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Through Big Data Management**

► **MAINTENANCE OF MICROCRACKING IN RIGID PAVEMENT WITH THE AID OF BACTERIA**

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►► Problem statement

Microcracking of rigid pavement usually occur due to the **shrinkage** of concrete after casting or throughout its **service life** due to **repeated compressive, tensile, and shear stress**.

Such cracking exhibit a **durability** problem.

►► Aim of the study

Self-healing techniques will be adopted with the aid of bacteria and healing agent to precipitate CaCO_3 on the formed micro-cracks.

A soil bacterium named *Bacillus subtilis* will be cultured in the laboratory, the concentration of bacteria cell of *B. subtilis* in normal saline (NaCl , 9 g/l) suspension was 10^6 cell/ml.


Impact of Healing of pre-cracked specimens with the aid of bacteria in the curing water on the strength properties of concrete will be monitored.

►► Methodology adopted

- Concrete specimens of various type (**cube** of 100x100x100 mm, **cylinder** of 100mm diameter and 200mm height, and **beam** of 100 x 100 x 500 mm) size will be prepared in the laboratory, then separated to three sets.
- The **first set** of specimens will be subjected to **controlled compression and flexure pre-cracking**, then subjected to **healing and curing** in a water bath which contains the prementioned bacteria at 20°C for 7 days.

The **second set** was the **control** specimens cured in water bath for 7 and 28 days at 20°C.

The **third set** of specimens will be subjected to **healing and curing** in a water bath which contains the prementioned bacteria at **20°C for 7 and 28 days** and then tested for compressive, indirect tensile, and flexure properties.

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- ▶▶ **Materials and testing methods**
 - **Ordinary Portland cement type 1.**
 - **Coarse and fine aggregates.**
 - **Water.**
 - **Culture of Bacteria Bacillus Subtilis.**

►► Chemical composition of cement

| Oxide | % by weight | Limit of Iraqi specification No.5/ 1984 |
|--------------------------------|-------------|---|
| CaO | 61.28 | ----- |
| SiO ₂ | 18.372 | ----- |
| Al ₂ O ₃ | 3.58 | ----- |
| Fe ₂ O ₃ | 5.02 | ----- |
| MgO | 1.39 | < 5.0 |
| SO ₃ | 2.02 | < 2.80 |
| C ₃ A | 0.988 | ----- |
| Loss on ignition | 2.85 | ≤ 4.0 |
| Insoluble residue | 1.07 | < 1.5 |
| Lime saturated Factor | 1.0148 | ≥ 0.66 ≤ 1.02 |

►► Physical properties of cement

| Physical properties | Test result | Limits of Iraqi specification |
|--|--------------|-------------------------------|
| Specific surface area, Blain's method m ² /kg | 394 | ≥ 230 |
| Soundness, Autoclave's Method, % | 0.03 | < 0.8 |
| Setting time, Vicat's method Initial setting hour: min Final setting hour: min | 2:15 3:30 | ≥ 45 min ≤ 10 hours |
| Compressive strength 3 days N/mm ² 7 days N/mm ² | 20.7 26.1 | ≥ 15 ≥ 23 |

►► Properties of Coarse and fine aggregates

| Type of aggregate | Bulk Specific Gravity | Density(kg/m ³) | Absorption % | SO ₃ % |
|-------------------|-----------------------|-----------------------------|--------------|-------------------|
| Crushed aggregate | 2.63 | 1646 | 1.167 | 0.034 |
| Fine aggregate | 2.5 | 1789 | 9.3 | 0.126 |

►► Combined aggregates gradation adopted

| Sieve size mm | 19 | 12.5 | 9.5 | 4.75 | 1.18 | 0.3 | 0.15 |
|----------------------------|-----|------|-----|------|------|-----|------|
| Percentage finer by weight | 100 | 85 | 70 | 59 | 46.4 | 16 | 3.3 |


►► Preparation, Isolation and Culture of Bacteria Bacillus Subtilis

- The isolation and culture of bacteria bacillus subtilis starts by collection of soil samples from Agricultural areas in Baghdad city and evacuation in glass bottles.
- The soil sample was then mixed with distilled water and shaken vigorously to ensure thorough mixing.
- Serial dilution was made by transfer of 1 ml of soil suspending to 9 ml of distilled water in test tube, this soil suspension is of 10^{-1} dilution.

This process was repeated until 10^{-6} dilution is obtained.

After all these steps, petri-dishes plates with nutrient agar media according to the bacteria requirement have been prepared.

The solution of concentration 10^{-4} to 10^{-6} was spread by cotton swab on the petri-dishes plate with nutrient agar media in it and incubated in 37°C to 24 hours in the incubator.

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- ▶▶ • After this, the different type of colonies on different plates were observed and the growth after 24 hours incubation was checked.
 - Afterword, Preparation of the concentration 10^6 of bacterial cell (*Bacillus subtilis*) starts by Transfer of loop full of single colonies of *Bacillus subtilis* that form urease enzyme from nutrient agar to brain heart broth media and incubation at 37°C on shaker at 150 rpm for 24 hours.

The harvest of bacterial cell was done by centrifuge the 24 hour's old grown culture (5000 rpm, 5 minutes).

The sediments (bacterial cell) was form after centrifuge and were re-suspended in normal saline (NaCl, 9 g/l).

The concentration of bacteria cell of bacillus subtilis in suspension was 10^6 cell/ml.

►► Isolation of the bacteria.



► Culture and dilution of the bacteria



►► Preparation of Concrete Mixture

Control concrete mixture was designed as per ACI 211.1-91 method, such mixture is usually used for rigid pavement construction.

The mix proportion is **(1:1.5:3.75) with 0.45 water cement ratio.**

Cube, beam and cylinder specimens have been prepared, covered with polythene sheets to retain the mixing water for curing for 24 hours. Specimens were immersed in **water with bacterial concentration of 10^6 cell/ml of water.**



Calcium lactate of 5% by weight of cement were added to the bacterial water.

Specimens were tested after **seven and 28 days to assess the impact of curing in bacterial water on the physical properties.**

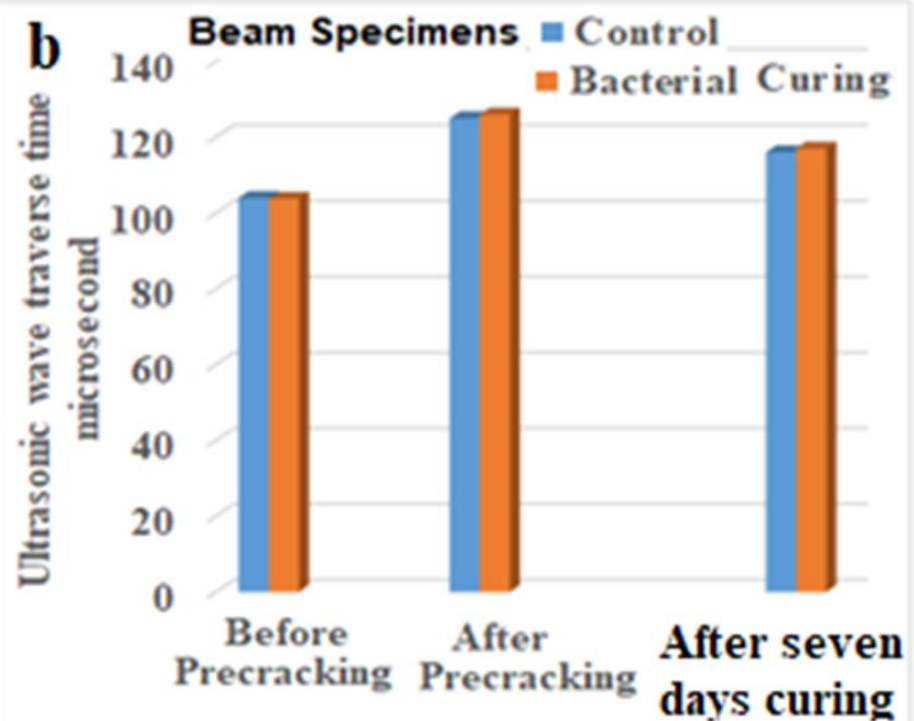
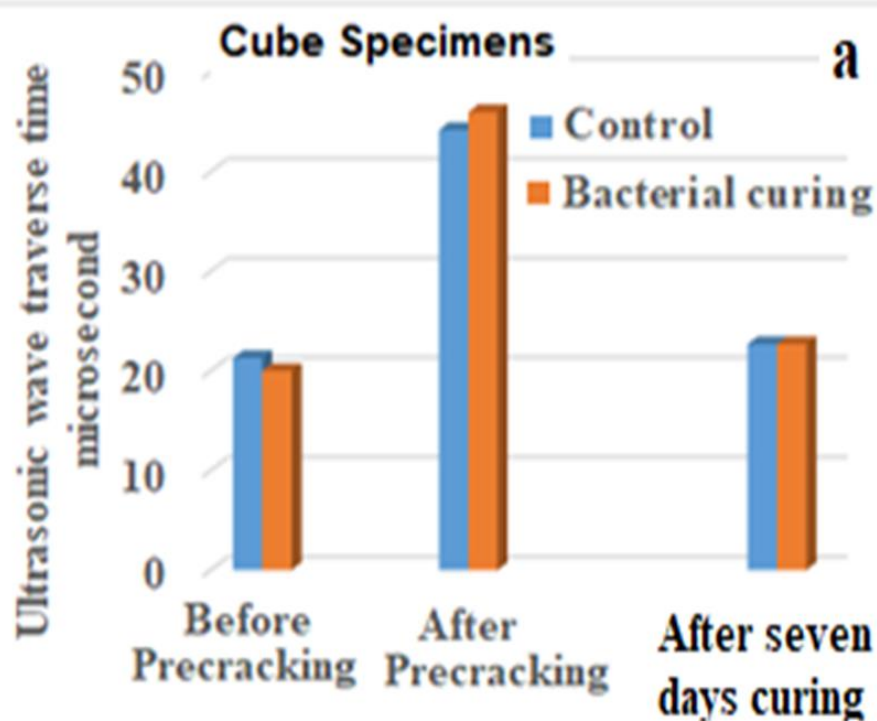
►► **Pre-cracking of specimens**

- Cube specimens were subjected to pre-cracking after 24 hours of curing, a load of 260 kN was applied for one minute then released.
- Cube specimens were subjected to ultrasonic wave traverse with the aid of pundit to determine the time required for the sound to traverse the specimen before and after the pre-cracking process.



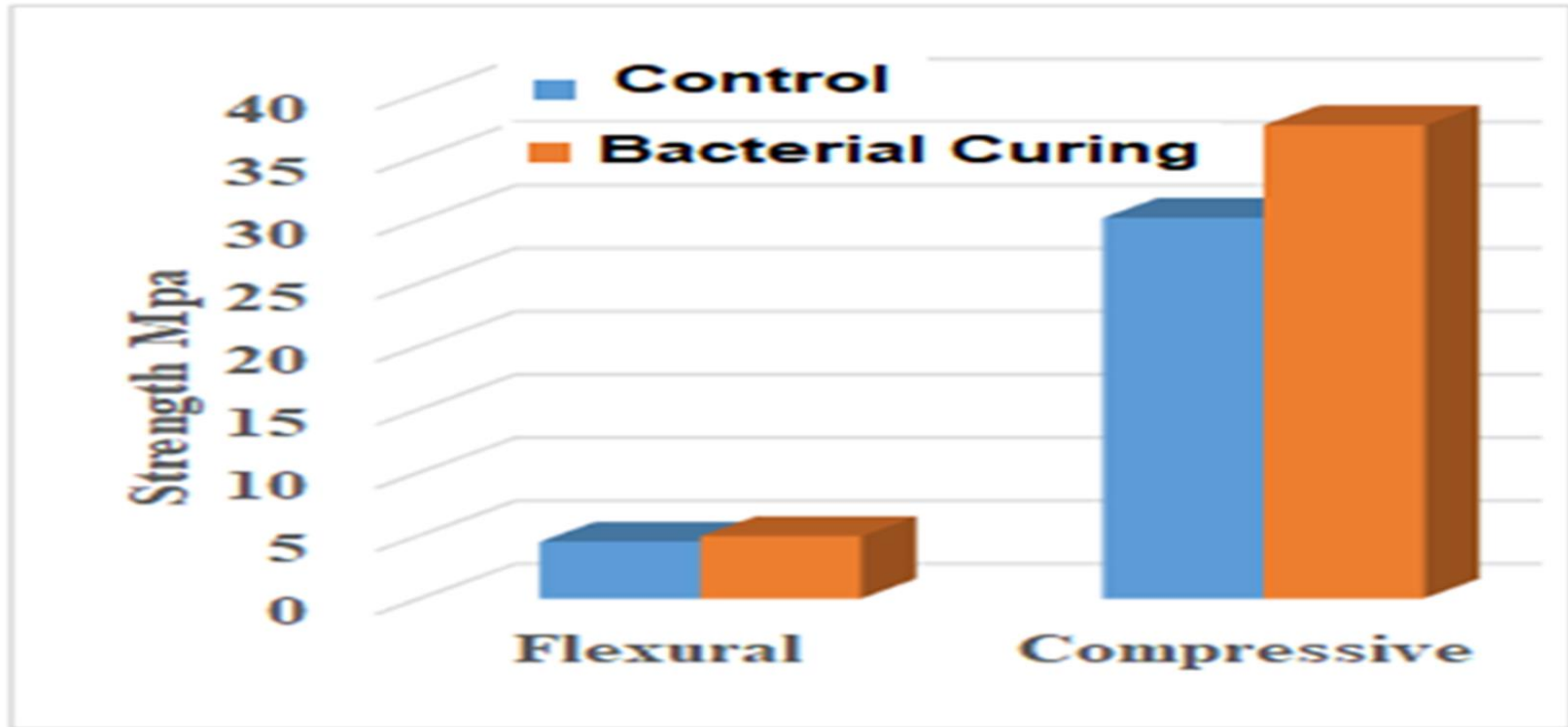
- For beam specimens, the pre-cracking was conducted using a flexure load of 5kN, while the ultrasonic wave traverse along the length of the beam.
- The time required for the sound to traverse the specimen before and after the pre-cracking process was recorded.

►► Ultrasonic wave traverse time for pre-cracked specimens

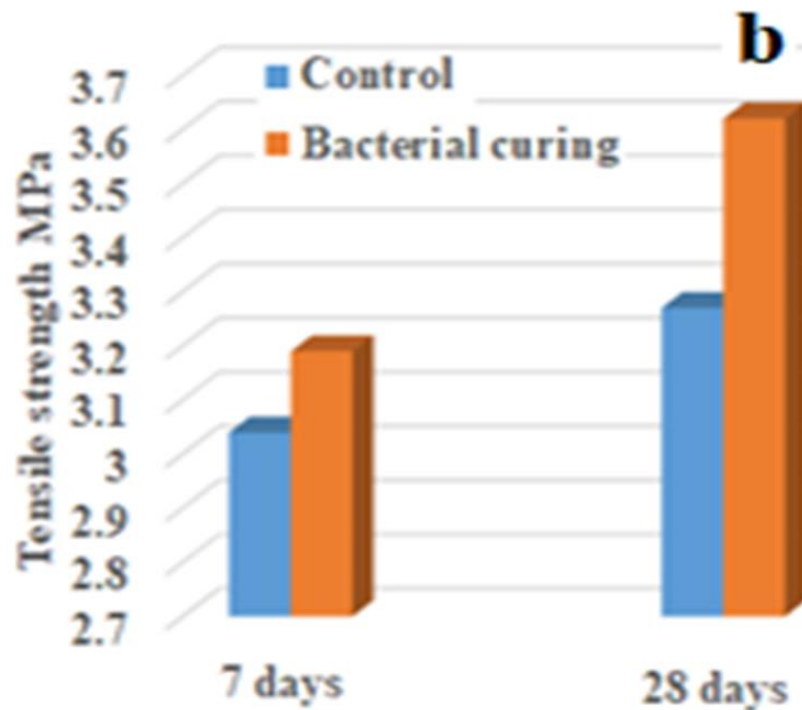
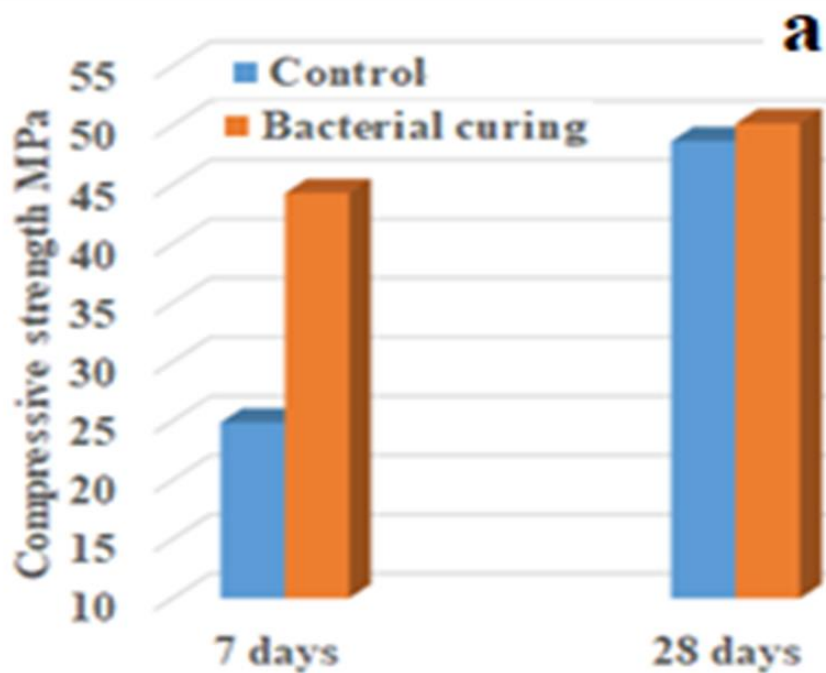


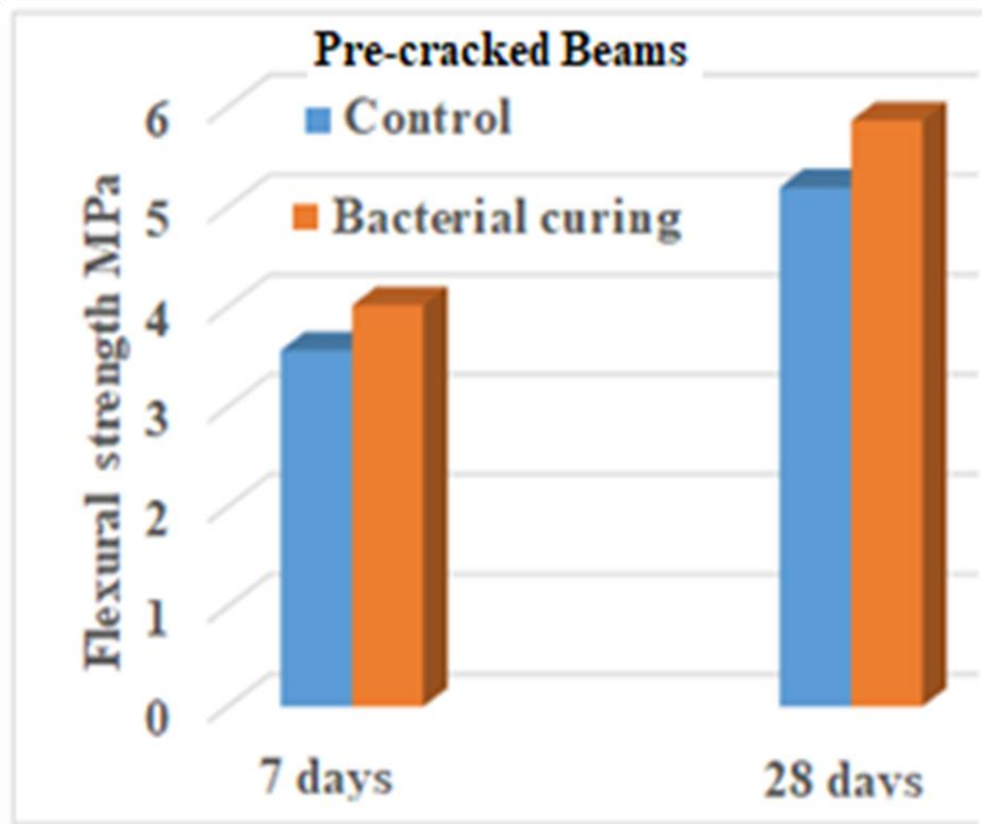


►► Strength properties after pre-cracking and curing for seven days











CONCLUSIONS

Implementation of bacteria (*Bacillus subtilis*) in the curing water exhibit a positive influence on the strength properties as well as crack healing of concrete.

Bacterial cured concrete exhibit high compressive strength of (107 and 23) % after 7 and 28 days of curing respectively when compared to that of control concrete. The increment in compressive strength after 28 days of curing was (95.5 and 18.4) % for control and bacterial Concrete respectively.

Bacterial cured concrete exhibit high tensile strength of (5 and 10.7) % after 7 and 28 days of curing respectively when compared to that of control concrete. The increment in tensile strength after 28 days of curing was (8.3 and 14.2) % for control and bacterial concrete respectively.

Bacterial cured concrete exhibit high flexural strength of (22.5 and 16) % after 7 and 28 days of curing respectively when compared to that of control concrete. The increment in flexural strength after 28 days of curing was (38 and 48.7) % for control and bacterial concrete respectively.



Specimens subjected to controlled pre-cracking exhibit improvement in strength properties after the healing process provided by the bacteria by (28 and 33) % for compressive and flexure strength respectively as compared to those of control mixture after 7 days of curing in bacterial water.



Recommendation

- It is recommended to implement bacterial water for curing of rigid pavement to assess the crack healing process.

Thank you