

# Barriers and Prospects of Carbon Sequestration in India

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Carbon sequestration is considered a leading technology for reducing carbon dioxide (CO<sub>2</sub>) emissions from fossil-fuel based electricity generating power plants and could permit the continued use of coal and gas whilst meeting greenhouse gas targets. India will become the world's third largest emitter of CO<sub>2</sub> by 2015. Considering the dependence of health of the Indian global economy, there is an imperative need to develop a global approach which could address the capturing and securely storing carbon dioxide emitted from an array of energy. Therefore technology such as carbon sequestration will deliver significant CO<sub>2</sub> reductions in a timely fashion. Considerable energy is required for the capture, compression, transport and storage steps. With the availability of potential technical storage methods for carbon sequestration like forest, mineral and geological storage options with India, it would facilitate achieving stabilization goal in the near future. This paper examines the potential carbon sequestration options available in India and evaluates them with respect to their strengths, weakness, threats and future prospects.

**Key words :** *Carbon sequestration, carbon capture and storage, carbon sources, CO<sub>2</sub> emission*

## 1 Introduction

India's economy is growing at the rate around 9% per annum, while the country is emitting significant amounts of greenhouse gases (GHGs) from fossil fuel use. The International Energy Agency (IEA) predicts that India will be one of the top three emitters of Greenhouse gases (GHG) in the world by 2030<sup>1</sup>. Therefore, India is a priority target for CO<sub>2</sub> reduction. However, almost half of India's households (56%) do not have electricity, and women and girls spend a total of 1 billion hours each year collecting firewood<sup>2</sup>.

The vast majority of India's population (70%) lives in rural areas and agriculture accounts for 35% of Gross National Product (GNP) and directly employs more than 60% of the Indian population<sup>1</sup>. According to the Intergovernmental Panel on Climate Change (IPCC), some of the most severe impacts of climate change will hit India's agriculture and natural resources<sup>3</sup>. For example, Himalayan glaciers are amongst the fastest retreating in the world; glacial melt water that feeds the major rivers on the sub-continent accounts for 37% of India's irrigated land, loss of this glacier melt water could cause water shortages for 500 million people<sup>3</sup>.

A high proportion of India's energy comes from coal, and the country's escalating fuel needs raise concerns of supply. Indian policymakers are taking a growing interest in promoting energy efficiency and renewable, as demonstrated by the National Action Plan on Climate Change<sup>4</sup>. The Indian government also plans to invest in several coal-fired Ultra-

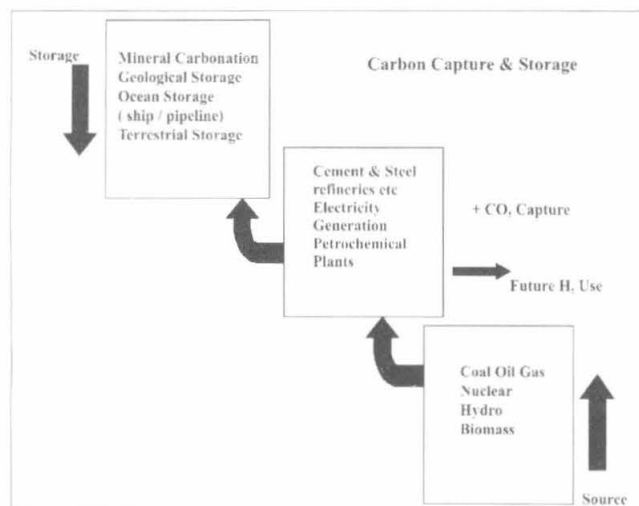
Mega Power Plants (UMPPs), with a power generating capacity of 4GW per unit accounting for 36GW overall. In 2006, the annual CO<sub>2</sub> emissions from large point sources in India was estimated to be 721 Metric tonne CO<sub>2</sub>, and a report by the IEAGHG<sup>5</sup> noted that this was roughly half of total emissions, which were approximately 1343 Metric tonne in 2004. Over half of India's current CO<sub>2</sub> emissions are from large point sources<sup>1</sup>. Such sources could be a suitable point for capturing emissions, transporting them, and then storing them in different forms like geological, oceanic, mineral and forest as a mitigation strategy.

CO<sub>2</sub> is emitted principally from the burning of fossil fuels, in large combustion units used for electric power generation and in smaller, sources such as automobile engines and furnaces used in commercial buildings. CO<sub>2</sub> emissions also result from some industrial processes, burning of forests and land clearance. Carbon Capture and Storage (CCS) is most likely to be applied to large point sources of CO<sub>2</sub>, which involves power plants and large industrial processes. These sources sometimes supply hydrogen as a fuel to the transportation, industrial and building, reducing emissions from the distributed sources. As illustrated in **Fig. 1**, there are three main components of the CCS process: capture, transport and storage. The capture step involves separating CO<sub>2</sub> from various gaseous products. The transport step involves carrying captured CO<sub>2</sub> to a suitable storage site located at a distance from the CO<sub>2</sub> source. Potential storage methods include injection into underground geological formations,

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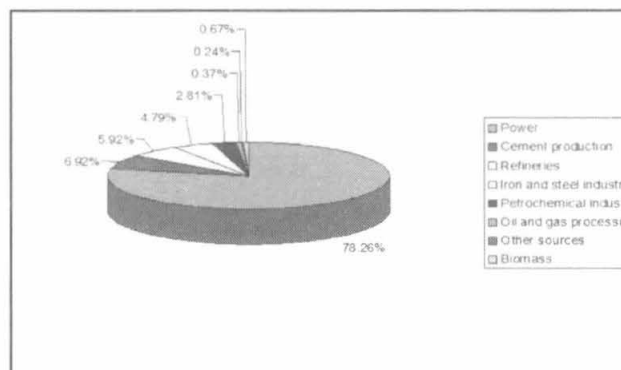
**Fig. 1 :** A schematic diagram of the possible carbon sequestration process, indicating the sources for which carbon sequestration might be relevant, as well as CO<sub>2</sub> transport and storage options

injection into the deep Ocean and industrial fixation. However, the technical maturity of specific CCS system components varies greatly.

### 1.1 Sources of CO<sub>2</sub>

The attractiveness of CO<sub>2</sub> source for capture depends on its volume, concentration and partial pressure, integrated system aspects, and its proximity to a suitable reservoir. Emissions of CO<sub>2</sub> arise from a number of sources, like fossil fuel combustion in the power generation, industrial, residential and transport sectors. Technological changes in the production and nature of transport fuels, however, may eventually allow the capture of CO<sub>2</sub> from energy use in this sector.

Global emissions of CO<sub>2</sub> from fossil-fuel use in the year 2000 totalled to about 23.5 Gega tonne CO<sub>2</sub> yr<sup>-1</sup> (6 Gega tonne C yr<sup>-1</sup>)<sup>1</sup>. Of this, close to 60% was attributed to large stationary emission sources (**Fig. 2**). Increased fuel utilization for electric power generation and transportation, resulted in energy related CO<sub>2</sub> emission. Increased demand for electricity for computers, electronics in homes and offices; demand for commercial lighting and cooling; substitution of electricity-intensive technologies, such as electric arc furnaces for steelmaking, in the industrial sector; and increased demand for transportation services have led to increased emissions of CO<sub>2</sub>. Though not all these sources are capable of CO<sub>2</sub> capturing. Coal which is the most dominant fuel in the power sector, accounts for 38% of total electricity generated in the year 2000, followed by hydro power (17.5%), natural gas (17.3%), nuclear (16.8%), oil (9%), and finally non-hydro renewable (1.6%)<sup>6</sup>. Fuel selection is greatly sector-specific



**Fig. 2 :** Profile by process or industrial activity of worldwide large stationary CO<sub>2</sub> sources with emissions of more than 0.1 million tonne CO<sub>2</sub> per year (Data taken from IPCC 2005<sup>6</sup>)

like the use of blast furnaces in the iron and steel industry using coal and coke as their primary fuel for the production of steel<sup>7-8</sup>. Refining industry on the other hand uses oil and gas as their fuel. Cement manufacturing industries use fossil fuels such as coal<sup>9</sup>. On the other hand, European cement manufacturing processes utilize non fossil fuel based fuels like tyres, sewage sludge and chemical-waste mixtures<sup>9</sup>. Other countries like Scandinavia and Brazil, use biomass as a fuel

The emission sources include mainly large stationary sources which involve the use of fossil fuel and biomass. From combustion activities, natural gas processing and industrial processes are some of the large stationary sources resulting in CO<sub>2</sub> emission. Carbon dioxide emissions also take place from a variety of industrial production processes involving transformation of materials by chemical, biological and physical processes.<sup>7, 9, 11, 12</sup>

### 1.2 CO<sub>2</sub> emission : future sources

The IPCC Special Report on Emission Scenarios (SRES) projects the future emissions of CO<sub>2</sub> on the basis of six illustrative scenarios where global CO<sub>2</sub> emissions range from 29 to 44 Gega tonne CO<sub>2</sub> per year in 2020, and from 50 to 84 Gega tonne CO<sub>2</sub> per year in 2050<sup>3</sup>. It is projected that in the South and East Asian countries the electric power and industrial sectors will account as significant CO<sub>2</sub> emission sources by 2050, whereas in Europe there may be a slight decrease in such sources. Great amount of CO<sub>2</sub> will be produced from higher number of plants installed making CO<sub>2</sub> sequestration process more feasible technology.

### 1.3 Energy and power sectors as major CO<sub>2</sub> sources in India

India depends largely on fossil fuel based energy input system in major industrial as well as domestic sectors. Coal is the main energy source in India dominating

current Indian energy sector<sup>1</sup>. India has the world's fourth largest hard coal reserves and is the third largest coal producer after China and the USA. The IEA<sup>1</sup> predicts that the overall share of imports in Indian primary coal supply would increase from 12% in 2005 to 28% in 2030. CCS could play a major role in this sector particularly as regards upcoming power plants.

In terms of installed power-generating capacity, Indian power sector is ranked fifth in the world. In 2005, power stations emitted an average 943g of CO<sub>2</sub> per kWh of electricity produced in 2005<sup>1</sup>. This accounted for nearly 60% of India's total CO<sub>2</sub> emissions for the year 2005. In 2006, the Ministry of Power launched an initiative to develop Ultra Mega Power Plants (UMPPs) having a capacity of 4 GW. These UMPPs account for 257 Metric tonne CO<sub>2</sub> emissions and 36000 MWe installed capacity within 7-8 years<sup>5</sup>. These represent an example of a massive point source of CO<sub>2</sub> emissions<sup>13</sup> which may be considered as a globally competitive opportunity for CCS. However, India is still seriously struggling to provide electricity to all its citizens.

In India, oil and natural gas are among the main energy sources. The total oil production in the country from 2003 to 2008 is shown in Fig. 3. The total oil production in 2008 for India was 880,500 bbl/day. The total oil consumption of India from 2003 to 2008 is shown in Fig. 4. The total oil consumption in 2008 for India was 2.722 million bbl/day. According to the IEA<sup>1</sup> by the end of 2006, the current reserves accounted approximately 0.4% of the world's reserves. As a result of which India has to import about 100 million tonnes of crude oil from the Middle East. Though India's current oil import dependence is about 70%, which is projected to rise to 94% by 2030<sup>1</sup>.

## 2.0 CO<sub>2</sub> capture and storage : general trends

CO<sub>2</sub> capture and storage process involves capturing CO<sub>2</sub> from large sources like power plants to produce a concentrated stream that can be easily transported to a CO<sub>2</sub>

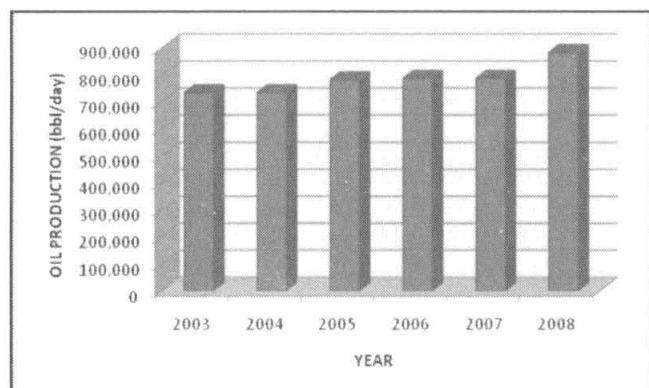


Fig. 3 : Total oil production in India <sup>42</sup>

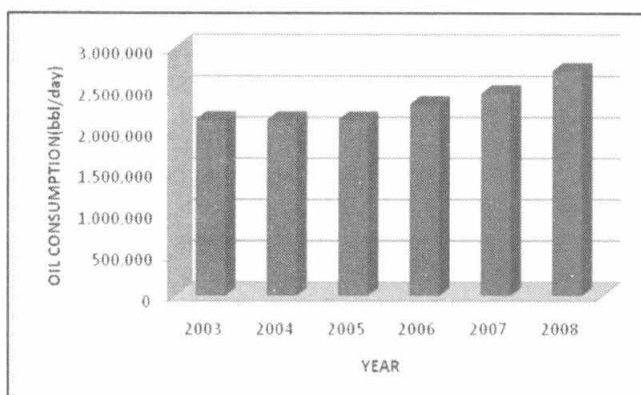


Fig. 4 : Total oil consumption in India<sup>42</sup>

storage site. This capturing technology is a way forward for development of new technology involving production of low-carbon or carbon-free electricity and fuels requirement for transportation, as well as for small-scale applications. Energy requirement for CO<sub>2</sub> capture operation systems reduces the overall efficiency of power generation leading to increase in the fuel requirements, solid waste generation and environmental impacts relative to the same type of plant without capture.

CO<sub>2</sub> is separated at some large industrial plants such as natural gas processing and ammonia production facilities, to meet process demands and not for storage. CO<sub>2</sub> capture has also been applied to several small power plants. There are mainly three main CO<sub>2</sub> capturing processes for industrial and power plant applications : (1) Post combustion systems which involve separation of CO<sub>2</sub> from the flue gases produced by combustion of primary fuel like coal, natural gas and oil in air; (2) Oxy-fuel combustion system utilizing oxygen for combustion and producing mainly H<sub>2</sub>O and CO<sub>2</sub>, commonly called syngas which can be readily captured; and (3) Pre-combustion systems process involving combustion of primary fuel in a reactor producing separate streams of CO<sub>2</sub> and H<sub>2</sub> which can be used as a fuel.

## 2.1 CO<sub>2</sub> storage systems

### 2.1.1 Geological storage

Geological storage of CO<sub>2</sub> is quite widely regarded as a potentially important mitigation option for various reasons like: (1) various commercial projects have been successfully undertaken and demonstrated increasing the level of confidence in the technology; (2) broader aspect of mitigation options is required; and (3) large CO<sub>2</sub> emission reduction in the atmosphere. The technique is safe, environmentally sustainable, and cost-effective and has broad applications. To store CO<sub>2</sub> geologically, it is normally

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compressed to a dense fluid state typically known as supercritical.

Geological storage of CO<sub>2</sub> is undertaken in a variety of geological settings such as oil fields, depleted gas fields, deep coal seams and saline formations are all possible storage formations as illustrated in Fig. 5. Subsurface geological storage is possible both onshore and offshore (continental shelf, some adjacent deep-marine sedimentary basins, storage in caverns, basalt and organic-rich shale's)<sup>14</sup>.

Injection of CO<sub>2</sub> has been done at a relatively small scale to significantly decrease emissions from the existing stationary sources. The injection rates would have to be at par with other injection operations at present. Unminable coal seams act as a potential source to capture and store because CO<sub>2</sub> adsorbs to the surface of coal. However, it depends largely on various geological and physical factors like the permeability of the coal bed. The process releases methane, and the methane can be recovered. Saline aquifers have common occurrence with large potential storage volume, however, little geological information is available about them as compared to oil fields. Major disadvantage is the leakage of CO<sub>2</sub> back into the atmosphere may be a problem in saline aquifer storage.

CO<sub>2</sub> storage is an integral part of the CCS chain. Attempts have been made at evaluating the geological storage potential in India. Unminable coal seams could store approximately 5 Gega tonne of CO<sub>2</sub>, depleted oil and gas reservoirs accounting for 7 Gega tonne of CO<sub>2</sub>, in deep saline aquifers accounting for 360 Gega tonne of CO<sub>2</sub>, and mineralization in basalt rocks accounting for 200 Gega tonne of CO<sub>2</sub><sup>15</sup>.

### 2.1.2 Ocean storage

Captured CO<sub>2</sub> can be injected into the ocean bed at great depth, where it would remain isolated from the atmosphere for many years. CO<sub>2</sub> is transported by pipeline or ship for its final release in the ocean bed. Small-scale field experiments along with various other theoretical calculations and laboratory modelling techniques for ocean storage of CO<sub>2</sub> have been studied, yet it has not been thoroughly tested. Several concepts have been proposed so far in respect to the disposal of CO<sub>2</sub> to the ocean/ sea bed like: (1) Lake deposits CO<sub>2</sub> directly onto the sea floor at depths greater than 3000 m, (2) Storage of CO<sub>2</sub> in solid clathrate hydrates<sup>16, 17</sup>, (3) Conversion of CO<sub>2</sub> to bicarbonates and (4) Dissolution injects CO<sub>2</sub> at depths of 1000 m or more where the CO<sub>2</sub> dissolves.

CO<sub>2</sub> is captured, compressed and transported to the deep ocean for release at or above the sea floor<sup>18,19</sup>. CO<sub>2</sub> then dissolves into the surrounding sea water and becomes a part

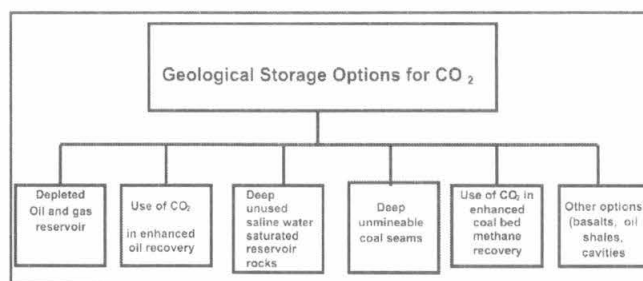


Fig. 5 : Options for storing CO<sub>2</sub> in deep underground geological formations

of the ocean carbon cycle<sup>20</sup>. Effectiveness of ocean storage depends on how long CO<sub>2</sub> remains isolated from the atmosphere. Release of CO<sub>2</sub> to the deep ocean results in eventually mixing of the gas throughout the oceans, finally affecting the concentration of CO<sub>2</sub> in the atmosphere. The main principle is the transfer of CO<sub>2</sub> to deep waters so that higher degree of isolation from the atmosphere could be achieved with an increase in the depth of the ocean.

#### 2.1.2.1 CO<sub>2</sub> storage by dissolution of carbonate minerals

Increased sea water acidity resulting from CO<sub>2</sub> addition will be greatly neutralized by the slow natural dissolution of carbonate minerals in sea-floor sediments. This allows the ocean to absorb more CO<sub>2</sub> from the atmosphere with less of a change in ocean pH and carbonate ion concentration<sup>21,22,23,24,25,26,27</sup>. Carbonate neutralization approaches attempt to promote reaction in which limestone reacts with carbon dioxide and water to form calcium and bicarbonate ions in solution<sup>23</sup>. For each mole of CaCO<sub>3</sub> dissolved there would be 0.8 mole of additional CO<sub>2</sub> stored in sea water<sup>23</sup>. The environmental effects of oceanic storage are: (1) Large concentrations of CO<sub>2</sub> kill ocean organisms, (2) Storage is not a permanent option as dissolved CO<sub>2</sub> will eventually equilibrate with the atmosphere, (3) Acidity of the ocean water increases with formation of carbonic acid on reaction of CO<sub>2</sub> with H<sub>2</sub>O, and (4) Effect on benthic life is poorly understood. Much more work is needed to define the extent of the potential problems.

#### 2.1.3 Mineral storage

Carbon sequestration by reacting naturally occurring Mg and Ca containing minerals with CO<sub>2</sub> to form carbonates is called mineral sequestration. This process occurs naturally and is similar to the weathering of rock over geologic time frame. Carbonates formed have a lower energy state than CO<sub>2</sub>, thus making the carbonation reaction thermodynamically favourable. The raw materials which are mainly magnesium and calcium based minerals are widely abundant and available. Carbonates are stable and thus the re-release of CO<sub>2</sub> into the atmosphere is not the matter of concern. Carbonation pathway

is normally a slow process under ambient temperatures and pressure conditions. Therefore, the primary objective is to address and investigate an industrially applicable and environmentally feasible/potential carbonation pathway that will allow mineral sequestration to be implemented at an economical platform. Mineral sequestration process involves reaction of CO<sub>2</sub> (exothermic reaction) with widely available and abundant metal oxides to produce a stable end product, carbonate.

The reaction rate can be accelerated by carrying out the reaction at higher temperatures, pressures, and by pre-treatment of the minerals. However this is an energy intensive method. According to IPCC, power plant equipped with Carbon Capture and Storage (CCS) utilizing mineral storage process would eventually require 60-180% more energy than a power plant without CCS.

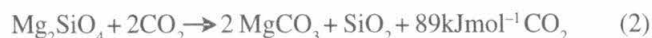
### 2.1.3.1 Chemistry of mineral carbonation

Reaction of CO<sub>2</sub> with metal oxides results in the formation of the corresponding carbonate and subsequent heat is released according to the chemical reaction as shown below.

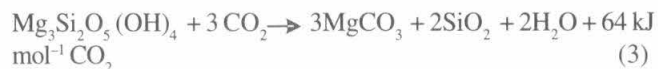


Where metal oxide is indicated as AO, A is the divalent metal, e.g., calcium, magnesium, or iron. Heat released during the reaction depends on the specific metal constituting the mineral as well as the mineral containing that metal oxide. The heat released during the combustion of elemental carbon is 393.8 kJ mol<sup>-1</sup>CO<sub>2</sub>. Following exothermic reaction takes place in few silicates mineral where the heat values are given per unit mol of CO<sub>2</sub>. The reaction takes place under temperature (25°C) and pressure conditions (0.1 MPa) as described below<sup>28</sup>.

#### Olivine:



#### Serpentine:



#### Wollastonite:



The reaction releases heat favouring the formation of carbonates at low temperature conditions. The reaction reverses under high temperature conditions favouring the calcinations process. Under ambient temperature and low partial pressure of CO<sub>2</sub> conditions, carbonation of minerals bearing the respective metal oxide occurs readily<sup>28,29</sup>. Silica or carbonate layers formation on the mineral surface during

carbonation hinders reaction<sup>30</sup> and rate of CO<sub>2</sub> uptake. The challenge for mineral carbonation is to find out ways to accelerate carbonation process for minimum energy and material losses.

India has about twenty-five major greenstone belts containing Ca, Fe and Mg silicate rich minerals such as olivine, serpentine, and pyroxene etc. The alkaline silicates may act as potential and feasible sinks to sequester CO<sub>2</sub> in the form of magnesium, iron or calcium carbonates.

### 2.1.4 Forests and carbon storage

Forests can store much carbon and their growth can be considered as a potential carbon sink. Fossil fuel emissions of carbon currently offset 310 million metric tons of U.S.<sup>31,32</sup>. A carbon balance of near zero could be achieved by large forested landscape over long period of time<sup>33,34</sup>.

In India the effects of climate on carbon reserves in cultivated soils has been studied in the past<sup>35</sup>. Several investigations and studies based on Indian forest carbon pools<sup>36,37,38</sup> and litter fall studies<sup>39</sup> are widely available and reported in literature. However, soil organic carbon (SOC) pool estimates based on Indian database with reference to forest types have not been undertaken for research so far. The total SOC stocks in India range from a wide range of 23.4–47.5 Pg C<sup>39,40</sup> and 5.4–6.7 Pg C<sup>36,39</sup>. However, India lack carbon inventories to monitor and enhance carbon sequestration potential of forests. Attempts at macro level have been made in India to carry out carbon sequestration studies<sup>36-41</sup>. However, attempts to assess the biomass and soil carbon sequestration at micro-level are still lacking.

## 3.0 Effectiveness of various sequestration options

CO<sub>2</sub> is released directly to the atmosphere mainly from human sources by burning of fossil fuels. Various anthropogenic sources release GHG to the atmosphere. By using a combination of several mitigation methods it is possible to achieve reduction of these emissions as illustrated in **Table 1**. The relationship between the concentration of CO<sub>2</sub> in the atmosphere and the storage of CO<sub>2</sub> can be clearly studied by understanding the relation between the stock of CO<sub>2</sub> stored and its flow in the respective reservoir.

The stock or amount of CO<sub>2</sub> stored over particular time frame in a given reservoir can be calculated by noting the difference between the current CO<sub>2</sub> stock and the rate at which CO<sub>2</sub> is being added or released. As far as the input storage rate of CO<sub>2</sub> in the reservoir exceeds the amount it is released back, CO<sub>2</sub> will certainly be accumulated in the given reservoir formation.

## 4.0 Summary

The various strengths, weaknesses, threats and future prospects of carbon sequestration technologies in

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**Table 1:** CO<sub>2</sub> storage potential of various options

Sino	Storage Option	CO <sub>2</sub> Stock	Storage Potential
1	Terrestrial biosphere	Stores and releases both natural and fossil fuel CO <sub>2</sub> through the global carbon cycle.	99% is stored for decades to centuries, although the average lifetime will be towards the lower end of that range.
2	Oceans storage	Absorb and release natural and fossil fuel CO <sub>2</sub> according to the dynamics of the global carbon cycle	Fraction retained by ocean storage at 3,000 m depth could be around 85% after 500 years
3	Geological storage	CO <sub>2</sub> has been in natural geological reservoirs for millions of years.	Fraction retained exceeds 99% over 1000 years
4	Mineral Carbonation	Carbonation reaction between CO <sub>2</sub> and mineral rock.	Fraction retained of early 100% for exceptionally long times in carbonate rock.
5	Converting carbon dioxide into other forms	Chemical reaction	Result in very small net storage of CO <sub>2</sub> . 99% of the carbon will be retained in the product for periods in the order of weeks to months, depending on the product.

**Table 2:** Overview of carbon storage technologies in India

Property	Terrestrial biosphere	Deep ocean	Geological reservoirs	Mineral Carbonation
<b>STRENGTHS</b>				
CO <sub>2</sub> sequestered or stored	Stock changes can be monitored over time.	Injected carbon can be measured	Injected carbon can be measured	Injected carbon can be measured.
Ownership	Stocks have discrete location	Stocks will be mobile and may reside in international waters.	Stocks may reside in reservoirs and differ from surface boundaries.	Stocks have a discrete location
Management decisions	Storage will be subject to continuing decisions about land use priorities.	Once injected, no further human decisions on maintenance.	Once injected, human decisions to influence continued storage involve monitoring and perhaps maintenance, unless storage interferes with resource recovery.	Once injected, human decisions to influence continued storage involve monitoring and perhaps maintenance, unless storage interferes with resource recovery.
Monitoring	Changes in stocks can be monitored.	Changes in stocks can be modelled.	CO <sub>2</sub> release might be detected by physical monitoring.	Changes in stocks can be monitored
Time scale with expected high values for fraction CO <sub>2</sub> retained	Decades	Centuries	Very small leakage	Centuries

Table 2 (...contd)

<b>THREATS</b>	1. Due to these horrible large-scale forest fires, the forest themselves become the sources of high CO <sub>2</sub> emissions in the atmosphere, thus adding to the problem of global warming. So it is certainly not very much exciting to rely upon the terrestrial ecosystems for a future reliable source of Carbon Sink	1. The injection of a few GtCO <sub>2</sub> would produce a measurable change in the ocean chemistry in the region of injection, whereas the injection of hundreds of GtCO <sub>2</sub> would produce larger changes in the region of injection and eventually produce measurable changes over the entire ocean volume. 2. Adding CO <sub>2</sub> can harm marine organisms. Effects of elevated CO <sub>2</sub> levels includes reduced rates of calcification, reproduction, growth, circulatory oxygen supply and mobility, as well as increased mortality over time.	1. Injection well failures or leakage up abandoned wells could create a sudden and rapid release of CO <sub>2</sub> , and can be dangers to human life and health. 2. Leakage may also occur through undetected faults, fractures or through leaking.. It may affect drinking- water aquifers and ecosystems 3. Groundwater can be affected both by CO <sub>2</sub> leaking 4. Acidification of soils and displacement of oxygen in soils.	1. The mineral carbonation process would require 1.6 to 3.7 tonnes of silicates per tonne of CO <sub>2</sub> to be mined, and produce 2.6 to 4.7 of materials to tonnes be disposed per tonne of tonne of CO <sub>2</sub> stored as carbonates. This would have environmental impact similar to surface mining operations. 2. Serpentine often contains chrysotile, therefore demands monitoring and mitigation measures 3. The environmental impact of mining, waste disposal and product storage could also limit potential. 4. Leaching could also cause containment problem.
<b>WEAKNESS</b>				
Cost	Low-cost	Costs for deep ocean disposal of liquid CO <sub>2</sub> are estimated at US\$40-80/ton.	High-cost	Low-cost
Time	Lifetimes of 10–100 years	The time it takes water in the deeper oceans to circulate to the surface has been estimated to be in the order of 1600 years, varying upon currents and other changing conditions.	CO <sub>2</sub> retained in appropriately selected and managed geological reservoirs is very likely to exceed 99% over 100 years and is likely to exceed 99% over 1,000 years.	Fraction of carbon dioxide stored through mineral carbonation that is retained after 1000 years is virtually certain to be 100%.
Source of CO <sub>2</sub>	Non- Point Source	Point /stationary Source	Point/stationary Sources	Point Source
<b>FUTURE PROSPECTS</b>	1. Terrestrial and agro-forestry modelling studies to enhance sequestration rate of CO <sub>2</sub> in ecosystem can be explored in future. 2. Regenerative carbon dioxide removal by chemical and biological	1. Iron Fertilization implies the introduction of iron to the upper ocean to increase productivity of marine food chain which in turn increases sequestration from atmosphere into the	1. Evaluation of Basalt Formations of India for environmentally safe and irreversible long time storage of CO <sub>2</sub> . 2. Scoping studies for CO <sub>2</sub> injection for EOR in various	1. Materials Research for Cost Effective Carbon Capture Process Development 2. Biomimetic sequestration implies the use of a particular aspect of biological process for resolving

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Table 2 (contd...)

<p>means are other areas that can be explored for further research.</p> <p>3. Micro-algae and Microbial Fixation of CO<sub>2</sub> from Industrial effluents as well as adaptive response of microalgae to high CO<sub>2</sub> coupled with high temperature and reduced pH posses a great potential in the future.</p> <p>4. Enhancing carbon storage in degraded dry lands could have direct environmental, economic and social benefits for local people. It could increase benefits for farmers as well as mitigate global warming, at least in the coming decades until alternative energy sources are developed.</p> <p>5. The wasteland available for forestation is quite large, 30 × 10<sup>6</sup> ha. An energy plantation can be used for power generation in applications, instance :Co-firing in fossil fuel fired power in agriculture and generation, Cogeneration forestry processing facilities, Stand alone grid connected biomass based power stations that consider energy plantation feed stocks along with other agriculture residues, like baggase, wood waste, rice hull, etc.</p>	<p>oceans. This technology proves to be beneficial in future.</p> <p>2. Marine plankton growth is enhanced by for physically distributing the iron particles in other wise nutrient rich but iron deficient ocean water using suitable delivering systems based on biomaterials.</p> <p>3. Studies of the response of biological systems in the deep sea to added CO<sub>2</sub>, including studies that are longer in duration and larger in scale than yet performed.</p> <p>4. Research facilities: Research facilities where ocean storage concepts can be applied and their effectiveness and impacts assessed in situ at small- scale on a continuing basis for the purposes of both scientific research and technology development.</p> <p>5. Investigation and development of technology for working and the development in the deep sea, of pipes, nozzles, diffusers which can be deployed in the deep sea with assured flow and be operated and maintained cost-effectively.</p> <p>6. Development of techniques and sensors to detect CO<sub>2</sub> plumes and their biological and geochemical consequences.</p>	<p>specific fields proves to be beneficial.</p> <p>3. Research on underground disposal and coal bed methane needs to be undertaken.</p> <p>4. Collaborative research on screening criteria development for Geological sequestration in Saline aquifers needs to be initiated.</p> <p>5. More research needs to be carried out in future geophysical sounding and deep resistivity surveys using Schumberger method for preparation of 2D and 3D models of subsurface lithography up to the depth of 800 m.</p> <p>6. The Central Ground Water Board and Geological Survey of India have established the presence of saline aquifers up to depths of ≥ 300m belowground level in the Ganga basin. Deep Resistivity studies carried out at different sites in various regions have shown the presence of saline aquifers at depths of 800m and beyond.</p> <p>7. The sale of the methane can be used to offset a portion of the cost of the CO<sub>2</sub> storage.</p> <p>8. There is strong evidence that storage of CO<sub>2</sub> in geological storage sites will be long term.</p>	<p>a specific non biological problem. The Carbonic Anhydrase (CA) is used as a catalyst for the conversion of CO<sub>2</sub> into bicarbonates and later to carbonates or amino acids. This new area can be explored for research</p> <p>3. The carbonates formed are stable and the disposal is therefore permanent. There is no chance of the carbon dioxide escaping into the atmosphere.</p> <p>4. Carbonate is the lowest energy state of carbon, not carbon dioxide.</p> <p>5. Mineral carbonation occurs naturally on a geological time scales and would eventually absorb all the additional carbon dioxide.</p> <p>6. The process is just speeding up one that occurs in nature. The minerals are readily accessible in locations near high- density power generation centres. There is potential to produce value-added by products.</p> <p>7. The process is compatible with both technologies under development and current power systems. Predicted cost is not unreasonable.</p> <p>8. No heat required reaction is exothermic.</p>
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Indian perspective are illustrated in **Table 2**. CCS technology involves various components like carbon capture and sequestration, transportation of CO<sub>2</sub> via pipeline and injection into geological formations. These are well proven in India but the number of such final CCS projects is very small as compared to what is currently required to achieve significant CO<sub>2</sub> emission reductions in the atmosphere. New technologies that make CCS much cheaper would be more helpful in getting it adopted quickly in terms of economics of the technology. India has low oil resources, therefore, enhanced oil recovery (EOR), and cannot be completely adopted for future sequestration in India.

In order to mitigate adverse affect of climate change, India needs to adopt a sustainable development approach by making use of environmental friendly technologies. Energy efficient technologies in conjugation with sustainable pathway will help to reduce the vulnerability of natural as well as other socio-economic system. In Indian context, an amalgamation of both geological and mineral sequestration technology should be promoted and considered for future mitigation of GHG emissions. However, a tailored approach for overcoming the adverse response of various anthropogenic emissions is needed for a large country size, like India. Though it is impossible to project how many Indian CCS plants will be installed on short term basis but in the long term, it can be realized if we start taking the right steps now.

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## Corrigendum

Please refer the article titled "Modeling and Simulation of Road Traffic Noise Using Artificial Neural Network and Regression" authored by M. Honarmand and S.M. Mousavi, published in Volume 56 No. 1, p. 1-6, January 2014 of *Journal of Environmental Science & Engineering*. In this context, the readers of the journal are requested to kindly read the revised article being published in the following pages.

The inconvenience caused is regretted.

- Editor, JESE