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Scaling green hydrogen and CCUS via cement–methanol co-production in China

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High costs of green hydrogen and of carbon capture, utilization, and sequestration (CCUS) have hindered policy ambition and slowed real-world deployment, despite their importance for decarbonizing hard-to-abate sectors, including cement and methanol. Given the economic challenges of adopting CCUS in cement and green hydrogen in methanol production separately, we propose a renewable-powered co-production system that couples electrolytic hydrogen and CCUS through molecule exchange. We optimize system configurations using an hourly-resolved, process-based model incorporating operational flexibility, and explore integrated strategies for plant-level deployment and CO₂ source-sink matching across China. We find that co-production could reduce CO₂ abatement costs to \$41–53 per tonne by 2035, significantly lower than approximately \$75 for standalone cement CCUS and over \$120 for standalone renewable-based methanol. Co-production is preferentially deployed at cement plants in renewable-rich regions, potentially reshaping national CO₂ infrastructure planning. This hydrogen–CCUS coupling paradigm could accelerate industrial decarbonization and scaling for other applications.

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Broader context

Industrial sectors such as cement and chemicals are central to economic development but account for a growing share of global CO₂ emissions. Green hydrogen and carbon capture, utilization and storage (CCUS) are widely viewed as essential options for deeply decarbonizing these “hard-to-abate” industries, yet their high costs and infrastructure requirements remain major barriers to deployment at scale. New opportunities emerge when green hydrogen production is coupled with oxy-fuel cement kilns and CO₂ capture, so that by-product oxygen and captured CO₂ are reused within an integrated co-production system. Here we show how industrial symbiosis between cement production and methanol synthesis can create bidirectional molecular flows and lower costs. We combine an hourly-resolved plant-level optimization model with CCS source–sink matching to explore system configurations, flexible operation under variable renewable electricity, and plant-by-plant deployment pathways for China’s cement fleet. Our results show that co-producing clinker and green methanol cuts costs significantly relative to decarbonizing cement and methanol separately, while reshaping the national CO₂ network. Beyond this specific case, the framework shows how coupling hydrogen, CO₂ and O₂ flows across sectors can inform infrastructure planning and circular-economy policy for industrial decarbonization.

1. Introduction

Delivering on global climate targets requires rapid and far-reaching emissions reductions across all sectors, including heavy industries.¹ Green hydrogen and CCUS are widely regarded as indispensable for decarbonizing sectors such as steel, cement, and chemicals, where fossil feedstocks, high-temperature heat, and process-related carbon dioxide (CO₂) emissions limit the effectiveness of electrification and efficiency improvements.^{2,3} In response, governments have announced increasingly ambitious policy targets, such as the EU’s goal of producing 10 Mt of renewable hydrogen and building 50 Mt of CO₂ injection capacity annually by 2030.^{4,5} However, real-world progress falls short of

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expectations and remains far from what net-zero pathways require due to high costs, infrastructure gaps, and uncertain market incentives.^{6,7}

Identifying and scaling early applications of electrolytic hydrogen and CCUS that are both economically viable and replicable can accelerate technological learning and enable broader adoption. One promising approach is to exploit industrial symbiosis, where distinct industrial processes are linked by exchange of mass and/or energy for mutual benefit.⁸ Existing forms can improve resource efficiency, lower costs, and reduce emissions, as illustrated by cogeneration of heat and power,⁹ utilization of petroleum coke for aluminum anodes¹⁰ and substitution of cement clinker with steel slag.¹¹ However, many of these have limited potential because they serve niche markets or operate only in highly specialized contexts. Symbiotic coupling of electrolytic hydrogen and CCUS through molecular exchange and process integration, as proposed here, could unlock novel and scalable applications of these decarbonization-enabling technologies. The oxygen (O₂) co-produced from water electrolysis can supply oxy-fuel combustion CCUS systems (e.g. high-temperature cement kilns),¹² thereby increasing flue-gas CO₂ concentration and reducing or even eliminating the need for air separation units (ASUs). Meanwhile, the captured CO₂ can serve as a carbon feedstock, which together with electrolytic hydrogen, can be used to synthesize methanol and other hydrocarbon-based chemicals, reducing or eliminating CO₂ transport and sequestration costs. In contrast to the predominantly one-way material or energy flows in most industrial symbiosis, coupling green hydrogen and CCUS could create bidirectional molecular linkages. Symbiotic coupling of cement and methanol production integrated with green hydrogen and CCUS is proposed here as a strategy for decarbonizing those industries while also supporting commercial development of green hydrogen and CCUS technologies.

In 2022, cement and methanol production globally emitted approximately 2.4 Gt CO₂¹³ and 0.26 Gt CO₂,¹⁴ respectively, together accounting for around 7% of energy related emissions globally. China accounted for 57% of cement and 37% of the methanol production (2020),^{15,16} with a large share of carbon-intensive coal-based routes. For cement, efficiency improvements, clinker substitution, and fuel switching can reduce fossil energy use, but roughly two thirds of emissions are from carbonate calcination and essentially unavoidable without capture, which makes CCUS indispensable.¹⁷ For methanol, dependence on fossil derived hydrogen and carbon feedstocks means that deep reductions require cleaner inputs, such as renewable hydrogen and carbon neutral CO₂ from biogenic sources or direct air capture.^{18,19} China is already piloting cement CCUS using oxy-fuel combustion²⁰ and renewable hydrogen-based methanol projects that source CO₂ from biomass or industrial flue gas.^{21,22} However, standalone deployment of CCUS in cement and renewable hydrogen-based methanol remain expensive. Equipment level improvements in cement capture systems and plant level source-sink optimization can reduce capture, transport and storage costs.^{23,24}

Renewable methanol pathways that pair electrolytic hydrogen with direct air capture and operational flexibility can lower leveled costs.^{25,26} Yet, the green premium is estimated at over 40% for cement and more than 100% for methanol compared to conventional pathways.^{26,27}

Linking cement and methanol, two historically unconnected sectors, through green hydrogen and CCUS could reduce green premiums for both. Previous studies on methanol production explored clinker kiln off-gases as a carbon source, including using captured CO₂ for catalytic hydrogenation with renewable hydrogen,¹⁹ employing electrochemical CO₂ reduction,²⁸ and reforming flue gases with natural gas.²⁹ For cement, prior work on cement oxy-fuel combustion evaluated the benefits of utilizing byproduct O₂ from water electrolysis to replace O₂ from ASU.³⁰ Our previous work proposed a co-located configuration that enables bidirectional exchange of O₂ and CO₂ between cement kilns and methanol synthesis units.³¹ We found synergistic economic and environmental benefits for both sectors and estimated that such co-production routes could account for 24% of China's clinker output and 55% of its methanol production by 2060. Despite these promising findings, two critical aspects of the co-production system warrant deeper investigation.

One is flexibility of operations. Traditionally, industrial facilities have been designed for continuous operation, relying on stable fossil energy supplies to maximize utilization and economic performance. Integrating variable renewable energy introduces variability and intermittency that require storage and flexible operation across units and time scales.^{32,33} In the co-production system, multiple process streams and inter-linked energy and material flows create a complex, tightly coupled multi-vector topology, calling for a unified optimization modeling framework that captures cross commodity flexibility and interdependent storage dynamics.

The second is facility-level deployment and system-level planning. The economic viability of co-production hinges on renewable quality and on proximity to CO₂ sequestration sites. Spatial heterogeneity in wind, solar and sequestration availability thus mandates a two-tiered approach: plant-specific techno-economic analysis using high-resolution renewable profiles, coupled with system-level geospatial CO₂ pipeline and storage network optimization to identify least-cost clusters for co-production roll-out.

In the present work, we develop a detailed process-based and hourly-resolved model that incorporates unit level flexibility under solar and wind supply and explicitly manages flows of H₂, O₂ and CO₂, as shown in Fig. 1. We represent the entire supply and utilization chain of commodities within the co-production system of cement fueled by biomass and methanol synthesized from green H₂ and captured CO₂, covering production, storage (in different states), consumption, and sequestration (if applicable), as shown in Fig. 7 (see Appendix). Then, we optimize capacity sizing, flexible operation, and storage across different cement-to-methanol stoichiometries (see Table 1). We conduct plant-level analysis of existing cement plants to evaluate the cost-effectiveness and spatial heterogeneity of





Fig. 1 Schematic diagram of the methodological framework.

Table 1 Scenario settings for cement–methanol system decarbonization. CO₂ emission reduction is defined as the proportion of CO₂ emissions avoided relative to incumbent coal-based technologies, specifically for the case where methanol is used as a shipping fuel. The accounting scope includes direct CO₂ emissions from cement and methanol production and direct CO₂ emissions from methanol combustion, assuming complete combustion. In the net-zero cement & methanol scenario, all process and uncaptured CO₂ emissions are fully balanced by geological sequestration, and the carbon embodied in methanol is entirely atmosphere-neutral, so that net CO₂ released into the atmosphere is zero within our accounting boundary (see details in Section 3 in the SI)

| Scenario | CO ₂ handling strategy | Cement-to-methanol stoichiometry | CO ₂ emission reduction (%) | Methanol synthesis flexibility | Notes |
|----------------------------------|--|----------------------------------|--|--------------------------------|---|
| Net-zero cement & methanol | Partial CO ₂ sequestration allowed | 9.76 | 100% | Flexible Inflexible | Aligned with certification schemes (e.g., ISCC sustainable marine fuels) that require methanol to use biogenic or atmospheric CO ₂ for long-term shipping fuel eligibility |
| No-CO ₂ sequestration | All captured CO ₂ used for methanol synthesis | 2.77 | 84% | Flexible Inflexible | Represents a closed-loop carbon pathway; no geological sequestration is allowed |
| Separate decarbonization | Cement and methanol decarbonized independently | 9.76 2.77 | 77% 89% | Flexible | Cement with standalone CCUS; methanol produced <i>via</i> standalone green H ₂ and CO ₂ sourced from DAC |

technology deployment. We integrate the process optimization model with a SimCCS-China model³⁴ to optimize CO₂ source–sink matching considering costs of pipeline construction across China. This enables generating comprehensive deployment strategies of co-production technologies under different CO₂ sequestration scenarios. We find that this co-production system could achieve a much lower overall abatement cost than decarbonizing these two sectors separately, primarily benefiting from more economical sources of O₂ and CO₂. Unlocking the flexibility of methanol synthesis and configuring molecular storage buffers offer additional but plant-specific economic benefits compared to steady and continuous production. Additionally, introducing co-production technology could have a significant impact on CO₂ transport network planning in China's cement industry, by prioritizing CCS implementation at cement facilities with abundant proximate renewable resources.

2. Results

2.1. Co-production reduces CO₂ abatement costs by lowering supply costs of O₂ and CO₂

We find that the co-production system can achieve a 38–65% reduction in CO₂ abatement costs compared to decarbonizing the two plants separately when considering a selected case-study facility in Inner Mongolia (Fig. 2a). Despite slightly higher H₂ costs (Fig. 2b), the cost savings primarily stem from lower-cost O₂ (Fig. 2c) and CO₂ (Fig. 2d). The increase in H₂ costs arises because O₂ consumption and CO₂ generation in cement kiln introduce inflexible elements to the system, limiting the ability of H₂ production to follow renewable availability as flexibly as in standalone H₂-based methanol. By diminishing the ASU, the system lowers both capital investment and energy consumption, resulting in a 71–95% decrease in O₂ costs (Fig. 2c). However, due to the temporal mismatch between variable O₂ production and fixed O₂ consumption, O₂ storage





Fig. 2 Cost structure of the co-production system and associated molecules for the selected plant in Inner Mongolia. (a) Cost breakdown of the co-production technology. CO₂ abatement cost is defined as the cost difference between the decarbonized technology and the incumbent dominant route without emissions abatement (*i.e.*, coal-based production and current grid electricity), divided by the corresponding emission reduction. In this case study, the CO₂ abatement cost for separate decarbonization is estimated at \$79 per tonne of CO₂ for cement and \$131 per tonne of CO₂ for methanol (see Section 3 in the SI for detailed calculations). The CO₂ abatement cost presented in this figure reflects the combined cost for cement and methanol under a given stoichiometric ratio. "Other costs" include labor and miscellaneous cement raw materials. "CO₂ seq." represents the costs associated with CO₂ compression, transport and sequestration, assuming the transport cost is \$8.7 per tCO₂ and storage cost is \$5.8 per tCO₂.³⁵ (b), cost breakdown of H₂ supply. (c) Cost breakdown of O₂ supply. (d) Cost breakdown of CO₂ supply. "Processing" in (c) and (d) refers to capital and operational costs associated with compressors (for gaseous storage) and liquefaction units (for liquid storage).

and its associated processes constitute the largest share of the O₂ cost structure. In the no-CO₂ sequestration scenario, the

amount of byproduct O₂ is 2.7 times the O₂ demand, eliminating the necessity for O₂ storage tanks. Capturing, processing and



storing CO₂ from cement production is 80–91% cheaper than costs of CO₂ from DAC, as shown in Fig. 2d. We use a relatively optimistic DAC cost at \$203.6 per tCO₂,²⁶ within the typical \$200–600 per tCO₂ range (see Section 3 in the SI for the literature review on DAC). Applying partial CO₂ sequestration could reduce CO₂ abatement costs by further leveraging the economically sourced O₂.

Also, unlocking the flexible operation of methanol synthesis could reduce overall production costs by 8–12% primarily due to lower H₂ costs, despite slight increases in O₂ and CO₂ costs. Harnessing demand-side flexibility in H₂ consumption can reduce the levelized cost of electricity (LCOE) from solar and wind sources and the required H₂ storage tank size, ultimately lowering the levelized cost of hydrogen (LCOH) by 22–24% (Fig. 2b). Inflexible methanol synthesis requires a stable, hour-by-hour supply of CO₂ and H₂. The fixed CO₂ extraction from clinker kilns aligns perfectly with the stable CO₂ demand, eliminating the need for CO₂ storage tanks. However, with flexible methanol synthesis scenarios, the flexible CO₂ demand and stable CO₂ supply create operational asynchrony and necessitate CO₂ storage tanks as a buffer. Similarly, O₂ supply is more compatible with a fixed demand, which reduces the cost of O₂ through downsized storage tanks and a smaller complementary ASU. Thus, maximizing the flexibility options could reduce overall system costs, but it requires more coordinated and complex system design and operations.

2.2. Optimized system configurations and flexible operations

By coupling flexibilities across processes, the variability of renewable generation cascades through all production and storage stages, with each component contributing to accommodating renewable fluctuations. We observe distinct patterns of optimal flexible operations across components within the co-production system under the net-zero cement & methanol scenario (Fig. 3). In the case-study plant in Inner Mongolia, electrolyzer loads closely follow the combined generation of solar and wind, operating primarily during the daytime while shutting down or reducing output at night. Methanol synthesis exhibits seasonal variations, operating at full load during most of the spring. Batteries, exhibiting an intra-day cycle, predominantly charge during the daytime and discharge at night. H₂ and gaseous O₂ storage tanks exhibit similar annual charge-discharge cycles, typically ranging from hours to days. Gaseous CO₂ tanks cycle 21 times per year, but with uneven temporal distribution across the year. In contrast, liquid O₂, liquid CO₂, and methanol storage function as long-duration storage, with the number of annual round-trip cycles ranging from 2.1 to 3.9. While liquid O₂ and methanol are charged in spring and summer when renewable generation is abundant and discharged in other periods, liquid CO₂ follows the opposite pattern.

The discrepancy in storage behavior among different commodities can be explained by unit capital cost considerations. Storage technologies with low capital costs per unit of stored volume (*e.g.*, liquid storage) tend to exhibit long-duration storage, despite the high capital and energy penalties

associated with charging and discharging, aligning with findings in a previous study.³⁶ Conversely, storage with higher volume capacity costs (*e.g.*, gaseous storage) is inclined to increase cycling frequency, diluting capital costs across higher turnover rates to improve cost efficiency per unit of stored commodity or energy. Overall, these operational patterns reflect the underlying variability of renewable resources, highlighting that the value of flexibility lies in enabling reliable integration of intermittent solar and wind.

In contrast to RE-rich weeks characterized by nearly full-load methanol synthesis, RE-scarce weeks feature frequent ramping, which drives distinct patterns in H₂, O₂, and CO₂ production, storage, and consumption. We analyze representative RE-scarce and RE-abundant weeks for the plant in Inner Mongolia, focusing on the balance of electricity, H₂, O₂, and CO₂, as shown in Fig. 4. Here, the RE-scarce week is characterized by persistently low renewable output, especially weak nighttime wind availability, whereas the RE-abundant week exhibits consistently higher wind and/or solar. Electrolyzers account for 85% of power consumption annually, with batteries discharging primarily at night to maintain the minimum electrolyzer load (5%). During RE-rich weeks, electrolyzers could operate overnight, powered by strong wind generation. H₂ storage is discharged at night to supplement the minimum H₂ supply required for methanol synthesis. Beyond real-time consumption, most excess O₂ is stored in gaseous form, supplementing demand at night. During RE-scarce weeks, liquid O₂ storage experiences lower charging and higher discharge, while ASU operates at a higher load compared to RE-rich weeks. Meanwhile, some O₂ may be vented in RE-rich weeks, indicating a trade-off between maximizing byproduct O₂ utilization and minimizing O₂ storage system costs. For CO₂ storage, gaseous CO₂ remains largely idle, while liquid CO₂ undergoes net discharge during RE-abundant weeks. Conversely, in RE-scarce weeks, gaseous CO₂ is charged at night and utilized on days with relatively higher renewable output, whereas liquid CO₂ is charged continuously without discharge.

2.3. Geographic differences in plant economic performance due to RE spatial heterogeneity

Given the estimated shrinking cement demand^{37,38} and the expanding methanol market³⁹ in China, we consider a scenario where new methanol facilities are co-located with existing cement plants (823 plants in total). We focus on greenfield methanol facilities because the current methanol fleet is predominantly coal-based and incompatible with low-carbon integration, whereas cement could involve retrofit. In the net-zero cement & methanol scenario (9.76 tonnes cement per tonne methanol), the spatial heterogeneity of renewable resources leads to variations in CO₂ abatement costs (see Fig. 5a, excluding CO₂ transport and sequestration costs) and results in diverse optimized system configurations and operational strategies (see Section 1 in the SI). Cement plants located in Northeast China and North-Central China, where renewable resources are abundant, achieve lower CO₂ abatement costs (excluding CO₂ transport and storage), approximately \$15–35



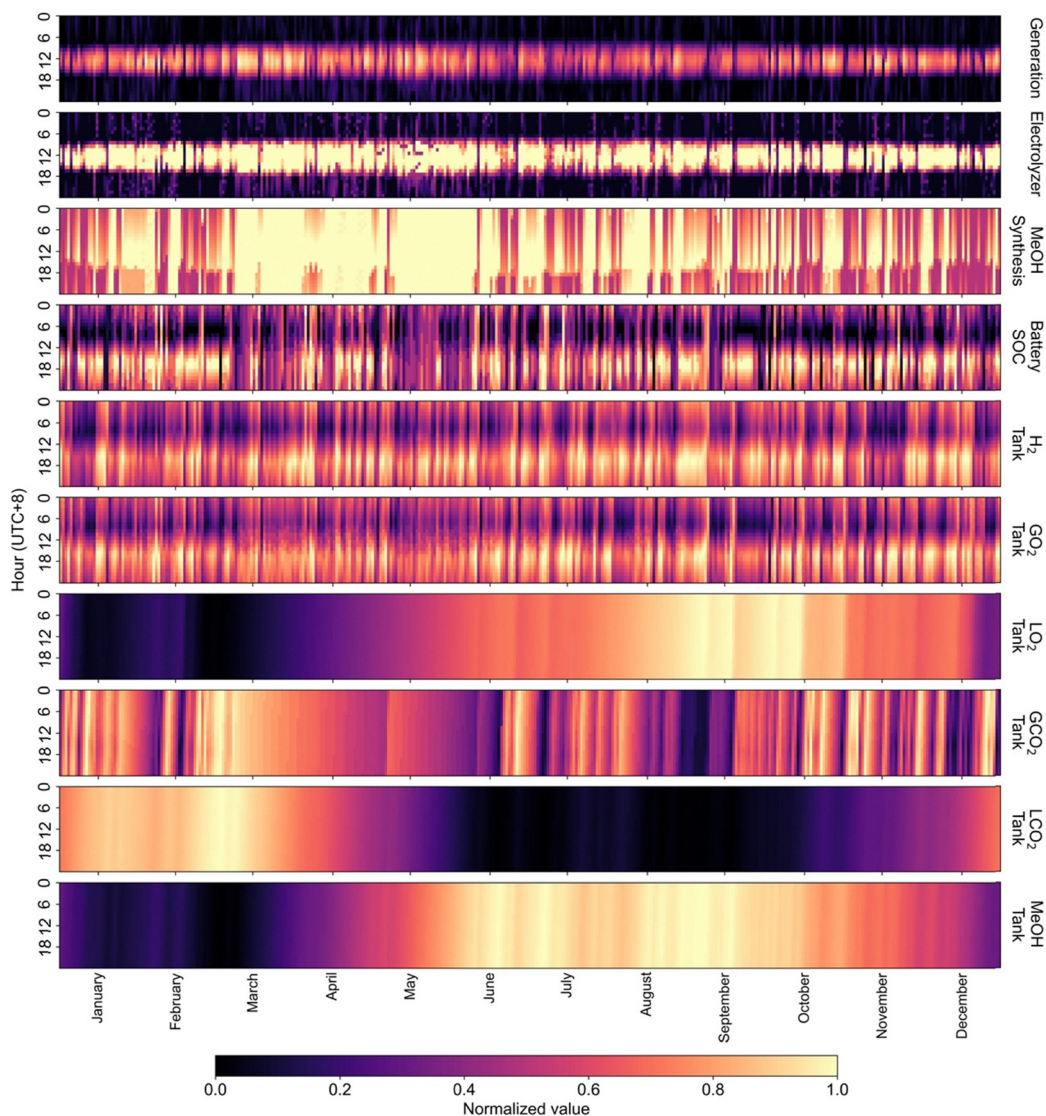


Fig. 3 Annual normalized operational profiles of the selected plant in Inner Mongolia under the net-zero cement & methanol scenario with flexible methanol synthesis. The x-axis represents 365 days over one year, and the heat maps illustrate hourly variations, with the y-axis indicating 24 hours per day. Values are normalized by dividing hourly absolute values by their annual maximum (linear normalization). “Generation” denotes the actual hourly power generation from solar and wind, normalized by the weather-dependent maximum available hourly generation. “Electrolyzer” and “MeOH synthesis” indicate hourly loads normalized by their respective maximum loads. “State of charge (SOC)” for battery, H₂ tank, gaseous O₂ (GO₂) tank, liquid O₂ (LO₂) tank, gaseous CO₂ (GCO₂) tank, liquid CO₂ (LCO₂) tank, and MeOH tank refer to the hourly SOC normalized by each component’s maximum SOC over the entire year. The operations under inflexible methanol synthesis are shown in Fig. 8 (see Appendix).

per metric ton. For example, the case plant in Inner Mongolia falls within this range, with an abatement cost of \$31 per metric ton. In contrast, plants located in central China exhibit higher abatement costs, often exceeding \$40 per metric ton of CO₂.

Unlocking the flexibility of methanol synthesis yields plant-specific cost reductions, with cement plants in China achieving co-production cost savings ranging from 3% to 17% (Fig. 5b). Generally, plants with lower baseline co-production costs under inflexible methanol synthesis exhibit smaller marginal benefits from adding flexibility. Nevertheless, enabling flexible methanol synthesis can still achieve over \$5 per metric ton in CO₂ abatement cost savings for each unit of added capacity following the ascending order of abatement cost (Fig. 5c).

Although CO₂ abatement costs for most plants are more sensitive to solar costs, the lowest-cost plants rely more heavily on high-quality wind resources and are therefore more sensitive to wind costs. For most plants, solar power accounts for a larger share of total system costs, averaging 11% compared with 5% for wind and 4% for electrolyzers. However, for plants with relatively low CO₂ abatement costs, wind cost parameters can emerge as the dominant sensitivity factor. For example, among the 50 lowest-cost plants, wind averages 9% of total costs compared with 6% for solar. These results suggest that while large-scale cost reductions depend on declining solar costs, achieving the lowest abatement costs might require prioritizing deployment in regions with high-quality wind resources.





Fig. 4 Operational profiles under renewable energy (RE)-scarce (left panel) and RE-abundant (right panel) weeks for the selected plant in Inner Mongolia under the net-zero cement & methanol scenario with flexible methanol synthesis. Positive values indicate the supply (including discharge) of each commodity, while negative values indicate demand (including charge). (a) Electricity balance. The dashed line indicates the aggregated electricity load; power generation exceeding the load is curtailed. “O₂ supply” includes electricity consumed by the ASU, compressors, and liquefaction units. “CO₂ processing” includes electricity consumed by compressors and liquefaction units. “H₂ supply” represents electricity consumption for water electrolysis and H₂ compression. (b) H₂ balance. (c) O₂ balance; “O₂ atmosphere” indicates vented O₂. (d) CO₂ balance; “CO₂ process” indicates CO₂ from carbonate decomposition, “CO₂ biogenic” indicates CO₂ from biomass combustion, and “CO₂ atmosphere” refers to CO₂ emitted to the atmosphere due to a less than 100% capture rate. The operations under inflexible methanol synthesis are shown in Fig. 9 (see Appendix).

2.4. Effect of co-production on planning of CO₂ source–sink matching

Although the abovementioned optimization model evaluates the economic performance based on renewable resource availability, incorporating post-capture (*i.e.*, transportation and storage) costs may alter the overall production cost distribution and influence the optimal plant selection strategy. Consequently, a co-optimization approach that integrates production and post-capture costs provides insights into the most cost-effective deployment of co-production facilities. To address this, we employ an extended version of SimCCS-China to

co-optimize onsite production costs, CO₂ pipeline network design, and CO₂ storage costs under different CO₂ sequestration amount scenarios (see the Methods section and Section 6 in the SI for details). As shown in Fig. 6a and Fig. S38a, the selected cement plants largely align with plants that have lower production costs shown in Fig. 5a. The “CO₂ sequestration 26.0 Mt per year” scenario corresponds to 7.06 Mt per year of methanol output under co-production, while the “CO₂ sequestration 311.5 Mt per year” scenario corresponds to 84.7 Mt per year. At the initial-deployment stage, the low-hanging fruit for adopting co-production technologies is found in Northeast





Fig. 5 Plant-by-plant economic performance and sensitivity analysis in China under net-zero cement & methanol scenarios. (a) CO₂ abatement cost by plant, defined as $(\text{cost}_{\text{decarbonized}} - \text{cost}_{\text{incumbent}}) / \Delta\text{CO}_2$ per 9.76 t cement + 1 t MeOH. Incumbent routes use coal-based production and current grid electricity; CO₂ transport and sequestration costs are excluded. Each dot marks an existing cement plant with a different capacity (size) and abatement cost (color). (b), The ratio of cost difference between flexible (Flex) and inflexible (Inflex) methanol synthesis, on the same 9.76 t cement + 1 t MeOH basis. (c) Cost–capacity curves sorted in ascending order of abatement cost versus cumulative cement capacity. The solid blue line is the Flex baseline; lighter blue lines show sensitivity to renewable-related cost parameters under Flex scenarios; the light blue shaded band is the Flex envelope (min–max). The sensitivity analysis is conducted by changing the cost parameters of solar, wind and electrolyzer by 20% one at a time. The solid orange line is the Inflex baseline; lighter orange lines show sensitivity of Inflex; the light orange shaded band is the Inflex envelope. It is assumed that local biomass resources are sufficient to meet the demand under the net-zero cement & methanol scenario, with supporting estimates and an availability analysis provided in Section 4 in the SI.

China, where the additional costs from CO₂ transport and sequestration are approximately \$15 per tonne. As CO₂ sequestration increases to large-scale deployment, CO₂ captured from some cement plants in North China, Northeast China, Southwest China, and East China is transported to nearby sequestration sites through a CO₂ pipeline network. However, compared to the plant distribution shown in Fig. 5a, some plants with low production costs are not connected to the network due to their long distances to suitable storage sites.

Meanwhile, the introduction of the co-production system significantly alters the CO₂ source–sink matching layout compared to decarbonizing China's cement sector alone (Fig. 6b and Fig. S38b). Under a cement-only decarbonization strategy, CO₂ pipelines connect fewer plants located in renewable-rich regions such as North China and Northeast China, but with

potential expansions into central China. CO₂ transport costs are higher when using co-production technologies, driven by lower capturable CO₂ volumes per plant and more pipeline installations required (Fig. 6c). This indicates that producing net-zero methanol through co-production systems involves a cost trade-off, where increased CO₂ transport requirements are offset by lower production-side abatement costs.

3. Discussion

Co-production, as proposed in this study provides substantial economic advantages over separate decarbonization of the cement and methanol sectors. By 2035, co-producing net-zero cement and methanol in China could lower CO₂ abatement





Fig. 6 Comparison of CO₂ source–sink matching under co-production *versus* cement-only decarbonization scenarios. (a) CO₂ network under co-production. (b) CO₂ network under cement-only. (c) The composition of CO₂ abatement costs under different scenarios. Compared to the case-study plant in Inner Mongolia, here we change the CO₂ transport and sequestration costs from fixed value per tonne of CO₂ into variable numbers (see the Methods section). “CO₂ sequestration 311 Mt per year” refers to the scenario where all incumbent methanol production (70.6 Mt) and an estimated share of marine fuel demand (14.1 Mt) are substituted with green methanol under co-production. “CO₂ sequestration 26 Mt per year” corresponds to a scenario with a 10% substitution of incumbent methanol production (7.06 Mt) by co-production (see Fig. S38). Green circles mark CO₂ point sources at cement plants, with size representing capturable emission volume; red triangles mark CO₂ storage sinks. Orange lines denote the proposed CO₂ pipeline network with different diameters reflected by line width. Additional CO₂ sequestration scenarios are presented in Section 7 in the SI.



costs to \$41–53 per tonne, markedly below approximately \$75 for independent CCS in cement and over \$120 for independent renewable-based methanol. With China's carbon price projected to reach \$30–70 per tCO₂ by 2035,⁴⁰ the co-production approach has promise to achieve cost parity with conventional technologies under plausible policy scenarios while achieving co-decarbonization. Spanning from detailed process modeling to system-level planning, this study offers critical insights into cost-effective and feasible decarbonization pathways for these two hard-to-abate sectors through optimized co-production systems.

Such industrial symbiosis reveals a transferable molecule coupling principle that can materially shape the cost and feasibility of integrated decarbonization systems. H₂ has attracted extensive attention for its potential as an industrial feedstock,⁴¹ spurring techno-economic analyses and lifecycle emission assessments.^{42,43} Conversely, the application potential of renewable-sourced O₂ and CO₂ as industrial feedstocks remains underexplored. Currently, industrial O₂ is primarily sourced through energy-intensive air separation for combustion, chemical synthesis, and metallurgy.⁴⁴ In a future industrial system deploying large-scale electrolytic hydrogen, high-purity O₂ generated from water electrolysis could be directed for oxy-fuel combustion, improving carbon capture efficiency and enhancing system-wide decarbonization economics by reducing costs associated with air separation.⁴⁵ Meanwhile, captured CO₂ can serve as a raw material for chemical synthesis where a carbon source is needed.⁴⁶ Using captured CO₂ is significantly more economical than direct air capture and less emission-intensive than conventional coal-based routes. Thus, the generalized co-production paradigm represents an innovative form of industrial symbiosis that integrates oxy-fuel combustion–CCUS with electrolytic hydrogen, opening new frontiers for industrial decarbonization and delivering mutual benefits for traditionally unconnected sectors.

Exploring the flexibility of industrial processes may further lower the costs of low-carbon technologies. Compared with most stable, continuously operated industrial facilities, a unique feature of the renewable-based co-production system is its integration of operational flexibility across different components and storage methods. Under the flexible methanol synthesis scenario, the temporal mismatch of supply and demand is characterized by: (1) H₂: flexible production and flexible consumption; (2) O₂: flexible production and fixed consumption; (3) CO₂: fixed production and flexible consumption. Different storage methods (*e.g.*, liquid *vs.* gaseous storage) operate at varying storage durations, shaped by their economic and technical parameters. The temporal asynchrony of molecular flows arising from renewable energy variability underscores the necessity and complexity of strategically coordinating operational modes and deploying buffering strategies to optimize system-wide adaptability. In practice, however, first-of-a-kind (FOAK) deployment is likely to operate more conservatively than the system-optimal schedules explored here. Flexible electrolysis and gas buffering are comparatively mature and have been demonstrated in industrial settings.⁴⁷

By contrast, flexible methanol synthesis remains largely at the early demonstration stage and is likely limited to narrower turndown ranges and slower ramping, given constraints related to catalyst performance, thermal management, and product quality.⁴⁸ Meanwhile, flexible operation may increase O&M burdens or accelerate equipment degradation.⁴⁹ We therefore interpret the modeled flexibility as an upper bound on integration value. The trade-off between flexibility benefits and operational penalties must be established through staged demonstration and deployment.

Implementing co-production technologies further demands customized deployment strategies to address the spatial heterogeneity of renewable resources, CO₂ pipeline costs, and sequestration site distributions. Our results indicate that optimal system configurations, economic performance, and potential savings from flexibility vary plant by plant, leading to plant-specific optimal operations across different components. Additionally, achieving net-zero emissions requires co-optimization of renewable-based production alongside CO₂ transport and sequestration planning. Compared to cement-only decarbonization, adopting co-production technologies results in different CO₂ pipeline layouts, both for initial-deployment and large-scale CO₂ sequestration scenarios. Given the substantial investments needed for CO₂ pipeline infrastructure, the deployment of co-production technologies must carefully assess the economic viability and technical feasibility of CO₂ network planning, and be coordinated with potential pipelines designed for the cement sector alone.

Introducing co-production technologies could also influence the decommissioning strategy for China's cement production capacity. In 2020, China produced 2395 Mt cement;⁵⁰ however, demand has started to decline, driven by a weakened real estate and infrastructure market, and is projected to decrease by 42–79% by 2060.^{37,38} Due to overcapacity and future demand decline, the criteria guiding decisions on cement plant retention are likely to prioritize factors such as plant efficiency rankings, availability of surrounding decarbonization resources (*e.g.*, CO₂ sequestration sites, alternative fuels), and regional cement demand.^{24,31,51} Co-production technologies might confer strategic advantages to cement plants located in regions with abundant renewable resources or even stimulate the construction of new capacity in these regions through capacity replacement policies. In contrast, China's methanol market remains robust, supported by both conventional chemical demand and emerging applications such as low-carbon fuels.⁵² Our analysis suggests that under the scenario of substituting all conventional methanol and methanol shipping fuel forecast in 2050, co-production technologies would necessitate utilizing approximately 41.3% of China's current cement capacity (see Section 7 in the SI). Therefore, dynamic planning of cement plant decommissioning and the incremental co-production deployment should be comprehensively evaluated for optimizing China's long-term decarbonization pathway.

However, successful deployment will require strong financial incentives, targeted policy support for demonstration projects, and coordinated corporate strategies bridging these



traditionally unrelated industries. In practice, co-production typically spans multiple owners and counterparties, including cement and methanol operators, renewable electricity suppliers or offtakers, and CO₂ transport and storage providers. Aligning incentives requires credible valuation of verified emissions reductions and transparent allocation of that value across participating actors. Bankable deployment will therefore depend on robust contracting arrangements, such as long-term renewable electricity procurement and low-carbon methanol offtake, with clear provisions on risk allocation, long-term liability, and revenue sharing. The limitations of this study (see Section 8 of the SI) reflect modeling abstractions on plant heterogeneity, siting feasibility, carbon-accounting scope, downstream logistics (see the estimated delivered-to-port cost in Section 9 of the SI), and technological constraints on alternative-fuel substitution.⁵³

Cement–methanol co-production could become increasingly attractive across a wider set of geographies beyond China. Cement and methanol are globally traded, geographically widespread foundation materials, enabling many opportunities for co-location or regional integration. Consistent with our China-wide sensitivity across renewable endowments, a key prerequisite is access to abundant low-cost renewables. Many regions offer wind and solar resources that are comparable to or stronger than China's (e.g., the U.S. Great Plains, Australia, Chile, and parts of the Middle East and North Africa, among others),⁵⁴ while biomass-rich regions (e.g., Brazil) may further ease low-carbon heat supply constraints.⁵⁵ Projected declines in the costs of wind, solar, and electrolysis energies would further improve the economics in the future. In parallel, policy and market frameworks that monetize emissions reductions are expanding, including carbon pricing (e.g., the EU ETS) and low-carbon fuel standards. For example, the average 2024 secondary market price of the EU ETS was around \$71 per tCO₂e,⁵⁶ versus approximately \$14 per tCO₂ for China's national ETS allowances.⁵⁷ However, national circumstances differ substantially in resource quality, infrastructure, and policy environments, and the quantitative outcomes therefore merit further country-specific assessment.

4. Methods

4.1. Optimization model

The process-detailed linear programming optimization model is designed to optimize capacity sizing, flexible operations, and storage strategies for each system component at an hourly resolution under constraints (see Section 2 in the SI). Electricity is sourced from a combination of dedicated local solar and wind sources. Locally-sourced biomass is used for clinker calcination, ensuring a carbon-neutral energy supply during the cement production stages. Alkaline electrolyzers generate H₂ and O₂ simultaneously with a flexible load. H₂ must be compressed to 150 bar to meet the pressure requirements for chemical production before being used in methanol synthesis or stored in gaseous H₂ tanks. O₂ can be utilized immediately,

stored in gaseous form, or liquefied for long-duration storage. Since the operating pressure of clinker kilns (~1 bar) is lower than the electrolyzer outlet pressure, O₂ can be used without compression. To manage temporal mismatches, O₂ can either undergo compression and be stored in gaseous tanks (lower energy penalty but higher capital cost per mass stored), or be liquefied and stored in liquid tanks (higher energy penalty but lower capital cost per mass stored). While byproduct O₂ utilization for oxy-fuel combustion offers economic benefits, an ASU is also included to supplement O₂ during temporal shortages or when elevating the stoichiometric balance between cement and methanol reduces byproduct O₂ availability. Cement production is divided into three stages: raw meal preparation, clinker production, and cement production. The output commodity of each stage can be easily stored since the solid outputs have simple storage requirements. Raw meal preparation and cement production involve grinding processes powered by renewable electricity and can operate flexibly. However, the clinker kiln must operate steadily due to the stability requirements of chemical reactions at more than 1500 °C.⁵⁸ CO₂ emissions from clinker kilns originate from biomass combustion (~1/3) and carbonate decomposition (~2/3). Captured CO₂ can be stored in liquid or gaseous form for future use, or compressed and transported for underground sequestration. For immediate consumption or gaseous storage, CO₂ is compressed to 50 bar, while for sequestration, it is compressed to 150 bar. Similar to O₂ storage, CO₂ storage options include gaseous and liquid storage. Methanol is synthesized from green H₂ and captured CO₂ and stored in liquid form. Methanol synthesis has the potential for flexible operation within a certain load range and ramping limits. We assume a fixed hourly supply for both cement and methanol. The resulting optimization problem involves hundreds of thousands of variables and constraints.

4.2. Input data

Technical and economic parameters were derived from the authors' process simulations and data from the literature. We use ASPEN Plus to simulate the detailed production processes for methanol synthesis, and O₂/CO₂ compression and liquefaction systems (Section 5 in the SI). The energy consumption for these processes is extracted from the simulations. Stoichiometric data for other processes are obtained from the literature and summarized in our previous work (see Table S2 in the SI). Flexibility-related parameters, including ramp rates and adjustable load ranges across all components, are also sourced from the literature (see Table S4 in the SI). The costs of renewable generation technologies, batteries, and electrolyzers are customized for China using 2035 projections. Economic data for other facilities are obtained from the literature (see Table S3 in the SI). Renewable energy profiles are sourced from Renewable.ninja, using 2019 weather data. The biomass cost is set at \$80 per tonne (dry matter). Costs are reported in 2020 USD.

4.3. Stoichiometric balance

The stoichiometric balance between cement and methanol is defined as the hourly supply mass ratio of these two products.



4.5.1. Input data preparation. The construction cost of CO₂ pipelines varies across regions due to geographic, topographic, and demographic factors. Our previous study on CO₂ source-sink matching in China's steel sector developed a geographic information system (GIS)-based cost surface with a 240 × 240 m resolution for pipeline construction.³⁴ In this study, we update the geographical coordinates of CO₂ sinks from prior research by incorporating new basins (see Section 6 in the SI). For CO₂ sources, we primarily evaluate the net-zero scenario by adopting the co-production strategy and compare it with the cement decarbonization-only scenario. The cement decarbonization-only scenario involves implementing CCS technology at selected cement plants without additional decarbonization measures (e.g., fuel switching, efficiency improvements), while achieving the same level of CO₂ sequestration as the net-zero scenario. Under the net-zero scenario, we derive plant-level CO₂ emissions and co-production costs (excluding CO₂ transport

and storage) from the process-detailed optimization model. In the cement decarbonization-only scenario, capturable CO₂ emissions at the plant level are assumed to be proportional to plant capacity, with cement production costs per tonne remaining constant (see Section 7 in the SI).

SimCCS is a decision-support software designed for the optimization and integration of CCUS technologies. One version of SimCCS generates the optimal pipeline layout under specified CO₂ sequestration targets. The input data includes geographical coordinates of CO₂ sinks and sources, the capturable CO₂ amount and capture costs by source, the sequestrable CO₂ amount and sequestration costs by sink, and the pipeline construction cost surface. The objective function minimizes the whole process cost, including capture, transport infrastructure, and sequestration costs, as defined in eqn (1), where i denotes the CO₂ source adopting CCS, and j represents the CO₂ sink utilized.

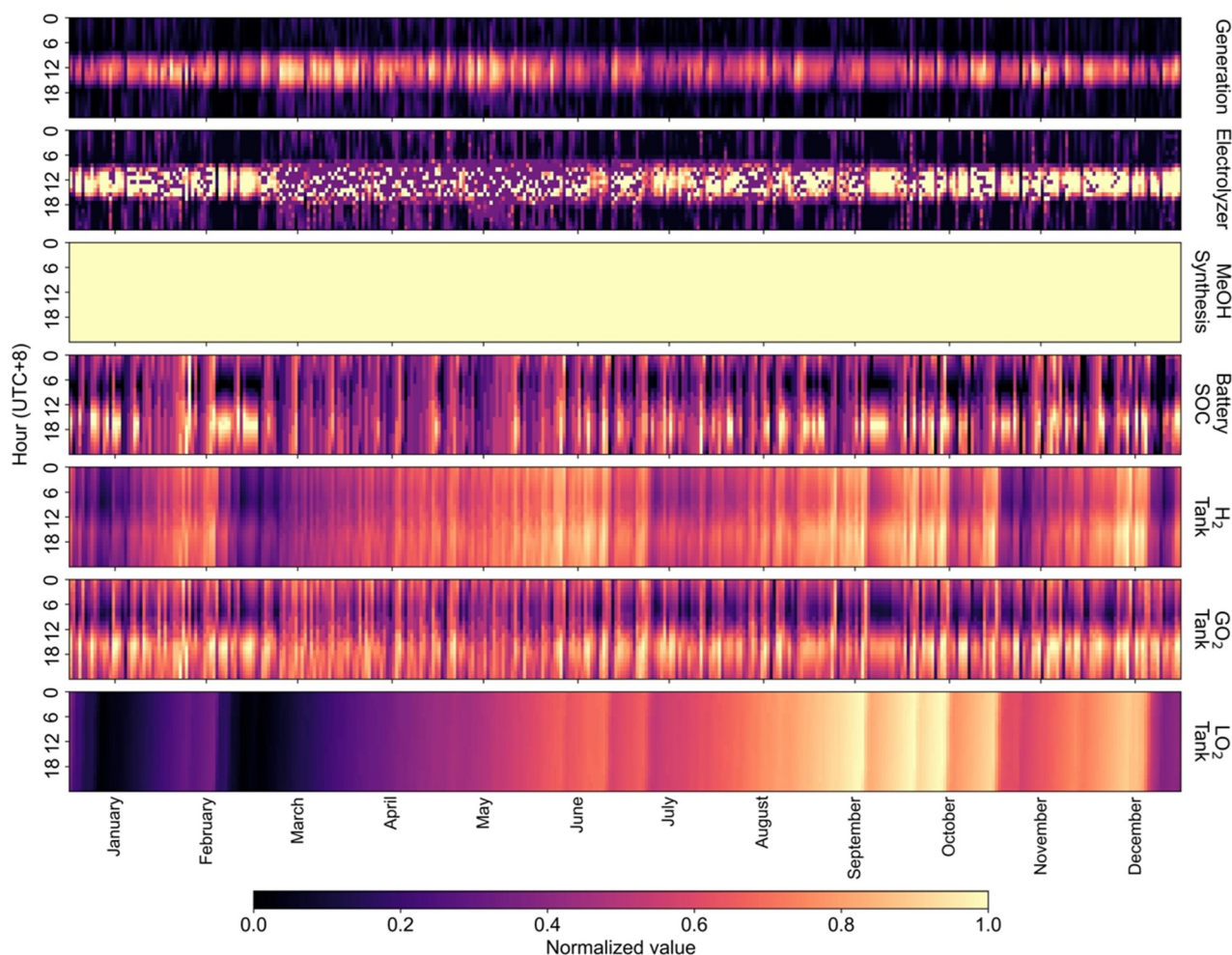


Fig. 8 Annual normalized operational profiles of the selected plant in Inner Mongolia under the net-zero cement & methanol and inflexible methanol synthesis scenario. The x-axis represents 365 days over one year, and the heat maps illustrate hourly variations, with the y-axis indicating 24 hours per day. Values are normalized by dividing hourly absolute values by their annual maximum (linear normalization). "Generation" denotes the actual hourly power generation from solar and wind, normalized by the weather-dependent maximum available hourly generation. "Electrolyzer" and "MeOH synthesis" indicate hourly loads normalized by their respective maximum loads. "State of charge (SOC)" for battery, H₂ tank, gaseous O₂ (GO₂) tank, and liquid O₂ (LO₂) tank refers to the hourly SOC normalized by each component's maximum SOC over the entire year.



In this study, we extend the SimCCS-China framework to optimize the overall CO₂ processing costs (equivalent to minimizing

the CO₂ abatement cost) under predefined targets. The objective function in eqn (2) replaces the capture cost in eqn (1) with the



Fig. 9 Operational profiles under renewable energy (RE)-scarce and RE-abundant weeks for the selected plant in Inner Mongolia under the net-zero cement & methanol and inflexible methanol synthesis scenario. The left panels illustrate operations during a representative RE-scarce week, while the right panels show a representative RE-abundant week. In each subfigure, positive values (above the x-axis) indicate the supply (including discharge) of each commodity, while negative values (below the x-axis) indicate demand (including charge). (a) Electricity balance. The dashed line indicates the aggregated electricity load; power generation exceeding the load is curtailed. "O₂ supply" includes electricity consumed by the ASU, compressors, and liquefaction units. "CO₂ processing" includes electricity consumed by compressors and liquefaction units. "H₂ supply" represents electricity consumption for water electrolysis and H₂ compression. (b) H₂ balance. (c) O₂ balance; "O₂ atmosphere" indicates discarded O₂. (d) CO₂ balance; "CO₂ process" indicates CO₂ from carbonate decomposition, "CO₂ biogenic" indicates CO₂ from biomass combustion, and "CO₂ atmosphere" refers to CO₂ emitted to the atmosphere due to less than 100% capture rate.



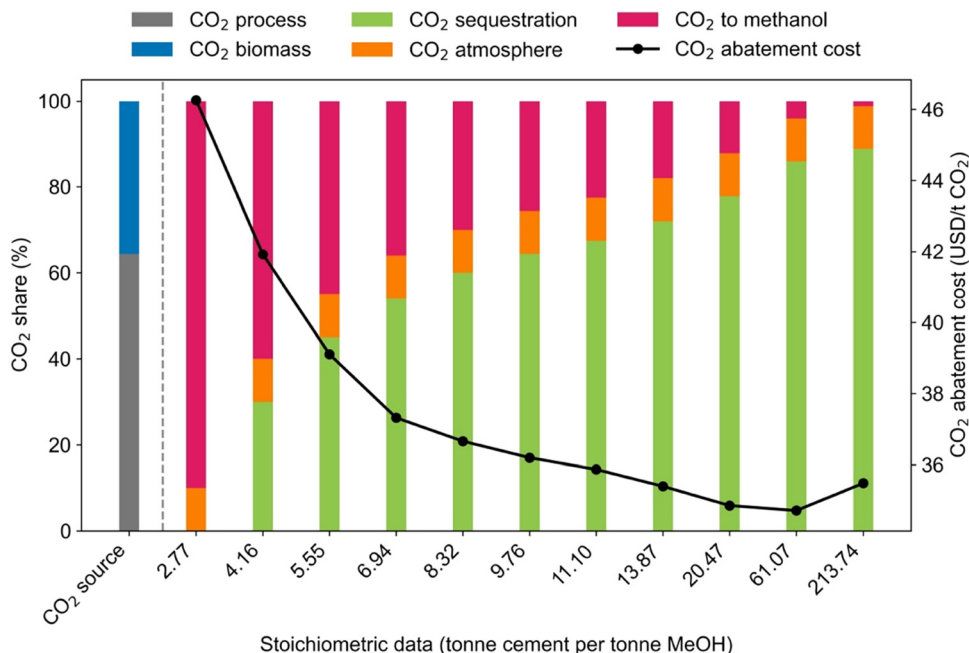


Fig. 10 CO₂ abatement cost and CO₂ share under different cement-to-methanol stoichiometric data. The leftmost bar shows the composition of the CO₂ source—exclusively from cement facilities—which remains the same across all scenarios. The subsequent bars indicate the share of CO₂ delivered to each final destination for the stoichiometric datasets on the x-axis. The line with markers represents the CO₂ abatement cost, calculated assuming CO₂ transport and storage costs of \$8.7 and \$5.8 per tonne of CO₂, respectively. As the ratio increases, CO₂ sequestration rises while CO₂ used for methanol synthesis correspondingly falls. This shift drives the CO₂ abatement cost down from \$46 per tCO₂ in the no-sequestration scenario to \$36 per tCO₂ under the net-zero methanol scenario (stoichiometric ratio = 9.76). Although abatement costs are minimized at stoichiometric ratios between 20.5 and 213.7, the incremental reduction relative to the net-zero methanol scenario (ratio = 9.76) is limited.

equivalent capture cost, which is derived by converting the co-production cost into an equivalent cost per unit of CO₂ sequestered (eqn (3)). At the optimum, the co-production cost, minus the cost of the coal-based benchmark technology and divided by the amount of CO₂ avoided, yields the system-level CO₂ abatement cost. We set different scales of green methanol output and convert these targets into CO₂ sequestration requirements based on the stoichiometric relationship between utilized and sequestered CO₂. Without considering CO₂ transport and sequestration costs, the co-production system is naturally favored in regions with abundant renewable energy. We incorporate post-capture (*i.e.*, transportation and storage) costs into the optimization to evaluate their impact on cost distribution and plant siting, and perform a geospatial assessment of CO₂ sequestration resources to enable source-sink matching for net-zero methanol and cement production.

$$\min \sum_{i,j} (\text{Capture_cost}_i + \text{Pipeline_cost}_{i,j} + \text{Sequestration_cost}_j) \quad (1)$$

$$\min \sum_{i,j} (\text{Eq.capture_cost}_i + \text{Pipeline_cost}_{i,j} + \text{Sequestration_cost}_j) \quad (2)$$

$$\begin{aligned} & \text{Eq.capture_cost} (\$ \text{ per tCO}_2) \\ & = \text{co-production cost} (\$ \text{ per (t MeOH + xt cement)}) / \text{stoichiometry}(x) / \text{CO}_2 \text{ sequestration factor} (\text{tCO}_2 / \text{t cement}) \end{aligned} \quad (3)$$

Author contributions

Y. H., H. L., P. L. and J. D. J. conceptualized the study and methodology. Y. H., H. L. and C. T. implemented the software, validated the model and analyzed the data, with contributions from Y. L. and Z. W. on data curation and visualization. Y. H. and Y. L. wrote the original draft, with major revisions from E. L., A. L., P. L., J. D. J., H. L., Y. F. and Z. L. E. L. and Z. L. acquired the funding. Z. L., P. L., E. L. and J. D. J. supervised the study.

Conflicts of interest

J. D. J. serves on the advisory boards of Eavor Technologies Inc., a closed-loop geothermal technology company, Rondo Energy, a provider of high-temperature thermal energy storage and industrial decarbonization solutions, Dig Energy, a developer of low-cost drilling solutions for ground-source heat pumps, and Karman Industries, a developer of advanced heat pumps for industrial applications and has an equity interest in each company. He also serves as a technical advisor to MUUS Climate Partners and Energy Impact Partners, both investors in early-stage climate technology companies.



Data availability

The data supporting this article have been included as part of the supplementary information (SI). Supplementary information is available. See DOI: <https://doi.org/10.1039/d5ee07379k>.

Appendix

Fig. 7–10 show the co-production framework, operational profiles under inflexible methanol synthesis, and CO₂ abatement cost and share.

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