

**Utilizing Ionic liquid-Nonthermal Plasma Combination for CO<sub>2</sub> Conversion**Pankaj Attri<sup>1\*</sup>, Takamasa Okumura<sup>1</sup>, Kunihiro Kamataki<sup>1</sup>, Kazunori Koga<sup>1</sup>, and Masaharu Shiratani<sup>1</sup>  
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Global energy consumption will increase by 28% between 2015 and 2040, according to a report by the U.S. Department of Energy's Energy Information Administration (EIA). The increased energy use will be matched by a 16% increase in energy-related carbon dioxide (CO<sub>2</sub>) emissions over that same period, with annual emissions rising from 33.9 billion metric tons in 2015 to 39.3 billion metric tons in 2040, according to EIA's report. Over the past decade, global CO<sub>2</sub> emissions increased by 50 % compared to the levels observed in 1990. The primary source of elevated CO<sub>2</sub> emissions is the use of fossil fuels for heat and power generation. Coal combustion as the major contributor is responsible for approximately 43 % of CO<sub>2</sub> emissions, while oil contributes around 37 %, and gas accounts for roughly 20 % of overall CO<sub>2</sub> emissions [1].

Therefore, there is an urgent need for prompt action from governments and industries across the globe to reduce the rising atmospheric concentrations of greenhouse gases. This emphasizes that there is a critical necessity to adopt negative emission technologies for achieving a net removal of greenhouse gases from the environment [2]. However, the transition of the energy infrastructure away from fossil fuels and toward renewable and nuclear energy resources is not an easy task. A potential solution lies in CO<sub>2</sub> capture from point sources, such as power plants and industries, which could play a crucial role in minimizing CO<sub>2</sub> emissions throughout this transition by integrating localized CO<sub>2</sub> capture and storage (CCS) technology [3]. The technology for capturing CO<sub>2</sub> from ambient air, known as "direct air capture" (DAC), has been advancing rapidly in recent years [4]. Despite significant advances in CCS technologies, there are still substantial limitations, such as high capital costs, reduced absorption, and desorption rates, solvent evaporation, etc. Ionic liquids (ILs) have recently gained considerable interest in CO<sub>2</sub> capture [5]. The absorption of CO<sub>2</sub> by ILs results from physical interactions between the anions and cations of the ILs and CO<sub>2</sub> molecules [6].

Plasma, known as the "fourth state of matter," is an ionized plasma made up of neutral ground-state molecules in addition to electrons, different types of ions, radicals, excited atoms, and molecules. The electrons may activate the CO<sub>2</sub> molecules and create new products without heating the entire gas, plasma is exciting for the conversion of CO<sub>2</sub> [7]. In our research, we have made significant progress by utilizing streamer plasma for the conversion of both CO<sub>2</sub> and N<sub>2</sub> into valuable plant

nutrients [8]. In another study, we have shown how humidity is utilized in CO<sub>2</sub> conversion [9].

Nevertheless, the simultaneous capture and conversion of CO<sub>2</sub> using NTP is of great interest. Hence, in this pioneering effort, we employed an IL for CO<sub>2</sub> capture and storage followed by the conversion of the captured CO<sub>2</sub> into CO using NTP. In this study, we utilized the 1-Butyl-3-methylimidazolium chloride [Bmim]Cl IL, to convert captured CO<sub>2</sub> to CO. We observed that integrates plasma and [Bmim]Cl IL for the simultaneous capture, storage, and conversion of CO<sub>2</sub>. Our experimental results indicate that, under atmospheric pressure and room temperature, the water + [Bmim]Cl solution can store CO<sub>2</sub>. Moreover, the release of CO<sub>2</sub> during plasma treatment produces CO. Our MD simulation supports our experimental findings, suggesting that CO<sub>2</sub> molecules easily transition from the gaseous phase into the water + [Bmim]Cl solution. In contrast, the penetration of CO molecules into the water + [Bmim]Cl solution is more challenging compared to water alone. This observation implies that once plasma produces CO, its solubility in the IL solution may be limited, showcasing the potential of this technology for efficient CO<sub>2</sub> capture and conversion.

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**References**

- [1] Y. Yang, et al, Journal of Cleaner Production 376, 134347 (2022).
- [2] P. Smith, et al, Nature Climate Change 2015 6:1 6 [1], 42 (2015).
- [3] J. L. Fan, et al, Nature Climate Change 13 [8], 807 (2023)
- [4] J. Sun, et al, Current Opinion in Green and Sustainable Chemistry 40, 100752 (2023).
- [5] W. Faisal Elmobarak, et al, Fuel 344, 128102 (2023).
- [6] W. U. Mulk, et al, Journal of CO2 Utilization 75, 102555 (2023)
- [7] R. Snoeckx and A. Bogaerts, Chemical Society Reviews, 2017, 46, 5805–5863
- [8] P. Attri, et al, Frontiers in Physics 11, 1211166 (2023).
- [9] P. Attri, et al, Plasma Processes and Polymers 21 [1], e2300141 (2024).