

Direct Air Capture: The Dire Climate Consequences of Capturing Carbon From the Atmosphere

Of all the technologies proposed to combat the climate crisis, perhaps none is as intuitively appealing as direct air capture (DAC). The premise of DAC (sometimes called “open air capture”) is simple: Machinery would capture carbon dioxide (CO₂) from the atmosphere, then compress, transport, and store it in underground reservoirs.

But despite decades of hype, DAC still plays no role in carbon removal. The recently announced proposal to build a facility in Texas — an \$811 million project that would be the largest direct air capture facility in the world — gives a sense of the scale of the problem. The plant owners promise to capture “up to” 500,000 metric tons per year, one hundredth of one percent of U.S. CO₂ emissions.¹

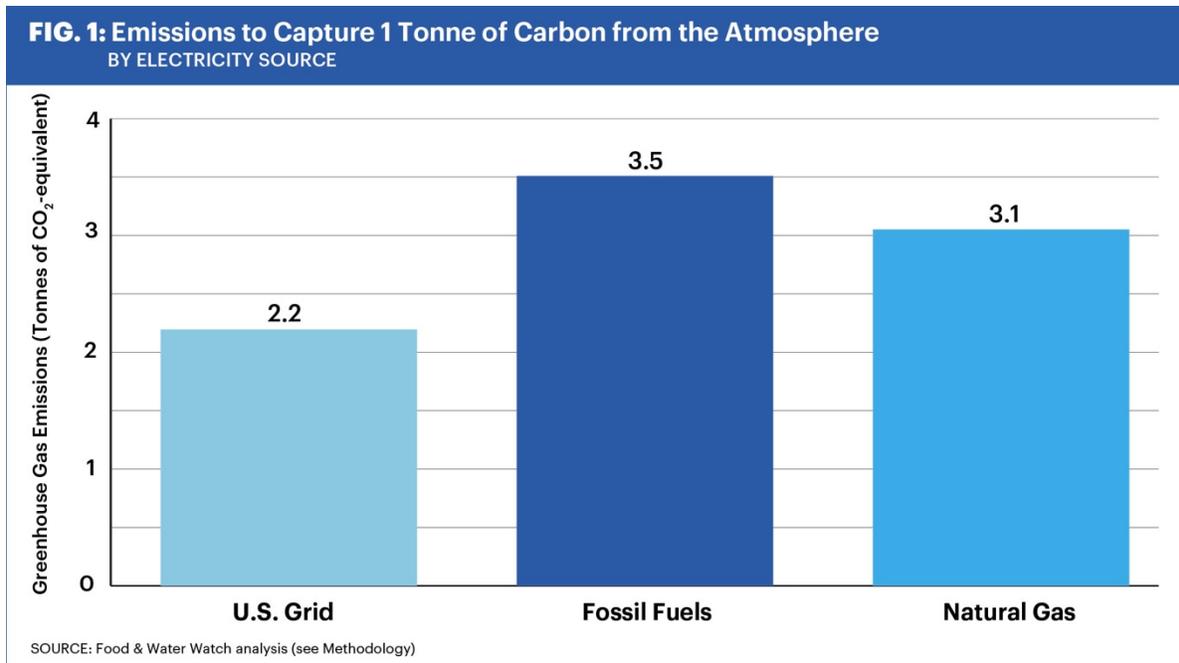
Direct air capture is prohibitively expensive and energy intensive; if it were ever to be successfully deployed, it would likely create more greenhouse gas emissions than it would capture. The promise of capturing CO₂ directly from the atmosphere not only delays the transition away from fossil fuels, it also powers and subsidizes increasing fossil fuel extraction and use. Replacing fossil fuels with renewables — rather than using fossil fuels to pull carbon out of the atmosphere — is a proven and significantly more cost-effective method of addressing climate change.

Direct Air Capture Cannot Be Scaled

Direct air capture is materially and energy intensive because of the low concentration of CO₂ in the air.² Building the technology that could pull carbon from the atmosphere would require considerable physical space — a DAC plant capable of capturing just one megaton of CO₂ per year takes up to 409,000 square feet of space.³ More importantly, direct air capture also consumes large quantities of water and electricity.⁴ Food & Water Watch (FWW) found that if all electricity generated in the United States were used to power DAC, it would capture only one-quarter of the carbon emissions generated by the country each year (see methodology section).

Generating this electricity would create more greenhouse gas emissions than the CO₂ captured at DAC facilities. According to calculations by FWW, capturing 1 ton of CO₂ from the atmosphere

using electricity from fossil fuels would create greenhouse gas emissions equivalent to 3.5 tons of CO₂. Using enough electricity from the U.S. grid to capture 1 ton of CO₂ from the air would create 2.2 tons of CO₂-equivalent emissions.

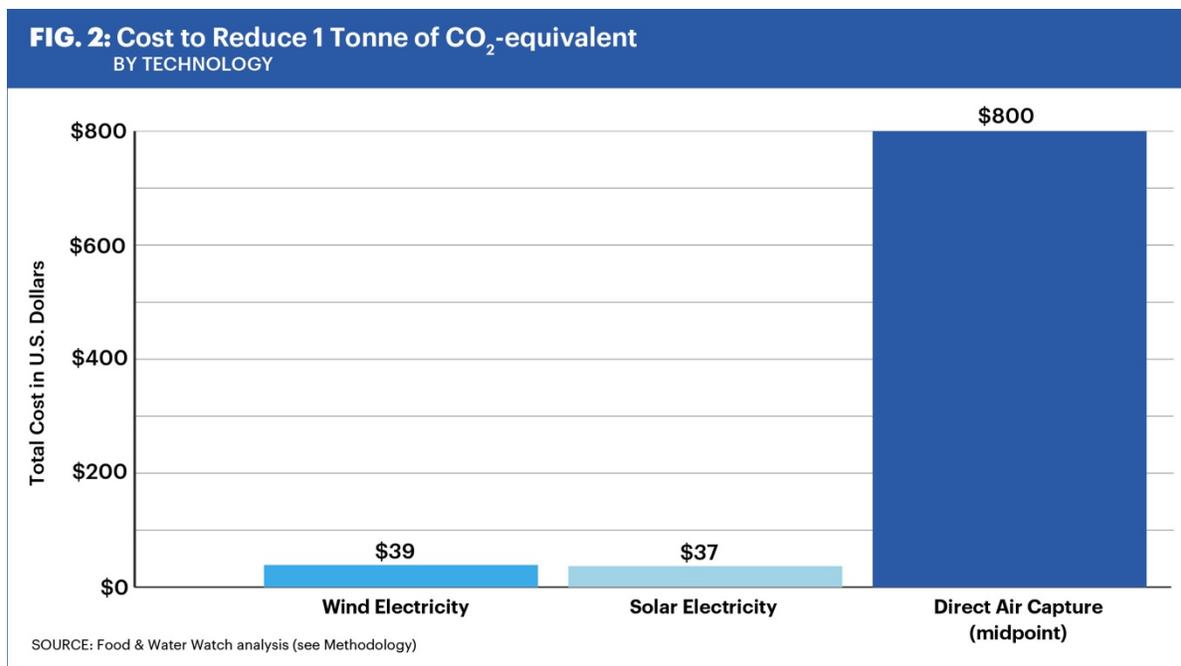


While using electricity from renewable energy could theoretically avoid this problem, that would involve building out a massive amount of clean energy capacity just to support the continuing extraction and combustion of fossil fuels. It is far more advantageous to simply shift our grid from fossil fuels to renewable energy. For instance, FWW found that replacing natural gas with renewables in the electrical grid avoids three times the amount of greenhouse gas emissions that could be captured by a DAC facility powered by renewable electricity. Additionally, the high temperature requirements of some common CO₂ capture systems make the process unable to use electricity (potentially generated from renewable sources) without additional technological progress.⁵

Direct Air Capture Is Absurdly Expensive

There is no way to turn a profit from capturing carbon pollution. To create an incentive for direct air capture, these facilities are eligible for a federal tax credit called 45Q. It is worth up to \$180 per metric ton for captured CO₂ that is sequestered underground, and \$130 per metric ton for CO₂ used for enhanced oil recovery (EOR).⁶ However, several cost estimates put the costs of direct air capture systems at \$600 to \$1,000 per ton of CO₂ captured. This suggests that to make the technology work, DAC will require more subsidies on top of the exorbitant 45Q tax credits.⁷

To illustrate the scale of the costs involved, consider that capturing one-quarter of annual U.S. CO₂ emissions from the atmosphere would cost between \$700 billion and \$1.2 trillion per year⁸ — and would be worth between \$153 billion and \$212 billion per year in 45Q tax credits.⁹



In comparison, FWW found that reducing the same quantity of greenhouse gas emissions with direct investments in wind and solar projects would cost \$45 billion, roughly one-fifteenth the cost of using DAC. Even assuming that 45Q subsidies are sufficient to finance DAC, there are significantly better technologies to subsidize. If 45Q credits were dedicated to making new renewable projects cheaper than operating the cheapest gas plant, the subsidies would be nearly ten times as effective at reducing greenhouse emissions, even ignoring the electricity demands of DAC.

All Carbon Capture Systems Rely on Toxic Solvents

In addition to being an ineffective means of reducing carbon pollution, DAC would create substantial environmental threats. The two most “promising” direct air capture solvent technologies are based on either aqueous hydroxide solutions (NaOH, KOH, etc.) or amine-modified solutions (such as monoethanolamine, or MEA).¹⁰ Industrial production of NaOH is very energy intensive and would massively increase production of chlorine gas, far exceeding utilization capacity.

Capturing one-quarter of U.S. CO₂ emissions would produce 245.5 million metric tons of chlorine gas, 3.2 times the current global use of chlorine gas.¹¹ Since current chlorine demand exceeds production, chlorine gas is used rather than disposed; there is not currently a large-scale disposal

system for chlorine gas.¹² Chlorine gas is highly toxic and corrosive, with both acute and long-term negative human health effects.¹³ Transport of industrial chlorine gas has resulted in fatal accidents, and chlorine gas has been used as a chemical weapon.¹⁴

Producing new MEA is associated with significant greenhouse gas emissions involving a similar supply chain to natural gas-based fertilizer production.¹⁵ When used to capture CO₂ emissions, MEA degrades, releasing chemical carcinogens such as nitrosamines that, according to one study, “are toxic and carcinogenic to humans even at extremely low levels.”¹⁶ At power plants, carbon capture operations using MEA can have air emissions of MEA up to 0.8 kilogram per metric ton of CO₂ captured. These emissions rates can exceed toxicity limits for drinking water and aquatic ecosystems.¹⁷

In addition to air emissions, MEA CO₂ capture systems generate large quantities of MEA-contaminated liquid waste.¹⁸ Historically, this waste was incinerated or landfilled, but neither of these disposal strategies are economically or environmentally sustainable, especially at much larger scales.¹⁹

Long-term Storage of CO₂ Is a Scam

Right now, around 95 percent of all captured carbon in the United States is used for enhanced oil recovery, a process of producing oil by injecting CO₂ mixed with other chemicals in existing wells to flush out additional oil.²⁰ This is far from a climate solution; EOR results in more carbon emissions than it stores. A ton of CO₂ produces two to three barrels of oil when injected; when burned, that oil emits around 1.2 tons of CO₂.²¹

The other option with captured carbon is to store it underground. But CO₂ sequestration also carries significant risks. CO₂ must be injected under sufficient pressure to displace existing fluids. In small spaces, this can create rapid pressure increases that fracture containment layers.²² Earthquakes from injection can also rupture storage seals, allowing CO₂ to leak.²³ Since many storage locations are in and around fossil fuel reservoirs, abandoned oil and gas wellbores provide a pathway for CO₂ leaking to the surface.²⁴ CO₂ can also slowly escape along well linings and has been shown to corrode materials used in well casings and seals.²⁵

Not only does CO₂ injection carry similar risks to wastewater injection (associated with technologies such as fracking), but reducing pressure to inject CO₂ may require extracting wastewater from the reservoir and reinjecting it elsewhere.²⁶ The precise constituents of the wastewater vary depending on the geology of the extraction site,²⁷ but it can contain salts (chlorides, bromides, and sulfides of calcium, magnesium, and sodium), metals (barium, manganese, iron, and strontium), oil, grease, dissolved organics (benzene and toluene), and

radioactive material (radium-226).²⁸ These chemicals can cause cancer, disrupt the endocrine system, affect the nervous, immune, and cardiovascular systems, and affect sensory organs and the respiratory system.²⁹ Underground injection of wastewater can put aquifers and drinking water at risk,³⁰ and extensive research links fluid injection and disposal to earthquakes.³¹

Conclusion

Instead of betting on speculative, dangerous, and ultimately climate-destroying carbon capture technologies, the United States must embrace real climate solutions that already work. Off-the-shelf technologies such as wind, solar, and batteries that generate and store carbon-free electricity are more than capable of addressing the climate crisis, at costs far lower than carbon capture.

Methodology

Food & Water Watch created an average emissions intensity for electricity generation in the United States using data from various sources. This includes 2019 electricity use and fossil fuel consumption data from the U.S. Energy Information Administration, CO₂ emissions data from the U.S. Environmental Protection Agency's Inventory of U.S. Greenhouse Gas Emissions and Sinks, and methane emissions data from peer-reviewed academic sources.³² Electricity requirements for DAC are from a plausible estimate presented in a peer-reviewed academic review.³³ Cost comparisons are based on levelized costs from Lazard.³⁴ Full calculations and treatment of small electricity sources (biomass, waste to energy, etc.) are on file with FWW.

Endnotes

- 1 1PointFive. [Press release]. "Occidental, 1PointFive to begin construction of world's largest direct air capture plant in the Texas Permian Basin." August 25, 2022; U.S. Environmental Protection Agency (EPA). "Inventory of U.S. Greenhouse Gas Emissions and Sinks: 1990-2020." EPA 430-R-22-003. April 2022 at ES-4; Mann, Joshua. "Occidental Petroleum eyes West Texas for largest-of-its-kind carbon capture project." *Texas Biz Journal*. March 23, 2021.
- 2 Qiu, Yang et al. "Environmental trade-offs of direct air capture technologies in climate change mitigation toward 2100." *Nature Communications*. Vol. 13, No. 3635. June 2022 at 2.
- 3 Fuss, Sabine et al. "Negative emissions — Part 2: Costs, potentials and side effects." *Environmental Research Letters*. Vol. 13. May 2018 at 18.
- 4 Qiu et al. (2022) at 3; Sekera, June and Andreas Lichtenberger. "Assessing carbon capture: Public policy, science, and societal need." *Biophysical Economics and Sustainability*. Vol. 5, Art. 14. October 2021 at 14.
- 5 Qiu et al. (2022) at 2.
- 6 Congressional Research Service (CRS). "Tax provisions in the Inflation Reduction Act of 2022 (H.R. 5376)." August 2022 at 8.
- 7 Fuss et al. (2018) at 18; Sabatino, Francesco et al. "Evaluation of a direct air capture process combining wet scrubbing and bipolar membrane electrodialysis." *Industrial & Engineering Chemistry Research*. Vol. 59, Iss. 15. January 2020 at 7007.
- 8 Fuss et al. (2018) at 18; Sabatino et al. (2020) at 7007; EPA (2022) at ES-4.
- 9 CRS (2022) at 8; EPA (2022) at ES-4.
- 10 Chatterjee, Sudipta and Kuo-Wei Huang. "Unrealistic energy and materials requirement for direct air capture in deep mitigation pathways." *Nature Communications*. Vol. 11, No. 3287. July 2020 at 1.
- 11 Food & Water Watch (FWW) analysis of *Ibid.* at 1.

- 12 Zhao, Yun et al. "Low-voltage gaseous HCl electrolysis with iron redox-mediated cathode for chlorine regeneration." *Chemical Engineering Journal*. Vol. 346, Iss. 15. August 2017 at 535; Treger, Yury and Mark Flid. "State of the art and problems of organochlorine synthesis." In Tundo, Pietro et al. (Eds.). (2016). *Chemistry Beyond Chlorine*. Cham: Springer International Publishing Switzerland at 533 to 536.
- 13 EPA. "R.E.D. Facts: Chlorine Gas." EPA-738-F-99-001. February 1999 at 2.
- 14 Hoyle, Gary W. and Erik R. Svendsen. "Persistent effects of chlorine inhalation on respiratory health." *Annals of the New York Academy of Sciences*. Vol. 1378, Iss. 1. August 2016 at 2.
- 15 Luis, Patricia. "Use of monoethanolamine (MEA) for CO2 capture in a global scenario: Consequences and alternatives." *Desalination*. Vol. 380. February 2016 at 93 and 94.
- 16 Zhang, Yuanyuan et al. "Health risk analysis of nitrosamine emissions from CO2 capture with monoethanolamine in coal-fired power plants." *International Journal of Greenhouse Gas Control*. Vol. 20. January 2014 at 37; Karl, Matthias. "Worst case scenario study to assess the environmental impact of amine emissions from a CO2 capture plant." *International Journal of Greenhouse Gas Control*. Vol. 5, Iss. 3. May 2011 at 439.
- 17 Karl (2011) at 439.
- 18 Botheju, Deshai et al. "Monoethanolamine biodegradation processes." *Proceedings of the 2nd Annual Gas Processing Symposium*. 2010 at 77.
- 19 *Ibid*.
- 20 Klinge, Naomi. "US representatives propose legislation that would exclude EOR from 45Q tax credits for CCS." *Upstream*. December 15, 2021.
- 21 Edwards, Ryan W. J. and Michael A. Celia. "Infrastructure to enable deployment of carbon capture, utilization, and storage in the United States." *PNAS*. Vol. 155, No. 38. September 2018 at E8817.
- 22 Holloway, S. "Storage capacity and containment issues for carbon dioxide capture and geological storage on the UK continental shelf." *Proceedings of the Institution of Mechanical Engineers, Part A: Journal of Power and Energy*. Vol. 223. December 2008 at 241.
- 23 Nicol, A. et al. "Induced seismicity and its implications for CO2 storage risk." *Energy Procedia*. Vol. 4. April 2011 at 3700.
- 24 Jahediesfanjani, Hossein et al. "Estimating the pressure-limited CO2 injection and storage capacity of the United States saline formations: Effect of the presence of hydrocarbon reservoirs." *International Journal of Greenhouse Gas Control*. Vol. 79. December 2018 at 14; Ajayi, Temitope et al. "A review of CO2 storage in geological formations emphasizing modeling, monitoring and capacity estimation approaches." *Petroleum Science*. Vol. 16, Iss. 5. October 2019 at 1029 and 1030; Carey, J. William. National Energy Technology Laboratory (NETL). "Probability Distributions for Effective Permeability of Potentially Leaking Wells at CO2 Sequestration Sites." NRAP-TRS-III-021-2017. April 27, 2017 at 19.
- 25 Qafoku, Nikolla P. et al. "Review of the impacts of leaking CO2 gas and brine on groundwater quality." *Earth-Science Reviews*. Vol. 169. June 2017 at 69; Ideker, J. H. et al. NETL. "Experimental and Numerical Modeling Approach to Elucidating Damage Mechanisms in Cement-Well Casing-Host Rock Settings for Underground Storage of CO2." NETL-TRS-4-2018. March 1, 2018 at 1.
- 26 Verdon, James P. and Anna L. Stork. "Carbon capture and storage, geomechanics and induced seismic activity." *Journal of Rock Mechanics and Geotechnical Engineering*. Vol. 8, Iss. 6. December 2016 at 929; Buscheck, Thomas A. et al. "Pre-injection brine production in CO2 storage reservoirs: An approach to augment the development, operation, and performance of CCS while generating water." *International Journal of Greenhouse Gas Control*. Vol. 54, Part A. November 2016 at 499.
- 27 Gallegos, Tanya J. et al. "Hydraulic fracturing water use variability in the United States and potential environmental implications." *Water Resources Research*. Vol. 51. 2015 at 5844.
- 28 U.S. Government Accountability Office. "Information on the Quantity, Quality, and Management of Water Produced During Oil and Gas Production." January 2012 at 12; Vengosh, Avner et al. "A critical review of the risks to water resources from unconventional shale gas development and hydraulic fracturing in the United States." *Environmental Science & Technology*. Vol. 48, Iss. 15. August 2014 at 8341 to 8342.
- 29 Colborn, Theo et al. "Natural gas operations from a public health perspective." *Human and Ecological Risk Assessment*. Vol. 17, Iss. 5. September 2011 at abstract.
- 30 Lustgarten, Abrahm. "Injection wells: The poison beneath us." *ProPublica*. June 21, 2012; Keranen, K. M. et al. "Sharp increase in central Oklahoma seismicity since 2008 induced by massive wastewater injection." *Science*. July 3, 2014 at 1 and 2.
- 31 Goebel, Thomas H. W. and Emily E. Brodsky. "The spatial footprint of injection wells in a global compilation of induced earthquake sequences." *Science*. Vol. 361, Iss. 6405. August 2018 at 899.
- 32 Howarth, Robert W. "Ideas and perspectives: Is shale gas a major driver of recent increase in global atmospheric methane?" *Biogeosciences*. Vol. 16, Iss. 15. August 2019 at 3038 to 3040; Worden, John R. et al. "The 2019 methane budget and uncertainties at 1 degree resolution and each country through bayesian integration of GOSAT total column methane data and a priori inventory estimates." *Atmospheric Chemistry and Physics*. Vol. 22, Iss. 10. May 2022 at 35.
- 33 Sekera and Lichtenberger (2021) at 17.
- 34 Lazard. "Lazard's Levelized Cost of Energy Analysis — Version 15.0." October 2021 at 6.