_3$ and CO$_2$ can easily be transported outside of cells, and react with H$_2$O as follows:

\[
2\text{NH}_3 + \text{CO}_2 + \text{H}_2\text{O} \rightarrow 2(\text{NH}_4)^+ + \text{CO}_3^{2-}
\]  

Figure 1. Microbial hydrolysis of urea, showing that first products are CO$_2$ and NH$_3$ inside the microbe’s cell and carbonates are produced outside of the cell.
It should be remembered that the enzyme [e] in Reaction (1) is not given appropriately in terms of quantity and quality. The species and strains of ureolytic microbes, their concentrations, and their micro-environments play an important role for the urease activity [7,13,15,25,26]. For example, the urease activity of biomass increases proportionally to the numbers of cells [25]. This implies that the concentration of microbes plays an important role in the generation rate of carbonate ions.

The product \((\text{NH}_4)_2 \text{CO}_3 \cdot \text{H}_2\text{O}\) increases the pH outside of the cell [8] and helps Reaction (3) in the presence of \(\text{Ca}^{2+}\). Reaction (3) indicates that 1 M \(\text{CaCO}_3\) can be produced when 1 M urea and 1M \(\text{CaCl}_2\) are used, as long as no inhibition of urease activity occurs in Reaction (1). Accordingly, the concentration of \(\text{Ca}^{2+}\) becomes a dominant factor for the \(\text{CaCO}_3\) precipitation rate.

\[
\text{Ca}^{2+} + \text{CO}_3^{2-} \rightarrow \text{CaCO}_3
\]  

(3)

The production of \(\text{CaCO}_3\) does not change the pH of the system. The \(\text{Ca}^{2+}\) is usually supplied in the form of calcium chloride, \(\text{CaCl}_2\). Then, \(\text{NH}_4^+\) and \(\text{Cl}^-\) may increase in the system, and the pH of the solution decreases during Reaction (3).

For the improvement of soils, rock, and concrete using microbial carbonate precipitation, there are basically three problems: the reliability, repeatability, and cost performance. This may result from the drawback in using microbes whose properties and characteristics are influenced by the change in cell viability. This problem becomes serious directly or indirectly when scale-up and commercial applications are considered [27]. In many studies, bacteria and media have been injected into soils for in situ cultivation. In those cases, bacteria cultivation has not always been well governed, and the design of the blending biocement solution (BCS) cannot be made. Under this condition, the reliability and repeatability are difficult to obtain. Therefore, the aim of this study is to solve the problem concerning the unknown quality of the BCS using stored frozen microbes.

2. Materials and Methods

2.1. Microbial Preparation

In this study, microbes were cultivated using a cell incubator (Life-Engineering, Shizuoka, Japan) and prepared with a continuous supply system. This may enable an optimum design of the blending of BCS. The culture solution is centrifuged, and the harvest is stored in a deep freezer (Life-Engineering, Shizuoka, Japan). With these processes, the volume reduction, the long-term-storage, easy and cheap transportation, etc., are achieved. At the same time, the cost performance is improved with the new development of cultivation techniques. The concentration of microorganisms drastically increased up to an optical density (\(\text{OD}_{600}\)) of 4.0 by developing a good medium for the bacteria. Therefore, a smaller cell incubator was enabled.

In 2008, the ureolytic microbes used in this study were isolated from the alluvial soils in Japan. The microbe strains were named NO-A10 [28]. The microbes were cultivated in a cell incubator with a volume of 1 m\(^3\). At present, the culture volume for a batch process is usually 300 L.

Centrifugation is also needed in the batch system. Usually, the 100 L culture solution is continuously centrifuged at 4 °C. The centrifugal concentration factors are usually 250 to 300. It took approximately 9 h to centrifuge the 100 L of the culture solution. A fifteen percent glycerin solution was added to the centrifuged microbes and mixed well. This was used to reduce damage to the microbes by freezing [29]. After mixing, the mixtures were packed in plastic bags or plastic tubes, and frozen in cooled ethanol containing the pieces of dry ice at below −40 °C for more than 20 min. The frozen mixtures were stored in a deep freezer at −80 °C until utilized, as illustrated in Figure 2.
Figure 2. From culture solution to biocement solution.

Under this situation, the quality of stored microbes is usually unknown because of many governing factors, such as the varied nature in the original culture solution, i.e., inherent properties, centrifugal effects, amounts of microbes in the supernatant during centrifuging, effects of added glycerin, aging, thawing effects, etc. Most of these factors reduce the number of viable cells. Accordingly, the urease activity in the BC solution (BCS) will decrease. To characterize the properties of BCS for engineering purposes, it is required to establish a technique to determine the quantity of microbes, including the carbonate precipitation rate (CPR). Herein, CPR is defined as the mass of precipitated carbonates, or the mass of precipitated carbonates for a given time.

In this study, the standard CPR was obtained using OD related to cell viability. Next, the calibration curves were introduced from the standard CPR curve, assuming that OD = Rcv OD*. Herein, OD* was tentatively used as the calculated OD value for the BCS, calculated by the dilution factor from the concentrated volume of microbes. Ultimately, the CPR curves with OD* were checked against the calibration curves, and the Rcv value was obtained. These are discussed later.

When it is expressed as the precipitation rate of CaCO₃, it is roughly divided into two meanings. One is to express the amount of precipitation by the quantity or percentage, and the other is to represent the precipitation rate. In this study, moles (M) were used to represent them by quantity, and percentages are used to represent them as a percentage. On the other hand, if it is expressed in the amount precipitated for a certain time, the precipitation speed was used.

The fundamentals of CPR were studied using the fresh culture in the laboratory. CPRs induced by two ureolytic microbes using 1.0 OD and various Ca²⁺ concentrations are shown in Figure 3. The two types of microbes were named NO-N10 and NO-A10. These are expressed as N10 and A10, respectively. In Figure 3, A10 shows the rapid increases in CPR. It is obvious that the CPRs are controlled by Ca²⁺. Thus, the limitation of Ca²⁺ clearly indicates that the chemical reactions occurred according to Reactions (1), (2), and (3), with partial constraints. It is noted that no precipitation occurred if the concentration of Ca²⁺ exceeded 1.5 M [28].
Concerning the reaction rate, it is common for the lower Ca\(^{2+}\) levels to cause the higher precipitation rate. This is discussed later.

The effects of OD using 1 M Ca\(^{2+}\) are shown in Figure 4. The various OD values were prepared by dilution of the 1.0 OD solution. The initial pH value used was approximately 9.5 due to the 20 mM ammonium buffer. The ambient temperature was 25 °C. The result showed that the CPRs were rapid at the greater OD. If 24 h is selected as the measured time, all of the curves except for 1.0 OD show that the CPRs at 24 h are nearly 1.0 M or greater than that. It is noted that the CPRs greater than 1.0 M can be obtained with greater OD values. This is because hydrated carbonates tend to be produced when the reaction rate is high [30]. For the CPR curve for OD = 0.1, the CPR is 0.65 M.

Considering that the hydrolysis of urea occurs inside of the cell, and that the ammonia and carbon dioxide react with water on the surface of the outside of the cell, molecular-sized carbonates will begin to cover the cell from the initial stage. Accordingly, the cell becomes a nucleus of the induced carbonate particles [31]. Okwadha and Li [32] concluded that the bacterial cell concentration, initial urea concentration, and Ca\(^{2+}\) concentration all influenced the amount of CaCO\(_3\) precipitated and CO\(_2\) sequestrated. Their results also indicated that a
greater amount of CaCO$_3$ would be precipitated with a greater concentration of urea, Ca$^{2+}$, and bacterial cells, so long as these quantities are economically feasible.

In general, the enzyme is not a reactive material. Therefore, it is not consumed in the hydrolysis of urea. In this sense, the hydrolysis continues if urea is supplied from the outside of cell. On the other hand, CaCO$_3$ precipitation continues around the surface of cell until Ca$^{2+}$ is consumed or the supply of CO$_2$ or CO$_3^{2-}$ or H$_2$O stops in Reactions (1) to (3). This condition may be due to the CaCO$_3$ amount per cell becoming too high and the precipitation of CaCO$_3$ being inhibited by the excess CaCO$_3$. Therefore, the hypothetical limitation of CaCO$_3$ precipitation per cell must be considered.

Counting the number of cells is a difficult task; simple methods include the optical density (OD) measurement of the cell suspension [33,34]. OD is a fast, cheap, and high-throughput measurement method widely used to estimate the density of cells in liquid culture [34].

Assuming that all of the cells are viable and under constant Ca$^{2+}$, the microbial CaCO$_3$ precipitation can be written as a function of OD and time. If the precipitation rate is defined at a given constant time, the precipitation rate at a given time is expressed as a function of OD only as follows:

$$\text{CaCO}_3 = f(\text{OD}) \quad (4)$$

It is assumed that Equation (4) has a limitation, such as the upper limit of CaCO$_3$ produced by unit cell. In this study, the limitations and interferences in Equation (4) are investigated experimentally. However, there is another problem: the cell viability reduces with age, temperature change (cold and heat shocks), freezing, and freezing–thawing cycles [29,35]. The drainage of the supernatant containing bacteria cells in centrifuging process causes a reduction in viable cells. These facts require a simple test method for obtaining OD values that can be a representative of the number of viable cells. Note that the conventional OD$_{600}$ cannot reflect the viable quantity of cells of BCS to be used.

2.3. Preparation of Sample for Obtaining Standard Precipitation Rate

The relationship between the OD value and precipitation rate for CaCO$_3$ was investigated based on the hypothesis described earlier. First, the OD of the fresh culture solution of A10 was adjusted to 1.0. The number of cells, i.e., colony forming unit (CFU), was measured using the agar culture technique. The cell viability corresponding to this CFU was assumed to be 100%. The solution was divided into three 100 mL flasks. The solution in the first flask was immediately used as fresh samples with 100% viability. Second flask was stored in the refrigerator at 4 $^\circ$C for one month, and the CFU was measured. Comparing the CFU value with that of 100% viability, the viability was estimated as 45%. Assuming the linear relationship between cell viability and OD value, the OD of the second solution was determined as 0.45. The third flask was kept at room temperature, 25 $^\circ$C, for one month. Like the second solution, CFU, viability, and OD were determined in order. As a result, the viability and OD for the third solution were 9% and 0.09, respectively. These three types of solutions were used for the following precipitation tests as the undiluted BCSs, respectively.

The samples for precipitation tests were prepared as follows.

For preparation of the OD value, the prescribed OD values were arranged in 10 mL BCS in test tubes, as indicated in Table 1. To carry this out, each 10 mL bio-solution was centrifuged using a centrifuge tube, and the supernatant was removed.
Table 1. Experimental data for determination of a standard precipitation curve.

| Flask No. | Temp. (°C) | Age | Spec. No. | Cell Viability (%) | OD  | CPR (M) |
|-----------|------------|-----|-----------|-------------------|-----|---------|
| 1         | 25         | 1 day | 1         | 100               | 1.0 | 1.06    |
|           | 2          |       | 2         | 0.75              | 1.04|         |
|           | 3          |       | 3         | 0.59              | 1.05|         |
|           | 4          |       | 4         | 0.37              | 1.00|         |
|           | 5          |       | 5         | 0.25              | 1.00|         |
|           | 6          |       | 6         | 0.10              | 0.66|         |
| 2         | 4          | 1 month | 7         | 45                | 0.45| —       |
|           |            |       | 8         | 0.36              | 1.06|         |
|           |            |       | 9         | 0.27              | 1.00|         |
|           |            |       | 10        | 0.18              | 0.93|         |
|           |            |       | 11        | 0.09              | 0.51|         |
| 3         | 25         | 1 month | 12        | 9                 | 0.09| 0.56    |
|           |            |       | 13        | 0.072             | 0.53|         |
|           |            |       | 14        | 0.054             | 0.42|         |
|           |            |       | 15        | 0.036             | 0.29|         |
|           |            |       | 16        | 0.018             | 0.15|         |

For the reactive solution,

1. Prescribed quantities of reagents, i.e., urea, calcium chloride, etc., were prepared;
2. Ten milliliter reactive solutions were added to each test tube containing the centrifuged microbes. Note that the volume of microbes is negligible, as the centrifugal concentration factor by volume is approximately 250–300;
3. The test tubes were left for 24 h, and the precipitated CaCO$_3$ was measured by dividing into suspended particles and adsorbed carbonates.

The results are indicated in terms of the carbonate precipitation rate (CPR) in Table 1 and Figure 5. It is seen that the quality of microbe changes with temperature, age, or dilution. In Table 1, it is seen that the precipitates for some samples exceed 1.0 M. This is possibly due to the precipitation of hydrated amorphous calcium carbonates (ACC) [30]. This unique relationship was obtained for the OD values under the controlled cell viability as the best performance. If the quality of microbes vary, the plots might deviate widely.

![Figure 5. Relationship between precipitation rates and OD related to cell viability, showing a standard precipitation curve.](image-url)
In this study, the CPR of the unspecified BCS was examined in terms of the estimated OD, which was calculated using only dilution effect. The line shown is defined as the standard precipitation curve, which is given as follows:

\[
\text{CaCO}_3 = \text{CPR} = 8.460 \text{ OD} - 17.633 (\text{OD})^2 \quad (M)
\]  

(5)

Equation (5) is a quadratic function of OD showing a convex at the top, and is illustrated in Figure 6. It seems that the first order term on the right side of Equation (5) shows the \( \text{CaCO}_3 \) precipitation rate with somewhat inherent properties of microbes, such as the activity of the unit cell. On the other hand, the second order term may be related to the interaction between microbes and \( \text{Ca}^{2+} \), and the competition of microbes. It may cause the delayed reaction or constrained reaction.

![Image of Equation (5)](image)

**Figure 6.** Illustration of Equation (5).

### 2.4. Calibration Curves

Considering the reduction in cell viability, the OD value is assumed as follows:

\[
\text{OD} = \text{Rcv} \text{ OD}^* 
\]  

(6)

where \( \text{Rcv} \) is the conversion factor and \( \text{OD}^* \) is the tentative optical density associated with the dilution factor used for sample preparation. The OD values of the test samples can be varied by diluting an undiluted BCS, which concerns the number of viable cells. Using the dilution technique, the CPR is measured in terms of various \( \text{OD}^* \) values. Introducing Equation (6) into Equation (5), the CPR as a function of \( \text{OD}^* \) and \( \text{Rcv} \) is as follows:

\[
\text{CaCO}_3 = 8.460 \text{ Rcv} \text{ OD}^* - 17.633 (\text{Rcv} \text{ OD}^*)^2 \quad (M)
\]  

(7)

In Equation (7), if the induced \( \text{CaCO}_3 \) is given, the value of \( \text{Rcv} \text{ OD}^* \) can be determined using the quadric formula. Therefore, if the \( \text{OD}^* \) is given, the corresponding \( \text{Rcv} \) is also determined. Experimentally, if the relationship between CPR and \( \text{OD}^* \) is obtained, the corresponding \( \text{Rcv} \) is determined by fitting the experimental data with calibration curves. The calibration curves consist of CPR-\( \text{OD}^* \) relationships with a constant \( \text{Rcv} \) value, as shown in Figure 7. In general, the \( \text{OD}^* \) value is given as an arbitrary experimental condition, provided that, in this study, it is defined according to a certain rule. The rule is needed to make \( \text{Rcv} \) meaningful, and is defined later.
when 100 mL BCS with 1.28 OD* is required, the volume of microbes $V_{mc}$ is needed as

$$V_{mc} = \frac{(100) (1.28)}{(4.0) (250)} = 0.128 \text{ (mL)}$$

(9)

where the initial optical density $OD_i$ and dilution factor $D_f$ (=concentration factor $C_f$) are known, and the volume of BCS $V_{bc}$ and desired tentative $OD^*$ are given. For example, when 100 mL BCS with 1.28 $OD^*$ is required, the volume of microbes $V_{mc}$ is needed as

$$V_{mc} = \frac{(V_{bc} \ OD^*)}{(OD_i \ D_f)}$$

(8)

Since the various curves with different $R_{cv}$ values are derived from the origin of $OD^*$ and CaCO$_3$ axes, the curves do not cross. Therefore, any point in the curve’s area on the coordinate has its own $R_{cv}$ value.

2.5. Materials

Materials used were microbes A10 strains, urea, calcium chloride, ammonia buffer solution, and water as solvent. The optimum blending of these materials is under consideration. Various ages of microbes were used to examine the ageing effects of microbes. The longest frozen time used was 1 year at $-80 \degree C$. The aspects of the samples were assumed as unknown, and the parameters used were the initial optical density of the culture solution of microbes, $OD_i$, and concentration factor, $C_f$.

2.6. Methods

The volume of microbes, $V_{mc}$, to be used was calculated as follows:

$$V_{mc} = \frac{(V_{bc} \ OD^*)}{(OD_i \ D_f)}$$

(8)

where $OD_i$ is usually 4.0, and $C_f$ and $D_f$ are usually around 250, as shown in Equation (9). If $V_{mc}$ was a very small amount, it was converted to the mass using a density of the concentrated microbes, i.e., $1.07 \text{ g/cm}^3$. Then, the microbes were weighed.

3. Results and Discussion

3.1. CPR Controlled by Ca$^{2+}$

If the BC solution was prepared with a relatively large density of viable cells, the CPR can be controlled by the Ca$^{2+}$ used. The hydrolysis of urea is completed by the microbe’s enzymes, and the carbonate ions required will be produced. Accordingly, Reactions (1) and (2) occur, provided that the Ca$^{2+}$ is arbitrary. This reaction process is easily examined using BCS. This is confirmed by the controlled precipitation by Ca$^{2+}$ as follows.

A hundred milliliters of BCS prepared by Equation (8) can be used for more than ten specimens with 10 mL. For the precipitation tests, the combination of Ca$^{2+}$ and $OD^*$ is selective. For a typical example, the BC solution with 1.0 M Ca$^{2+}$ and 1.0 $OD$ can be
diluted with dilution factors of 0.7, 0.5, 0.3, and 0.1, respectively. Distilled water or a saline solution was used for dilution. In the case of this dilution, the prepared BC solutions have the same Ca\textsuperscript{2+} /OD* ratio, i.e., 0.78 M, and both Ca\textsuperscript{2+} and OD* decrease with the similar dilution factors.

The used Ca\textsuperscript{2+}-OD* and the induced CaCO\textsubscript{3}-OD* relationships are shown in Figure 8, which clearly indicates that the two relationships are parallel straight lines. This indicates that the CPRs were controlled by Ca\textsuperscript{2+}. In Figure 8, the molar concentrations of the induced CaCO\textsubscript{3} are slightly higher than that of Ca\textsuperscript{2+}. This is due to the hydration of carbonates, as described earlier. If the carbonate precipitations for all samples occurred under control of Ca\textsuperscript{2+}, it means that the number of cells was sufficient to complete the reactions. Accordingly, the Rcv values for the BCSs cannot be specified, but the minimum Rcv is evaluated, as shown in the Figure. It indicates that, if BCS has a Rcv value greater than 0.18, the precipitation result is like Figure 8. Thus, in those cases, a proper Rcv cannot be evaluated. However, in the design of the blending of BCS, the minimum value of Rcv provides significant information.

![Figure 8](image_url)

**Figure 8.** CPR controlled by Ca\textsuperscript{2+} levels, which shows the fact that the Ca\textsuperscript{2+} used was all consumed, and the calibration curve with minimum Rcv required for obtaining the results of precipitation tests (at 24 h).

In Figure 9, microscopic photos for 1.0, 0.5, 0.3, and 0.1 M CaCO\textsubscript{3} corresponding to Figure 8 are demonstrated. The calcium carbonates fully precipitated on the inner walls of the test tubes were dependent on the used levels of Ca\textsuperscript{2+}. The precipitation time was approximately 24 h. It is noted that, in general, the minerals precipitated are not always the same, but vary with temperature, pH, reaction rates, etc. In general, minerals induced by microbial precipitation are hydrated amorphous calcite (ACC), calcite, aragonite, vaterite, etc. [36,37], whose size and shape (crystals) often vary with time. However, it was reported that microbes and urease do not affect the morphology of carbonates to be induced [38]. This may be true because microbes become nuclei for carbonate formation and urease only has a role as an enzyme for catalysis in the hydrolysis of urea. There is no direct evidence that microbes and urease affect the morphology of carbonates.
It was confirmed that these particles are dissolved by 3N HCl. The carbonate content was measured by mass analysis using HCl. The particles were also observed by a microscope as amorphous or crystalline calcites, and vaterite. Calcite is known to exhibit many morphological aspects, i.e., many crystalline shapes.

3.2. Effects of Increasing Ca\(^{2+}\)/OD*  
For the next tests, the Ca\(^{2+}\)/OD* ratio was increased to 1.72. The BCSs used were diluted with factors of 0.7, 0.5, 0.3, and 0.1. The results of the precipitation tests are shown in Figure 10. It shows that, in comparison to Figure 8, the higher precipitation occurred for the higher Ca\(^{2+}\)/OD* ratio. However, the CPR at the point P for OD* = 0.58 was lower than expected. It is considered that the precipitation indicated by point P at OD* = 0.58 was delayed because the number of cells was not enough under 1.0 M Ca\(^{2+}\). This is examined later.

The CPRs for the OD* lower than 0.4 are expressed by a straight line, so the precipitation was probably controlled by the Ca\(^{2+}\) used. Therefore, the calibration curve for this
case can be fitted by \( R_{cv} = 0.3 \), as shown in Figure 10. It is a key point in this case that the calibration curve passes between the \( \text{Ca}^{2+} \) used and the lower measured CPR at a given OD* value. Note that the OD is evaluated by 0.3 OD* using \( R_{cv} = 0.3 \) because of the definition in Equation (6). Therefore, if 0.4 OD*, i.e., 0.12 OD is used, it is found that approximately 0.7 M \( \text{CaCO}_3 \) is obtained.

To see this phenomenon in detail, the \( \text{Ca}^{2+}/\text{OD}^* \) ratio was increased again. The results of the precipitation tests at a \( \text{Ca}^{2+}/\text{OD}^* \) ratio of 2.5 is indicated in Figure 11. It shows the relatively high CPR below OD* = 0.2 and the inhibitory effect at a higher OD*. The CPR at OD* = 0.4 was only 12% of the applied \( \text{Ca}^{2+} (=1.0 \text{ M}) \). This decreasing effect is possibly due to the shape of the calibration curve depending on the second order term of OD in Equation (5) or (7). In fact, this second order term can be understood as the delayed reaction from 24 h for the growth of mineral grains.

![Figure 11](image-url)

**Figure 11.** Relatively high CPRs at lower OD* and over-loading effects at higher OD* under the condition of relatively high \( \text{Ca}^{2+}/\text{OD}^* \).

The effect of the delayed reaction on the engineering properties of biocement has not been well studied. It may give good results in terms of cementation effects. However, considering the various cases in the lab and field conditions, the inhibitory phenomena of CPR with \( \text{Ca}^{2+} \) is defined as the over-loading effect and is distinguished from the standard precipitation rate in this study.

In Figure 11, the over-loading effect also seems to occur at OD* = 0.28. Thus, it is found that, with increasing \( \text{Ca}^{2+}/\text{OD}^* \), the CPR increases at a lower OD*, whereas the over-loading effect occurs at a higher OD*.

### 3.3. Over-Loading Effect

In Figure 12, the CPRs for the highest OD* values at various \( \text{Ca}^{2+}/\text{OD}^* \) ratios are indicated as the over-loading 1M \( \text{Ca}^{2+} \) line. It indicates that higher \( \text{Ca}^{2+} \) causes the over-loading effect. On the other hand, for the lower OD*, a relatively high CPR corresponding to the \( \text{Ca}^{2+} \) used is obtained.
Over-loading effects due to relatively high Ca\(^{2+}\).

3.4. Over-loading Effects for Low OD Values

If the OD value is unknown, it is not known if the over-loading effect occurs. Another example of the over-loading effect is obtained using different samples, as shown in Figure 13. The test conditions are presented in terms of Ca\(^{2+}\), OD*, ODi, and Df for the undiluted BCS.

The undiluted OD* value used is already adjusted to the OD value. Therefore, the diluted OD* values are also equivalent to the OD value. The calculated Ca\(^{2+}\)/OD ratios for the specimens ranged from 7.7 M to 83 M. The result is very similar to that shown in Figure 12. Thus, the results assured the repeatability of the over-loading effect. If the over-loading effect is known, lower Ca\(^{2+}\) levels are selected, i.e., the CPR of the standard precipitation curve can be obtained in 24 h.

![Figure 12](image-url)  
**Figure 12.** Over-loading effects due to relatively high Ca\(^{2+}\).

![Figure 13](image-url)  
**Figure 13.** Over-loading effect under high Ca\(^{2+}\)/OD* ratios, where OD* \(\equiv\) OD.

3.5. Intersection of Constant Ca\(^{2+}\) Line and Calibration Curve

It was described earlier that the CPR cannot exceed the Ca\(^{2+}\) line, except for the occurrence of hydrated amorphous carbonate (ACC). Figure 14 shows the case where the constant Ca\(^{2+}\) line intersects with the calibration curve. The CPR line at 24 h increases along the calibration curve (Rcv of 0.077) from the lower OD* value until it is controlled by Ca\(^{2+}\), as shown in Figure 14. Near the Ca\(^{2+}\) line, the CPR is controlled by the amount of Ca\(^{2+}\). If ACC forms, the CPR may exceed the Ca\(^{2+}\) line because of the effect of hydration, as was described. It is noted that the OD values are estimated by Equation (6), i.e., Rcv OD*.
ACC forms, the CPR may exceed the Ca$^{2+}$ line because of the effect of hydration, as was described. It is noted that the OD values are estimated by Equation (6), i.e., $R_{cv}$ OD is greater than that of CaCO$_3$.

To investigate time dependence on the over-loading effect, the precipitation rates were measured at a longer elapsed time, under the condition of 1.0 M Ca$^{2+}$. The test conditions used are indicated in Table 2. Note that the No. 1 specimen consisted of five aliquots, whereas four aliquots were used for No. 2. The CaCO$_3$ measurements of each sample were made at a different date. Herein, CPR is defined for CaCO$_3$ precipitation at any elapsed time. However, it does not mean the precipitation rate, but the amount.

3.6. Example of Translation from CPR-OD* to CPR-OD Relations

If the $R_{cv}$ value is obtained, the corresponding OD value is determined from Equation (6). The conversion from OD* to OD or OD to OD* is shown in Figure 15. The horizontal axis is used for both the OD and OD* values. As can be seen, the precipitation rates were converted by the coordinate conversion from OD* to OD. The agreement between the standard curve and the converted points of the precipitation rates indicates that the value of the selected $R_{cv}$ was appropriate. It shows that, if the molar concentration of used Ca$^{2+}$ is greater than that of CaCO$_3$ given by the standard curve, the actual precipitation of CaCO$_3$ will decrease from the value predicted by the calibration, as shown in the figure. This is discussed later.

3.7. CPR after Over-Loading Effects

To investigate time dependence on the over-loading effect, the precipitation rates were measured at a longer elapsed time, under the condition of 1.0 M Ca$^{2+}$. The test conditions used are indicated in Table 2. Note that the No. 1 specimen consisted of five aliquots, whereas four aliquots were used for No. 2. The CaCO$_3$ measurements of each sample were made at a different date. Herein, CPR is defined for CaCO$_3$ precipitation at any elapsed time. However, it does not mean the precipitation rate, but the amount.
The precipitated amounts measured are shown in Figure 16. The precipitation seems to end at certain Ca\(^{2+}\) values. For the No. 1 specimen, the precipitation rate at 24 h was approximately 0.25 M, which was approximately a half of the standard value. The CPR increased by approximately 0.5 M at 4 days. After 18 days, the precipitation decreased to approximately 0.45 M but slightly increased again, as shown in Figure 16.

For the No. 2 specimen (four aliquots), the precipitation rate was approximately 0.63 M at 24 h. The value is smaller than the standard at 24 h. The precipitation exceeded 1.0 M 4 days after and then decreased 12 days after. However, it seems to be slightly increased again, as shown in Figure 16. The decrease in volume of CaCO\(_3\) after a few days is common. It may be due to the metamorphism of carbonate crystals, which may be due to the dehydration of hydrated carbonates. It is considered that such a slight change in CaCO\(_3\) precipitation may be due to the combined effects of hydration, dehydration, metamorphism, delayed formation of carbonates, etc.

For specimens No.1 with an OD value of 0.068, the 24 h precipitation was approximately half that of the standard curve at 24 h, but exceeded the standard curve after 4 days. The measured CaCO\(_3\) after 18 days is plotted as shown in Figure 17. For the No. 2 specimen, the precipitation at 0.12 OD exceeded the standard curve (24 h) on the second day. After 3 days, the CaCO\(_3\) production reached almost 1.0 M. The measured value of 1.03 M after 23 days is plotted, as shown in Figure 17.

From the observation of a long time CaCO\(_3\) precipitation, it was found that the amount of CaCO\(_3\) exceeded the standard value. Therefore, the second term of the right side in Equation (5) is considered to be a delayed effect due to the over-loading effect of Ca\(^{2+}\). It means that, given the unlimited time, the second term limited by 24 h can be ignored. As a result, the standard curve without time limitation is represented by a linear relationship of the following equation, leaving only the first term.

\[
\text{CaCO}_3 = 8.46 \text{OD (M)}
\] (10)

Equation (10) indicates that the maximum amount of precipitation is proportional only to the OD value (representative of the viable cell number). It was a hypothetical starting point in this study on the blending design of bio-cement.

| No. | OD\(^*\) | Rcv | OD | Ca\(^{2+}\) | Ca\(^{2+}\)/OD\(^*\) | Ca\(^{2+}\)/OD |
|-----|---------|-----|----|------------|-----------------|---------------|
| 1   | 0.180   | 0.38| 0.068| 1.0        | 5.56            | 14.62         |
| 2   | 0.310   | 0.38| 0.118| 1.0        | 3.23            | 8.47          |

Table 2. Precipitation tests after over-loading effects.
Figure 17. Precipitation rates after the over-loading effect of Ca\(^{2+}\), showing that ultimate precipitation of carbonates can be presented as a linear function of OD value.

4. Conclusions

The carbonate precipitation induced by ureolytic microbes was investigated to develop the optimum engineered blending method of BCS. Based on the hypothesis that CPR is governed by OD with viable cells, the unique standard precipitation curve for A10 strains, a basic calibration curve was determined. To apply the unique standard precipitation curve, the tentative OD* value was defined. The OD* is associated with OD = Rcv OD*, where Rcv is defined as the conversion factor concerned with cell viability. Herein, the CPR-OD* relationship can be converted to the CPR-OD relationship in terms of Rcv. This concept was experimentally examined. As a result, it was found that Rcv is predictable compared to the unique standard CPR-OD and the experimental CPR-OD* relationship. It was also found that the OD value was predictable using Rcv and OD*. With this, it became possible to evaluate the quality of microbes uniformly, without measuring OD\(_{600}\). Furthermore, multiple achievements were obtained through this study. The main achievements are summarized as follows.

1. The microbial strain (NO-A10) is mainly examined in this study. Note that the quantitative results presented are dependent on microbes and strains;
2. The CPR desired can be blended in terms of OD and Ca\(^{2+}\);
3. CPR is governed by OD or Ca\(^{2+}\). If the OD is dominant, the CPR (24 h) is expressed as CPR = 8460 OD − 17.633 OD\(^2\);
4. If Ca\(^{2+}\) is dominant, CPR = a CaCO\(_3\) (M/24h) for the case of NO-A10 strains and a M Ca\(^{2+}\);
5. If Ca\(^{2+}\) is high and OD is relatively low, the over-loading effect, which causes exceptionally low CPR, occurs. However, the carbonate precipitation increases according to the following extension of the concept with time;
6. Extending the concept of the unique standard precipitation rate regarding time, the CPR-OD relationship was simply expressed by CPR = l OD, where l was 8.46 M for NO-A10 strains. Note that the CPR does not necessarily mean CaCO\(_3\) precipitated for 24 h.

The study is based on the BCS, but not on the soil-BCS system. However, the microscopic photos presented in this paper indicate the carbonate precipitated on the surfaces of glass like minerals. The effect of the adsorption between microbes and soil particles may influence the Ca\(^{2+}\) and OD* values. It may depend on the soil depth or flow distance associated with the injection of the solution. Accordingly, the adsorption properties and behavior of Ca\(^{2+}\) and microbes on soil particles can be important for future study. The most important contribution in this research is the determination that CPR is controlled by the terms of Rcv, OD*, and Ca\(^{2+}\).
Author Contributions: Conceptualization, M.F. and Z.L.; methodology, M.F. and Z.L.; software, M.F. and K.E.; validation, Z.L., C.N.M., Y.F. and K.E.; formal analysis, M.F.; investigation, M.F., Y.F. and K.E.; resources, M.F.; data curation, not applicable.; writing—original draft preparation, M.F.; writing—review and editing, M.F., Z.L. and C.N.M.; visualization, K.E.; supervision, M.F.; project administration, M.F.; funding acquisition, not applicable. All authors have read and agreed to the published version of the manuscript.

Funding: This research received no external funding.

Institutional Review Board Statement: Not applicable.

Informed Consent Statement: Not applicable.

Data Availability Statement: Data available on request.

Acknowledgments: Bacteria NO-A10 and -N10 strains used for the experiment were supplied by Life Engineering, Co., Japan.

Conflicts of Interest: The authors declare no conflict of interest.

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