

CO₂ sequestration derived calcium carbonate and methane formation in deep subsurface porous media: A State-of-art review to decoding microbes-minerals crosstalk and molecular bio mineralization process

Abstract

The concentration of carbon dioxide has been rising extensively due to human activities and it reached dangerous levels that had not been seen in the previous 3 million years. In 2022, carbon dioxide emissions from the transport sector increase by 250Mt CO₂ which is 3% more than in 2021(8Gt). Natural sources of carbon dioxide emissions include decomposition, forest fires, ocean release, volcanic eruption, and ocean release. Each year globally 2.6 billion tons of carbon dioxide are released from natural forest fires. Ocean absorbs 30% of carbon dioxide which is released by human activities and holds 60 times more carbon than the atmosphere. Microbial communities play a major role in carbonate rock formation through bio mineralization within subsurface porous media. Moisture content in subsurface porous media significantly impacts the carbonate bio mineralization process during CO₂ sequestration. It influences mineral dissolution, microbial activity, water transport, CO₂ solubility, and transport efficiency. Environmental factors like temperature, pH, and nutrient availability can influence urease activity. Microbial community dynamics also impact ureolysis efficiency. Understanding these aspects is essential for applications like CO₂ sequestration and environmental remediation. To stop CO₂ leakage in the context of CO₂ sequestration, the subsurface may be covered with a layer of cap rock, or impermeable rock. The formation of carbonate minerals induced by MICP can strengthen the integrity of the cap rock by sealing potential pathways for CO₂ migration and filling fractures. Maintaining sustainable CO₂ sequestration and bio-methane production requires striking a balance between the advantages of MICP and its effects on subsurface processes. This review examines how geochemical and environmental controls affect microbial activity and bio mineralization, emphasizing the importance of sustainability and long-term stability.

Keywords

CO₂ sequestration, microbial communities, subsurface porous media, bio mineralization, MICP.

Introduction

Since the Industrial Revolution, the concentration of carbon dioxide has been rising extensively due to human activities and it reached dangerous levels that had not been seen in the previous 3 million years¹. Increased concentrations of carbon dioxide in the atmosphere are due to anthropogenic activities such as deforestation, as well as the burning of fossil fuels. Although anthropogenic sources of carbon emission are much smaller than natural emissions, they disturb the natural balance that was before human influence. Researchers estimated that the world's emission of carbon dioxide exceeded 40 billion tons in 2023 and nearly 37 billion tons of emissions are from fossil fuels².

Every year due to deforestation more than 1.5 billion tons of carbon dioxide are released into the atmosphere. Global carbon dioxide emissions from fossil fuel burning are about 34 billion tons (Gt) per year³. Industrialization and energy consumption had a positive impact on carbon emissions, a 1% increase in industrialization leads to a 0.54% rise in CO₂ emissions while a 1% rise in energy consumption produces a 0.61% increase in emissions. In 2022, carbon dioxide emissions from the transport sector increase by 250Mt CO₂ which is 3% more than in 2021(8Gt). Air travel contributed to 70% of emissions in 2019. Between 2001 and 2023 global forest fires release 35.9 tons of carbon dioxide. While agriculture contributes to 10% of global CO₂ emissions⁴.

Furthermore, flavonoids and phenolic also being explored for pharmaceutical and medical applications⁵. A range of natural compounds derived from plants has been explored for their potential health benefits⁶. Chlorogenic Acid from *Equisetum arvense* L. is a phenolic compound that possesses antioxidant properties and has been studied for its antiviral, antibacterial, and anti-inflammatory effects⁷. Methotrexate from *Plantago lanceolata*, on the other hand, is a versatile phenolic drug used to treat various conditions, including cancer and autoimmune diseases⁸. Among the flavonoids, Tamoxifen, Raloxifen, and Anastrozole are derived from *Urtica dioica*, showing promise when combined with radiotherapy and adjuvants in improving the survival rates of breast cancer patients⁹. Additionally, Genistein, being a flavonoid, is being studied in clinical trials for prostate cancer treatment. Crofelemer serve various medical purposes, including cancer and tumors¹⁰, while Silibinin, a flavonolignan, and is used for treating toxic liver damage, cirrhosis, and hepatitis. It's crucial to consult with healthcare professionals and rely on scientific research and clinical trials when considering the use of these compounds to ensure their effectiveness and safety¹¹.

Natural Sources

Carbon emissions from natural sources are a vital part of Earth's carbon cycle. Natural sources of carbon dioxide emissions include decomposition, forest fires, ocean release, volcanic eruption, and ocean release. The decomposition of leaves and plants releases 60 pentagrams of carbon every year into the atmosphere¹². Oceans release 10 billion tons of carbon every year into the atmosphere but oceans quickly absorb 3 billion tons of these emissions. Each year globally 2.6 billion tons of carbon dioxide are released from natural forest fires. Due to volcanic eruptions globally 94 million tons of carbon dioxide are released each year¹³.

Carbon sequestration methods

The removal, capturing, and storage of carbon dioxide from the earth's atmosphere is called carbon sequestration. Two basic forms of carbon sequestration are biological and global carbon sequestration. Three main carbon sequestration methods are described below:

1. Biological Carbon sequestration

Biology carbon is stored in the natural environment. Major carbon sinks in nature are forests, grasslands, soil, oceans, and other bodies of water. Forest plays a significant role in natural/biological carbon sequestration¹⁴. 25% of global carbon emissions are sequestered by

forests and other vegetative forms such as rangelands or grasslands (FAO) ¹⁵. Forest stores double carbon as they emit. In the fight against climate change, soil plays a very important role as healthy soil helps remove carbon from the atmosphere and store it in soil (IPCC) ¹⁶. Globally 1500- 2500 billion tons of carbon can be sequestered by soil (USDA) ¹⁷. Aquatic bodies are the greatest absorbers of CO₂, as they absorb 25% of carbon dioxide emitted by the earth's atmosphere ¹⁸. 50% of the oxygen we need is generated by the ocean while it absorbs 25% of all carbon emissions and oceans capture 90% of heat emitted by these emissions. Ocean absorbs 30% of carbon dioxide which is released by human activities and holds 60 times more carbon than the atmosphere¹⁹.

2. Geological Carbon Sequestration

The storage of carbon in underground geological formations or rocks is called geological carbon sequestration ²⁰. It is an effective way of neutralizing the carbon emissions due to human activities such as construction or manufacturing²¹. Recent technological innovations show that carbon can be sequestered on a larger scale more effectively. These technological innovations include:

Graphene production

Carbon dioxide is used as a raw material in grapheme production. It is heavily utilized in certain industries for the production of tech devices such as computer processors and smartphones ²².

Engineered molecules

By capturing carbon from the air, scientists can change the shape of molecules which helps in the formation of new compounds. While reducing atmospheric carbon could be a systematic way of creating raw material²³.

3. Mechanical carbon sequestration

It is a method of capturing and storing carbon dioxide from the atmosphere by the usage of mechanical processes such as enhanced weathering or air capture. In this method, carbon dioxide is physically removed from the air and then stored in oceans, geological formations, or other long-term storage site²⁴. Direct capture from the air is the best strategy in the fight against climate change and it is designed to remove 4000 tons of carbon dioxide per year. DAC is not limited to national emissions it can be done anywhere in those areas that have good wind speeds²⁵. In the case of exhaust stack capture it usually be carried out closer to CO₂ burial sites. While mechanical trees absorb carbon more effectively, when planted in clusters of 12 they capture a metric ton of CO₂ per day. A large farm with 120,000 trees in the working phase, can capture 10,000 metric tons of carbon per day which would be equal to the CO₂ emissions of 800,000 cars²⁶.

The most stable and permanent form of CO₂ sequestration in geological formation is mineralization. Different researchers conducted experiments on this method and they revealed that the reaction takes decades to sequester CO₂. The process takes place directly or indirectly when dissolved CO₂ and brine form carbonic acid²⁷. In the geologic formation, there is a reaction that takes place after a long time between carbonic acid and minerals which leads to the improvement of precipitates of carbonate minerals and after that results in solid precipitates of carbon dioxide. It is attractive because it can permanently store CO₂. It is the most secure trapping mechanism in

the CO₂ sequestration process²⁸. Trapping speed is slow but if certain parameters are completely understood then there is the possibility to speed up the trapping process. Some factors such as brine pH, brine composition, temperature, pressure, leakage rate, microorganism interaction, and CO₂ dissolution kinetics have a major effect on the mineral trapping of CO₂ in brines²⁹.

Rock formation via bio mineralization

A process by which living organisms facilitate mineral formation is called bio mineralization. During this process, microorganisms interact with injected CO₂ and the subsurface environment. High concentrations of carbon dioxide are injected underground, where they interact with minerals found in the rock formation³⁰. The aim of this process is also referred to as mineralization or mineral trapping it turns gaseous CO₂ into stable mineral form to prevent its release into the atmosphere. Various factors can contribute to rock formation through bio mineralization in the context of deep industrial subsurface porous media during the injection of high concentrations of CO₂³¹. Here is a general overview of how this might happen:

Mineral Activity

Bacteria and other microorganism can be very important to bio mineralization. Certain bacteria can modify the local geochemical conditions to catalyze mineral precipitation reactions. They can produce metabolic byproducts that alter the subsurface chemical composition³².

Ureolysis and calcium carbonate precipitation

Certain bacteria are known for their ureolytic activity, which allows them to hydrolyze urea and produce carbonate ions. PH and carbonate ion availability increase as a result of this process which encourages the precipitation of calcium carbonate. It is a mineral that can help to form a solid rock by consolidating the porous media³³.

Sulfide mineral precipitation

Sulfide ions are produced when certain microbes participate in the sulfate reduction process. Iron sulfide (pyrite, greigite) and other sulfide minerals may precipitate as a result of this process. As a result of these minerals, the properties of subsurface rocks may change³⁴.

Biofilm formation

On rock surfaces, microorganisms frequently form biofilms. Biofilms can precipitate minerals and serve as nucleation sites for the synthesis of new minerals. A mineralized matrix that strengthens the rock structure may result from this³⁵.

Nutrient availability

Carbon, nitrogen, and phosphorus availability are examples of nutrients that can affect microbial activity and in turn bio mineralization. These nutrients may be utilized by microorganisms for metabolic activities that contribute to the formation of minerals. It's important to note the details of bio mineralization in the deep industrial subsurface during CO₂ injection can differ based on the

site's geological and geochemical features, the composition of CO₂ injected, as well as the microbiological community that is present³⁶.

Molecular mechanism of carbonate rock formation

As discussed, earlier rock formation takes place by the process of bio mineralization when a high concentration of CO₂ is injected into subsurface porous media. Although the specific microbiological species and reactions may differ, the general procedure usually entails the following steps³⁷.

Injection of CO₂

Subsurface porous media that contains minerals and water is injected with CO₂³⁸. A favorable atmosphere for bio mineralization is created by the high concentration of carbon dioxide³⁹.

Microbial Activity

Some bacteria can catalyze bio mineralization; these bacteria are commonly found in genera like Bacillus and Sporosarcina⁴⁰. Through metabolic processes, these bacteria help in the formation of carbonate minerals⁴¹.

Uptake of CO₂ by microorganisms

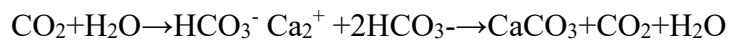
CO₂ serves as a source of carbon for the metabolism of microorganism. The CO₂ is transformed into bicarbonate ions (HCO₃⁻) through enzymatic processes such as carbonic anhydrase activity⁴².

Production of carbonate ion

The calcium ions (Ca₂⁺) in the subsurface porous media react with the bicarbonate ions produced by microbial activity. As a result, calcium carbonate (CaCO₃) is produced⁴³

Biochemical reaction

The bio mineralization process involves many biochemical reactions. One simplified example of a reaction is:



In these reactions, CO₂ is converted to bicarbonate ions, and then the bicarbonate ions react with calcium ions to form calcium carbonate, which precipitates as solid mineral phases⁴⁴.

Types of microbial communities involved in carbonate rock formation

Microbial communities play a major role in carbonate rock formation through bio mineralization within subsurface porous media. Microorganisms that affect the mineralization and precipitation of carbonate minerals are involved in this process⁴⁵. A variety of microbial communities are involved in this process of bio mineralization

Cyanobacteria

Through bio mineralization, these photosynthetic bacteria can help in the formation of carbonate minerals⁴⁶. These bacteria have the capability of fixing carbon dioxide and precipitating carbonate minerals such as calcium carbonate in their extracellular polymeric substance (EPS). Cyanobacteria create biofilms in subsurface environments, which leads to the precipitation of carbonate minerals and the formation of microbialite structures⁴⁷.

Sulfate-reducing bacteria

Through microbial sulfate reduction, certain sulfate-reducing bacteria are known to encourage the precipitation of carbonate minerals⁴⁸. Formation of calcium carbonate takes place when these bacteria react with dissolved calcium ions to produce sulfide (S_2^-) from reduced sulfate (SO_4^{+2}).

Iron-Reducing Bacteria

By encouraging the reduction of iron minerals, iron-reducing bacteria can affect the precipitation of carbonate minerals. This process contributes to the precipitation of carbonate minerals by releasing bicarbonate ions and dissolved iron into the surrounding environment⁴⁹.

Ureolytic Bacteria

Ureolytic bacteria, including some Bacillus species, can produce urease enzymes, which hydrolyze urea and release ammonium and carbonate ions. Carbonate minerals, especially calcium carbonate, may precipitate as a result of this rise in carbonate ions⁴⁹.

Denitrifying Bacteria

By modifying the chemical environment in their surroundings, denitrifying bacteria indirectly affects carbonate precipitation. During denitrification, nitrate (NO_3^-) is reduced to nitrogen gas (N_2), which increases pH and encourages the precipitation of carbonate minerals⁵⁰.

Microbial Consortia

Complex microbial consortia in which various microbial species work synergistically, are frequently involved in the formation of carbonate rocks. For example, a group of fungi, bacteria, and archaea may help in the general biomineralization process in subsurface settings⁵¹.

Biofilm Formation

In subsurface porous media, microorganisms frequently create biofilms on the surface. The environment that creates these biofilms is favorable for microbial activity, which includes the precipitation of carbonate minerals. For carbonate mineral growth EPS which is produced by biofilm-forming microorganisms can act as nucleation sites. In addition to subsurface geology, these microbial processes affect environmental remediation and carbon sequestration⁵².

Factors affecting carbonate biomineralization process

Abiotic factors

Texture of soil sub-strata or rock types

The texture of the rock types and soil substrata in porous media strongly influences the carbonate biomineralization process during CO₂ sequestration. Contact between CO₂ and minerals are made easier by the soil or rock porosity and permeability, which control fluid movement⁵³. The mineral compositions of different types of rock vary, which influences the availability of ions necessary for carbonate mineralization in mineral reactivity with injected CO₂ and other solution components⁵⁴. The chemical composition of pore fluids also influences the carbonate biomineralization process⁵⁵. Carbonate biomineralization may be impacted by secondary minerals created by earlier geological processes. The subsurface fluid movement of injected fluids is influenced by hydraulic conductivity⁵⁶.

Amount of CO₂ concentration in subsurface porous media

The carbonate biomineralization process during CO₂ sequestration is strongly influenced by the concentration of CO₂ in subsurface porous media. Carbonic acid is created when CO₂ dissolves in water and reacts with the minerals in the porous medium to produce carbonate mineral precipitation⁵⁷. Increased carbonic acid formation from higher dissolved CO₂ concentrations encourages mineral dissolution and releases the ions required for carbonate precipitation. The concentration of dissolved CO₂ affects the equilibrium between carbonic acid, bicarbonate ions, and carbonate ions. The concentration of reactants, such as dissolved CO₂, affects the rate of carbonate mineralization reactions⁵⁸. The ideal mineralization depends on the pH of the pore fluids, which is influenced by the CO₂ concentration as well. Limitations on mass transfer, like flow dynamics and injection rate, also affect the mass transfer of CO₂.

Level of O₂ concentration in subsurface porous media

During CO₂ sequestration, carbonate biomineralization may be impacted by the oxygen content of subsurface porous media. Microbial activity and respiration can be impacted by the oxidation of organic carbon, which is facilitated by oxygen. Additionally, oxygen affects redox conditions, which in turn affects the possibility of mineralization reactions⁵⁹. Maintaining a stable pH is necessary to encourage carbonate biomineralization. Anaerobic processes can be inhibited by oxygen, which lowers the amount of carbonate ions available for precipitation⁶⁰. The total amount of oxygen available for biomineralization can vary depending on oxygen consumption. Oxygen can also catalyze oxidative reactions, allowing certain minerals to undergo oxidative dissolution, releasing ions that contribute to carbonate mineral precipitation⁶¹. Oxygen plays a complicated role in carbonate biomineralization. Understanding the possible influence of oxygen on the overall success of carbonate mineralization in porous media requires site-specific evaluations⁶².

Temperature of subsurface porous media

In the carbonate biomineralization process during CO₂ sequestration, temperature is critical. It affects microbial activity, thermodynamics, and reaction kinetics all of which are critical for the mineralization of carbonate⁶³. Higher temperatures increase the kinetic of reactions, which causes minerals to dissolve and precipitate more quickly. Lower temperatures increase CO₂ solubility in water, which is temperature-dependent⁶⁴. The thermodynamics of mineral precipitation are influenced by temperature as well; certain carbonate minerals are more stable at higher temperatures. The metabolic rates of the microorganisms involved in the carbonate

biomineralization process are also influenced by temperature. Maintaining the ideal conditions for carbonate precipitation requires careful pH control⁶⁵. The physical and chemical characteristics of water are also influenced by temperature, which affects how it interacts with minerals. Carbonate mineralization is also influenced by thermal gradients and temperature distribution⁶⁶.

Moisture of subsurface porous media

Moisture content in subsurface porous media significantly impacts the carbonate biomineralization process during CO₂ sequestration⁶⁷. It influences mineral dissolution, microbial activity, water transport, CO₂ solubility, and transport efficiency. Elevated moisture content causes CO₂ to dissolve more readily, raising the concentration of carbonic acid⁶⁸. Water transport ensures uniform contact between reactants and minerals, and microbial activity supports the growth and metabolic activity of microorganisms involved in mineralization reactions⁶⁹. Moisture content also affects the transport and dissolution of mineral⁷⁰. Determining the dynamics of the carbonate biomineralization process requires an understanding of these interactions. Thus, keeping the moisture content at a suitable level is crucial to the success of carbonate biomineralization during CO₂ sequestration.

pH of subsurface porous media

During CO₂ sequestration, the carbonate biomineralization process is greatly impacted by the pH of subsurface porous media. Carbonic acid formation, mineral solubility, saturation state, bicarbonate and carbonate ion availability, microbial metabolism, buffering capacity, chemical equilibria, stability of silicate minerals, and injection material corrosion are all impacted by pH. Higher concentrations of bicarbonate and carbonate ions result from lower pH values, which encourages the formation of more carbonic acid⁷¹. The availability of bicarbonate and carbonate ions, which are necessary for carbonate precipitation reactions, is also influenced by pH. The metabolic activity of microorganisms involved in carbonate biomineralization is also influenced by pH levels. To ensure the long-term stability of carbonated formations and to promote carbonate biomineralization, pH conditions must be optimized⁷².

ORP level of subsurface porous media

Subsurface porous media's Oxidation-Reduction Potential (ORP) level has a major effect on the carbonate biomineralization process during CO₂ sequestration. Higher values to ORP indicate oxidizing conditions, while lower values indicate reducing conditions. ORP is a measure of redox conditions. The overall efficiency of carbonate biomineralization can be impacted by anaerobic processes being inhibited by positive ORP conditions. To guarantee the integrity of injection materials and infrastructure during CO₂ sequestration, ORP can also indicate corrosion potential⁷³.

Biotic factors

Organic carbon situation in subsurface porous media

Carbonate biomineralization is strongly influenced by the organic carbon state of subsurface porous media, particularly in the context of carbon capture and storage operations. This is brought on by the metabolic activity of ureolytic bacteria, which create urease enzymes by using organic

carbon as a substrate⁷⁴. This raises pH levels and produces carbonate ions. Carbonate biomineralization is facilitated by the energy that organic carbon provides to microbial communities. Organic acids that are released when microorganisms break down organic carbon can have an impact on the formation of carbonate minerals by regulating pH. By acting as substrates for microbial metabolism, promoting ureolysis, and raising the availability of carbonate ions, organic carbon amendments can improve microbial activity and encourage carbonate biomineralization⁷⁵.

Microbial composition in subsurface porous media

Carbonate biomineralization is greatly influenced by the microbial composition of subsurface porous media, particularly in carbon capture and storage initiatives. Sporosarcina, Bacillus, and certain strains of Clostridium are examples of ureolytic bacteria. These bacteria produce urease enzymes, which hydrolyze urea to produce carbonate ions and raise pH⁷⁶. Utilizing sulfate as a terminal electron acceptor, sulfate-reducing bacteria (SRB) produce sulfide ions, which combine with carbonate ions and metal cations to form carbonate minerals. PH and redox conditions are impacted by the reduction of ferric ions to ferrous ions by iron-reducing bacteria. Acetate is produced by acetogenic bacteria and provides other microbial communities involved in carbonate biomineralization with an organic carbon source. Methane produced by methanogenic archaea modifies redox conditions, which in turn affects carbonate precipitation indirectly⁷⁷. Autotrophic bacteria use CO₂ as a carbon source, whereas heterotrophic bacteria obtain their energy and growth from organic carbon sources. The formation of biofilms, competition, and adaptation to harsh environments all affect carbonate biomineralization⁷⁸.

Enrichment of carbonate biomineralization involving functional genes within the microbial community in subsurface porous media

Metabolism and certain functional genes promote carbonate biomineralization in subsurface porous media. The enrichment of carbonate minerals is facilitated by genes that encode urease, carbonic anhydrase, alkaline phosphatase, polyphosphate kinase, sulfate-reducing, methanogenic, organic carbon metabolism, biofilm-forming, and metal-related functions⁷⁹. Urease-producing bacteria release carbonate ions and elevated pH, promoting carbonate mineral precipitation. Phosphate ions are released by alkaline phosphatase enzymes during hydrolysis which influences the formation of carbonate minerals. By adjusting redox conditions, methanogenic archaea indirectly help in the biomineralization of carbonate. It is essential to comprehend the genetic and metabolic capacities of microbial communities to forecast and improve carbonate mineral precipitation in processes such as CO₂ sequestration⁸⁰.

Microbial activity (especially urase enzyme) in subsurface porous media

Microbial activity, particularly urease enzymes, is crucial in carbonate biomineralization in subsurface porous media. Urease enzymes catalyze the hydrolysis of urea, releasing carbonate ions and ammonium ions⁸¹. This process increases pH, which is favorable for carbonate mineral precipitation. Urease activity directly contributes to the production of carbonate ions, which are essential building blocks for carbonate minerals⁸². Urease-mediated carbonate biomineralization has implications for carbon capture and storage strategies. Environmental factors like temperature,

pH, and nutrient availability can influence urease activity. Microbial community dynamics also impact ureolysis efficiency. Understanding these aspects is essential for applications like CO₂ sequestration and environmental remediations⁸³.

EPS (Extracellular polymeric substances) production rate in subsurface porous media

Microorganisms produce complex, high-molecular-weight polymers known as extracellular polymeric substances (EPS), which are essential for subsurface porous media, including carbonate biomineralization. Carbonate minerals use EPS as nucleation sites to form biofilms, which offers a safe site for microbial activity⁸⁴. They serve as a barrier of defense, encouraging cell survival and improving the uptake and retention of nutrients. EPS also contributes to microbial adhesion to minerals, which will change the permeability and pore structure⁸⁵. It also controls redox and pH levels, which influence the microenvironment surrounding microbial cells. The rate and degree of carbonate biomineralization in subsurface porous media can be influenced by EPS-mineral interactions. Understanding the relationship between EPS production rate and carbonate biomineralization is vital for optimizing microbial processes for applications like CO₂ sequestration⁸⁶.

Carbonate Mineral formation through MICP process

Microorganisms cause the precipitation of calcium carbonate minerals in the subsurface through a biomineralization process known as Microbially Induced Calcium Carbonate Precipitation (MICP)⁸⁷. This process can impact CO₂ sequestration efficiency by decreasing subsurface porous media's porosity in several ways:

Porosity Reduction

The physical filling of the pore spaces in subsurface media is caused by the precipitation of calcium carbonate minerals in these spaces⁸⁸. The media's overall porosity decreases as carbonate minerals, such as calcite, build up and sometimes fill the pore spaces that were once filled with gas or water.

Cementation of Sediments

The carbonate minerals formed by MICP can bind sediment particles together naturally like cement. A more compact and solid subsurface structure is formed as a result of the cementation process, which helps to consolidate loose sediments⁸⁹.

Pore Throat Occlusion

Smaller pore throats in the subsurface media may become obstructed as a result of the precipitation of carbonate minerals. As a result, the pore network's connectivity decreases, which restricts the flow of fluids including CO₂ through the porous media⁹⁰. The occlusion of the pore throat can improve CO₂ trapping below the surface.

Increase in Mechanical Strength

The subsurface media's mechanical strength increases as a result of porosity reduction and particle cementation. This increased strength lowers the possibility of subsurface deformations or collapses, which is especially important for stability concerns⁹¹.

Enhanced Caprock Integrity

To stop CO₂ leakage in the context of CO₂ sequestration, the subsurface may be covered with a layer of caprock, or impermeable rock. The formation of carbonate minerals induced by MICP can strengthen the integrity of the caprock by sealing potential pathways for CO₂ migration and filling fractures⁹².

CO₂ Trapping within Minerals

When CO₂ is dissolved in pore water, it can react with calcium ions to precipitate as carbonate minerals, which is a long-term storage medium for sequestered CO₂. This chemical trapping of CO₂ within the mineral strength adds an extra layer of permanence to the sequestration process⁹³.

Controlled Release of CO₂

Although immobilizing carbon dioxide is the main objective of CO₂ sequestration, it's also important to take into account the possibility of controlled CO₂ release from the carbonate minerals⁹⁴. Changes in pressure or pH could cause the CO₂ that has been stored to leak back into the solution⁹⁵.

Enhanced Sealing of Faults and Fractures

MICP-induced carbonate precipitation can help seal faults and fractures in geologic formations. This sealing improves the containment of injected CO₂ within the target storage zone and lowers the possibility of CO₂ migration along these preferred flow paths⁹⁶.

It is important to note that although MICP presents potential advantages for CO₂ sequestration, the process-specific efficacy is contingent upon several variables, such as the selection of microorganisms, the accessibility of reactants, and the subsurface geological features⁹⁷. To maximize MICP for real-world application of CO₂ sequestration projects, research is ongoing to maximize⁹⁸.

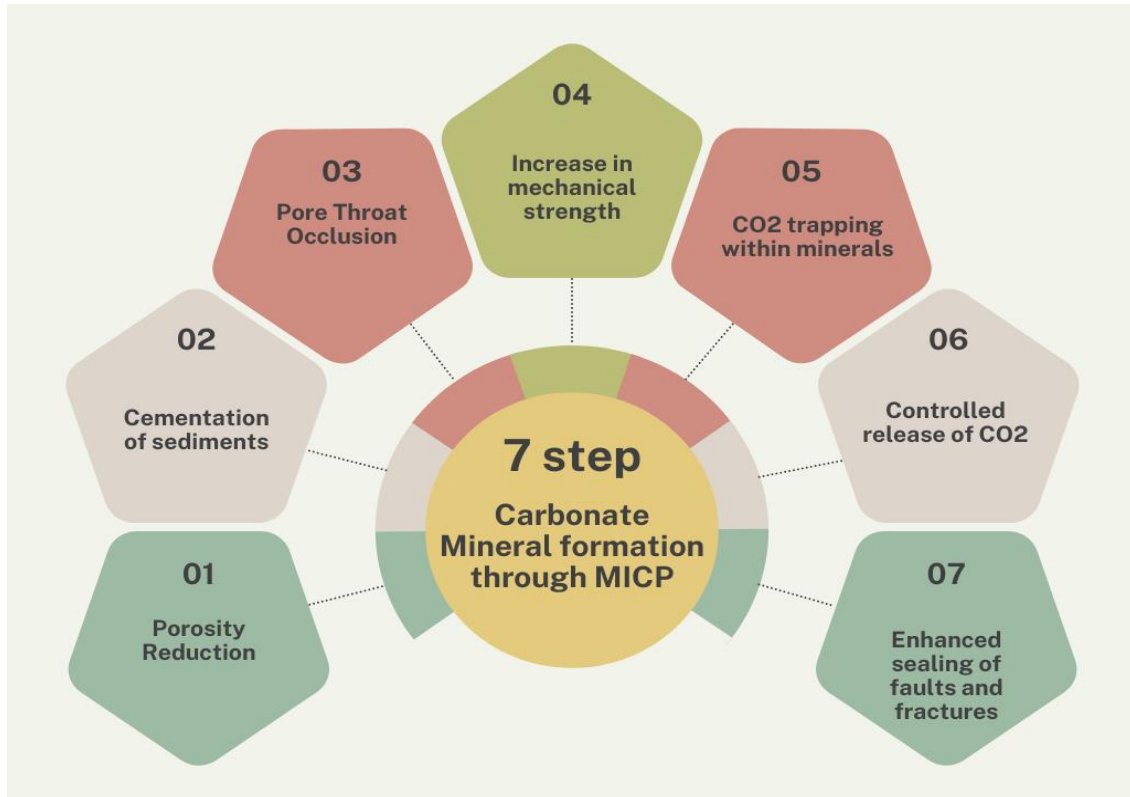


Fig: Carbonate Mineral Formation through MICP.

Methane gas generation

Through the metabolic activities of microorganisms, a process known as MICP precipitates calcium carbonate (CaCO_3). This process is widely applied in several contexts, such as CO_2 sequestration, where it improves the storage of carbon dioxide in geological formations⁹⁹. One of the less common outcomes of MICP is the direct generation of methane gas in subsurface porous media. The following steps are involved in the MICP process:

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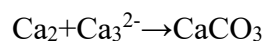
Carbon dioxide (CO_2) uptake

Certain types of bacteria and other microorganisms commonly involved in MICP absorb CO_2 from their environment¹⁰⁰.

Calcite Precipitation

The microorganisms transform the absorbed CO_2 into carbonate ions (CO_3^{2-}). Following their reaction with calcium ions (Ca^{2+}) in the surrounding water, these carbonate ions produce calcium carbonate (CaCO_3), which precipitates as solid particles¹⁰¹.

The following reaction represents the overall process:



Now, methane (CH₄) is not directly generated through the MICP process. Complex biochemical processes are usually involved in the formation of methane, and this process is usually linked to the anaerobic breakdown of organic matter by certain microorganisms called methanogens. Methane can be produced in subsurface porous media through processes like microbial methanogenesis if organic matter is present and the environment is conducive to methanogenic activity¹⁰². This is a different kind of microbiological process from MICP that breaks down organic materials into carbon dioxide and methane¹⁰³.

In conclusion, MICP is not a direct source of methane in subsurface porous media, even though it helps in the formation of carbonate minerals during CO₂ sequestration¹⁰⁴. Methane production is more strongly linked to particular microbial processes involved in the anaerobic breakdown of organic materials¹⁰⁵.

Discussion

In MICP, urease-producing bacteria are usually the driving force behind the formation of carbonate minerals through microbial activity. This process affects porosity, subsurface CO₂ sequestration, and the production of biomethane, but it also has potential uses in soil stabilization and groundwater remediation. In addition to changing fluid flow patterns and causing cementation and pore blockage, these effects may have an impact on the efficiency of methane production and CO₂ injection¹⁰⁶. MICP lowers subsurface formation porosity, which affects the effectiveness of CO₂ sequestration. Reduced permeability, pressure accumulation, and possible seismicity may result from this. To avoid unforeseen consequences, it is essential to monitor changes in pressure. To stop the dissolution and re-release of CO₂, biogenically precipitated carbonate minerals must remain stable over the long term. Reduced porosity and permeability in subsurface formations can have an impact on fluid transport, microbial communities, and methane migration limitation¹⁰⁸. Porosity variations can impact methanogenic bacteria's access to nutrients, substrates, and electron acceptors. Carbonate mineralization-induced pore blockage may restrict the production of sustainable biomethane. Mitigation techniques include maximizing MICP conditions, keeping an eye on and modeling subsurface conditions, and investigating technical fixes like smart injection techniques or permeability enhancers. Maintaining sustainable CO₂ sequestration and bio-methane production requires striking a balance between the advantages of MICP and its effects on subsurface processes¹⁰⁹.

Future Direction

Studying the formation of methane and calcium carbonate from CO₂ sequestration in deep subsurface porous media has enormous potential to advance science and improve energy and environmental applications. The subsequent path of investigation encompasses comprehensive characterization of microbial communities, integration of omics, extended stability of biomineralized products, modeling and simulation, engineering tactics, biotechnological implementations, field-scale exhibits, interdisciplinary cooperation, and surveillance and risk evaluation¹¹⁰. These areas will center on addressing climate change, creating efficient and sustainable technologies for CO₂ sequestration and methane generation, and comprehending the complex interactions between microbes and minerals in deep subsurface porous media¹¹¹.

Incorporating omics data, evaluating the long-term stability of biomineralized products, creating sophisticated modeling and simulation techniques, investigating engineering tactics, and carrying out field-scale demonstrations are further areas of investigation for this research.

Conclusion

The intricate interactions between bacteria, minerals, and molecular mechanisms involved in biomineralization are highlighted in this review on the formation of methane and calcium carbonate derived from CO₂ sequestration in deep subsurface porous media. It draws attention to how microbial activity affects how methane and calcium carbonate are formed in these conditions. The review deciphered molecular biomineralization processes, shedding light on enzymes, and metabolic pathways used by microorganisms during CO₂ sequestration and methane generation. Additionally, it highlights how extracellular polymeric substances help in microbial interactions and biomineralization. This review examines how geochemical and environmental controls affect microbial activity and biomineralization, emphasizing the importance of sustainability and long-term stability. To fully comprehend CO₂ sequestration and methane generation in deep subsurface environments, the review also emphasizes the significance of field-scale demonstrations and interdisciplinary collaborations.

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