



Article Optimization of Bio-Foamed Concrete Brick Strength via Bacteria Based Self-Healing and Bio-Sequestration of CO₂

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1. Introduction

Global carbon dioxide (CO_2) emissions have increased in recent decades, in line with increases in anthropogenic activities. Therefore, a great deal of research has been conducted to reduce the impacts of the catastrophic environmental issues due to CO_2 emission such as global warming, rising sea levels, and climate change [1]. Fossil fuel combustion and cement manufacturing are the biggest contributors to CO_2 emissions, representing around 88% [2]. The process of producing one ton of cement emits around 900 to 1000 kg of CO_2 due to the energy required to burn limestone [3,4]. Between 2005 and 2015, cement production increased worldwide by 79.5%, i.e., from 2284 to 4100 Mt/yr [2,4]. For this reason, various studies on concrete technology have focused on reducing cement production by using alternative materials [5,6]. However, the demand and production of cement continue to increase, resulting in increased global emissions of CO_2 . Furthermore, most of the replacement materials used in concrete reduce its strength.



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Copyright: © 2021 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (https:// creativecommons.org/licenses/by/ 4.0/). Due to the ability of bacteria to improve the physical and mechanical properties, particularly compressive strength, of concrete, so-called bio-concrete has become popular worldwide [7,8]. Many researchers have realized the potential of bacteria to increase the strength of bio-concrete through a self-healing process. Bacteria are typically used in different types of concrete such as normal, fly ash, and rice husk ash. Different types of bacteria have been used for this purpose, such as *Bacillus pasteurii*, *Pseudomonas aeruginosa*, *B. alkalinitrilicus*, *B. sphaericus*, *B. subtilis*, *Enterococcus faecalis*, *Shewanella* sp., *S. pasteurii* and *Ureolytic* [7]. However, the use of bacteria in foamed concrete has not been reported in previous studies, according to the latest update of the Scopus database in 2019. Bacteria have the potential to be more effective in foamed concrete than in normal concrete due to the high level of porosity and availability of oxygen formed by the foaming agent in the former [9].

Challenges to the use of bacteria in bio-concrete, such as overcoming the highly alkaline and anaerobic conditions [10], have given rise to the use of silica gel, capsules, and adaptation media [11]. Furthermore, the ability of bacteria to produce enzymes such as urease or carbonic anhydrase (CA) is critical to the precipitation of calcium carbonate (CaCO₃) on the surface, which results in self-healing of the concrete pores. The healing of bio-concrete pores occurs due to the reaction of urease or CA enzymes with the available cement-based calcium ions (Ca⁺) on the surface of the bacteria. Typically, the reaction results in the precipitation of CaCO₃, which improves the compressive strength [12,13]. The reaction of urease and CA enzymes enhances the natural carbonation reaction of bio-concrete with bicarbonate (CO₃²⁻), which further reacts with the calcium in cement to increase CaCO₃ precipitation and accelerates carbonation [14,15].

According to Rafat Siddique (2011), bacteria concentration is one of the most critical factors affecting the performance and compressive strength of bio-concrete [16]. While increasing the bacteria concentration increases compressive strength, exceeding the optimum value can have a negative effect [17]. Furthermore, higher cell concentrations disrupt the integrity of the matrix due to excessive microbial activity. The various factors that affect the compressive strength of bio-concrete may be categorized the solid physical properties, material chemical properties, and external conditions of the environment [18]. Therefore, the effect of bacteria and other factors must be investigated to optimize the compressive strength of bio-concrete.

This research aims to use accelerated CO_2 production via *Bacillus tequilensis* (*B. tequilensis*) to optimize the strength of bio-foamed concrete brick B-FCB, using a 2^k factorial design and response surface methodology (RSM) using the Minitab 18 software to complete the analysis. *B. tequilensis* concentration (B), the density of B-FCB (D), CO_2 concentration (CO₂) and temperature (T) of curing in the chamber were carefully investigated before optimization of the compressive strength of B-FCB.

2. Materials and Methods

In this study, the materials used to prepare the foamed concrete were cement, sand, water and a foam agent, while *B. tequilensis* was added to produce a new type of concrete, namely, bio-foamed concrete. The two methods used to optimize the compressive strength of B-FCB were 2^k factorial design and RSM. Details regarding the materials and methods used in this study are provided in the following subsections.

2.1. Materials

2.1.1. Cement

Ordinary Portland cement (OPC) manufactured by Cement Industries of Malaysia Berhad (CIMA), type I, MS 522 according to American society for testing and materials (ASTM), was used in this study. The composition and specifications of the OPC are defined in BS 197-1:2000, as shown in Table 1. The quantity of cement was adjusted according to the density used in the foamed concrete mixture with and without *B. tequilensis*.

Chemical Compound	Concentration (%)
SiO ₂	20.6
Al_2O_3	5.4
Fe ₂ O ₃	4.2
SO_3	2.2
K ₂ O	0.6
CaO	64.8
MgO	2.2

Table 1. Summary of the chemical composition of the OPC.

2.1.2. Sand

River sand was sieved using a plate passing 1 mm, according to IS 383:1970 [19,20]. The particle size distribution is presented in Figure 1. The sieved sand was placed in an oven at 100 °C for 24 h to dry and to remove any microorganisms present. It was then cooled at room temperature before use in the concrete mixture.



Figure 1. Grading curve for river sand.

2.1.3. Water

Water was used with a foamed concrete mix to produce the FCB. The same water was used with *B. tequilensis* added in powdered form to produce B-FCB; the water containing *B. tequilensis* was agitated to ensure that the powder was evenly distributed.

2.1.4. Bacteria

The *B. tequilensis* was isolated from five samples of cement kiln dust (CKD), which has an extreme pH value and provides anaerobic conditions. The isolation of *B. tequilensis* was subjected to several tests, namely, CA and urease assays, growth in theioglycollate, a candle jar test and growth in a bio-foamed concrete medium [21]. Then, the most resilient bacteria were isolated from the five samples, and tested for the following properties: ability to produce CA and urease enzymes, facultative anaerobic, and capable of growth in high concentration of CO_2 and in a bio-foamed concrete simulation medium. After that, a powdered form of *B. tequilensis* was produced to control its concentration in the concrete mixture: a bacteria pellet was placed into a freeze dryer with the following settings: $-40 \,^{\circ}C$ and 0.133 mbar of pressure for 96 h [21].

In this research *B. tequilensis* was selected as one of the factors used to optimize the compressive strength of B-FCB. The concentration of *B. tequilensis* was suggested by the 2^k factorial and RSM methods shown in Table 2.

Level	Bacteria Concentration (Cell/mL)	<i>B. tequilensis</i> Concentration (g/L)	Value Used in (RSM)
Low (-1)	$3 imes 10^5$	0.001	5
Centre (0)	$3 imes 10^6$	0.01	6
High (1)	$3 imes 10^7$	0.1	7

Table 2. *B. tequilensis* concentration as the main factor in (RSM).

2.1.5. Foam Agent

A synthetic type CF 500 foaming agent with a density between 45–65 kg/m³ and an expansion ratio of 1:20 was used in this study to produce air bubbles in the foamed concrete mixtures with and without bacteria. The foaming agent was diluted with water at a ratio of 1:20 and aerated to a density of 65 kg/m³, according to ASTM C796 [20,22].

2.2. Methods

2.2.1. Design of Experiments

In this study, the optimization of the compressive strength of B-FCB involved two stages. The first stage, called screening experiments, was carried out according to the 2^{k} factorial design method, consisting of 11 runs with three center runs added for the curvature test analysis. The 2^k factorial design is the most commonly used technique in design of experiments (DOE), and was very helpful in determining the important factors of the experiment. Due to time and funding constraints, the 2^k factorial design was prioritized. The experiment was run based on the 2^3 full factorial design and was performed to study the effects of the four process parameters shown in Table 3. The low- and high-level settings of several factors, particularly the *B. tequilensis* concentration (B), density of the concrete (D), temperature (T) and CO_2 concentration (CO_2), were input and analyzed using Minitab software (version 18, Pennsylvania State University, State College, PA, USA) and analyzed. A reasonable range for each factor was selected according to previous research findings [2], as presented in Table 3. The next stage used to optimize the compressive strength of B-FCB by RSM analysis involved adding eight axial and two additional runs at the center points. Hence, a total of 21 experiments were performed, comprising eight factorial runs, eight axial runs, and five center runs. This certainly increased the accuracy levels of the empirical models deduced from the Analysis of Variance (ANOVA) data, as shown in Equation (1).

$$Y = \beta_0 + \beta_1 x_1 + \beta_2 x_2 + \beta_{11} x_1^2 + \beta_{22} x_2^2 + \beta_{12} x_1 x_2 \tag{1}$$

where

Y is the response (output function)

 x_1, x_2, \ldots, x_k are the factors (Input variables)

 β_0 is the grand average of all observations

 β_1 is half of the *A* effect

 β_2 is half of the *B* effect.

Table 3. Design scheme and levels of the process parameters.

Factor Symbol	D (T In: 1		
	Parameter	Low (-1)	Centre (0)	High (+1)	- Unit
CO ₂	CO ₂ Concentration	10	15	20	(%)
В	<i>B. tequilensis</i> concentration	$3 imes 10^5$	$3 imes 10^6$	$3 imes 10^7$	(cell/mL)
Т	Temperature	27	33.5	40	(°C)
D	Density of concrete	1300	1550	1800	(kg/m^3)

ANOVA was also used to rank the main effects and analyze the interactions between the input factors. ANOVA is considered essential for structured analyses of results in the 2^k design.

2.2.2. Mixture Design

The mixture design of each run was based on the suggested densities and *B. tequilensis* concentration. The mass of the solid materials (cement/sand) was distributed in the ratio of 1:1.35 according to ACI 523.3R, using the trial method of mix design [20]. Details of the experimental runs are shown in Table 4. The first 11 runs were suggested by the 2^k factorial design for the screening stage, whereas the last 10 runs were added to complete the RSM analysis using the Minitab 18 software.

Table 4. Concrete mixture and RSM design for the optimization experiment.

Run No.	Density (kg/m ³)	Cement (kg/m ³)	Fine Sand (kg/m ³)	Water (L/m ³)	Т (°С)	CO ₂ (%)	B (Cell/mL)
1	1300	553.2	746.8	276.6	27	10	$3 imes 10^5$
2	1800	766	1034	383	40	10	$3 imes 10^5$
3	1300	553.2	746.8	276.6	40	20	$3 imes 10^5$
4	1800	766	1034	383	27	20	$3 imes 10^5$
5	1300	553.2	746.8	276.6	40	10	$3 imes 10^7$
6	1800	766	1034	383	27	10	$3 imes 10^7$
7	1300	553.2	746.8	276.6	27	20	$3 imes 10^7$
8	1800	766	1034	383	40	20	$3 imes 10^7$
9	1550	659.5	890.4	329.7	33.5	15	$3 imes 10^6$
10	1550	659.5	890.4	329.7	33.5	15	$3 imes 10^6$
11	1550	659.5	890.4	329.7	33.5	15	$3 imes 10^6$
12	1550	659.5	890.4	329.7	33.5	10	$3 imes 10^6$
13	1550	659.5	890.4	329.7	33.5	20	$3 imes 10^6$
14	1550	659.5	890.4	329.7	33.5	15	$3 imes 10^5$
15	1550	659.5	890.4	329.7	33.5	15	$3 imes 10^7$
16	1550	659.5	890.4	329.7	27.0	15	$3 imes 10^6$
17	1550	659.5	890.4	329.7	40.0	15	$3 imes 10^6$
18	1300	553.2	746.8	276.6	33.5	15	$3 imes 10^6$
19	1800	766	1034	383	33.5	15	$3 imes 10^6$
20	1550	659.5	890.4	329.7	33.5	15	$3 imes 10^6$
21	1550	659.5	890.4	329.7	33.5	15	$3 imes 10^6$

2.2.3. Specimen Preparation and Curing

FCB and B-FCB specimens were prepared with the following dimensions: 215 mm \times 100 mm \times 75 mm. The total number of specimens for each run was six, comprising three 3 without (as controls) and three with *B. tequilensis*. The specimens were demolded for 24 h before oven drying at 50 °C for 72 h to avoid water evaporation, which has the potential cause chemical reactions with the available CO₂ in the chamber, resulting in a decrease in the CO₂ concentration during the process. The FCB and B-FCB specimens were subjected to curing in the chamber under different run conditions (T and CO₂) for 28 days, as suggested by the 2^k factorial and RSM methods, as shown in Table 4.

3. Results and Discussion

3.1. Screening Stage Analysis (2^k Factorial Design)

The results of the compressive strength experiments during screening are presented in Table 5. For one replicate design, the internal error could not be estimated. Therefore, all high-order interactions were neglected and the mean squares of the omitted factors were combined to estimate the error. The percentage of R^2 was 96.08%, which reflects the highly significant effect of these factors on the compressive strength of B-FCB. This proved that the experimental results could be reproduced with high repeatability and accuracy.

Std. Run No	Input Variables					
5tu: Kuir 10. –		Uncoded Value				
	Density (kg/m ³)	Bacteria (Cell/mL)	Temperature (°C)	CO ₂ (%)	Compressive Strength of B-FCB (MPa)	
1	1300	$3 imes 10^5$	27	10	0.90	
2	1800	$3 imes 10^5$	27	20	7.23	
3	1800	$3 imes 10^7$	27	10	8.38	
4	1300	$3 imes 10^7$	27	20	1.02	
5	1800	$3 imes 10^5$	40	10	6.72	
6	1300	$3 imes 10^5$	40	20	0.78	
7	1300	$3 imes 10^7$	40	10	0.61	
8	1800	$3 imes 10^7$	40	20	6.13	
9	1550	$3 imes 10^6$	33.5	15	2.72	
10	1550	$3 imes 10^6$	33.5	15	2.63	
11	1550	$3 imes 10^6$	33.5	15	2.60	

Table 5. Compressive strengths of all samples in the screening stage.

The results of the screening stage presented in Tables 5 and 6 indicate that all the selected factors significantly influenced the response; as such, they were retained for the RSM analysis. The discussion of the results is divided into two parts to assist in the interpretation of the ANOVA of the DOE, optimization, and modelling of the compressive strength of B-FCB.

Table 6. ANOVA analysis of the compressive strength of B-FCB.

Source	DF	Adj SS	Adj MS	F-Value	<i>p</i> -Value	Effect	Ranking
Model	8	84.964	10.620	2723.220	0.000	-	-
Linear	4	79.991	19.997	5127.640	0.000	-	-
CO ₂	1	0.316	0.316	81.030	0.012	-0.397	5
В	1	0.017	0.017	4.390	0.171	0.092	7
Т	1	1.470	1.470	377.080	0.003	-0.857	2
D	1	78.187	78.187	20,048.080	0.000	6.252	1
2-Way Interactions	3	1.062	0.354	90.840	0.011	-	-
CO ₂ *B	1	0.546	0.546	140.000	0.007	0.522	3
CO ₂ *T	1	0.070	0.070	18.030	0.051	0.187	6
CO ₂ *T	1	0.446	0.446	114.490	0.009	-0.472	4
Curvature	1	3.910	3.910	1002.660	0.001	-	-
Error	2	0.007	0.004	-	-	-	-
Total	10	84.972	-	-	-	-	-

3.2. Compressive Strength of B-FCB (2^k Factorial Analysis)

3.2.1. ANOVA Analysis, Main Effect and Interaction Plots

This section presents the effects of each factor, i.e., B, D, T and CO₂, on the compressive strength of B-FCB. The compressive strength of each run was analyzed to identify the effects of and interactions among the factors.

The results in Table 6 show the effects of and interactions among the factors on the compressive strength of the B-FCB specimens after 28 days under controlled curing conditions in a chamber. CO₂, T and D significantly influenced the compressive strength of B-FCB, as evidenced by the low *p*-values (<0.05) 0.012, 0.003 and 0.000, respectively). B was insignificant because its *p* > 0.05; however, a significant interaction with CO₂ was observed.

The main plot shown in Figure 2a presents the effect of each factor on the compressive strength of B-FCB. The values of the main factors of D, T, CO_2 and B were 6.2525, 0.8575, 0.3975 and 0.0925, respectively, as presented in Table 6. As observed, factor D had the highest effect on the compressive strength, whereby an increase in D led to greater compressive

strength, as presented in Figure 2a. However, the compressive strength of B-FCB was higher with a lower level of T, i.e., an increase in T had a negative effect on the strength of B-FCB. In addition, an increase in CO₂ concentration during curing in the chamber caused a reduction in compressive strength. According to the ANOVA results, and compared to other factors, *B. tequilensis* alone was less likely to influence the compressive strength.



Figure 2. Results of factorial analysis with curvature test: (**a**) Main effect plot of compressive strength, and (**b**) Interaction plot of compressive strength.

The interactions between CO_2*B , CO_2*T , and CO_2*D had *p*-values < 0.05, as shown in Table 6 and Figure 2b, which means that all interactions had a significant effect on compressive strength; however, the highest interaction was for CO_2*B with P, with effect values of 0.008 and 0.4800, respectively. The strength of B-FCB increased with a high level of B and low level of CO_2 , but decreased with a high level of CO_2 . This finding confirmed the strong relationship between B and CO_2 and the reactions with CA and urease enzymes to accelerate CO_2 sequestration in the B-FCB pores to form $CaCO_3$. The cumulative formation of $CaCO_3$ led to the high compressive strength of B-FCB. In contrast, the increase or decrease of CO_2 with a low level of B did not have a strong effect on compressive strength. The strength of B-FCB improved with a low level of T and CO_2 , while it decreased with higher levels of T and CO_2 . The CO_2*T interaction can be interpreted as follows: a high level of CO_2 during the curing period may restrict the activities of bacteria enzymes, resulting in reduced $CaCO_3$ formation. This finding is supported by the steep line of CO_2 performance shown in Figure 2a. The interaction between CO_2*D shows a drastic change in the compressive strength of B-FCB when D is at a high level and CO_2 at a low level. However, at the same level of D, an increase in CO_2 to 20% lowered the compressive strength. This result indicated that by increasing the CO_2 to 20%, the restriction of enzymatic reactions suppressed the ability of the bacteria to generate CaCO₃ on the surface and heal the pores of B-FCB.

The ANOVA results confirmed that the factors and interactions had a significant effect on the compressive strength of B-FCB. A ranking of each factor is presented in Table 6.

3.2.2. Cube Plot of B-FCB Compressive Strength

The relationship between the factors and response (compressive strength) can be analyzed from the cube plots shown in Figure 3. The predicted values for each combination of factor levels, split by different densities of 1300 kg/m³ and 1800 kg/m³ D, are depicted in the corners of the plots. The maximum compressive strength was 8.38 MPa, which appeared with a high level of D (1800 kg/m³) and B when T and CO₂ were set at low levels. The compressive strength with a low level of D (1300 kg/m³), CO₂ and T was 1.74 when B remained at a high level. It can therefore be concluded that in order to obtain the highest compressive strength, factor B must be set to a high level, while the CO₂ and T should remain low, regardless of the density setting. By applying these parameters, favorable conditions may be established for *B. tequilensis* to become more reactive to CA and urease enzymes; this will accelerate the sequestration of CO₂ from the atmosphere and greatly improve the healing process of B-FCB via the formation of abundant CaCO₃.



Figure 3. Cube plot of compressive strength.

The increase in compressive strength due to an increase in B has already been confirmed in previous research works [23,24]. Typically, an increase of bacteria resulted in an increase in CaCO₃ formation on the surface that healed the pores in the concrete [23]. Therefore, it can be inferred that the healing process in bio-concrete pores leads to an increase in compressive strength. However, higher bacteria concentrations occasionally led to excessive bacteria activity, which disrupted the matrix integrity and ultimately decreased the compressive strength [17].

3.3. Analysis of Response Surface Methodology (RSM) Results

The 2^k factorial analysis proved that all the factors in this study had significant effects on the responses, as discussed in Section 3.2. RSM analysis was applied to optimize the compressive strength of B-FCB after the screening stage. The effect and correlation of the four selected factors were further analyzed using RSM. The Minitab 18 software suggested that 10 more runs be added to the previous eleven runs shown in Table 7, according to central composite design (CCD). Two out of ten runs were assigned to center points, and therefore, the full RSM design comprised five center points. The optimization results for compressive strength are discussed in detail in the following sections.

Std. Run	Input Variables					
No.		Uncoded Value				
	Density (kg/m ³)	Bacteria (Cell/mL)	Temperature (°C)	CO ₂ (%)	Compressive Strength of B-FCB (MPa)	
1	1300	3×10^5	27.0	10	0.9	
2	1800	$3 imes 10^5$	27.0	20	7.23	
3	1800	$3 imes 10^7$	27.0	10	8.38	
4	1300	$3 imes 10^7$	27.0	20	1.02	
5	1800	$3 imes 10^5$	40.0	10	6.72	
6	1300	$3 imes 10^5$	40.0	20	0.78	
7	1300	$3 imes 10^7$	40.0	10	0.61	
8	1800	$3 imes 10^7$	40.0	20	6.13	
9	1550	$3 imes 10^6$	33.5	15	2.72	
10	1550	$3 imes 10^6$	33.5	15	2.63	
11	1550	$3 imes 10^6$	33.5	15	2.60	
12	1550	$3 imes 10^6$	33.5	10	2.55	
13	1550	$3 imes 10^6$	33.5	20	2.56	
14	1550	$3 imes 10^5$	33.5	15	2.16	
15	1550	$3 imes 10^7$	33.5	15	2.31	
16	1550	$3 imes 10^6$	27.0	15	2.54	
17	1550	$3 imes 10^6$	40.0	15	2.12	
18	1300	$3 imes 10^6$	33.5	15	0.81	
19	1800	$3 imes 10^6$	33.5	15	7.2	
20	1550	$3 imes 10^6$	33.5	15	2.54	
21	1550	$3 imes 10^6$	33.5	15	2.59	

Table 7. Results of the full design of RSM for compressive strength and carbonation depth.

3.3.1. Optimisation of Compressive Strength for B-FCB

RSM was used to optimize the compressive strength of B-FCB using residual plot, ANOVA analyses, along with the surface, contour, and optimization plots, as discussed in detail in the following sections.

Residual Plot for B-FCB Compressive Strength

The residual plots of the DOE analysis helped us to evaluate the accuracy of the data. The effects of the nuisance factors were not included in the analysis, except when error measurements were performed. The data distribution must be normal and independent from the zero mean value and constant variance to ensure that the F_o ratio follows the (F) distribution [25]. The distribution of the compressive strength data of B-FCB is present in the residual plots of the normal probability shown in Figure 4.

The presented data indicate that the model meets the assumptions of the analysis. Furthermore, most points follow a straight line, with only three drifting away from the central data. In addition, the residual versus fitted plots presented randomly dispersed data around zero with no clear observable pattern. However, three points exceeded the limit, which confirmed that the error is normal in the results regarding compressive strength. The permissible error to conclude the findings was below 10%, which indicated a high level of accuracy in the data analysis. The fine segregation of the points around the normal probability line demonstrated a precise prediction of the B-FCB strength.



Figure 4. Residual plot for compressive strength of B-FCB.

Analysis of RSM-ANOVA for Compressive Strength of B-FCB

The ANOVA results of compressive strength were analyzed using RSM once all the insignificant factors had been removed, as presented in Table 8.

D, T, and CO₂ were found to be the most significant factors in the analysis, as indicated by p < 0.05, whereas the B was insignificant due to p > 0.05. Furthermore, the square of density (D*D) also showed a highly significant effect, which confirmed that a strong relationship exists between density and compressive strength in B-FCB. In contrast, interactions CO₂*B and CO₂*D were shown to be highly significant, i.e., p < 0.05. As observed, the effect of factor B alone was insignificant. However, the CO₂*B interaction demonstrated a highly significant effect with a *p*-value equal to 0.003, as shown in Table 8. This finding corroborates the theory that the CA and urease enzymes secreted by the bacteria chemically reacted with atmospheric CO₂, resulting in the formation of CaCO₃ on the surface [26,27]. Therefore, it can be inferred that the CO₂*B interaction considerably affected the compressive strength by healing the pores of B-FCB by CaCO₃ precipitation [28].

Table 8. ANOVA results of compressive strength (after backward elimination).

Source	Degrees of Freedom	Sum of Squares	Mean Square	F-Value	<i>p</i> -Value
Model	7	113.265	16.180	413.290	0.000
Linear	4	100.355	25.088	640.820	0.000
CO ₂	1	0.250	0.249	6.380	0.025
В	1	0.027	0.027	0.690	0.421
Т	1	1.482	1.482	37.860	0.000

Source	Degrees of Freedom	Sum of Squares	Mean Square	F-Value	<i>p</i> -Value
D	1	98.596	98.596	2518.370	0.000
Square	1	11.918	11.917	304.400	0.000
D*D	1	11.918	11.917	304.400	0.000
2-Way Interaction	2	0.993	0.496	12.680	0.001
CO ₂ *B	1	0.546	0.546	13.950	0.003
CO ₂ *D	1	0.447	0.446	11.400	0.005
Error	13	0.509	0.039	-	-
Lack-of-Fit	10	0.500	0.050	16.570	0.020
Pure Error	3	0.009	0.003		
Total	20	113.774	-	-	-
Standard deviation = $(R^2 = 99.55\%)$ R ² adjusted = 99.31% Predicted R ² = 98.06%).197865				

Table 8. Cont.

Response Surface and Contour Plots for the Compressive Strength of B-FCB

The response surface plots shown in Figure 5a,c,e depict the effects of the various parameters on the compressive strength of B-FCB. The contour plots played a significant role in the creation of the response surface analysis, which includes layers with different graded colors indicating possible independence of factors to a response, as depicted in Figure 5b,d,f. Both the surface and contour plots graphically depict the relationship between each two process factors and compressive strength, while the other two associated factors are maintained at the center value.



Figure 5. Response surface and contour plots for compressive strength of B-FCB; (**a**,**b**) between B and CO₂, (**c**,**d**) between B and D and (**e**,**f**) between B and T.

Optimization Plots for Compressive Strength of B-FCB

The optimization plot shows how different experimental settings affected the predicted compressive strength of B-FCB. Figure 6 shows that the single desirability (d) and response (y) were 0.77806 and 8.2245, respectively. The red solid lines indicated the values of each factor that led to the highest compressive strength. Meanwhile, the dotted blue lines represent the predicted compressive strengths.



Figure 6. Optimization plot for the compressive strength of B-FCB.

A decrease in CO₂ and T during curing process increased the compressive strength of B-FCB, while the opposite result was observed with higher levels of B and D. The resulting predicted values of D and T were in accordance with data from a previous study, whereby the compressive strength improved when D was set at a high level but T was below 30 °C [31]. The highest predicted response for compressive strength was obtained under the following conditions: CO₂, B, T and D were set at 10%, 30×10^7 cell/mL, 27 °C, and 1800 kg/m³, respectively.

An empirical model was developed via RSM analysis after optimizing the compressive strength of B-FCB. The model derived from the ANOVA results indicated a clear relationship between the independent variables (significant terms) and compressive strength response. The final regression equation in uncoded units of compressive strength is given in Equation (2).

Compressive strength (MPa) = $34.05 + 0.575 \text{ CO}_2 + 0.836 \text{ B} - 0.05923 \text{ T} - 0.05942 \text{ D} + 0.000024 \text{ D*D} - 0.0522 \text{ CO}_2 \text{*B} - 0.000189 \text{ CO}_2 \text{*D}$ (2)

3.3.2. Comparison of Compressive Strength between FCB and B-FCB





The previous statement illustrates that B-FCB has a higher compressive strength compared to FCB at different D levels. In fact, the higher compressive strength was demonstrated by runs with higher levels of D. However, the improvements between FCB and B-FCB were achieved in runs using lower levels of D. Typically, *B. tequilensis* plays a vital role in improving the compressive strength of B-FCB by precipitating high quantities of CaCO₃, which heals pores on its surface. Thus, the pore-healing process was more evident in specimens with low D of B-FCB compared to those with high D. This was because the specimens with low D were characterized by high porosity compared to those with high D.

These findings reveal the trend of compressive strength results, and confirm that compressive strength is affected by numerous factors. Thus, considering only one factor may yield defective FCB and B-FCB. Consequently, in this study, 21 runs were used in RSM to analyze and optimize the compressive strength of B-FCB according to four main variables and their interactions.

4. Microstructure Analysis

A microstructure analysis was conducted by SEM, EDX, and XRD to investigate the healing of specimen pores, weight changes in chemical elements, and comparative precipitation of $CaCO_3$ in B-FCB and FCB.

4.1. Healing Process and Porosity Determination

SEM images were used to demonstrate the healing process and determine the porosity of the FCB and B-FCB specimens, as presented in Figure 8. In general, the healing process of B-FCB pores was clearly present, compared to the FCB pores. The healing process of the pores was due to the formation CaCO₃ on the surface in the B-FCB pores. This occurs due to the chemical reaction between the sequestrated CO₂ and the CA and urease enzymes of *B. tequilensis* during the curing process.



Figure 8. SEM images with specimens with 1300 kg/m³ density of (a) FCB and (b) B-FCB at 28 days.

However, some pores did not fully heal, either due to the high level of porosity in the specimens or inadequate curing time to precipitate more $CaCO_3$ in the pores, as shown in Figure 8. This implies that the effectiveness of the high level of CO_2 sequestrated in B-FCB is responsible for the increase in $CaCO_3$ precipitation that accelerates the healing process. In addition, the relationship between the healing of the B-FCB pores and the high level of CO_2 during curing confirms the ability of the bacteria to accelerate the CO_2 sequestration process. This can be considered a future direction for CO_2 sequestration technology [2]. The results also confirmed that the high compressive strength performance of B-FCB compared to FCB was due to the healing of B-FCB pores. The inference is that the high performance of the compressive strength observed in B-FCB compared to FCB was due to the healing of CaCO_3. Conversely, the availability of CA and urease enzymes in the *B. tequilensis* used in B-FCB helped accelerate CO_2 sequestration and $CaCO_3$

formation on the *B. tequilensis* surface, which gave rise to the healing process observed in Figure 8 [32].

4.2. Elemental Analysis of FCB and B-FCB





Figure 9. Comparative EDX analyses between (a) FCB and (b) B-FCB at 28.

It was also observed that the wt.% of oxygen increased in B-FCB compared to FCB due to the microbial activities, which released oxygen during the precipitation of CaCO₃. In this study, oxygen was also available in both FCB and B-FCB from to the foaming agent used in the concrete mixture. In a study by Zhang [34], the availability of oxygen in concrete was shown to help the bacteria to precipitate CaCO₃. Thus, to provide oxygen for microbial

CaCO₃ precipitation inside the bio-concrete, bacteria and oxygen-releasing compounds were compressed together as the core material of the microcapsule.

The results of the EDX analyses showed that the composition of the primary elements Ca, O and C confirmed the presence of CaCO₃. The increase in C and decrease of Ca in B-FCB confirmed the formation of CaCO₃ on the surface of *B. tequilensis*, which improved its compressive strength [36].

4.3. Crystallinity Analysis of FCB and B-FCB

The XRD technique was used in this study to observe and compare calcite formation in FCB and B-FCB. The results indicated that the D of FCB and B-FCB played the most significant role in the increase or decrease of the XRD intensity. The calcite intensity of the runs at different densities, i.e., 1300 kg/m³, 1550 kg/m³ and 1800 kg/m³, of FCB and B-FCB are demonstrated in Figure 10. The intensity decreased when the density of FCB and B-FCB decreased. Therefore, the highest intensity occurred at 1800 kg/m^3 , and the lowest at 1300 kg/m³. At a low density of FCB and B-FCB, air bubbles were generated due to the high reactivity of the foaming agent compared to the high density. As a result, the cement content reduced, resulting in diminished Ca content, which is the key factor in the formation of CaCO₃. Therefore, the results of the XRD analysis and the figures plotted were classified based on the densities used in this study. Figure 10a,b show that the highest intensity of the FCB specimens was 5500 at 1300 kg/m³, while it was 5900 for B-FCB with a However, the highest intensities for both types of FCB and B-FCB with 1550 kg/m^3 were 6900 and 8200, as shown in Figure 10c,d, respectively. Figure 10e,f show that at 1800 kg/m^3 , the highest intensity of FCB was almost 11,000, while for B-FCB, it was almost 13,500 with the same level of D.



Figure 10. Cont.



Figure 10. XRD analysis comparison between FCB and B-FCB at 28 days, (**a**) Specimens with 1300 kg/m³ of density, (**b**) The highest peak at specimen with 1300 kg/m³ of density, (**c**) Specimens with 1550 kg/m³ of density, (**d**) The highest peak at specimen with 1550 kg/m³ of density (**e**) Specimens with 1800 kg/m³ of density, (**f**) The highest peak at specimen with 1800 kg/m³ of density.

The results suggest that the crystallinity intensity in B-FCB is higher than that of FCB, which reflects the role of *B. tequilensis* in precipitating CaCO₃. Moreover, some peaks appeared on the specimens of B-FCB, but were absent in FCB [37], due to the reactions of the *B. tequilensis* enzymes, which integrated the natural sequestration of CO₂ (resulting in higher CaCO₃ yield) and the healing process of the specimen pores [37].

5. Conclusions

 T and CO₂*B were the most significant factors affecting the compressive strength results of B-FCB. The highest compressive strength of B-FCB was 8.22 MPa at 28 days, which occurred when CO₂, B, T and D were at the following levels: 10%, 3×10^7 cell/mL, 27 °C and 1800 kg/m³, respectively. In addition, the strong interaction effects between CO₂*B and compressive strength reflected the relationship between sequestrated CO₂ and *B. tequilensis* concentration. This, in turn, enhanced the precipitation of CaCO₃, healed the B-FCB pores, and improved the compressive strength. Therefore, the compressive strength of B-FCB was higher than that of FCB.

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