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Realizing CO₂ emission reduction in lime and soda ash manufacturing through anion exchange[†]

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Lime (CaO) and soda ash (Na₂CO₃) are two foundational chemicals for modern civilization, and the CO₂ emissions from their production processes are challenging to reduce. Furthermore, decarbonization of the lime industry could also reduce the CO₂ emissions associated with cement production, for which lime is the key precursor. In this paper, we show that an anion exchange process to co-produce CaO and Na₂CO₃ from CaCO₃ and NaOH can reduce the carbon footprint of both chemicals through industrial symbiosis. Heating energy and NaOH production are the major contributing factors towards the cost and CO₂ emissions of this process, which can supply the global annual soda ash demand (~65 Mt) and co-produce ~50 Mt of lime in an economically sustainable manner (16% gross margin) while immediately reducing global CO₂ emission by 37 Mt compared to current production methods. Using electrified industrial heat sources and heat pumps to reuse heating energy would further reduce the cost and CO₂ emissions of the anion exchange process.

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1. Lime and soda ash production belong to the “hard-to-abate” sector, where reduction of CO₂ emissions is difficult. In this paper, we have shown that an anion-exchange process to co-produce lime and soda ash can reduce global CO₂ emissions by 37 Mt per year in an economically feasible manner.
2. Lime and soda ash are fundamentally essential chemicals for today’s human civilization. They are used for carbon capture, cement, detergent, and glass manufacturing, and they are critical for energy storage applications, including lithium and sodium ion batteries, and renewable energy generation, such as solar panels.
3. Integrating soda ash and lime manufacturing is a pivotal step in decarbonizing the inorganic bulk chemical sector, which can have a ripple effect on decarbonizing other hard-to-abate sectors.

Introduction

Lime (CaO) and soda ash (Na₂CO₃) are historically essential chemicals for human civilization. Soda ash has been used for manufacturing glass since 5000 years ago by the Egyptians,¹ and in more modern times, soda ash demand is continuously growing as it is used for producing powdered detergents, scrubbing polluting emissions from flue

gas emissions, and as a buffering agent in different processing technologies, including lithium ion battery production, a rapidly expanding industry. Soda ash is extracted from mined trona (“natural soda ash”) or in regions without trona availability, such as in Europe, it is produced using synthetic processes, mostly the Solvay² or the modified Solvay or Hou’s process (“synthetic soda ash”). Soda ash production from trona mining has a CO₂ footprint of 0.3–0.7 kg of CO₂ per kg of soda ash, whereas synthetic soda ash production, which accounts for two-thirds of global soda ash production, has a footprint of 1 kg of CO₂ per kg of soda ash produced.³ Lime, on the other hand, was also used as early as 7000 BCE at Aşıklı Höyük in Turkey,⁴ and is currently utilized majorly in cement production as well as in soil stabilization,⁵ wastewater treatment,⁶ remediation of heavy metal pollution,⁷ and CO₂ capture.⁸ Lime is generally manufactured from the calcination of limestone with a CO₂ footprint of ~1–1.8 kg of CO₂

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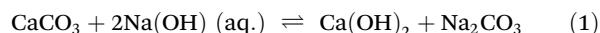
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per kg of lime produced.⁹ Both lime and soda ash are globally produced at a high volume (65 Mt of soda ash and 430 Mt of lime annually).¹⁰ Furthermore, the calcination of limestone is an essential step in conventional cement production (4.1 Gt annual global production),¹⁰ which contributes about 8% of anthropogenic CO₂ emissions,¹¹ and a significant reduction in carbon emissions from lime production could be key to reducing cement's CO₂ footprint. Recently, a concurrent recycling of steel and concrete using electrical energy without any direct CO₂ emissions has been proposed;¹² however, this process cannot supply the current global cement demand, and other complementary solutions are necessary.

Both lime and soda ash production are considered part of hard-to-abate sectors^{13,14} and are expected to require radical transformation in their production processes to make them carbon-neutral. Recently, a novel chemical process has been developed to simultaneously produce Ca(OH)₂ and Na₂CO₃ from CaCO₃ and NaOH through anion exchange without direct CO₂ emission (eqn (1)).¹⁵



Eqn (1) is a reversible anion-exchange reaction, and the output is always a mix of CaCO₃, NaOH, Ca(OH)₂, and Na₂CO₃. However, the conversion efficiency of CaCO₃ into Ca(OH)₂ (eqn (2)) can be as high as 96%, depending on the initial amount of CaCO₃, NaOH, and water (H₂O).

$$\text{Conversion efficiency}(\%) = 100 - \frac{\text{Unreacted CaCO}_3}{\text{CaCO}_3 \text{ in mix}} \times 100 \quad (2)$$

The different compounds are separated by leveraging the high solubility of NaOH in alcohol (methanol or ethanol) and Na₂CO₃ in water. This anion exchange reaction occurs at ambient pressure and temperature and is successful even when utilizing industrial sources of limestone containing various impurities such as Si, Al, and Fe.¹⁶ Furthermore, it is not highly sensitive to reaction temperature.¹⁷ Although this anion-exchange process shows promise in reducing carbon emissions, a thorough and systematic techno-economic analysis has not been performed to evaluate its potential for scaling up within the timeframe necessary to achieve climate goals.

In this work, we have evaluated the energy, cost, and carbon footprint of executing the processes previously suggested,¹⁵ along with proposing a new process for better techno-economic opportunities. The aim of our analysis was to identify the process and the initial amounts of CaCO₃, NaOH, and water in eqn (1) that would provide the lowest cost and CO₂ footprint, as well as to compare them to the current global emission and cost values to evaluate the potential of this technology for successful scaling up in the short term.

Methodology

Chemical processes and techno-economic assessment

The anion exchange process has CaCO₃ and NaOH as input materials and Na₂CO₃ and Ca(OH)₂ as output materials. If the anion exchange of Ca(OH)₂ production is scaled up to supply the current global cement clinker production (annually 3.8 Gt),¹⁰ a total of ~2.47 Gt of CaO would be required, assuming that clinker contains 65% CaO on average. Production of the 2.47 Gt of CaO would be accompanied by a production of 4.68 Gt of Na₂CO₃ and a consumption of 3.53 Gt of NaOH, assuming 100% conversion efficiency. These amounts are much higher than the current global annual production of NaOH and Na₂CO₃ (82 Mt (ref. 18) and 65 Mt,¹⁰ respectively), and the techno-economic assessment of this process must carefully consider the cost and CO₂ footprint of these materials.

The production of Ca(OH)₂ and Na₂CO₃ from CaCO₃ and NaOH can involve various steps depending on the order of separation and purification after the anion exchange reaction (Fig. 1). In this paper, two processes, process A and process B, proposed by Hanein *et al.*,¹⁵ would be considered. Process A involves removing the unreacted NaOH using methanol, and then separating Na₂CO₃ from the mixture of Ca-containing compounds, Ca(OH)₂ and CaCO₃. In process B, the Ca-containing compounds are first separated after adding excess water to dissolve NaOH and Na₂CO₃.

For a techno-economic assessment, a set of 71 observed experimental results¹⁵ that evaluated the conversion rate of CaCO₃ into Ca(OH)₂ as in eqn (1), when different amounts of CaCO₃, NaOH, and water are mixed, are considered. As the chemical reaction described in eqn (1) is never 100% complete and depends on a complex equilibrium, the Ca-containing product will always be a mixture of different compositions of Ca(OH)₂ and CaCO₃, and for comparison purposes, the mixture of Ca(OH)₂ and CaCO₃ is here assumed to be heated at 1000 °C using natural gas to form CaO as the final product.

For the title study, two types of NaOH, (1) "commercial NaOH" (obtained from the currently available supply chain) and (2) "renewable NaOH" (produced using 100% renewable green electricity), have been considered. In terms of output materials, two cases for the Na₂CO₃ generated were also considered: (1) "commercialized Na₂CO₃", where the Na₂CO₃ is sold in the market, or (2) "recycled Na₂CO₃", where the Na₂CO₃ produced is used to regenerate NaOH (input in eqn (1)). Thus, a total of 4 cases were considered for each process, process A and B – (1) commercial NaOH, commercialized Na₂CO₃, (2) renewable NaOH, commercialized Na₂CO₃, (3) commercial NaOH, recycled Na₂CO₃, and (4) renewable NaOH, recycled Na₂CO₃. A detailed description of the cost and CO₂ footprint of these materials is provided below and in the ESI (Tables 1 and S1†).

Process model

A simulation model was built in Aspen Plus 11, based on electrolyte equilibria for the prediction of carbonate speciation reactions and E-NRTL¹⁹ as a thermodynamic model. The

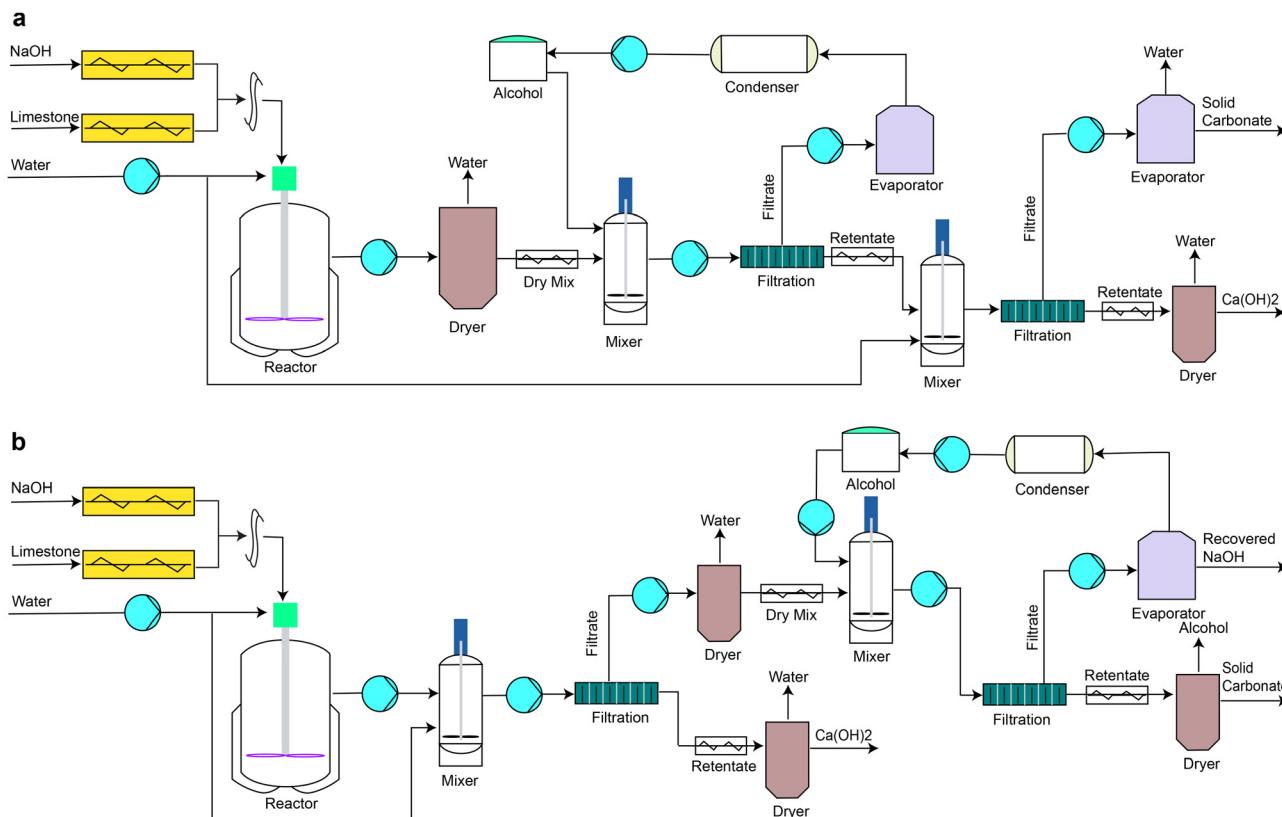


Fig. 1 Flow diagrams of the two processes (a: process A, b: process B). In process "A", the unreacted NaOH after the anion exchange reaction is separated first before separating the Na_2CO_3 from the mixture of Ca-containing compounds, $\text{Ca}(\text{OH})_2$ and CaCO_3 . In process "B", the Ca-containing compounds are separated from the mixture after adding excess water to remove/dissolve both NaOH and Na_2CO_3 .¹⁵

model takes the composition of the input reactants (*i.e.* calcite, water, and sodium hydroxide) and the observed conversion from the experimental results¹⁵ and scales the process up to 1000 tons of calcite per day. The modeling was performed for separation and purification routes A and B.

Evaporators were calculated using an adiabatic flash block to estimate the heat requirements for drying and evaporation at 150 °C. The rest of the processes (mixers, reprecipitations and/or redissolutions) were estimated from solubility data, while the thermal balance of each block was estimated using Aspen Plus. The two options (A and B) were integrated into the same simulation flowsheet with a stream multiplicator to calculate the mass and energy balance for all the 142 potential cases (71 experimental results through 2 different processing options) in one simulation run. These calculations and the solubilities were integrated into a worksheet calculator that includes the 71 case studies developed by Hanein *et al.*¹⁵ and feeds to the Aspen model the conversion of calcite and the amount of required methanol or water for reprecipitation/redissolution.

The mass and energy balance results are then calculated through a sensitivity analysis for all the laboratory results reported by Hanein *et al.* (Table S4†).¹⁵ Methanol was assumed to be recovered in a closed cycle, as for conventional solvents. However, a 5% loss per cycle was considered to estimate the

potential impact of various losses in an industrial plant. Evaporated water is not recovered, while electricity needs, calculated for pumping and pressure-driven filtrations, are lower than 1 kW h per ton of input calcite for options A and B.

CO₂ footprint

The carbon footprint is calculated using a cradle-to-gate LCA approach following the methodology described in the ISO 14040 standard.²⁰ In terms of the system's functionality, the desired function is to fully convert calcium carbonate in limestone to calcium oxide and CO₂. So, the functional unit of the assessment is the treatment of 1 ton of calcium carbonate with sodium hydroxide to obtain 0.56 tons of CaO. The system boundaries include all the impacts from raw materials and energy source supply (background system) and the proposed limestone treatment system (foreground system) but do not include further use of CaO, such as in clinker making or any other further processing since its comparison with the production of CaO through conventional thermal calcination of limestone would suffice to estimate emission savings. The CO₂ emissions from the final calcination step were considered in the GHG balance. The construction of the plant was excluded from the boundaries of the LCA system.

The system is multifunctional since the by-product consists of a solid composed mainly of sodium carbonate that may be

Table 1 Main techno-economic considerations assumed in the study

| Feature | Value/calculation/comment |
|--|---|
| Base year | 2019 |
| Currency | USD |
| Geographical scope | Europe |
| Annual total costs | Variable costs + fixed costs |
| Variable costs | RM + U + C&A |
| Raw materials (RM) | NaOH: USD 219 per t Renewable NaOH: USD 218 per t Methanol: USD 254 per t (ref. 26) Sodium carbonate: USD 223 per t Recycled NaOH: USD 363 per t Recycled renewable NaOH: USD 362 per t CaO: USD 140 per t (ref. 10) Natural gas: USD 8.5 per GJ (ref. 27) |
| Utilities (U) | Power: Grid – USD 16.3 per GJ (ref. 28) Renewable (wind) – USD 15.5 per GJ (ref. 29) Water: USD 3.7 per m ³ 5% of M |
| Consumables and auxiliaries (C&A) | OL&S + IN + M Based on European labour costs for manufacturing sectors (USD 32.0 per h) ³⁰ + 15% overhead |
| Fixed costs | OL&S + IN + M |
| Labour, supervision, laboratory and overheads (OL&S) | Based on European labour costs for manufacturing sectors (USD 32.0 per h) ³⁰ + 15% overhead |
| Insurance (IN) | 1% of TIC |
| Maintenance (M) | 3% of TIC |

RM: raw materials, U: utilities, C&A: consumables and auxiliaries, OL&S: Labour, supervision, laboratory and overheads, IN: insurance, M: maintenance.

recycled to make NaOH (case – “recycled Na₂CO₃”) or commercialized (case – “commercialized Na₂CO₃”). In order to evaluate the influence of the by-product on the carbon footprint, for the case of “commercialized Na₂CO₃”, the system was expanded in order to include the avoidance of the production of commercial sodium carbonate and lime. For the case “recycled Na₂CO₃”, sodium carbonate was assumed to be recycled back to sodium hydroxide through an electrolytic system; pure CO₂ would then be produced, which would be stored underground and not considered in the GHG balance.

The background system inventories were taken from ecoinvent 3.8,²¹ except for sodium hydroxide, for which the inventory model is modified to account for realistic and current electricity market-derived emissions during the electrolytic process of sodium hydroxide. For the foreground system, *i.e.* the production of CaO through the reaction of sodium hydroxide and calcite, the mass and energy balances for the two design options described above (A and B) were taken from the mass and energy balances produced in Aspen Plus. Energy supply was modeled considering an average, from the latest available European inventory for natural gas heat and the European mix for electricity. Average water, limestone, and methanol supplies in Europe were accounted for from ecoinvent 3.8, maximizing its representativeness.

Economic modelling

The economic assessment of the processes was carried out assuming a 50-year design life of the plant, and the total

investment cost was calculated based on the exponential model dependent on the size parameter, as per Lang’s method.²² The different equipment and sizes used in evaluating the total investment cost are shown in the ESI (Table S2).† The annual operating costs of the plant are composed of variable costs, which are dependent on the annual production, and fixed costs, which are independent of the annual production but dependent on the design capacity. Fixed costs such as maintenance, insurance or others were generally calculated as a percentage of the investment cost (Table 1).

Table 1 also summarizes the main economic assumptions used in the calculations and shows all the prices and costs already converted into USD₂₀₁₉.

Cost data were all actualized to December 2019, *i.e.*, a pre-pandemic situation and without the current (2024) price volatility, by using the Producer Price Index (PPI)²³ for raw materials and the Chemical Engineering Plant Cost Index (CEPCI)²⁴ for investment costs unless originally sourced from 2019 (Table S3†). For consistency, as most cost references for raw materials were found on a USD basis, we utilized the PPI reported by the United States Bureau of Labor Statistics. The yearly exchange rates USD/EUR were taken as the average December exchange rates each year.

Sodium hydroxide costs were estimated from the weighted average flow of import and export values reported in the UN Comtrade database.²⁵ The average was weighed considering the suppliers’ traded flow covering 90% of the total traded flow and then converted into USD₂₀₁₉ per ton of NaOH, which is taken as the “Commercial NaOH” price. The price of “Renewable NaOH” was calculated based on the “Commercial NaOH” price and the difference between European average electricity prices for non-household consumers and European average electricity prices from wind turbines.

From this value, the estimate for the price of sodium hydroxide manufactured from 100% renewable electricity (Re-NaOH) was calculated using eqn (3).

$$\begin{aligned}
 & \text{Price (Re-NaOH)}(\text{USD}_{2019} \text{ per kg NaOH}) \\
 &= \text{Price}(\text{USD}_{2019} \text{ per kg commercial NaOH}) \\
 &\quad - \text{Electricity Demand (kW h per kg NaOH)} \\
 &\quad \times (\Delta \text{Electricity Price (USD}_{2019} \text{ per kW h)}).
 \end{aligned} \quad (3)$$

The heat requirement of the processes was generated using a natural gas heater.

Results

Carbon footprint of the anion exchange process

The carbon footprint of the process was expressed in terms of net carbon footprint per ton of CaCO₃ processed, as the output from the process was dependent on the input material proportions. The net carbon footprint of the system was calculated by subtracting the carbon footprint of the output materials from the gross carbon footprint of the anion exchange process. The CO₂ footprints of lime and soda vary greatly depending on the production process, and three



different CO₂ footprints (low, high, and representative) of CaO and Na₂CO₃ were used in this study. The CO₂ footprints of lime were 1.0, 1.8, and 1.2 t CO₂ per t CaO produced for the low, high, and representative cases, respectively, based on the different kiln types used.⁹ Similarly, the CO₂ footprints of Na₂CO₃ were 0.3 and 1 t CO₂ per t Na₂CO₃ produced for the low and high cases, corresponding to the Na₂CO₃ production from high-quality trona in the USA and the synthetic Solvay

process, respectively.³ The representative CO₂ footprint of Na₂CO₃ was 0.67 t CO₂ per t Na₂CO₃ produced, as calculated from ecoinvent based on Na₂CO₃·10H₂O.³¹

The net carbon footprints of the anion exchange system are shown in Fig. S2† and Fig. 2 for processes A and B, respectively. The net carbon footprint was dependent on the NaOH type (commercial vs. renewable) and the fate of the Na₂CO₃ produced (recycled vs. commercialized). The net carbon foot-

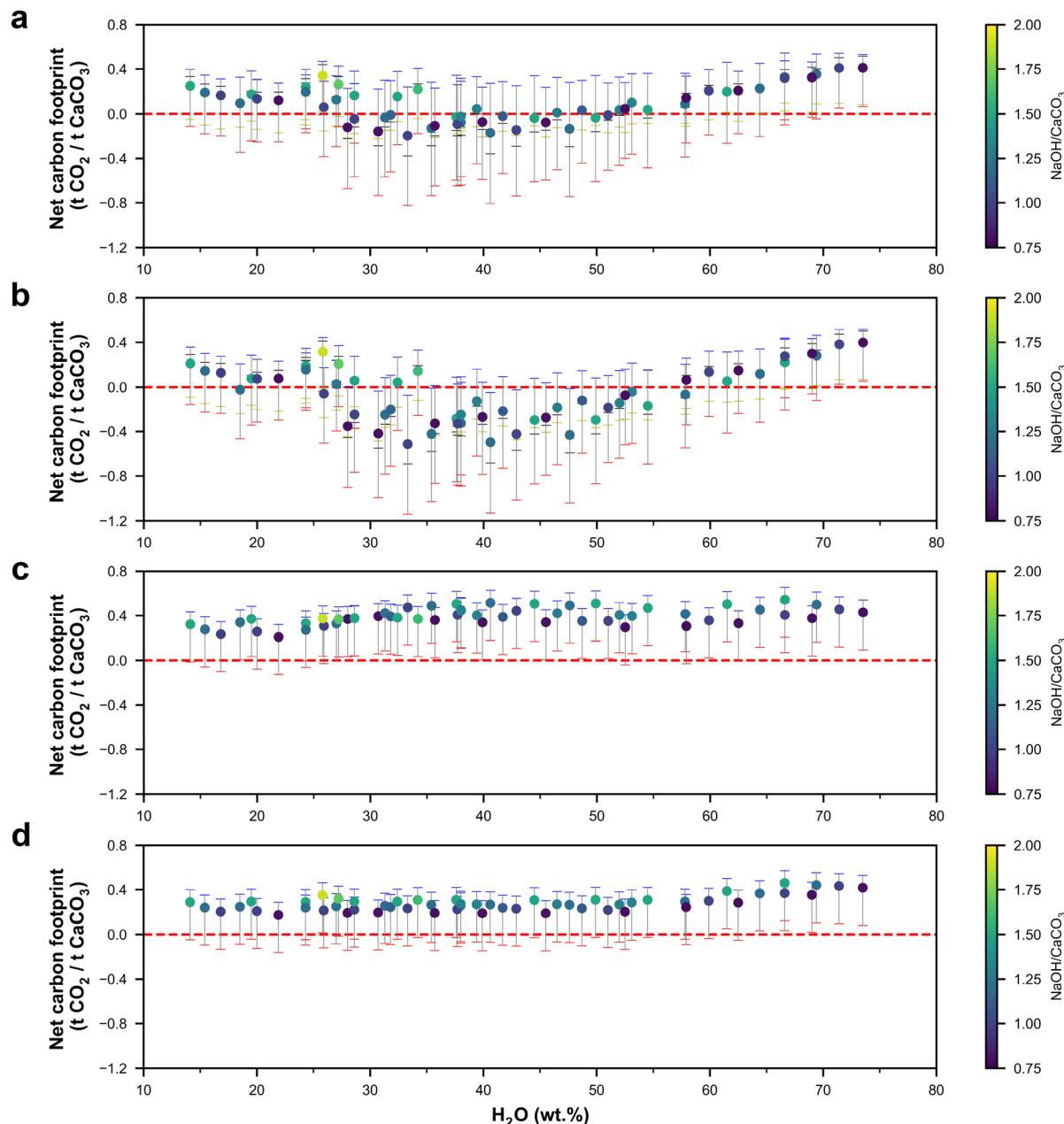


Fig. 2 Net carbon footprint of process B expressed in t CO₂ per t CaCO₃ processed compared to current CaO and Na₂CO₃ production scenarios. The four cases of materials are plotted: (a) commercial NaOH and commercial Na₂CO₃, (b) renewable NaOH and commercialized Na₂CO₃, (c) commercial NaOH and recycled Na₂CO₃, and (d) renewable NaOH and recycled Na₂CO₃. The solid circles show the representative net carbon footprint. The colored dashes on the error bar represent the following types of carbon footprint used while calculating the net carbon footprint: both CaO and Na₂CO₃ minimum carbon footprint – blue, both CaO and Na₂CO₃ maximum carbon footprint – red, CaO minimum and Na₂CO₃ maximum carbon footprint – black, and finally, CaO maximum and Na₂CO₃ minimum carbon footprint – yellow. The color of the points in all the figures represents the NaOH and CaCO₃ ratio in the mix design, as shown in the color bar.

print of the system was in the following order: commercial NaOH and recycled Na_2CO_3 > renewable NaOH and recycled Na_2CO_3 > commercial NaOH and commercialized Na_2CO_3 > renewable NaOH and commercialized Na_2CO_3 . Recycling Na_2CO_3 increased the carbon footprint of the system significantly, and apart from a few cases where it was assumed that the CaO produced replaced existing CaO production with a high carbon footprint, the net carbon footprint was positive. Recycling Na_2CO_3 to regenerate NaOH has a carbon footprint of 0.92 t CO_2 per t NaOH regenerated, which is higher than the CO_2 footprint of the currently used NaOH production process (0.78 and 0.31 t $\text{CO}_{2\text{eq}}$ per t NaOH for commercial and renewable NaOH, respectively). Furthermore, recycling Na_2CO_3 also eliminates the CO_2 savings from replacing current Na_2CO_3 production (0.67 t CO_2 per t Na_2CO_3), making the recycling of Na_2CO_3 to regenerate NaOH an unattractive method.

For the cases where Na_2CO_3 is commercialized, the net representative carbon footprint for process A could be as low as -0.15 t CO_2 per t CaCO_3 processed for commercial NaOH, and -0.47 t CO_2 per t CaCO_3 processed for renewable NaOH. Similarly, when Na_2CO_3 was commercialized for process B, a

minimum net representative carbon footprint of -0.20 and -0.50 t CO_2 per t CaCO_3 processed was achieved for commercial NaOH and renewable NaOH, respectively. When Na_2CO_3 was commercialized, the net representative carbon footprint was lower when the water content in the mix design was within ~25 to 50%, compared to that for very high or low water content. The net representative carbon footprint decreased with an increase in the conversion efficiency of CaCO_3 in the anion exchange process (Fig. 3a), as a lower conversion efficiency implies that a higher fraction of unreacted CaCO_3 is later calcined to form CaO, which directly releases 0.44 t CO_2 per t CaCO_3 calcined.

In terms of effects of NaOH type and Na_2CO_3 usage compared to the control case (*i.e.*, commercial NaOH and commercialized Na_2CO_3), using renewable NaOH reduced the net carbon footprint, and the reduction increased linearly with increasing conversion efficiency, as an increased efficiency implies that an increasing amount of NaOH is consumed per ton of CaCO_3 processed (Fig. 3b). Similarly, the recycling of Na_2CO_3 increased the net carbon footprint linearly with increasing conversion efficiency as Na_2CO_3 generation

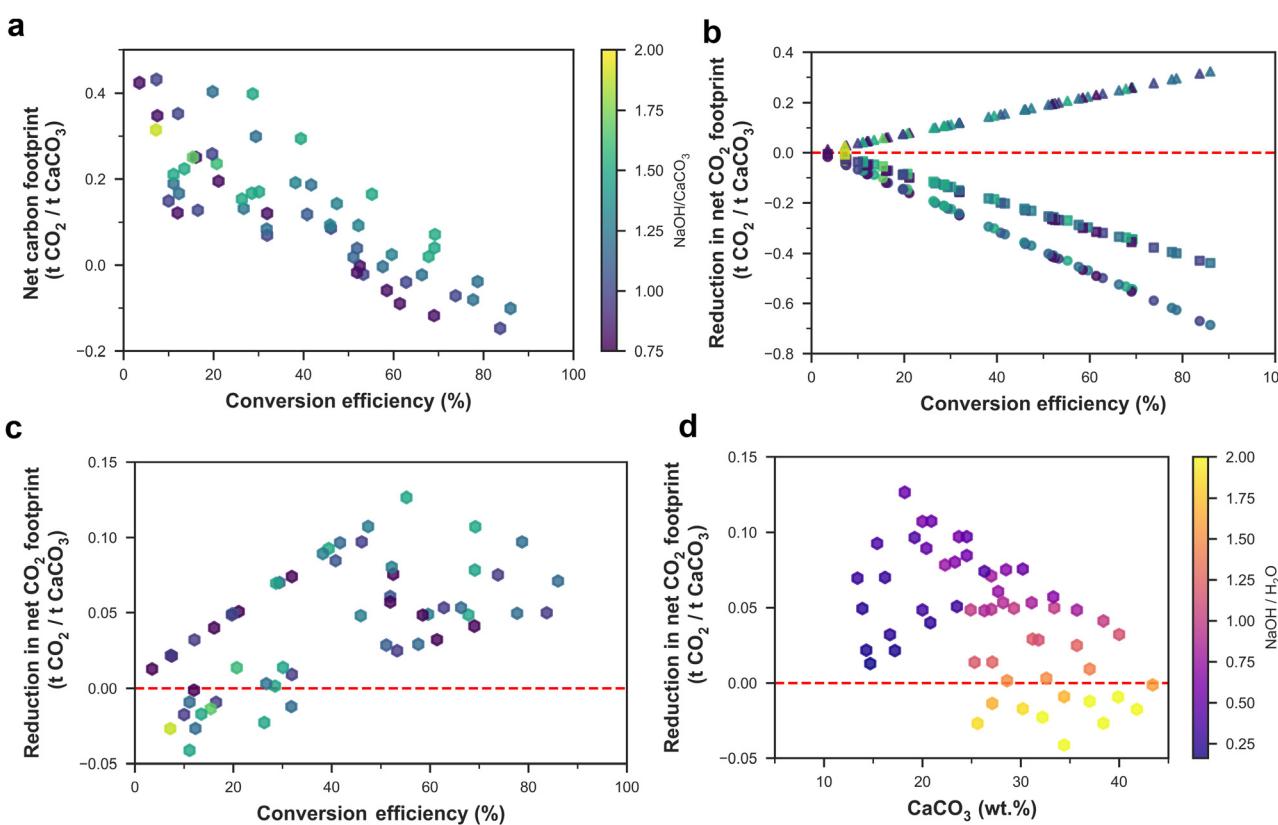


Fig. 3 Effect of raw material types, mix design, and process type on the representative net carbon footprint of the anion exchange process. The effects of conversion efficiency on the net representative carbon footprint for process A and the control case (NaOH-commercial, Na_2CO_3 -commercialized) are shown in (a). The changes in the net representative CO_2 footprint by using renewable NaOH, recycled Na_2CO_3 , and a combination of both are compared with the control case for processes A and B, as shown in (b). The reduction in the net representative CO_2 footprint by using process B compared to process A with respect to conversion efficiency and CaCO_3 content in the mixture is shown in (c) and (d), respectively. The color bar in (a) represents the ratio of NaOH to CaCO_3 (in wt. %) for (a) to (c). In all the panels, hexagons represent the case of control raw materials; triangles represent NaOH-renewable, Na_2CO_3 -commercialized; circles represent NaOH-commercial, Na_2CO_3 -recycled; and finally squares represent NaOH-renewable and Na_2CO_3 -recycled.

increased with conversion efficiency. As recycling Na_2CO_3 made the net carbon footprint positive for both processes A and B, we will not further consider these cases for techno-economic assessment. Using process B instead of process A, in general, reduced the net carbon footprint, but process B had a higher net carbon footprint when the ratio of NaOH to H_2O was higher than 1.5, similar to the trend observed in the net heat energy requirement in the next section (Fig. 4), as a higher amount of excess water had to be added to dissolve both Na_2CO_3 and NaOH to separate the Ca-containing compounds (Fig. 3d). In terms of mix design, the mix with water, NaOH, and CaCO_3 contents of 33.3, 33.4, and 33.3 wt%

(Experiment ID 6) provided the lowest net carbon footprint for both processes and NaOH types.

Energy requirements

The anion exchange process is energy-intensive, and the consumed energy requirements of this process mostly arise from drying/evaporation requirements in various steps and from calcining the unreacted CaCO_3 into CaO . The heating energy requirements were expressed per ton of CaCO_3 processed, and the heating energy requirements decreased with increasing CaCO_3 content in the mix design (Fig. 4a and b) because almost all the experimental data points had CaCO_3 as the limiting

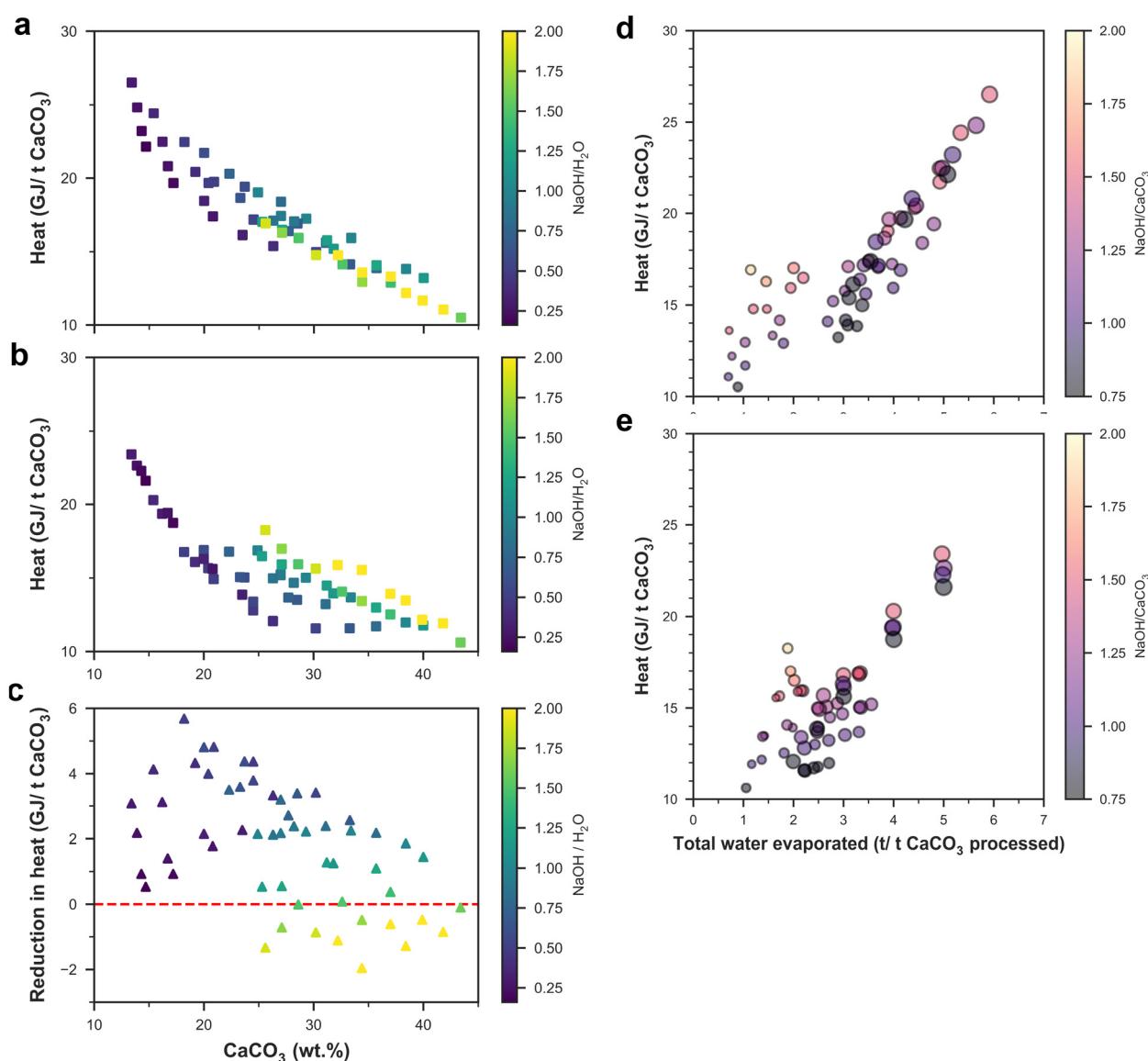


Fig. 4 Heating energy requirement of the anion exchange process. Total heating energy requirements from natural gas for the drying process and converting unreacted CaCO_3 to CaO for processes A (a) and B (b) are shown as a function of CaCO_3 content in the mix design, with the color of the scatterplot being a function of the NaOH to H_2O ratio in the mix design. The reduction in heat requirement in process B compared to process A is shown in (c). The effects of total water evaporated during the anion exchange process on the total heat requirement for processes A (d) and B (e) are also plotted. The color of the scatterplot depends on the NaOH to H_2O ratio in the mix design, whereas the sizes of the scatter points are proportional to the water content in the mix design.



reagent (*i.e.*, excess NaOH) as the goal of the anion exchange process is to convert all CaCO_3 into Ca(OH)_2 . The heating energy requirements of process B were lower than those for process A for most of the mix designs except for the few cases when the ratio of NaOH to H_2O was higher than 1.5 (Fig. 4c). A higher NaOH to H_2O ratio meant that a higher amount of excess water (which was evaporated in subsequent steps) had to be added to dissolve Na_2CO_3 and NaOH to separate the Ca-containing compounds. Evaporating the water consumes a significant amount of energy as the latent heat of water evaporation (2.26 GJ per ton) is comparable to the heat of reaction for calcining the unreacted CaCO_3 to CaO (1.78 GJ per ton). However, the amount of water evaporated during the whole anion exchange process is much higher than unreacted CaCO_3 as water is added to dissolve the Na_2CO_3 , and the total heat energy required in the process mainly depends on the total amount of water evaporated (Fig. 4d and e).

Economic feasibility of the anion exchange process including carbon credits

The total cost of the anion exchange process was calculated as the sum of operating expenses (OpEx) and capital expenses (CapEx) distributed over the 50-year lifetime of the processing plant, which can process 1000 tons of CaCO_3 per day. However, different relative quantities of starting materials produce different amounts of Na_2CO_3 and lime, and to

compare the different processes and types of NaOH, we calculated the profit per ton of CaCO_3 processed after selling the Na_2CO_3 and CaO produced. In general, the profit increased as the amount of CaCO_3 in the mix design and conversion efficiency increased and was the highest for a medium NaOH to H_2O ratio (Fig. S3 and S4†). None of the mix designs turned out to be profitable for process A, whether using commercial NaOH or renewable NaOH. On the other hand, process B turned a maximum profit of \$3.6 per t CaCO_3 and \$4.2 per t CaCO_3 when using commercial and renewable NaOH, respectively (Fig. S3c and S3d†). In terms of mix design, the mix with water, NaOH, and CaCO_3 contents of 30.7, 30.9, and 38.4% (Experiment ID 19) provided the highest profitability for both processes and NaOH types. In most of the mix designs, the anion exchange process would not turn a profit highlighting the importance of choosing an efficient mix design, and further experiments may be performed to fine-tune the mix designs for process B to obtain higher profitability.

The net carbon footprint also generally decreased with increasing profitability (Fig. S5†), as both parameters showed a strong correlation with conversion efficiency. Although the anion exchange process is profitable, the gross profit margin is relatively low (~2% for process B). However, selling carbon credits (\$80 per t CO_2) arising from the net carbon footprint reduction would make this process more profitable (Fig. 5),

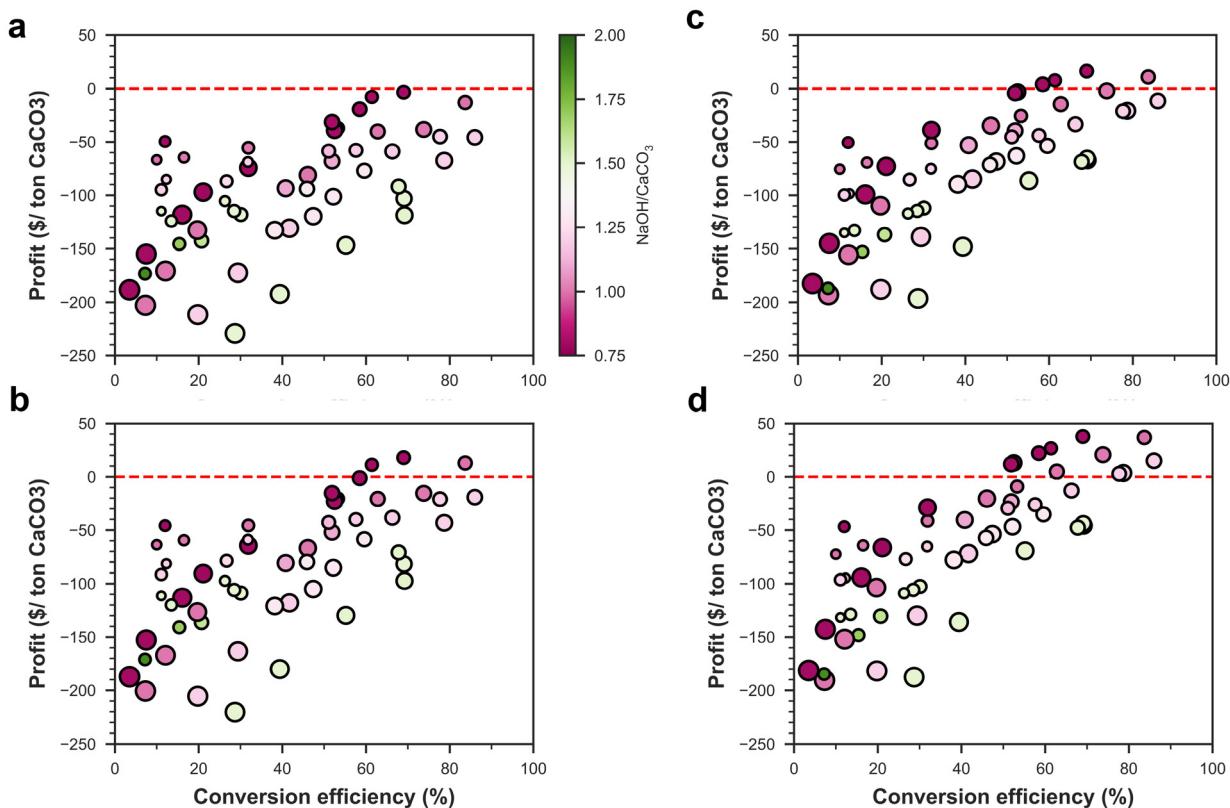


Fig. 5 Profitability vs. net carbon footprint of the anion exchange process after selling carbon credits. Process A – (a) commercial NaOH and commercialized Na_2CO_3 , and (b) renewable NaOH and commercialized Na_2CO_3 . Process B – (c) commercial NaOH and commercialized Na_2CO_3 , and (d) renewable NaOH and commercialized Na_2CO_3 . The color of the scatterplot depends on the NaOH to CaCO_3 ratio in the mix design, whereas the sizes of the scatter points are proportional to the water content in the mix design.



with a gross margin of 16% for renewable NaOH and 6% for commercial NaOH. Even after selling carbon credits, the mix design with Experiment ID 19 remains most profitable for both processes and NaOH types. This mix design had a CaCO_3 conversion efficiency of 69%, and thus, for 1-ton CaO production, 1.3 ton of Na_2CO_3 is generated and 0.49 kg of NaOH is consumed. In terms of major factors affecting the costs (Fig. 6), we found that heating energy and NaOH both contribute to almost ~50% of the total cost each for both commercial and renewable NaOH usage, whereas selling carbon credits, and selling CaO and Na_2CO_3 were the major sources of revenue. The CO_2 emissions had another major contributor along with NaOH and heating energy – the direct CO_2 emission arising from calcining the unreacted CaCO_3 to form CaO . Thus, the economic viability of the process is sensitive to changes in the price of carbon credit, natural gas, NaOH, Na_2CO_3 , and CaO .

Integration with the production of cementitious materials

The anion exchange process could ideally be integrated with a cement plant because the unreacted NaOH from the anion exchange reaction can be used to capture CO_2 produced during Portland cement manufacturing (Fig. 7) and has been named process C. Process B was only considered for integration with the production of cement clinker as it was more economical and had a lower CO_2 footprint than process A for optimized mix designs. In process C, the unreacted NaOH after the anion exchange process would react with CO_2 to form Na_2CO_3 , which can then be commercialized. This integration removes the need for lime transportation to the cement plant, reducing costs and CO_2 emissions. In some cases, the amount of CO_2 needed to convert all the unreacted NaOH into Na_2CO_3 may exceed the CO_2 produced by the cement plant using the lime from the anion exchange process. In such instances,

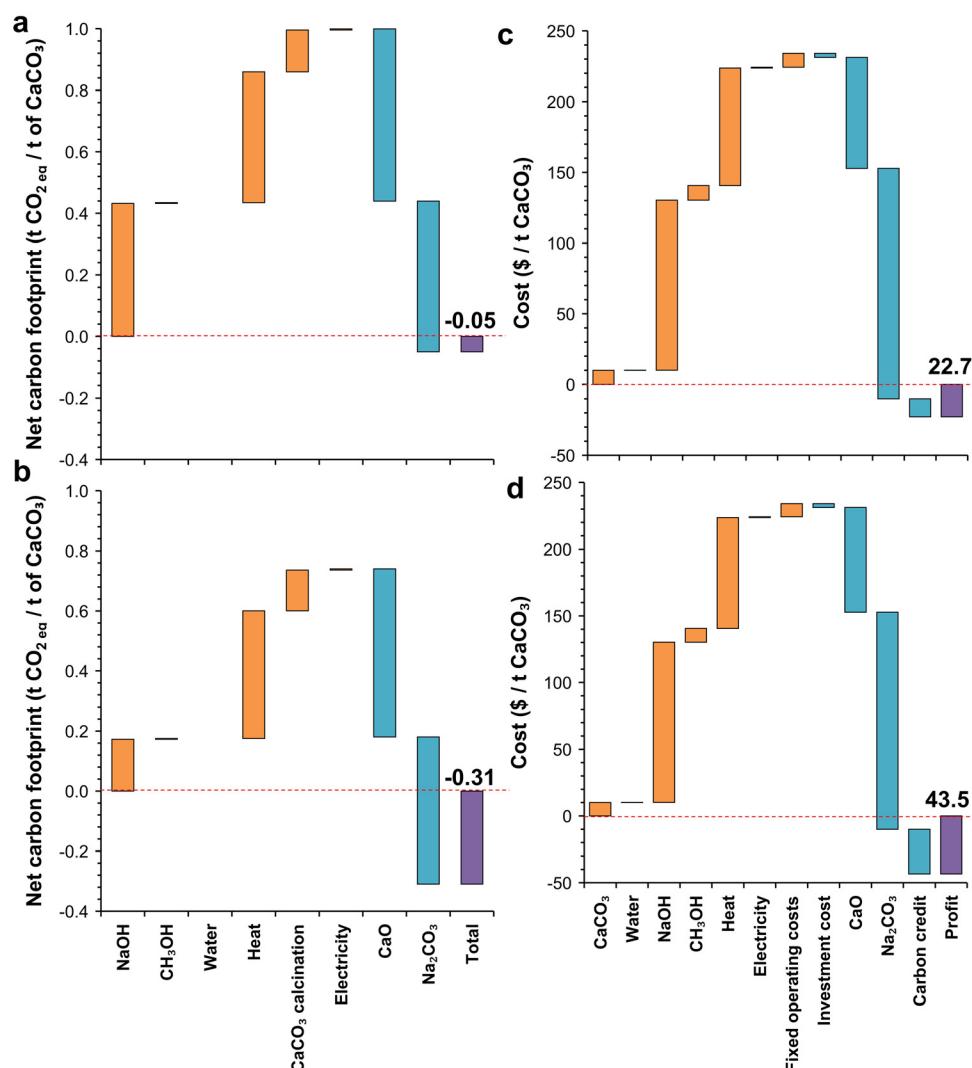


Fig. 6 Different components of carbon footprint (a and b) and costs (c and d) of the anion exchange process B for the mix design Experiment ID 19 (water: 30.7, NaOH: 30.9, and CaCO_3 : 38.4 wt%). The cases for commercial NaOH are shown in (a) and (c), whereas renewable NaOH cases are shown in (b) and (d).



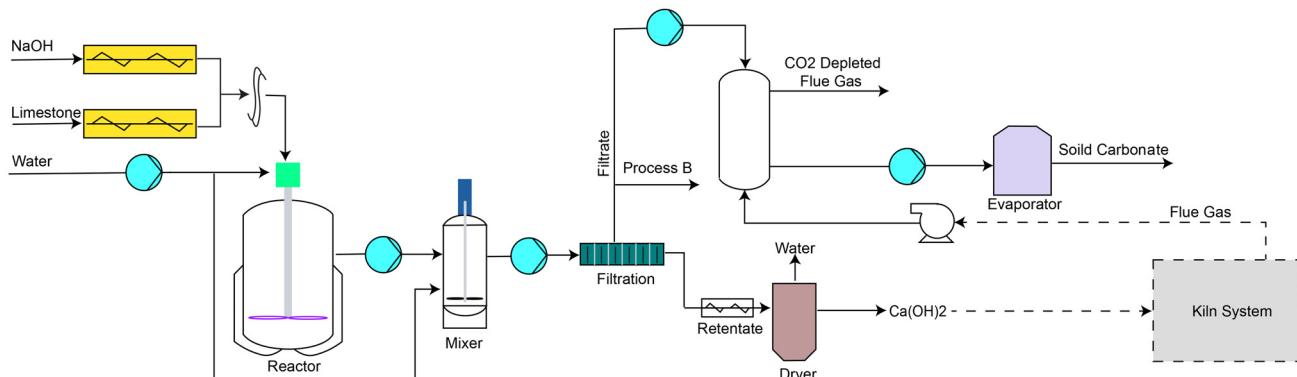


Fig. 7 Schematic diagram showing the integration of anion exchange process B with the cement kiln system.

some of the products from the anion exchange reaction can be separated using the original process B. The amount of CO_2 captured by NaOH to form Na_2CO_3 depends on the pH and temperature of the NaOH solution.³²

Considering Experiment ID 19 mix design with renewable NaOH, we found that the amount of CO_2 produced by the

cement plant using the lime from the anion exchange process was sufficient to convert all the unreacted NaOH into Na_2CO_3 . Integrating the anion exchange process with a cement plant for Experiment ID 19 mix design would reduce the net carbon footprint by 0.15 t CO_2 per t CaCO_3 with a final net footprint of -0.45 t CO_2 per t CaCO_3 . However, as recycling Na_2CO_3 to

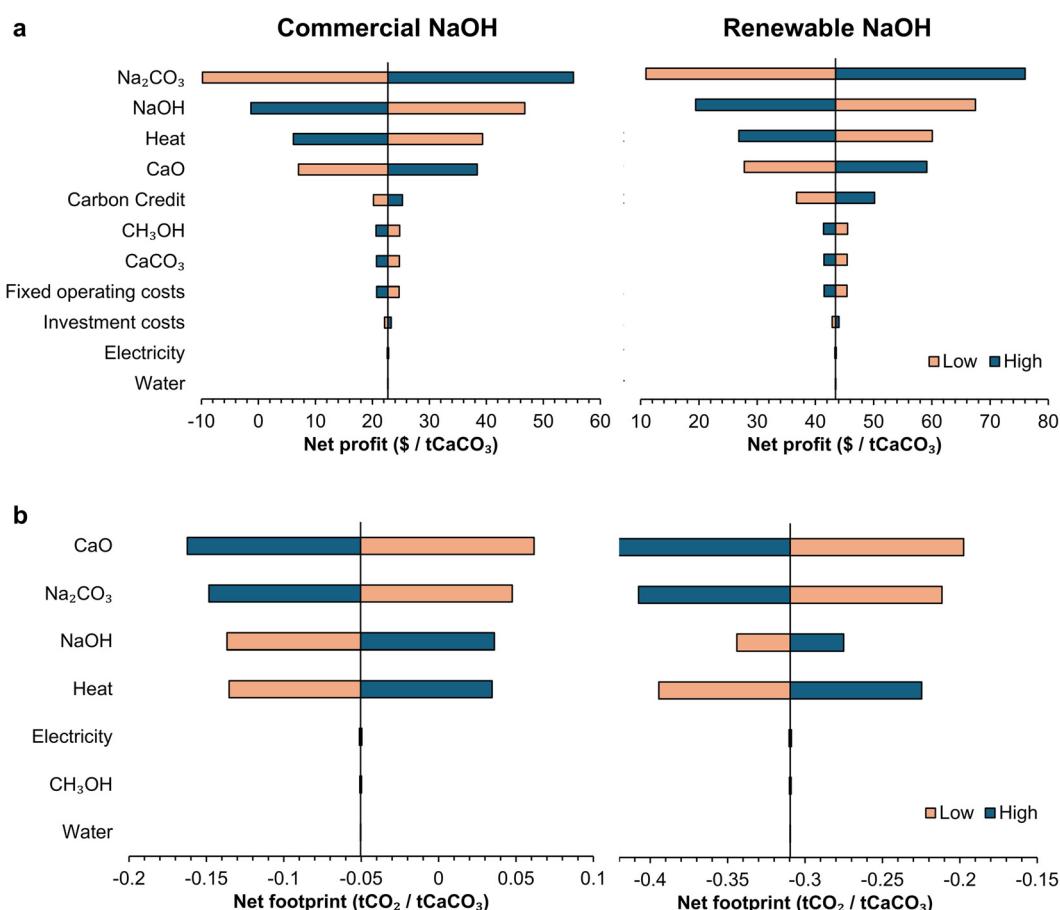


Fig. 8 Effects of different factors on the profitability (a) and CO_2 emissions (b) of the anion-exchange process. The sensitivity of the profit and CO_2 footprint was calculated when the cost and CO_2 footprints were lowered ("low") or increased ("high") by 20%. The analysis was performed for process B for the mix design Experiment ID 19 (water: 30.7, NaOH: 30.9, and CaCO_3 : 38.4 wt%) for both renewable and commercial NaOH cases.

generate NaOH was not environmentally friendly, modifying a significant portion of global cement production capacity with this anion exchange process in an economically sustainable way would not currently be possible without finding a new avenue for utilizing the significant amount of Na_2CO_3 co-produced or realising advances in NaOH production that may allow the storage of Na_2CO_3 to be economical. Assuming that the annual global consumption of Na_2CO_3 stays unchanged (65 Mt), this anion exchange process could produce ~ 50 Mt of lime, immediately reducing the global CO_2 emission by 37.3 Mt per year using process B.

Challenges and opportunities of scaling up the anion exchange process

The anion exchange process should be ideally used with renewable NaOH, and if sufficient supply is not available, new production facilities of NaOH may need to be constructed. Renewable NaOH production requires the availability of renewable electricity, and constructing a new NaOH production facility for the anion exchange process may improve economics through vertical integration at the cost of increased initial investment. One of the ways to decrease the initial investment cost would be to re-purpose equipment currently used for the Na_2CO_3 production process, such as filtration columns, distillation columns, pumps, *etc.*

Currently, the Na_2CO_3 and $\text{Ca}(\text{OH})_2$ production processes are decoupled, as the modified Solvay process (or Hou's process) no longer uses CaO to regenerate NH_3 , the catalyst used in the original Solvay method. Adopting co-production of Na_2CO_3 and $\text{Ca}(\text{OH})_2$ would change the current global material flow, which may affect the costs and CO_2 footprint of transporting these materials from the production location to the customers. The limited global demand for Na_2CO_3 compared to that for $\text{Ca}(\text{OH})_2$ (including potential demand for Portland cement production) is one of the bottlenecks of scaling-up the anion-exchange process beyond satisfying the global Na_2CO_3 demand. Potential avenues of increased usage of Na_2CO_3 are in glass production, water purification, and the construction industry, including activating secondary raw materials (slags) with Na_2CO_3 (~ 5 wt% of slag)^{33,34} to reduce the global clinker demand and thus reduce global CO_2 footprint.³⁵

To understand the effect of pricing volatility and changing carbon footprint associated with the materials involved in the anion exchange process, we performed a sensitivity analysis of the anion exchange process for the mix design Experiment ID 19 (water: 30.7, NaOH: 30.9, and CaCO_3 : 38.4 wt%), when the various factors were changed by 20% (Fig. 8). The economic feasibility of this anion-exchange process is highly sensitive to the market price of the major input and output raw materials, *i.e.*, Na_2CO_3 , NaOH, and CaO, apart from CaCO_3 , as CaCO_3 is inexpensive compared to the other materials. Heating energy is also one of the key parameters – and the high volatility in the price of natural gas (or any other fuel) needs to be considered when implementing this process. The initial investment cost and fixed operating expense do not influence the economic feasibility of this simple process. Similarly, the net

CO_2 footprint of the process is also primarily sensitive to the CO_2 footprint of the raw materials other than CaCO_3 and heating energy. The CO_2 emission and cost of the anion exchange process could be further lowered by electrifying the industrial process heat used.^{36,37}

Conclusions

We show that the co-production of CaO and Na_2CO_3 using the anion exchange process could reduce the carbon emissions economically today through the industrial symbiosis of the soda ash and lime manufacturing, especially if NaOH produced from renewable energy is used. Separating the Ca-containing compounds (CaCO_3 and $\text{Ca}(\text{OH})_2$) first from the mixture of reactants and products after the anion exchange reaction seemed to be more economically and environmentally efficient than separating the Na-containing compounds at the beginning. The most efficient mix design (water: 30.7, NaOH: 30.9, CaCO_3 : 38.4 wt%) had a net profit of \$43.5 per t CaCO_3 processed and reduced the carbon footprint by 0.31 t CO_2 per t CaCO_3 compared to the current production process.

Decarbonizing a significant portion of the cement industry using this technology requires new avenues for utilizing the Na_2CO_3 produced and significant developments in global NaOH production, as recycling Na_2CO_3 to regenerate NaOH was not economically and environmentally favorable. The proposed process is highly sensitive to the CO_2 emissions and market price associated with NaOH and Na_2CO_3 , and the supply of heating energy.

We show that this anion exchange technology, even when using natural gas to supply heating energy, can be used immediately to decarbonize the lime and soda ash industries. When NaOH produced using renewable energy is utilized to produce soda ash and lime employing process B, an annual reduction of 37.3 Mt CO_2 emission can be achieved compared to the current production process.

Data availability

The data supporting the findings of this study are included within the main article and the ESI.†

Conflicts of interest

Theodore Hanein is a shareholder in United earth4Earth Holding.

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