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PERSPECTIVE

The technological and economic prospects for CO₂ utilisation and removal

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Capturing and utilising carbon dioxide to create valuable products might lower the net costs of reducing emissions or removing carbon dioxide from the atmosphere. Here we review ten such pathways. Pathways involving chemicals, fuels, and microalgae might reduce carbon dioxide emissions but have limited potential for carbon dioxide removal; pathways involving construction materials can both utilise and remove carbon dioxide. Land-based pathways can increase agricultural output and remove carbon dioxide. Our assessment suggests that each pathway could scale to over 0.5 Gt carbon dioxide utilisation annually, although barriers remain substantial and resource constraints prevent the simultaneous deployment of all pathways.

CO₂ utilisation is receiving increased interest from the scientific community¹, partly due to climate change considerations and partly because using CO₂ as a feedstock can be a cheaper or cleaner production process than using conventional hydrocarbons². CO₂ utilisation is often promoted as a way to reduce net costs – or increase profits – of reducing emissions or removing carbon dioxide from the atmosphere, and therefore to help scale mitigation or removal efforts³. CO₂ utilisation is also seen variously as a stepping stone towards⁴ or a distraction away from⁵ the successful implementation of carbon capture and storage (CCS) at scale.

In most of the literature, including the IPCC 2005 Special Report on CCS⁶, the term 'CO₂ utilisation' refers to the use of CO₂, at concentrations above atmospheric levels, directly or as a feedstock in industrial or chemical processes, to produce valuable carbon-containing products⁶⁻¹¹. Included in this conventional definition is the industrial production of fuels using, e.g., amines to capture and concentrate the CO₂ from air, potentially with solar energy. However, excluded is the production of an identical fuel from the same essential inputs, but where the CO₂ is captured by plant-based photosynthetic processes.

Here, we consider CO₂ utilisation to be a process in which one or more economically valuable products are produced using CO₂, whether the CO₂ is supplied from fossil-derived waste gases, captured from the atmosphere by an industrial process, or – in a departure from most (but not all^{12,13}) of the literature – captured biologically by land-based processes. Biological or land-based forms of CO₂ utilisation can generate economic value in the form of, e.g., wood products for buildings, increased plant yields from enhanced soil carbon uptake, and even the production of biofuel and bioderived chemicals. We employ this broader definition deliberately; by thinking functionally, rather than narrowly about specific processes, we hope to promote dialogue across scientific fields, compare costs and benefits across pathways, and consider common techno-economic characteristics across pathways which could potentially assist in identifying routes to climate change mitigation.

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In this perspective, we consider a non-exhaustive selection of ten CO₂ utilisation pathways. We provide a transparent assessment of the potential scale and cost for each pathway, namely: (i) CO₂-based chemical products, including polymers; (ii) CO₂-based fuels; (iii) microalgae fuels and other microalgae products; (iv) concrete building materials; (v) CO₂ enhanced oil recovery (CO₂-EOR); (vi) bio-energy with carbon capture and storage (BECCS); (vii) enhanced weathering (EW) (viii) forestry techniques including afforestation/reforestation (AF/RF), forest management and wood products; (ix) land management via soil carbon sequestration (SCS) techniques; and (x) biochar.

These ten CO₂ utilisation pathways (described in Table 1 and in the Supplementary Materials and depicted by the thin arrows in Figure 1) can also be characterised as 'cycling', 'closed' and 'open' utilisation pathways. For instance, many (but not all) conventional industrial utilisation pathways, such as CO₂-based fuels and chemicals, tend to be 'cycling': they move carbon through industrial systems over timescales of days, weeks or months. Such pathways do not provide net CO₂ removal from the atmosphere but can reduce emissions via industrial CO₂ capture that displaces fossil fuel use. In contrast, 'closed' pathways involve utilisation and near-permanent CO₂ storage, such as in the lithosphere (via CO₂-EOR or BECCS), deep oceans (via terrestrial EW), or in mineralised carbon in the built and natural environments. Finally, 'open' pathways tend to be biologically-based and characterised by large removal potentials and storage in 'leaky' natural systems such as biomass and soil with risks of large-scale flux back to atmosphere.

Our pathways include some novel or emerging pathways, such as a CO₂-based fuel economy, for which current flows are near-zero, as well as established pathways such as CO₂-EOR and afforestation/reforestation. Pathways were selected based upon discussions at a joint meeting of the US National Academy of Sciences and the UK Royal Society¹; each is relatively well studied to date and has an acknowledged potential to scale. Many other pathways meet our definition but are not reviewed here (Supplementary Materials).

This perspective is structured as follows. First, the ten utilisation pathways are presented in the context of the scale of CO₂ stocks and flows on Earth. Second, the potential scale and economics of each pathway is assessed. Third, a selection of key barriers to scaling are identified. Fourth, we assess the outlook for CO₂ utilisation, and conclude with future research and policy priorities.

CO₂ utilisation pathways and the carbon cycle

Carbon dioxide utilised by a pathway (CO_2u) is not necessarily the same as carbon dioxide removed (CO_2r) or carbon dioxide stored (CO_2s); does not necessarily reduce emissions ($CO_2\rho$); and does not necessarily deliver a net climate benefit, once indirect and other effects are accounted for. The various concepts overlap and relate to each other, but are distinct (Figure S1, Supplementary Materials). Some carbon capture and utilisation (CCU) processes achieve carbon dioxide removal (CDR) from the atmosphere. Some CCU processes involve carbon capture and storage (CCS). CCS itself can contribute either to mitigation (e.g. reducing net emissions from a gas-fired power plant) or to atmospheric removals (e.g. direct air carbon capture and storage, or DACCS) – CCS does not necessarily imply CDR. Further, CCS and CDR can fail to deliver a climate benefit. For instance, perverse indirect effects such as land-use change resulting from $BECCS^{14}$ could increase net atmospheric CO_2 concentrations.

CO₂ utilisation does not necessarily contribute to addressing climate change, and careful analysis is essential to determine the overall impact. Identifying the counterfactual (what would have happened without CO₂ utilisation) is important but often particularly challenging, and the impact of a given CO₂ utilisation pathway on the mitigation of climate change varies as a function of space and time (Box 1).

Box 1: Concepts: CO₂ utilisation, removal, storage, reduced emissions and net climate benefit

Does CO_2 utilisation (CO_2 u) leads to a climate benefit? It might reduce emissions (CO_2 p), or remove CO_2 (CO_2 r) from atmosphere and/or store it (CO_2 s). But various direct and indirect effects over the relevant life-cycle must be considered and compared to a plausible baseline or 'counterfactual' – what would have happened without CO_2 utilisation⁸⁴. Assiduously calculating direct impacts in one place, and one time, is of little use if there is a 'waterbed effect' (also referred to as a 'rebound' or 'leakage') and emissions occur somewhere else, or later.

For instance, a barrel of oil derived via CO_2 -enhanced oil recovery (CO_2 -EOR) utilises CO_2 which can remain in the oil formation rather than being re-emitted into the atmosphere. Assuming the CO_2 does not return to atmosphere, the CO_2 utilised is equal to the CO_2 emissions stored, i.e. $CO_2u = CO_2s$, but whether $CO_2r \ge 0$ depends upon the source of the CO_2 ; if it is from a fossil power station, there is no net removal of CO_2 from atmosphere. Emissions have been reduced, and $CO_2\rho = CO_2u = CO_2s > 0$, even though $CO_2r = 0$.

To see this, consider a 'reference' scenario in which 1 t CO_2 is emitted from a fossil power plant, and 1.5 t CO_2 are emitted from oil use, such that total emissions are 2.5 t CO_2 . Compare this to a 'utilisation' scenario, in which the CO_2 from the power plant is used for CO_2 -EOR instead, i.e., CO_2 u = 1 t CO_2 . Total emissions in this 'utilisation' scenario comprise the 1.5 t CO_2 from the consumption of the CO_2 -EOR oil. $CO_2 \rho = 2.5 - 1.5 = 1.0$ t $CO_2 \rho$ which is identical to the CO_2 u, but net CO_2 r = 0 because the CO_2 came from fossil power plant, rather than the atmosphere.

In reality, the emissions from the baseline barrel of oil that was displaced by the CO₂-EOR oil might be higher or lower, depending on its origin and production process. If the CO₂-EOR oil displaces the use of renewable electricity in an electric vehicle, CO₂-EOR generates a net increase in emissions. If CO₂-EOR is to offer net removals, the CO₂ must be captured from the atmosphere, and more carbon must injected into the well than extracted.

Life-cycle analyses (LCAs) on some industrial CO_2 utilisation pathways suggests that the potential for net emission reductions ($CO_2\rho$) is much larger than for net removals (CO_2r), which appears very modest⁹⁵. Up to 3 t CO_2 emissions may be avoided for every 1 t CO_2 utilised in polycarbonate polyols², even though no CO_2 is removed from atmosphere. Nearly 4 t CO_2 emissions may be avoided for each tonne of dry wood utilised that displaces concrete-based materials⁹⁶.

Other LCAs have found neutral or negative impacts of CO_2 utilisation on $CO_2\rho^{75,97-99}$. For instance, CO_2 utilisation pathways requiring energy inputs that are not decarbonised may result in net lifecycle increases in CO_2^{97-100} .

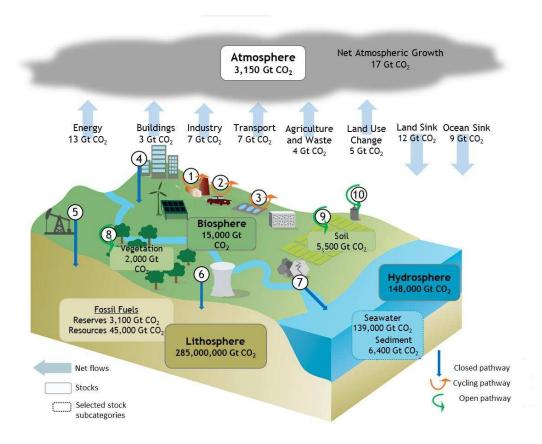


Figure 1 | **Stocks and net flows of CO₂ including potential utilisation and removal pathways.** Orange, blue and green arrows (numbered 1-10, as described in Table 1) represent cycling, closed and open pathways for utilisation and removal. Grey block arrows represent annual flows to and from the atmosphere, with estimates averaged over the 2008-2017 period ^{15,92}. Solid boxes represent stocks in the Earth's spheres, with all estimates based on IPCC estimates ¹⁶ except where noted, and converted from C to CO₂. Carbon stocks in the hydrosphere comprise of seawater, sediment, and dissolved organic carbon (not shown, ~2,600 Gt CO₂). The vast majority of carbon stocks in the lithosphere are locked in the Earth's crust⁹³, with estimated accessible fossil fuel reserves and resources of >45,000 Gt CO₂²⁵. Atmospheric stocks are converted from the 2017 estimates of atmospheric CO₂ of 405 ppm⁹⁴ using a conversion factor of 2.12. Carbon stocks in the biosphere include those stored in permafrost and wetlands (not shown, ~7,500 Gt CO₂), vegetation, and soils. Soil stocks to 1m depth have been recently estimated at 5,500 Gt CO₂⁶³. Illustration by Jillian Ditner and Ella Adlen.

For CO₂ utilisation to contribute usefully to reducing atmospheric CO₂ concentrations, the scale of the pathways must be meaningful in comparison with the net flows of CO₂ in Figure 1. The flux of carbon from fossil fuels and industry to the atmosphere (34 Gt CO₂ yr⁻¹)¹⁵) is dwarfed by the gross flux to land via photosynthesis in plants (440 Gt CO₂ yr⁻¹)¹⁶. However, only 2–3% of this photosynthetic carbon remains on land (12 Gt CO₂ yr⁻¹, Figure 1), and only for decades; the remainder is re-emitted by plant and soil respiration. If soil carbon uptake could be increased by 0.4% per annum, this would contribute to achieving net zero emissions, as per the '4 per mille' initiative¹⁷, but this is challenging¹⁸. Five of the ten pathways in Table 1 leverage our ability to perturb these land-based fluxes.

The other five conventional industrial CO₂ utilisation pathways could also perturb the net flows of CO₂. The production of plastics and other products creates a demand for so-called 'socioeconomic carbon'¹⁹ (~2.4 Gt CO₂ yr⁻¹, of which around two-thirds are wood products) that could be met in part through CO₂ utilisation. The total stock of carbon accumulated in products (such as wood products, bitumen, plastic and cereals) has been estimated at 42 Gt CO₂ in 2008, of which 25 Gt CO₂ is in wood products¹⁹. Up to 16 Gt CO₂ has been sequestered in human infrastructure as mineralised carbonates in cement from 1930-2013, at current rates of ~1 Gt CO₂ yr^{-1 20,21}.

Table 1 | Ten CO₂ utilisation and removal pathways

Table 1 Ten CO ₂ utilisation and removal pathways							
Pathway (Fig 2#)	Removal and/or Capture (Figure 2 step A, B and/or C)	Utilisation product [Figure 2 yellow boxes]	Storage ^a (Figure 2 step D, E, or F) and likelihood of release (high/low)	Emission on use (Figure 2 step G) / Release during storage (Figure 2 steps H, I or J)	Figure 2 Cycle examples		
(1) Chemicals	Catalytic chemical conversion of CO ₂ from flue gas or other sources into chemical products	CO ₂ -derived platform chemicals such as methanol, urea and plastics	Various chemicals (days/decades) - high	Hydrolysis or decomposition	«KCLG»		
from CO ₂					«KCLF»		
					«ALFJ»		
					«ALG»		
(2) Fuels from	Catalytic hydrogenation	CO ₂ -derived fuels	Various fuels	Combustion	«KCLG»		
CO ₂	processes to convert CO ₂ from flue gas or other sources into fuels	such as methanol, methane, and Fischer-Tropsch- derived fuels	(weeks/months) - high		«ALG»		
(3) Products from microalgae	Uptake of CO ₂ from the atmosphere or other sources by microalgae biomass	Biofuels, biomass, or bioproducts such as aquaculture feed	Various products (weeks/months) - high	Combustion (fuel) or consumption (bioproduct)	«KCLG» «BG»		
(4) Concrete	Mineralisation of CO ₂	Carbonated	Carbonates	Extreme acid	«KCLF»		
building materials	from flue gas or other sources into industrial waste materials, and CO ₂ curing of concrete	aggregates or concrete products	(centuries) - low	conditions	«ALF»		
(5) CO ₂ -EOR	Injection of CO ₂ from flue gas or other sources into oil reservoirs	Oil	Geological sequestration (millennia) - low b	N/A	«KCD»		
(6) Bioenergy with carbon capture and storage (BECCS)	Growth of plant biomass	Bioenergy crop biomass	Geological sequestration (millennia) - low ^b	N/A	«BCD»		
(7) Enhanced weathering (EW)	Mineralisation of atmospheric CO ₂ via the application of pulverised silicate rock to cropland, grassland and forests	Agricultural crop biomass	Aqueous carbonate (centuries) - low	Extreme acidic conditions	«BE»		
(8) Forestry techniques	Growth of woody biomass via afforestation, reforestation or sustainable forest management	Standing biomass, wood products	Standing forests and long-lived wood products (decades to centuries) - high	Disturbance, combustion or decomposition	«BFJ»		
(9) Soil carbon sequestration (SCS) techniques	Increase in soil organic carbon content via various land management practices	Agricultural crop biomass	Soil organic carbon (years to decades) - high	Disturbance or decomposition	«BFJ»		
(10) Biochar	Growth of plant biomass for pyrolysis and application of char to soils	Agricultural or bioenergy crop biomass	Black carbon (years to decades) - high	Decomposition	«BFJ»		

 $^{^{}a}$ Storage durations represent best-case scenarios. For instance, in CO₂-EOR if the well is operated with complete recycle, the CO₂ is trapped and can be stored on a centuries timescale or more 22 This is also relevant only for conventional operations.

^b Release during geological storage is usually a consequence of engineering implementation error.

The flow of CO₂ through the different utilisation pathways can be represented by a combination of different steps (labels A to L, ure 2 and Table 1). Utilisation pathways often (but not always) involve removal (**«A»** or **«B»**) and storage (**«D»**, **«E»** or **«F»**), but the permanence of CO₂ storage varies dramatically from one utilisation pathway to another, with storage timeframes ranging from days to millennia. Permanence partially depends upon where the carbon ends up (Figure 1): (i) the lithosphere, by geological sequestration into reservoirs such as saline aquifers or depleted oil and gas reservoirs, or by mineralisation into rocks; (ii) the biosphere, in trees, soils and the human built environment; or (iii) the hydrosphere, with storage in the deep oceans. Geological storage, when executed correctly, is considered to be more permanent²² than storage in the biosphere, which is shorter and subject to more human and natural disturbances²³ like wildfires and pests, as well as changes in climate²⁴. However, even 'closed' pathways do not offer completely permanent storage over geological timescales (>100,000 years²⁵), giving rise to intergenerational ethical questions²⁶.

In the short run, products created with concentrated CO_2 by «L» (albeit, CO_2 conversion is not a necessary requirement for utilisation) could leverage industrial capture of flue gases following the extraction and combustion of fossil fuels «KC»²⁷. In the longer term, the CO_2 loop will need to be closed to achieve net zero emissions, implying that CO_2 will need to be sourced from the atmosphere, potentially via direct air capture (DAC) «A» or land-use based uptake by photosynthesis or mineralisation «B». For instance, net zero CO_2 -based fuels must shift the current flows of carbon, from lithosphere to atmosphere «KCLG», to an atmosphere to atmosphere cycle «ALG» (Figure 2).

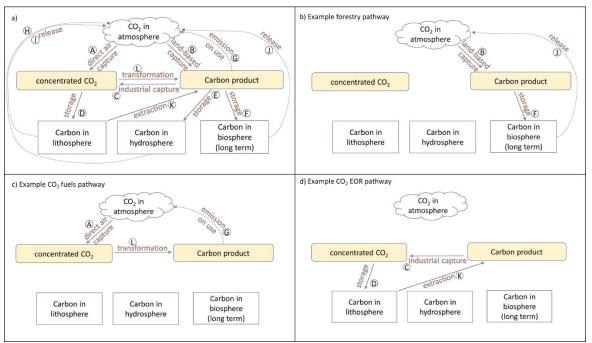


Figure 2| **Carbon dioxide utilisation and removal cycle.** Utilisation pathways are represented as a combination of steps, A-L. Panel a) represents the full cycle, with panels b), c) and d) showing three potential routes through the cycle. Conventional fossil fuel extraction, combustion and CO₂ emission to the atmosphere traces «KG». Panel b) uses the example of an open pathway such as forestry (pathways 8-10 in Table 1 and Figure 1). Panel c) uses the example of CO₂ fuels (pathway 2 in Table 1 and Figure 1) using DAC to acquire CO₂. Panel d) is an example of CO₂-EOR (pathway 5 in Table 1 and Figure 1). Cycling pathways (with the exception of polymers) end with G; closed pathways end with D, E or F; and open pathways end with J. See Table 1 for further description. All flows are net of process emissions. **E**

Scale and economics of utilisation pathways

We assess the peer-reviewed literature comprising over 11,000 papers on the ten pathways. For the conventional pathways, our scoping review covered over 5,000 papers, a minority (n=186) of which provide cost estimates. Estimates of potential scale were informed by a structured estimation process and an expert opinion survey. For the non-conventional, land-use-based utilisation pathways, we build upon existing CO_2 removal estimates (also derived from a scoping review²⁸ of over 6,000 papers, of which n=927 provide usable estimates, and an expert judgement process) and identify preliminary published research on the relationship between CO_2 removal and CO_2 utilisation to offer estimates of scale and cost of CO_2 utilisation.

Where possible, breakeven costs in 2015 \$/t CO₂ for each pathway are calculated. The breakeven CO₂ cost represents the incentive per tonne of CO₂ utilised that would be necessary to make the pathway economic (see Supplementary Materials, S1.2). This can be thought of as the breakeven (theoretical) subsidy per tonne of CO₂ utilisation, although we are not recommending such a subsidy.

Conventional utilisation pathways

Depending on a multitude of technological, policy and economic factors that remain unresolved, utilisation potential in the conventional pathways – chemicals, fuels, microalgae, building materials and CO_2 -EOR – might each achieve around 0.5 Gt CO_2 yr⁻¹ or more in 2050 respectively. Of that, 0.2 to 3.2 Gt CO_2 yr⁻¹ could be simultaneously removed and stored in the lithosphere or the biosphere for centuries or more.

[Chemicals] CO_2 can be transformed efficiently into a range of chemicals, but only a few of these technologies are economically viable and scalable. Some are commercialised²⁹, such as urea³⁰ and polycarbonate polyols³¹. Some are technically possible but not widely adopted, such as CO_2 -derived methanol in the absence of carbon monoxide^{c 32}. Breakeven costs per tonne of CO_2 calculated from the scoping review for urea (circa -\$100) and polyols (circa -\$2,590) reflect currently profitable markets. The estimated utilisation potential for CO_2 in chemicals is ~0.3 to 0.6 Gt CO_2 yr⁻¹ in 2050, with a scoping review interquartile range of breakeven costs of -\$80 to \$320/t CO_2 .

The largest scale chemical utilisation pathway is for urea production. Urea yield boosting is a large existing market (current global production 200 Mt yr^{-1 33}) for CO₂. Urea is produced from ammonia (produced by the energy-intensive Haber-Bosch process $3H_2 + N_2 \rightarrow 2NH_3$) and CO₂ according to $2NH_3 + CO_2 \rightleftharpoons CO(NH_2)_2 + H_2O$, usually with coal or natural gas providing the necessary energy. Within a matter of days of being applied as fertilizer, the carbon in urea is released to the atmosphere. For urea to be net zero carbon, it would require its carbon to be sourced from the atmosphere, e.g., using direct air capture, and a renewable energy source. All nitrogen-based fertilisers produce N₂O, a greenhouse gas around 300 times more potent than CO₂ over a 100-year time horizon³⁴. Increasing urea inputs may thus have a negative impact on climate³⁵.

For polymers, the utilisation potential is estimated to be ~ 10 to 50 Mt CO₂ yr⁻¹ in 2050. In the current market structure, around 60% of plastics have applications in sectors other than packaging, including durable materials for construction, household goods, electronics, and in vehicles. Such products have lifespans of decades or even centuries³⁶.

[Fuels and microalgae] CO₂-derived fuels are argued to be an attractive option in the decarbonisation process^{37,38} as they can be employed within existing transport infrastructure. CO₂-derived fuels might play a role in harder-to-decarbonise sectors, such as aviation³⁹, given hydrocarbons have energy densities orders of magnitude above that of present-day batteries³². The long-term use of carbon-based energy carriers in a net zero emissions economy relies upon their production with renewable

 $[^]c$ Methanol is a platform chemical for a multitude of other reaction pathways, including to fuels. The dominant present day manufacturing method for methanol is the hydrogenation of a mixture of CO and around 1-2% CO₂, added to boost methanol yield.

energy and low-cost scalable clean hydrogen production e.g. by electrolysis of water or novel alternatives.

Here we consider products such as methanol; methane; dimethyl ether (DME); and Fischer-Tropsch fuels as potential CO₂ energy carriers for transportation. The wide range of estimated potential for the scale of CO₂ utilisation in fuels, from ~1 to 4.2 Gt CO₂ yr⁻¹, reflects uncertainties in potential market penetration. The high end represents a future where synfuels have sizeable market shares, due to cost reductions and policy drivers. The low end, which is itself considerable, represents very modest market penetration in methane and fuels markets, but it could also be an overestimate if CO₂-derived products do not become cost competitive with alternative clean energy vectors, or with direct sequestration.

A CO₂-to-methanol plant operates in Iceland, and various power-to-gas plants operate worldwide. However, these may represent special cases that are difficult to replicate because they are exploiting geographic advantages, such as cheap geothermal energy. Whilst producing more complex hydrocarbons is energetically and therefore economically expensive¹¹, rapid cost-reductions could potentially occur, particularly with policy support, given that the cost of renewable energy is a large proportion of the total. The US Department of Energy's target for hydrogen production costs, \$2/kg-H₂, is roughly equivalent to \$2 per gasoline-gallon equivalent, and would require carbon free electricity to cost <US\$0.03/kWh⁴⁰ (accounting for kinetics and other losses to the enthalpy of electrolysis-based hydrogen production ~40 kWh/kg-H₂). Several wind and solar power auctions around the world have been won in recent years with prices below US\$0.03/kWh⁴¹.

The interquartile range for breakeven costs for CO₂ fuels from our scoping review was \$0/t CO₂ to \$670/t CO₂. Negative breakeven costs appear in studies that model particularly beneficial scenarios, such as low discount rates, free feedstocks, or low-cost or free renewable electricity.

For pathways with high capital costs, the benefits of economies of scale and learning could be considerable⁴². This is particularly relevant for algal pathways requiring photobioreactors⁴³ and fuel synthesis pathways requiring electrolysers⁴⁴. Microalgae are a subject of long-standing research interest given their potential to fix CO₂ efficiently (up to 10%, vs 1 to 4% for other biomass⁴⁵), as well as produce a range of products such as biofuels, high-value carbohydrate and proteins, and plastics⁴³. The estimated 2050 potential for microalgae is again wide, given complex production economics, from 0.2 to 0.9 Gt CO₂ yr⁻¹ with a breakeven cost interquartile range from the scoping review of \$230 to \$920/t CO₂.

[Concrete building materials] We estimate that CO_2 utilisation pathways in concrete building materials may remove, utilise and store ~ 0.1 to 1.4 Gt CO_2 yr $^{-1}$ over the long term (with the CO_2 sequestered well beyond the lifespan of the infrastructure itself) at interquartile breakeven costs of ~ 30 to ~ 70 /t CO_2 . This estimate mostly comprises the use of CO_2 as a curing agent in cement, and the high end might reflect a scenario (amongst other possibilities) whereby the technology is fully adopted by the precast concrete market and has a 70% penetration into pourable cement markets. We also consider aggregates produced from carbonated industrial wastes such as cement and demolition waste, steel slag, cement kiln dust, and coal pulverised fuel ash,.

Cement requires the use of lime (CaO), which is produced by the emissions-intensive calcination of limestone, so unless calcination is paired with carbon capture and sequestration, it is difficult for building-related pathways to deliver CO₂ emissions reductions on a life-cycle basis. Several commercial initiatives seek to replace the lime-based ordinary Portland cement (which currently dominates the global market) with alternative binders (such as steel-slag based systems⁴⁶) or geopolymers made from aluminosilicates⁴⁷ in the end product completely, but regulatory barriers curently prevent these approaches from scaling.

[CO₂-EOR] Enhanced oil recovery using CO₂ currently provides ~5% of total US crude oil production⁴⁸. Conventionally, operators aim to maximise both the oil and the CO₂ recovered (rather than CO₂ stored) per tonne of CO₂ injected, with between 1.1 and 3.3 barrels (bbl)) of oil produced per tonne of CO₂ injected under conventional operation and within the constraints of natural reservoir heterogeneity⁴⁹. However, CO₂-EOR can in principle (and depending on operating conditions and project type) be operated such that, on a life cycle basis, more CO₂ is injected than produced upon consumption of the final oil product⁵⁰.

More than 90% of the world's oil reservoirs are potentially suitable for CO₂-EOR⁵¹, implying that as much as 140 Gt CO₂ could be utilised and stored in this way⁵. We estimate a 2050 utilisation rate of ~0.1 to ~1.8 Gt CO₂ yr⁻¹. If EOR was deployed to maximise CO₂ storage, rather than oil output, then CO₂ reduced could approximately equal CO₂ utilised, depending on the emissions-intensity of the counterfactual and relevant inefficiencies (Box 1).

At oil prices of approximately \$100/bbl, EOR is economically viable if CO₂ can be sourced for between \$45 and \$60/t CO₂^{51,52}, implying a breakeven cost of CO₂ of -\$60 to -\$45/t CO₂. These cost estimates are specific to the United States, where the business model is mature, but may not hold for the rest of the world.

Non-conventional utilisation pathways

The five non-conventional utilisation pathways reviewed comprise BECCS, enhanced weathering, forestry techniques, land management practices and biochar. Prior reviews^{18,28,53-55} demonstrate that these pathways offer substantial removal potential; based on a recent substantive scoping review²⁸, 0.5 to 3.6 Gt CO₂ yr⁻¹ for afforestation/reforestation, 2.3 to 5.3 Gt CO₂ yr⁻¹ for land management, and 0.3 to 2 Gt CO₂ yr⁻¹ for biochar and 0.5 to 5 Gt CO₂ yr⁻¹ for BECCS. Enhanced weathering offers a removal potential of 2 to 4 Gt CO₂ yr⁻¹ at costs of ~\$200/t CO₂²⁸. Not all of this potential involves utilisation of carbon dioxide resulting in economic value, but the approximate scale of CO₂u described below is potentially considerable. The breakeven costs per tonne of CO₂u estimated here are low and frequently negative.

[BECCS] BECCS involves the biological capture of atmospheric carbon by photosynthetic processes, producing biomass used for the generation of electricity or fuel, before CO₂ is captured and removed. Although there is substantial uncertainty regarding the total quantity of available biomass⁵⁶ (particularly in the light of concerns over land use competition with food crops) 100–300 EJ yr⁻¹ of primary energy equivalent of biomass could be deployed by 2050.

BECCS provides two distinct services, bioenergy and atmospheric CO_2 removal. Whilst a number of cost estimates exist in the literature, e.g. ~\$200/t CO_2^{28} , these typically assign all costs to the CO_2 removal service, and thus implicitly assume that no revenue is received for the bioenergy services generated. Approximating those revenues using a basket of wholesale electricity prices across countries suited to host BECCS systems⁵⁷, we estimate breakeven costs of between ~\$60 and ~\$160/t CO_2 utilised.

[EW] For terrestrial enhanced weathering (EW) employed on croplands, a yield-enhancing impact is anticipated²⁸. This yield enhancement effect is unlikely to originate directly from soil carbon improvements, but to nutrient uptake facilitated by pH effects⁵⁸; however under our broad definition there may still be an as-yet-unquantified CO₂u associated with the increase in net primary productivity (NPP).

[Forestry techniques] In sustainable forestry, atmospheric CO₂ is removed via photosynthesis into carbon in standing forests, and a portion of that carbon enters production processes and (after minor energetic losses) becomes wood products. Both wood products and standing forests provide economic value and can be seen as CO₂ utilisation (standing forests provide ecosystem services, not

quantified here). CO₂u in wood products will occur additionally to CO₂r directly by forests under certain highly specific circumstances; sustainable harvesting can maintain carbon stocks in forests while providing a source of renewable biomass^{59,60}.

Our estimate is that of the volumes of CO_2 sequestered via afforestation/reforestation in 2050, 0.07 to 0.5 Gt CO_2 u yr⁻¹ may flow into industrial roundwood products, at approximate breakeven costs of between ~-\$40 and ~\$10/t CO_2 utilised. An optimistic scenario might also consider the volumes of wood products that are sustainably harvested from existing forests and plantations. Yearly inflows of carbon utilised as wood products is estimated to be ~1.8 Gt CO_2 in 2050. An additional 0.6 Gt CO_2 utilised may arise from the portion of those flows that are industrial roundwood products sustainably harvested for use in the construction industry (Supplementary Materials), leading to a top-end estimate of 1.1 Gt CO_2 u yr⁻¹ from AF/RF and sustainable forestry techniques.

Wood products have the potential to be long-term stores of carbon, particularly when used in long-lived buildings, wherein lifespans can be conservatively estimated at 80-100 years⁶⁰. We estimate that around half of the carbon in the wood product pool may continue to be stored beyond their usable product life (the non-decomposed fraction (\sim 77%) of the portion of wood presently committed to landfill (\sim 60%)⁶¹). The remainder will return to atmosphere as a fraction (\sim 0.5 Gt CO₂ yr⁻¹) of the 5 Gt CO₂ yr⁻¹ land use change flux portrayed in Figure 1.

[SCS & biochar] CO₂ in land management and biochar pathways is properly considered to be 'utilised' if it enhances economically valuable agricultural output. The CO₂ taken up by land is ultimately either CO₂u (with increased output) or CO₂r (stored in soils), but not both. We estimate that ~0.9 to 1.9 Gt CO₂ yr⁻¹ may be utilised by SCS techniques on croplands and grazing lands by 2050, at approximate breakeven costs of between -\$90 and -\$20/t CO₂ utilised, due to yield increases associated with increases in soil organic carbon stock. We tentatively estimate that ~0.2 to 1 Gt CO₂ yr⁻¹ may be utilised via yield increases following the application of biochar on managed lands at approximate breakeven costs of between -\$70 and -\$60/t CO₂ utilised. These estimates are based on currently reported yield increases (of 0.9% to 2% associated with SCS techniques^{62,63} and 10% associated with biochar⁶⁴) from sparse literature, using crop production as a proxy for net primary productivity. Yield impacts are likely to be highly variable, e.g. according to climatic zone⁶⁵. Crop productivity increases are important not only for economic returns for operators but also for land-use requirements. For instance if tropical biomass yield increases following biochar application reached 25%, the associated reduction of land requirements would equate to 185m ha, and would result in a cumulative net emission benefit from those increased yields of 49 Gt C to 2100⁶⁶.

Table 2 presents breakeven cost ranges and estimated volumes of CO₂ utilised or removed per annum in 2050.

Table 2 | Range estimates of the potential for CO₂ utilisation and present-day breakeven cost

	Professional Chalance and Chalance and Chalance of the Control of					
Pathway	Global gross removal	Global gross utilisation	Breakeven cost of CO ₂			
	potential in 2050	potential in 2050	utilisation			
	(Mt CO ₂ r yr ⁻¹)	(Mt CO ₂ u yr ⁻¹)	(2015 \$/tCO ₂ u)			
CONVENTIONAL						
UTILISATION						
CHEMICALS	~10 to 30	300 to 600	-\$80 to \$320			
FUELS	0	1,000 to 4,210	\$0 to \$670			
MICROALGAE	0	200 to 900	\$230 to \$920			
CONCRETE BUILDING	100 to 1,400	100 to 1,400	-\$30 to \$70			
MATERIALS						
ENHANCED OIL	100 to 1,800	100 to 1,800	-\$60 to -\$45			
RECOVERY						
NON-CONVENTIONAL						
UTILISATION						
BECCS	500 to 5,000	500 to 5,000	\$60 to \$160			
ENHANCED	2,000 to 4,000	N/C	<\$200*			
WEATHERING						
FORESTRY	500 to 3,600	70 to 1,100	-\$40 to \$10			
TECHNIQUES						
LAND MANAGEMENT	2,300 to 5,300	900 to 1,900	-\$90 to -\$20			
BIOCHAR	300 to 2,000	170 to 1,000	-\$70 to -\$60			

The breakeven cost is the cost in 2015 \$/tCO2 adjusted for revenues, by-products, and any CO2 credits or fees. A breakeven cost of zero represents the point at which the pathway is economically viable without governmental CO2 pricing (e.g. a subsidy for CO2 utilisation). Breakeven costs presented as a range represent either (for conventional pathways with the exception of EOR) 25th and 75th percentile estimates as calculated *via* the scoping review of the academic literature (where the magnitude of difference reflects the diversity of technological and economic assumptions available within and across each sub-pathway) or (for land-use based pathways) top-down estimates of revenues that may accrue (where the uncertainty of the accuracy of the estimation is high). Breakeven costs presented with a * are calculated unadjusted for revenues and by-product credits. To get the utilisation potential high and low values for conventional pathways, we averaged the interpolated expert opinions with a co-author estimate. For non-conventional utilisation pathways, estimated utilisation potential ranges are based on estimates of additional realised yield of carbon in vegetation (for soil carbon sequestration and biochar, additional yield approximates to net primary productivity (NPP), and for afforestation/reforestation, it approximates to wood products). These are first rough estimates based on preliminary but sparse published research reporting relationships between C storage and additional C that can be utilised.

Techno-economic barriers to scaling

Numerous challenges exist for scaling CO₂ utilisation. Here we consider cost, technology and energy related issues. Although market penetration can be facilitated by cost competitiveness there is no certainty that the cheapest CO₂ utilisation pathways will scale up. Geographical, financing, political and societal considerations are briefly addressed in the Supplementary Materials, but further investigation of these issues is warranted particularly in light of the UN Sustainable Development Goals (SDGs).

Cost and performance differentials

The breakeven cost per tonne of CO_2 is one way to assess the economics of utilisation. The impact of CO_2 utilisation on the price and value-add proposition of the end product is also important, particularly for CO_2 utilisation processes where the final price differential is immaterial, while small differences in key properties may be important. For instance, prices for a fuel product made using CO_2 currently considerably exceed market prices (Table 3).

Pathway	Cost of product made with CO ₂ utilisation	Selling price of product (\$/t product)		Anticipated cost relative to incumbent in 2050	Anticipated direction of cost relative to incumbent in 2050
	(\$/t product) Median, scoping	Present day	% difference	Summary, expert opinion survey & author judgement	Summary, expert opinion survey & author judgement
Polymers	review 1440	2040	-30%	Likely cheaper	
Methanol	510	400	+30%	Insufficient consensus	
Methane	1740	360	+380%	Likely more expensive	*
FT fuels	4160	1200	+250%	Likely more expensive	*
DME	2740	660	+320%	Insufficient consensus	-
Microalgae	2680	1000	+170%	Likely more expensive	Insufficient consensus
Aggregates	21	18	+20%	Insufficient consensus	_
Cement Curing	56	71	-20%	Likely cheaper	-
CO2-EOR	N/A	N/A	N/A	Likely more expensive	7

Table 3 | Costs of utilisation compared with product costs, scoping review

Median cost estimates for products made with CO₂ utilisation are derived from the backward-looking scoping review. References for the selling prices are set out in more detail in the Supplementary Materials, Table S4. The costs and cost trends anticipated in 2050 are derived from a forward-looking expert opinion survey and coauthor judgement.

Many of the other pathways – in particular products in construction and plastics – have economics that are driven not just by price alone but by the performance characteristics of the end product. There may be trade-offs between product quality and mitigation value, or synergies between the two.

Our cost estimates for conventional pathways, because they are based on a backward-looking scoping review, do not capture current unpublished innovations and advances in the industrial arena. Our expert opinion survey, which included sources from both academia and industry reflected great uncertainty about future costs. Participants from industry were particularly likely to express confidence that costs in pathways that are already economic (for instance in CO₂ cement curing and polyols) would continue to decrease relative to incumbent product costs.

Energy requirements

Some CO₂ utilisation pathways involve chemical transformations with substantial energy inputs (Figure S2). Some require energy to increase CO₂ concentrations from 0.04% towards 100% ⁶⁷. Lifecycle emissions and costs depend upon the source of the energy used. Land-based natural processes use solar energy, harnessed by photosynthesis, to transform CO₂ and water into carbohydrates. While photosynthesis is an inefficient process (average efficiency is around 0.2% globally ⁶⁸), biological pathways are not necessarily more expensive. In industrial processes, hydrogen often serves as feedstock. 'Brown' hydrogen is currently primarily and most cheaply generated by reforming methane, ⁶⁹ with associated CO₂ emissions. These emissions could be captured and stored, producing 'blue' hydrogen. Production of 'green' hydrogen by electrolysis of water also has potential, and the ultimate technology choice for hydrogen generation will depend on the rates of cost reduction, ⁷⁰ among other factors.

The outlook for CO2 utilisation

Our high-end and low-end scale and cost estimates in Table 2 are drawn as cost curves in (A) low and (B) high scenarios in Figure 3. These curves are constructed using currently available (and often sparse) data in the peer-reviewed literature, or (where data is not available) approximations, and should be considered as a speculative first pass at envisioning future scenarios. The curves should not be interpreted as comprehensive assessments of costs, do not represent nth of a kind costs, and are incompatible with other sequestration or abatement cost curves. The limitations of cost curves particularly with regards to exogenous costs such as establishment costs have been previously described⁷¹ and remain relevant here. An important caveat is that individual potentials cannot be arbitrarily summed: some access the same demand, for instance for transport, which may or may not be filled by a process that utilises CO₂. For instance, the putative success of CO₂-fuels may reduce the demand for oil, thus also reducing the potential of CO₂-EOR. Furthermore, land availability means that choosing one land-based pathway (e.g. BECCS) might preclude the application of another at scale (e.g. biochar).

Notwithstanding the many caveats, the potential scale of utilisation could be considerable. Much of this potential CO₂ utilisation – notably in 'closed' and 'open' pathways - may be economically viable without dramatic shifts in prices. The specific assumptions of the low scenario, which do not account for potential overlaps in utilisation volumes between pathways, imply an upper bound of over 1.5 Gt CO₂ yr⁻¹ at well under \$100/t CO₂u. For policymakers interested in climate change, these figures demonstrate the theoretical potential for correctly designed policies to incentivise the displacement of fossil fuels or the removal of CO₂ from the atmosphere.

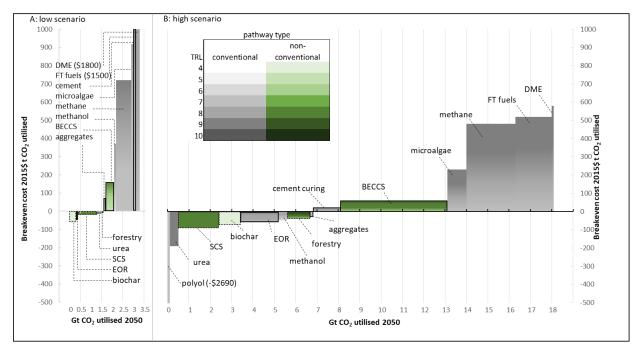


Figure 3 | Estimated CO₂ utilisation potential and breakeven cost of different sub-pathways in a (A) low and (B) high scenario. The breakeven cost is the incentive, measured in 2015\$/t CO₂, required to make the pathway economic. Negative breakeven costs indicate that the pathway is already profitable, without any incentive to utilise CO₂ (such as a tax on CO₂ emissions in the case where utilisation avoids emissions, or a subsidy for CO₂ removed from atmosphere in the case where utilisation removes CO₂). Utilisation estimates are based on 2050 projections. Many technologies are at very early stages of research and cost optimization via research and development could dramatically change these estimates. Colour shadings reflect the level of technology readiness levels (TRL) of the pathways, which again vary dramatically within each pathway. Solid borders indicate that the pathway offers CO₂ storage durations of centuries or more; dotted borders that the pathway offers storage durations of decades; and no borders that the pathway offers storage durations of days or months. See Supplementary Materials for further details.

Figure 3 also highlights some of the economic and technological challenges faced by the pathways. The cycling pathways often (other than urea and polyols) must compete with lower-cost incumbents. The four closed pathways, except for CO₂-EOR, are at low technology readiness levels (TRLs). Open pathways, while both theoretically profitable and implementable, often incur additional operating costs such as implementation, transaction, institutional, and monitoring costs, which can be high⁷².

Each of the potentially large scale, low-cost pathways also face challenges as mitigation strategies. CO₂-EOR utilises and (with correct policy) stores CO₂ at scale, but may not yield any net climate benefit and may even be detrimental. BECCS has a range of well-articulated problems, including major land-use change emissions⁷³. Land management, biochar and forestry all suffer risks of permanence and large-scale flows of CO₂ back to the atmosphere²³. The chemicals pathways may reduce net emissions by displacing fossil fuel use but will not contribute to net removals, unless paired with DAC in a net zero world. Building materials face a challenging route to market penetration due to regulatory barriers which may take decades to surmount. In general, low TRLs will also challenge pathways' ability to scale rapidly enough and within the desired timeframe for mitigation⁵. The uncertainty in future outcomes is relatively large, and very few industries globally involve over 1 Gt yr⁻¹ of material flows.

The net climate impact of the CO₂ utilisation pathways will in many cases depend upon the emissions intensity of prevailing processes⁷⁴. For instance, CO₂-EOR might currently contribute to an overall reduction in atmospheric CO₂, compared to business-as-usual⁴⁹. As decarbonisation proceeds, the climate benefit of CO₂-EOR is reduced. At some point before full decarbonisation, EOR without DAC will result in a net increase in CO₂ emissions⁷⁵. Conversely, in an economy with high supply chain emissions, the climate benefit from BECCS is low⁷³. In a decarbonised world, those supply-chain emissions will be close to zero so the climate benefit from BECCS will be amplified.

Each of the utilisation pathways described here should be seen as a part of the cascade of mitigation options available. For instance, using recycled organic matter to reduce fertiliser use and emissions is a priority, followed by more efficient user of fertiliser⁷⁶, followed by increasing urea yields to reduce total emissions (via more efficient use of NH₃)³⁰. Eventually, fertilizers derived from fossil fuel free ammonia⁷⁷ should be used to supplement fertilizers derived from organic materials. Similarly, a robust finding in the integrated assessment modelling literature is that the electricity sector should be decarbonized first, which then facilitates decarbonisation in other more difficult sectors⁷⁸. In terms of climate impact per kWh of electricity use, available renewable electricity is more efficiently directed towards e-mobility and heat pumps rather than hydrogen-based CCU technologies in the chemical industry⁷⁴.

Future priorities for CO₂ utilisation

An important caveat for policymakers and practitioners is that scaling up CO_2 utilisation will not necessarily be beneficial for climate stability; policy should not aim to support utilisation per se, but instead seek to incentivise genuine emission reductions and removals on a life cycle basis, and thus provide incentives for the deployment of CO_2 utilisation that is climate beneficial. The depth and quality of much of the literature underlying the scoping reviews that support this article is in many senses insufficient in this respect.

Nonetheless given the slow nature of the innovation process, and the urgency of the climate problem, prioritisation should be given to the most promising and least developed options so that early and effective adoption of a portfolio of techniques can be achieved. For the pathways with apparently negative cost (i.e. those that should be profitable in the absence of a theoretical CO_2 subsidy), the challenge – particularly for the open pathways – is to identify and overcome other barriers to adoption.

Conventional utilisation pathways

The emissions-reduction potentials of the three cycling pathways would be facilitated by declines in the costs of CO_2 capture. New sorbents could reduce the cost of energy-intensive separation of CO_2 from flue gases and industrial streams^{40,79}. In the longer term, cheaper direct air capture (based on clean energy) would support the scale up of these pathways⁸⁰. The cost of DACCS has recently been assessed as being between \$600 and \$1000/t CO_2 for the first-of-a-kind plant, with nth-of-a-kind costs potentially on the order of \$200/t CO_2^{80} .

Research on materials and catalysts for CO₂ reduction could enable the efficient and lower-cost transformation of CO₂ into a broader range of products⁷⁹. This includes developing catalysts for efficient production of syn gas via dry reforming of methane with CO₂, efficient photo/electro-catalysts to release hydrogen from water, along with photo/electro-catalysts that can reduce CO₂, or new high-temperature, reversibly reducible metal oxides⁷⁹ to produce syn gas using concentrated sunlight. New membrane materials that can separate miscible liquids, e.g., methanol and water, have a role to play⁸¹. Catalytic processes can be optimised to increase CO₂ emission reductions or reduce energy consumption⁸². One important research challenge is to produce materials showing the highest material property profiles, in particular temperature stability and wider operating or processing temperature windows. Rigorous, realistic techno-economic analyses of these scientific advances could determine their contribution to valuable cost reductions.

Given the rapid rate at which human societies are urbanising⁸³, there is an urgent one-time opportunity to deploy new building materials (including wood, discussed below) that utilise and store CO₂ and displace emissions-intensive Portland cement. In this area, as others, progress would be helped by techno-economic analyses and lifecycle analyses with clearer system boundaries, counterfactuals, and accounting for co-products,⁸⁴ and integrated modelling frameworks that can co-assess background system changes⁸⁵.

Non-conventional utilisation pathways

Figure 1 and Figure 3 imply that land-based biological processes offer a large opportunity to utilise, remove and store more CO₂. Progress here is partly dependent upon field-based trials to improve understanding of the system-wide impacts of different pathways on plant yields and impacts on water, food and water systems, and other resources. Such research might fruitfully prioritise multiple-land-use approaches, such as agro-forestry plantations; rice straw as biomass; low-displacement bioenergy strategies like crassulacean acid metabolism (CAM) plants on marginal land; or nipa palm in mangroves. A better understanding of soil carbon dynamics and improved phenotypic and genotypic plant selection will also help⁸⁶.

Biochar is at a low TRL, with associated uncertainties, but if these can be overcome, its position low on the cost curve in both low and high scenarios suggests that this pathway may have considerable potential. The key challenge is to improve variations in yield effects that will likely be a hindrance in the economic decisions on the part of farmers to use biochar applications⁸⁷, and to find ways to secure potential revenue streams.

Increased forestation, where land availability and biodiversity constraints allow, and the greater use of wood products in buildings, are strategies that appear worth pursuing. Whilst our estimates consider the scale up of existing industrial roundwood use via afforestation and reforestation, new wood-based products such as cross-laminated timber and acetylated wood⁸⁸ aimed at new markets may also have potential. Specification, quality and safety measures for these products are approaching comparability for many concrete structures⁸⁹, and current manufacturing scale-up suggests this may be a market with strong growth prospects.

Cross-cutting efforts

Broad policy and regulatory changes that may support the appropriate scale-up of CO₂ utilisation include creating carbon prices of around US\$40-80/tCO₂, rising over time, to penalise CO₂ emissions⁹⁰ and to incentivise verifiable CO₂ emissions reductions and removals from atmosphere. We do not advocate a direct subsidy for utilisation. Instead, subsidies for CO₂ removals and reductions (or penalties for emissions) are justified and these will support CO₂ utilisation where it is beneficial for the climate. For instance, our analysis suggests closed pathways with scalability, such as BECCS and building materials, would be sensitive to a subsidy for CO₂ removals. Changes to standards, mandates, procurement policies and research and development support to close gaps in knowledge across a portfolio of pathways,⁹¹ are also desirable. Financing and managing the emergence of a globally important new set of CO₂ utilisation industries will likely need clear direction and industrial support from government. An enabling 'net zero' legislative regime – such as that in Sweden and the UK and proposed in New Zealand – can provide clarity about the necessary scale of industries that reduce and remove CO₂, including the pathways examined here.

Collaboration between scholars, public officials and business leaders to ensure accurate comparisons between different alternatives, including directly comparing CCU, CDR and CCS pathways, could facilitate the blending of advantageous features of the ten pathways described here, the exploration of pathways not addressed here, and the identification of novel CO₂ utilisation pathways to accelerate emissions reductions and removals.

 CO_2 utilisation is not an end in itself, and these pathways solely or even collectively will not provide a key solution to climate change. Nevertheless, there is a substantial societal value in continued efforts to determine what will and will not work, where the climate will benefit from CO_2 utilisation and where it will not, and how expensive it will be.

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Author contributions

JB conceived of the paper. CH and EA conducted the analysis and drafted the paper, with extensive input from NMD, and critical input on estimates, methodology and drafting from JB, EC, SF, JM, PS and CW.

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References

- 1 The Royal Society & National Academy of Sciences. Dealing with Carbon Dioxide at Scale. (Sackler Forum, London, 2017).
- von der Assen, N. & Bardow, A. Life cycle assessment of polyols for polyurethane production using CO₂ as feedstock: insights from an industrial case study. Green Chemistry 16, 3272-3280, doi:10.1039/C4GC00513A (2014).
- Ampelli, C., Perathoner, S. & Centi, G. CO₂ utilization: an enabling element to move to a resource- and energy-efficient chemical and fuel production. Philosophical Transactions of the Royal Society A: Mathematical, Physical and Engineering Sciences **373**, doi:10.1098/rsta.2014.0177 (2015).
- 4 The Royal Society. The Potential and Limitations of Using Carbon Dioxide. (London, 2016).
- Mac Dowell, N., Fennell, P. S., Shah, N. & Maitland, G. C. The role of CO₂ capture and utilization in mitigating climate change. Nature Climate Change **7**, 243-249, doi:10.1038/nclimate3231 (2017).

This paper assesses the potential for CO₂-derived fuels and chemicals to be a fraction of that possible via CO₂-EOR.

Metz, B., Davidson, O., De Coninck, H., Loos, M. & Meyer, L. IPCC Special Report on Carbon Dioxide Capture and Storage. Prepared by Working Group III of the Intergovernmental Panel on Climate Change. (Cambridge, United Kingdom and New York, NY, USA, 2005).

This IPCC report provides an overview of the technology and expected costs of carbon capture and sequestration, and provides a key definition of CO₂ utilisation.

- Aresta, M., Dibenedetto, A. & Angelini, A. Catalysis for the valorization of exhaust carbon: from CO₂ to chemicals, materials, and fuels Chemical Reviews **114**, 1709-1742, doi:10.1021/cr4002758 (2014).
- Quadrelli, E. A., Centi, G., Duplan, J. L. & Perathoner, S. Carbon dioxide recycling: emerging large-scale technologies with industrial potential. ChemSusChem 4, 1194-1215, doi:doi.org/10.1002/cssc.201100473 (2011).
- 9 Mikkelsen, M., Jorgensen, M. & Krebs, F. C. The teraton challenge. A review of fixation and transformation of carbon dioxide. Energy & Environmental Science 3, 43-81, doi:10.1039/B912904A (2010).
- Markewitz, P. et al. Worldwide innovations in the development of carbon capture technologies and the utilization of CO₂. Energy & Environmental Science **5**, 7281-7305, doi:10.1039/C2EE03403D (2012).
- Bushuyev, O. S. et al. What should we make with CO₂ and how can we make it? Joule, doi:10.1016/j.joule.2017.09.003 (2018).
- Majumdar, A. & Deutch, J. Research opportunities for CO₂ utilization and negative emissions at the gigatonne scale. Joule **2**, 805-809, doi:10.1016/j.joule.2018.04.018 (2018).

This high-level commentary proposes, using industrial methods, harnessing of the natural biological cycle and a systems approach for industrial CO₂ utilisation at scale.

- Bennett, S. J., Schroeder, D. J. & McCoy, S. T. Towards a framework for discussing and assessing CO₂ utilisation in a climate context. Energy Procedia **63**, 7976-7992 (2014).
- Harper, A. B. et al. Land-use emissions play a critical role in land-based mitigation for Paris climate targets. Nature Communications **9**, 2938, doi:10.1038/s41467-018-05340-z (2018).
- 15 Le Quere, C. et al. Global Carbon Budget 2018. Earth System Science Data 10, 2141-2194, doi:10.5194/essd-10-2141-2018 (2018).
- 16 IPCC. Climate Change 2014 Mitigation of Climate Change. Contribution of Working Group III to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change. (New York, 2014).
- 17 Minasny, B. et al. Soil carbon 4 per mille. Geoderma **292**, 59-86, doi:10.1016/j.geoderma.2017.01.002 (2017).
- Smith, P. et al. Biophysical and economic limits to negative CO₂ emissions. Nature Climate Change **6**, 42-50, doi:10.1038/nclimate2870 (2016).

This paper quantifies potential global impacts of various Negative Emissions Technologies in the context of biophysical resource constraints.

- Lauk, C., Haberl, H., Erb, K.-H., Gingrich, S. & Krausmann, F. Global socioeconomic carbon stocks in long-lived products 1900–2008. Environmental Research Letters 7, 034023, doi:10.1088/1748-9326/7/3/034023 (2012).
- Xi, F. et al. Substantial global carbon uptake by cement carbonation. Nature Geoscience 9, 880-883, doi:10.1038/ngeo2840 (2016).
- 21 Maries, A., Tyrer, M. & Provis, J. L. Sequestration of CO₂ emissions from cement manufacture. Proceedings of the 37th Cement and Concrete Science Conference (2017).
- Alcalde, J. et al. Estimating geological CO₂ storage security to deliver on climate mitigation. Nature Communications **9**, 2201, doi:10.1038/s41467-018-04423-1 (2018).
- Baccini, A. et al. Tropical forests are a net carbon source based on aboveground measurements of gain and loss. Science, doi:10.1126/science.aam5962 (2017).
- Allen, C. D. et al. A global overview of drought and heat-induced tree mortality reveals emerging climate change risks for forests. Forest Ecology and Management **259**, 660-684, doi:10.1016/j.foreco.2009.09.001 (2010).
- Scott, V., Haszeldine, R. S., Tett, S. F. B. & Oschlies, A. Fossil fuels in a trillion tonne world. Nature Climate Change 5, 419, doi:10.1038/nclimate2578 (2015).
- Gardiner, S. M. A Perfect Moral Storm: Climate Change, Intergenerational Ethics and the Problem of Moral Corruption. Environmental Values **15**, 397-413, doi:10.3197/096327106778226293 (2006).
- Naims, H. Economics of carbon dioxide capture and utilization—a supply and demand perspective. Environmental Science and Pollution Research 23, 22226-22241, doi:10.1007/s11356-016-6810-2 (2016).

This paper analyses CO₂ supply and demand scenarios to conclude that the business case for CO₂ utilisation is technology-specific and that in the long term capture costs of up to 65 €/t will be needed.

Fuss, S. et al. Negative emissions—Part 2: Costs, potentials and side effects. Environmental Research Letters 13, doi:10.1088/1748-9326/aabf9f (2018).

This paper estimates via a large scoping review that afforestation and reforestation, BECCS, biochar, enhanced weathering, DACCS and soil carbon sequestration all have multi-gigatonne sequestration potentials in 2050, and that costs vary widely.

- Otto, A., Grube, T., Schiebahn, S. & Stolten, D. Closing the loop: captured CO₂ as a feedstock in the chemical industry. Energy Environ. Sci. **8**, 3283-3297, doi:10.1039/C5EE02591E (2015).
- Pérez-Fortes, M., Bocin-Dumitriu, A. & Tzimas, E. CO₂ utilization pathways: Techno-economic assessment and market opportunities. Energy Procedia **63**, 7968-7975, doi:doi.org/10.1016/j.egypro.2014.11.834 (2014).
- Langanke, J. et al. Carbon dioxide (CO₂) as sustainable feedstock for polyurethane production. Green Chemistry **16**, 1865-1870, doi:10.1039/C3GC41788C (2014).
- 32 Shih, C. F., Zhang, T., Li, J. & Bai, C. Powering the future with liquid sunshine. Joule, doi:10.1016/j.joule.2018.08.016 (2018).
- Jarvis, S. M. & Samsatli, S. Technologies and infrastructures underpinning future CO₂ value chains: A comprehensive review and comparative analysis. Renewable and Sustainable Energy Reviews 85, 46-68, doi:10.1016/j.rser.2018.01.007 (2018).
- Myhre, G. et al. Anthropogenic and Natural Radiative Forcing. Climate Change 2013: The Physical Science Basis: contribution of Working Group 1 to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change. 714 (2013).
- Luo, J., Ledgard, S. & Lindsey, S. Nitrous oxide emissions from application of urea on New Zealand pasture. New Zealand Journal of Agricultural Research **50**, 1-11, doi:10.1080/00288230709510277 (2007).
- Geyer, R., Jambeck, J. R. & Law, K. L. Production, use, and fate of all plastics ever made. Science Advances 3, e1700782, doi:10.1126/sciadv.1700782 (2017).
- Jiang, Z., Xiao, T., Kuznetsov, V. L. & Edwards, P. P. Turning carbon dioxide into fuel. Philosophical Transactions. Series A, Mathematical, Physical, and Engineering Sciences 368, 3343-3364, doi:10.1098/rsta.2010.0119 (2010).
- Olah, G. A. Beyond oil and gas: the methanol economy. Angewandte Chemie International Edition **44**, 2636-2639, doi:doi.org/10.1002/anie.200462121 (2005).
- National Academies of Sciences Engineering and Medicine. Commercial Aircraft Propulsion and Energy Systems Research: Reducing Global Carbon Emissions. (National Academies Press, 2016).
- 40 Secretary of Energy Advisory Board. Letter Report: Task Force on RD&D Strategy for CO₂ Utilisation and/or Negative Emissions at the Gigatonne Scale. (US Department of Energy, Washington, DC, 2016).
- De Luna, P. et al. What would it take for renewably powered electrosynthesis to displace petrochemical processes? Science **364**, eaav3506, doi:10.1126/science.aav3506 (2019).

This paper reviews the potential for and costs of using renewable energy for electrochemical synthesis of concentrated CO₂ into formic acid, carbon monoxide, ethylene and ethanol, and compares biocatalytic and FT routes to long-chain chemical production.

- Dimitriou, I. et al. Carbon dioxide utilisation for production of transport fuels: process and economic analysis. Energy & Environmental Science **8**, 1775-1789, doi:10.1039/C4EE04117H (2015).
- 43 IEA Bioenergy. State of Technology Review Algae Bioenergy. (IEA, Paris, 2017).
- Brynolf, S., Taljegard, M., Grahn, M. & Hansson, J. Electrofuels for the transport sector: A review of production costs. Renewable and Sustainable Energy Reviews, 1-19, doi:10.1016/j.rser.2017.05.288 (2017).
- Williams, P. J. l. B. & Laurens, L. M. Microalgae as biodiesel & biomass feedstocks: review & analysis of the biochemistry, energetics & economics. Energy & Environmental Science 3, 554-590 (2010).
- 46 Mahoutian, M. & Shao, Y. Production of cement-free construction blocks from industry wastes. Journal of Cleaner Production 137, 1339-1346, doi:10.1016/j.jclepro.2016.08.012 (2016).
- 47 Provis, J. L. & Bernal, S. A. J. Geopolymers and related alkali-activated materials. Annual Review of Materials Research 44, 299-327, doi:10.1146/annurev-matsci-070813-113515 (2014).
- Dai, Z. et al. CO₂ accounting and risk analysis for CO₂ sequestration at enhanced oil recovery sites. Environmental Science & Technology **50**, 7546-7554, doi:10.1021/acs.est.6b01744 (2016).
- 49 Lipponen, J. Storing CO₂ through Enhanced Oil Recovery: Combining EOR with CO₂ storage (EOR+) for Profit. (International Energy Agency, 2015).
- 50 Stewart, R. J. & Haszeldine, R. S. Can producing oil store carbon? Greenhouse gas footprint of CO₂EOR, offshore North Sea. Environmental Science & Technology **49**, 5788-5795, doi:10.1021/es504600q (2015).
- 51 Godec, M. L. Global Technology Roadmap for CCS in Industry: Sectoral Assessment CO₂ Enhanced Oil Recovery. (United Nations Industrial Development Organisation, 2011).
- 52 IEA. Storing CO₂ through Enhanced Oil Recovery. (International Energy Agency, Paris, 2015).
- Griscom, B. W. et al. Natural climate solutions. Proceedings of the National Academy of Sciences 114, 11645-11650 (2017).
- Smith, P. Soil carbon sequestration and biochar as negative emission technologies. Global Change Biology 22, 1315-1324, doi:10.1111/gcb.13178 (2016).
- Minx, J. C. et al. Negative emissions—Part 1: Research landscape and synthesis. Environmental Research Letters 13, 063001, doi:10.1088/1748-9326/aabf9b (2018).
- Slade, R., Bauen, A. & Gross, R. Global bioenergy resources. Nature Climate Change 4, 99, doi:10.1038/nclimate2097 (2014).
- Vaughan, N. E. et al. Evaluating the use of biomass energy with carbon capture and storage in low emission scenarios. 13, 044014, doi:10.1088/1748-9326/aaaa02 (2018).
- Beerling, D. J. et al. Farming with crops and rocks to address global climate, food and soil security. Nat Plants 4, 138-147, doi:10.1038/s41477-018-0108-y (2018).

- 59 Pingoud, K., Ekholm, T., Sievänen, R., Huuskonen, S. & Hynynen, J. Trade-offs between forest carbon stocks and harvests in a steady state A multi-criteria analysis. Journal of Environmental Management **210**, 96-103, doi:10.1016/j.jenvman.2017.12.076 (2018).
- Lippke, B. et al. Life cycle impacts of forest management and wood utilization on carbon mitigation: knowns and unknowns. Carbon Management **2**, 303-333, doi:10.4155/CMT.11.24 (2011).
- 61 FAOSTAT. FAOSTAT statistical database. (http://ref.data.fao.org, 2018).
- Lal, R. Enhancing crop yields in the developing countries through restoration of the soil organic carbon pool in agricultural lands. Land Degradation & Development 17, 197-209, doi:10.1002/ldr.696 (2006).
- Soussana, J.-F. et al. Matching policy and science: Rationale for the '4 per 1000-soils for food security and climate' initiative. Soil and Tillage Research, doi:10.1016/j.still.2017.12.002 (2017).
- Jeffery, S., Verheijen, F. G., Van Der Velde, M. & Bastos, A. C. A quantitative review of the effects of biochar application to soils on crop productivity using meta-analysis. Agriculture, ecosystems & environment **144**, 175-187, doi:10.1016/j.agee.2011.08.015 (2011).
- Jeffery, S. et al. Biochar boosts tropical but not temperate crop yields. Environmental Research Letters 12, doi:10.1088/1748-9326/aa67bd (2017).
- Werner, C., Schmidt, H. P., Gerten, D., Lucht, W. & Kammann, C. Biogeochemical potential of biomass pyrolysis systems for limiting global warming to 1.5 °C. Environmental Research Letters 13, doi:10.1088/1748-9326/aabb0e (2018).
- Darton, D. & Yang, A. Removing carbon dioxide from the atmosphere assessing the technologies. Chem Eng Transactions **69** (2018).
- 68 Barber, J. Photosynthetic energy conversion: natural and artificial. Chemical Society Reviews 38, 185-196, doi:10.1039/B802262N (2009).
- 69 Izquierdo, U. et al. Hydrogen production from methane and natural gas steam reforming in conventional and microreactor reaction systems. International Journal of Hydrogen Energy 37, 7026-7033, doi:10.1016/j.ijhydene.2011.11.048 (2012).
- Kuckshinrichs, W., Ketelaer, T. & Koj, J. C. Economic analysis of improved alkaline water electrolysis. J Frontiers in Energy Research 5, 1, doi:doi.org/10.3389/fenrg.2017.00001 (2017).
- Kesicki, F. & Strachan, N. Marginal abatement cost (MAC) curves: confronting theory and practice. Environmental Science & Policy 14, 1195-1204, doi:10.1016/j.envsci.2011.08.004 (2011).
- Viana, V. M., Grieg-Gran, M., Della Mea, R. & Ribenboim, G. The costs of REDD: lessons from Amazonas. International Institute for Environment and Development (IIED): London, UK (2009).
- Fajardy, M. & Mac Dowell, N. Can BECCS deliver sustainable and resource efficient negative emissions? Energy & Environmental Science 10, 1389-1426, doi:10.1039/C7EE00465F (2017).
- Kätelhön, A., Meys, R., Deutz, S., Suh, S. & Bardow, A. Climate change mitigation potential of carbon capture and utilization in the chemical industry. Proceedings of the National Academy of Sciences 116, 11187-11194, doi:10.1073/pnas.1821029116 (2019).
- Jaramillo, P., Griffin, W. M. & McCoy, S. T. Life cycle inventory of CO₂ in an enhanced oil recovery system. Environmental Science & Technology **43**, 8027-8032, doi:10.1021/es902006h (2009).
- Gerber, J. S. et al. Spatially explicit estimates of N₂O emissions from croplands suggest climate mitigation opportunities from improved fertilizer management. Global Change Biology 22, 3383-3394, doi:10.1111/gcb.13341 (2016).
- 77 Chen, J. G. et al. Beyond fossil fuel–driven nitrogen transformations. Science **360**, doi:10.1126/science.aar6611 (2018)
- 78 Luderer, G. et al. Residual fossil CO₂ emissions in 1.5–2 °C pathways. Nature Climate Change 8, 626-633, doi:10.1038/s41558-018-0198-6 (2018).
- Senftle, T. P. & Carter, E. A. The holy grail: chemistry enabling an economically viable CO₂ capture, utilization, and storage strategy. Accounts of Chemical Research **50**, 472-475, doi:10.1021/acs.accounts.6b00479 (2017).
- 80 Keith, D. W., Holmes, G., St. Angelo, D. & Heidel, K. A process for capturing CO₂ from the atmosphere. Joule **2**, 1573-1594, doi:10.1016/j.joule.2018.05.006 (2018).
- Mahmood, A., Bano, S., Kim, S.-G. & Lee, K.-H. Water–methanol separation characteristics of annealed SA/PVA complex membranes. Journal of Membrane Science **415**, 360-367, doi:10.1016/j.memsci.2012.05.020 (2012).
- Xiao, T. et al. The Catalyst Selectivity Index (CSI): a framework and metric to assess the impact of catalyst efficiency enhancements upon energy and CO₂ footprints. Topics in Catalysis **58**, 682-695, doi:10.1007/s11244-015-0401-1 (2015).
- 83 Seto, K. C., Güneralp, B. & Hutyra, L. R. Global forecasts of urban expansion to 2030 and direct impacts on biodiversity and carbon pools. Proceedings of the National Academy of Sciences 109, 16083-16088, doi:10.1073/pnas.1211658109 (2012).
- Zimmermann, A. et al. Techno-Economic Assessment & Life-Cycle Assessment Guidelines for CO₂ Utilization. (2018).
- Arvesen, A., Luderer, G., Pehl, M., Bodirsky, B. L. & Hertwich, E. G. Deriving life cycle assessment coefficients for application in integrated assessment modelling. Environmental Modelling & Software **99**, 111-125, doi:10.1016/j.envsoft.2017.09.010 (2018).
- Scharlemann, J. P. W., Tanner, E. V. J., Hiederer, R. & Kapos, V. Global soil carbon: understanding and managing the largest terrestrial carbon pool. Carbon Management 5, 81-91, doi:10.4155/cmt.13.77 (2014).
- Dickinson, D. et al. Cost-benefit analysis of using biochar to improve cereals agriculture. Gcb Bioenergy 7, 850-864 (2015).
- Song, J. et al. Processing bulk natural wood into a high-performance structural material. Nature **554**, 224, doi:10.1038/nature25476 (2018).

- 89 Ramage, M. H. et al. The wood from the trees: The use of timber in construction. Renewable and Sustainable Energy Reviews **68**, 333-359, doi:10.1016/j.rser.2016.09.107 (2017).
- 90 High-Level Commission on Carbon Prices. Report of the High-Level Commission on Carbon Prices. (Washington, DC, 2017).
- 91 Hepburn, C., Pless, J. & Popp, D. Encouraging innovation that protects environmental systems: five policy proposals. Review of Environmental Economics and Policy, doi:10.1093/reep/rex024 (2018).
- 92 Muntean, M. et al. Fossil CO₂ emissions of all world countries 2018 report. (Luxemboug, 2018).
- 93 Sundquist, E. & Visser, K. The geologic history of the carbon cycle. Treatise on Geochemistry **8**, 682, doi:10.1016/B0-08-043751-6/08133-0 (2003).
- 94 Blunden, J., Derek, S., Hartfield, G. State of the Climate in 2017. Special Supplement to the Bulletin of the American Meteorological Society, **99** (2018).
- Cuéllar-Franca, R. M. & Azapagic, A. Carbon capture, storage and utilisation technologies: A critical analysis and comparison of their life cycle environmental impacts. Journal of CO₂ Utilization **9**, 82-102, doi:10.1016/j.jcou.2014.12.001 (2015).

This paper compares environmental impacts of CO₂ utilisation and CCS technologies by reviewing the literature of life cycle assessment studies.

- Sathre, R. & O'Connor, J. Meta-analysis of greenhouse gas displacement factors of wood product substitution. Environmental Science & Policy 13, 104-114, doi:10.1016/j.envsci.2009.12.005 (2010).
- van der Giesen, C., Kleijn, R. & Kramer, G. J. Energy and climate impacts of producing synthetic hydrocarbon fuels from CO₂. Environmental Science & Technology **48**, 7111-7121, doi:10.1021/es500191g (2014).
- 98 Sternberg, A., Jens, C. M. & Bardow, A. Life cycle assessment of CO₂-based C1-chemicals. Green Chemistry **19**, 2244-2259, doi:10.1039/C6GC02852G (2017).
- Abanades, J. C., Rubin, E. S., Mazzotti, M. & Herzog, H. J. On the climate change mitigation potential of CO₂ conversion to fuels. Energy & Environmental Science 10, 2491-2499, doi:10.1039/C7EE02819A (2017).
- Sternberg, A. & Bardow, A. Life cycle assessment of power-to-gas: syngas vs methane. ACS Sustainable Chemistry & Engineering 4, 4156-4165, doi:10.1021/acssuschemeng.6b00644 (2016).