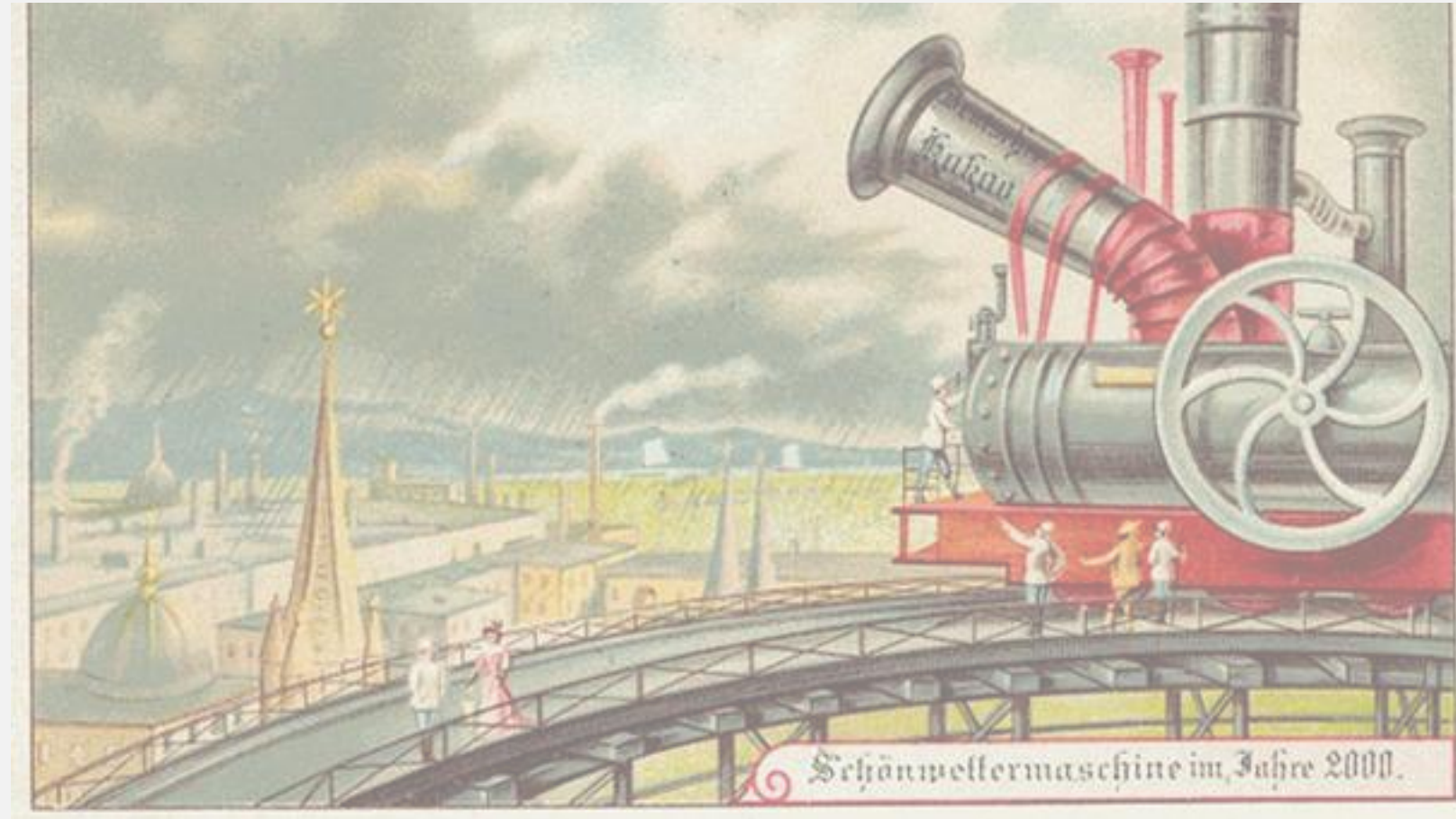


HighTechXL

PFAS & CO2 capture



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From demonstration to Realisation



Expectations



Innovation
Trigger



ureau of
luctivity



Out of the valley, into the light

Some key Updates and Projects 2025:

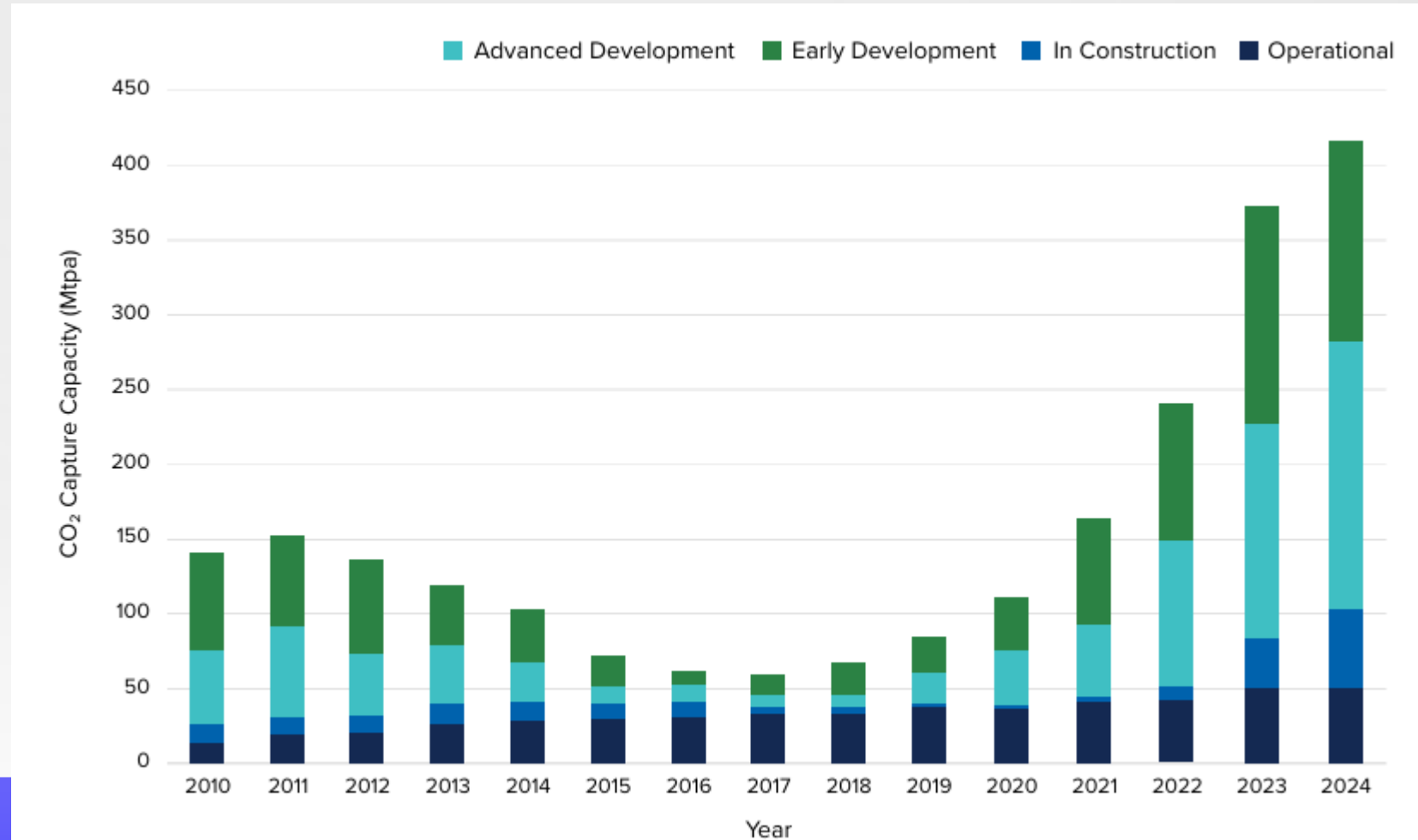
Netherlands: Twence's Modular Carbon Capture Plant (SLB Capturi)

United Kingdom: enfinium's CCS Pilot Programs

Canada: Alberta's Waste-to-Energy CCS Project → \$300 million plant will process 150,000 tonnes mun. waste

European Union: Major Funding for WtE CCS:

- Öresundskraft AB's Innozero project will receive €54 million to build a CCS facility capable of capturing 200,000 tonnes of CO₂ per year by 2026.
- Herambiente's Ferrara WtE plant in Italy will receive €24 million to install enzymatic CCS technology, targeting a 90% capture rate (64,000 tonnes of CO₂ annually), using renewable geothermal heat



Pipeline of CCS projects capture capacity from the Global Status of CCS Report 2024

PFAS & CO2



1. How much and which PFAS is going to the capture plant?
2. What is the effect of this PFAS on the capture plant itself
3. How much PFAS will be in the treated flue gas?
4. How will PFAS influence CO2 quality?

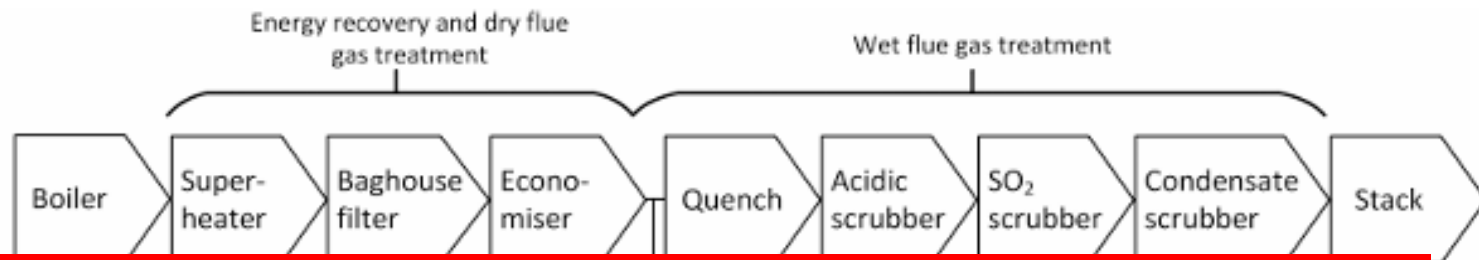
PFAS entering the capture plant

PFAS detected in flue gas, ash, condensate, and other residues

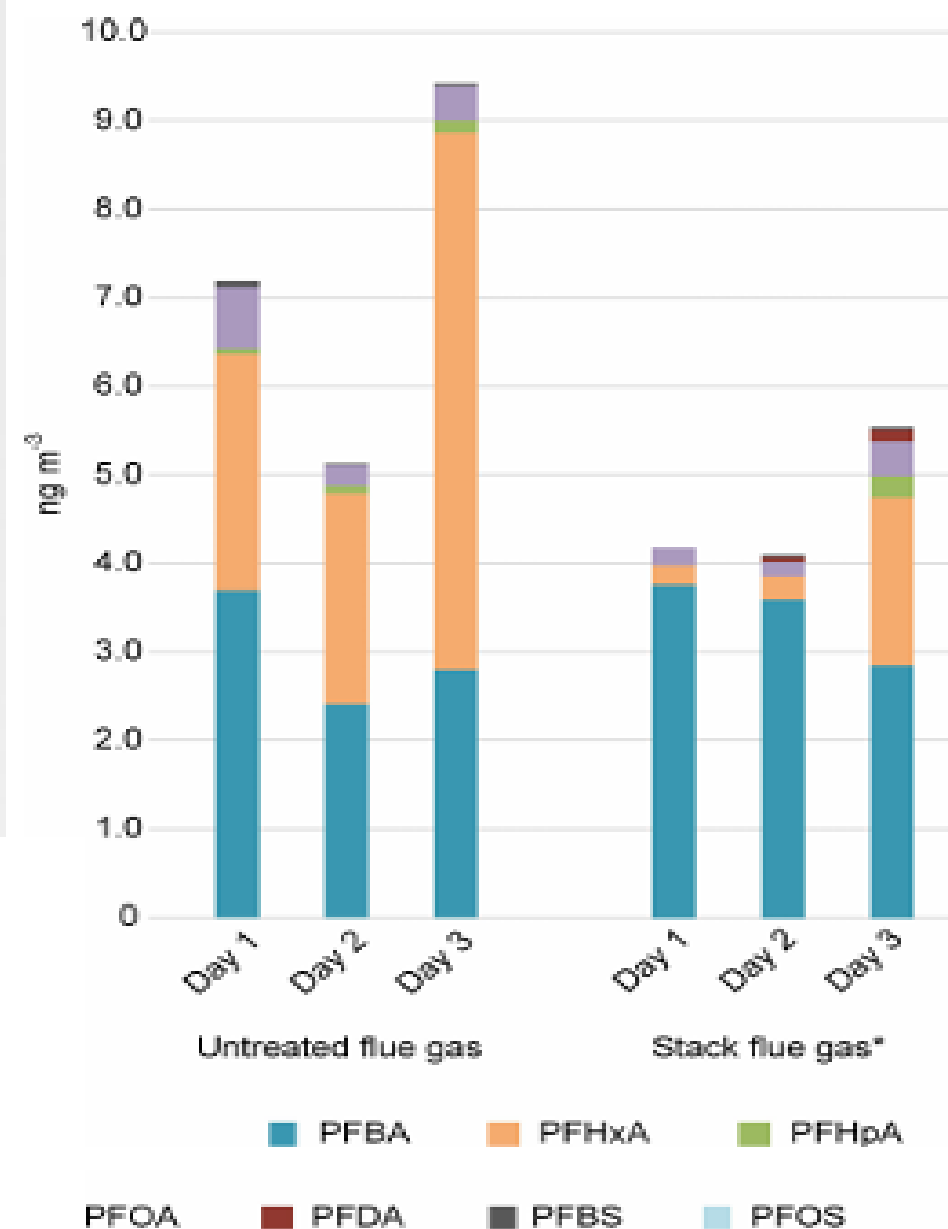
Typical concentrations: **4–20 ng/m³ in flue gas**; higher in condensate and ash

Wet flue gas treatment can reduce PFAS levels by ~35%, but efficiency varies by compound
Some PFAS (e.g., PFHxA) are efficiently captured; others (e.g., PFBA) are not

PFAS entering the capture plant



Perfluoroalkanes (fully fluorinated alkanes like CF₄, C₂F₆) are known to be thermally stable and can be present in flue gas, but their specific concentrations are generally not reported separately in these studies.

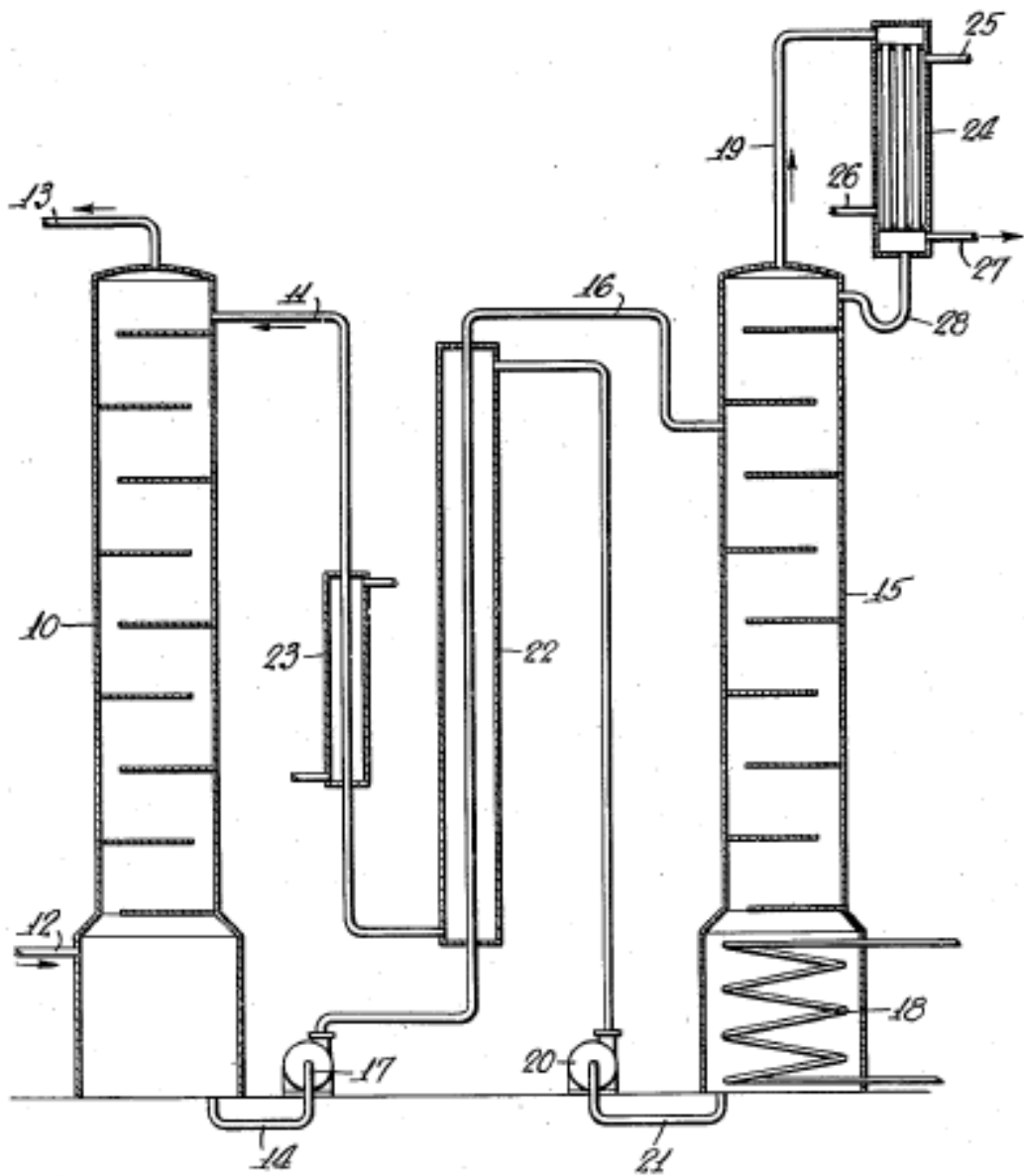


Sofie Björklund, Eva Weidemann, and Stina Jansson*

Cite This: *Environ. Sci. Technol.* 2024, 58, 8457–8463

[Read Online](#)

A worldclass example: AVR DUIVEN



PFAS expected to enter the capture plant as gaseous components and as solids (eg flue ash)

PFAS effect on the capture plant

Most PFAS are expected to be removed during CO₂ recovery, but some polar, non-dissociating PFAS may end up in the captured CO₂ stream or in the treated flue gas

Total PFAS (ng/g)
in solid streams

Solid							ng g ⁻¹
Boiler ash	n.d.	n.d.	n.d.	0.67	0.40	1.2	
Filter ash	0.28	0.79	0.70	0.75	0.13	0.37	
APCR†*	n.d.	n.d.	n.d.	1.3	0.99	1.1	
Bottom ash*	0.16	n.d.	0.54	1.4	0.81	1.5	
Gypsum*	n.d.	n.d.	n.d.	0.31	0.17	0.26	

PFCAs like PFHxA and occasionally PFOA are the main PFAS species captured on fly ash, with some variability in PFSA presence (e.g., PFOS) depending on specific plant conditions and waste feedstock.

PFAS effect on the capture plant

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Solvent reclaiming

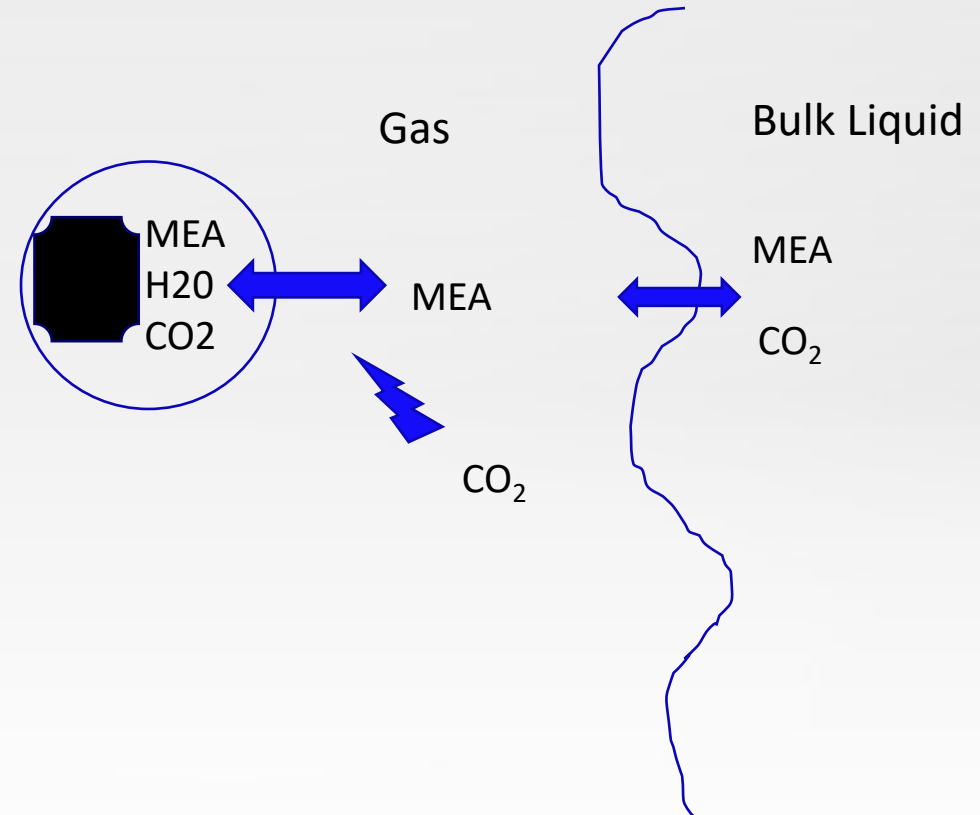
MEA visual change in 1000 operating hours



PFAS effect on the capture plant

Most PFAS are expected to be removed during CO₂ recovery, but some polar, non-dissociating PFAS may end up in the captured CO₂ stream or in the treated flue gas

Aerosol based emission



PFAS in treated flue gas

Short-chain PFAS, fluorotelomer alcohols, and certain substituted perfluoroalkane sulphonamido ethanols are most likely to persist through incineration and gas cleaning

- Short-chain perfluorocarboxylic acids (PFCAs)
PFBA (perfluorobutanoic acid, C4) and PFHxA (perfluorohexanoic acid, C6)
- Fluorotelomer alcohols (FTOHs)
3-5:2 FTOH (fluorotelomer alcohols with 3–5 fluorinated carbons) exhibit high volatility and low adsorption affinity, enabling them to bypass wet scrubbing systems
- Substituted perfluoroalkane sulfonamido ethanols
MeFASEs and EtFASEs (methyl and ethyl derivatives) persist due to their non-dissociating, polar nature, allowing them to partition into recovered CO₂ streams
- Perfluoroalkanes (PFCs)
Fully fluorinated alkanes like PFHx (perfluorohexane) show low reactivity and high thermal stability, surviving incineration and gas cleaning

PFAS effect on CO₂ quality

No published measurements of PFAS in recovered CO₂ to date

Standard fluoride tests are not suitable for PFAS; specific PFAS analysis or total organic fluorine (TOF) methods are needed

Detection limits for PFAS in flue gas: 0.1–0.5 ng/m³; for CO₂: estimated 10 ng/m³

Upgrading via adsorption Media

Eg Granular activated carbon (GAC):

Less effective for short-chain PFAS (e.g., PFBA, PFHxA) due to lower adsorption capacity. Breakthrough occurs 2–3x faster than for long-chain compounds like PFOS

Functional group impact: Sulfonate-containing PFAS (e.g., PFOS) adsorb better than carboxylates (e.g., PFOA) of equivalent chain length

When
are we
happy?

Conclusions & Recommendations

PFAS are not fully destroyed during waste incineration and can be found in flue gas, ash, and condensate.

Wet flue gas treatment can reduce but not eliminate PFAS; capture efficiency depends on PFAS type.

Most PFAS are expected to be removed during CO₂ recovery, but some may contaminate the captured CO₂ stream.

There is a critical need for improved measurement, monitoring, and treatment technologies to address PFAS in WtE and CO₂ capture processes



Zero
doesn't
exist

Great to have you on board of the change



[Let's energise innovation together](#)

Questions?

