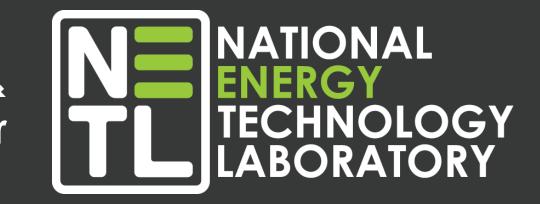
Review of Sensors for In-Situ Amine Degradation Monitoring in Post-Combustion Carbon Capture

Research & **Innovation Center**



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Introduction

- Reducing the CO₂ emissions is paramount to meet the decarbonization goal of net-zero emission by 2050
- Post-combustion carbon capture offers a variety of advantages:1-4
 - Retrofitted to existing coal fired power plants
- Suitable for natural gas fired power plants
- Power generation can be achieved even if the carbon capture process is down for maintenance unlike the pre-combustion process
- Chemical absorption is a widely used post-combustion method¹⁻⁴
 - The most common chemical absorbers are amine-based solvents
 - These solvent system degrades losing its carbon capture efficiency over time
 - Monoethanolamine (MEA) being the most studied
- □ Objective
 - In situ real-time monitoring of amine degradation will optimize operational control, carbon capture efficiency, and reduce the overall cost

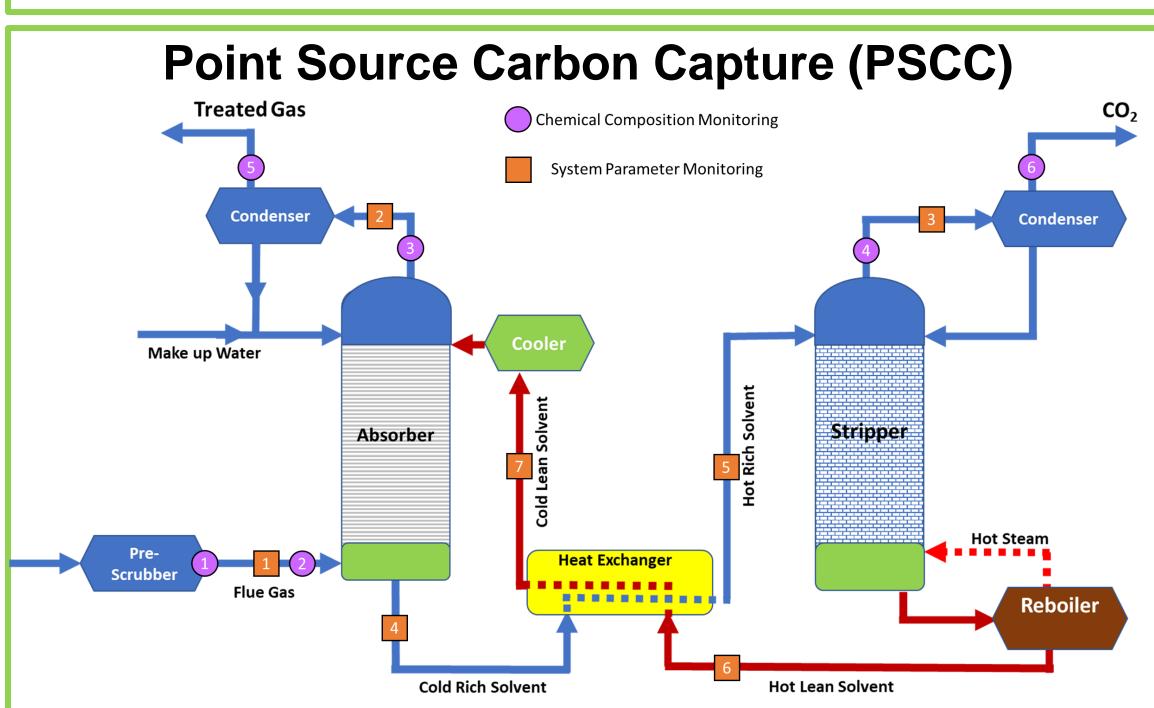


Figure 1. Pictorial representation of PSCC system with system parameter and chemical composition monitoring locations indicated

Amine Degradation Mechanisms^{1,5-9}

- Oxidative: absorber, cross exchanger
- Thermal: stripper
- Caused by flue gas contaminants

Problem Statement: 1) Solvent degradation is hindering large-scale deployment of amine-based carbon capture. Amine solvent degradation associated costs can be significant compared with the cost to monitor. 2) Existing monitoring methods usually involve sampling from the process lines and sending samples to laboratories for analysis using expensive instruments.

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Industry Monitoring Necessities

Table 1. Industrial monitoring requirements both current and future

Current Monitoring Requirements	Emerging Monitoring Requirements
 Density of Solvent Viscosity of Solvent Temperature CO₂ Capture Efficiency Chemical Composition Pressure pH Amine Concentration 	 Water Mass Balance < 1 % Nitrosamine Concentration NH₃ Concentration in Water Wash SO₃ Impact on Aerosol Ammine Carryover Trace Metal: Mercury, Arsenic, Selenium, Chromium pH Changes Following Distinct Functional Groups Color Change of Solvent

State of the Art Monitoring

Electrochemical Changes

Physical Parameters

Table 2. Physical monitoring parameters for PSCC²⁻⁶

Location	Equipment	System Parameter Monitoring
1,2,3	Pressure Gauge	Pressure of Gas and Liquids
1,2	Volumetric Flow Rate	Rate of Gaseous Flow
4,5,6,7	Viscosity	Flow Rate of Solvent
4,5,6,7	Temperature	Temperature of Solvent

Monitoring locations for Table 2 & 3 are indicated in Figure 1.

Cost of analysis instrument

degradation products

☐ Sensitivity to low-concentration

☐ Lack of monitoring of trace toxic

Periodic sampling

Point sensing

metals

Chemical Parameters

Table 3. Chemical monitoring parameters for PSCC²⁻⁶

Location	Equipment	Chemical Composition Monitoring
1	pH Meter	Basicity
1	UV	SO ₂ , NO ₂
1	Total Organic Carbon Analyzer	CO ₂
2,5,6	FTIR	CO ₂ , H ₂ O, NH ₃ , NO, NO ₂ , SO ₂ , CH ₂ O, C ₂ H ₄ O, Amines
2,5,6	NDIR	CO ₂
2	Paramagnetic	O_2
3,4	GC/MS	CO ₂ , O ₂ , N ₂ , H ₂ O
3,4	LC/MS	CO ₂ , O ₂ , N ₂ , H ₂ O
2,4	Electric Conductivity	O ₂ content
5,6	Single Ion Monitoring	Mass Spectrometry
5,6	Electric Low-Pressure Impactor	Aerosol Measurements (Size Distribution and Count)

Technology Gap

Table 4. Potential equipment cost for PSCC monitoring¹⁰

Equipment	Cost (\$)
pH Meter/ Automatic Titrator	\$ 3,000
UV Gas Sensor	\$ 10,000
Total Organic Carbon Analyzer	\$ 3,000
Fourier-Transform Infrared Spectroscopy (FTIR)	\$ 100,000
Nondispersive Infrared Sensor (NDIR)	\$ 20,000
Paramagnetic O ₂ Analyzer	\$ 8,000
Gas Chromatography–Mass Spectrometry (GC/MS)	\$ 100,000
Liquid Chromatography–Mass Spectrometry (LC/MS)	\$50,000
Electric Conductivity	\$ 1,000
Single Ion Monitoring	< \$ 50,000
	Aerosol
Electric Low-Pressure Impactor	Measurements (Size
Lieunic Low-Fressure impactor	Distribution and
	Count)

Key Parameters for Amine Degradation Monitoring

Direct Monitoring

Amine Solvent Color Change⁹

Amine degradation leads to color changes

Amine Concentration in Water^{5,8}

pH Change¹¹

Indicates CO₂ loading; CO₂ dissolution into water; heat stable salt neutralization

Degradation Products Detection⁸

Nitrate, sulfate salts, nitrosamine, ammonia gas

Indirect Monitoring

Temperature Monitoring⁸

Related to thermal degradation

O₂ Monitoring

- Oxidative: absorber, cross exchanger
- O₂ concentration: 5-10 ppm in solvents

Monitoring of Flue Gas Contaminants

• SOx, NOx, etc.

Toxic Trace Metal Ion Monitoring

• Trace Metals: Hg, As, Se, Cr



Figure 2. Examples of an amine solvent system degradation over time. 9

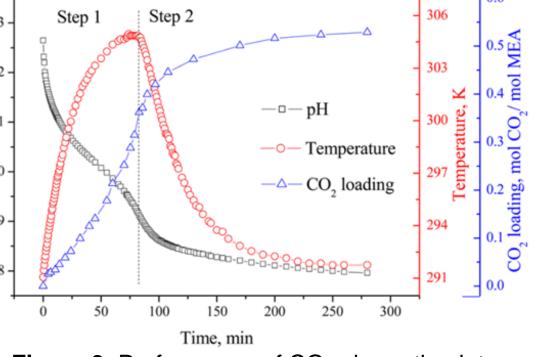


Figure 3. Performance of CO₂ absorption into MEA solution over time.¹¹

NETL Sensor Capabilities

Long-distance Distributed Optical Fiber Sensors

Imperfections in fiber lead to Rayleigh backscatter

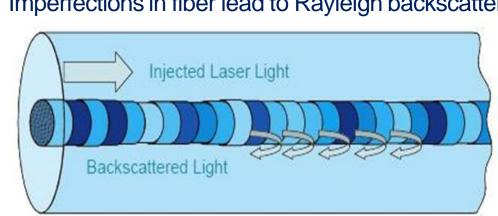


Figure 4. fiber optic representation of Rayleigh backscattering

Electrochemical Sensors

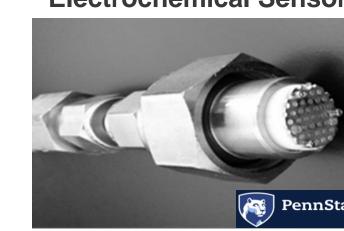
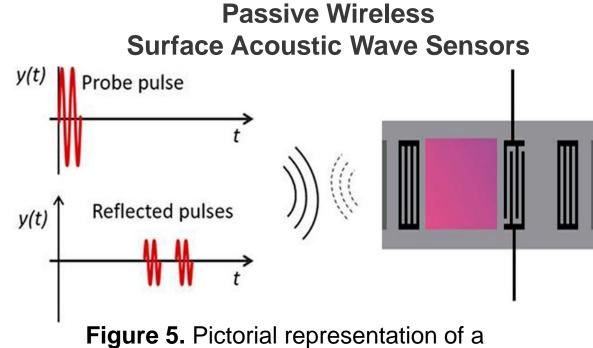


Figure 6. Picture of electrochemical sensor



surface acoustic wave sensor

LIBS: Laser Induced Breakdown Spectroscopy

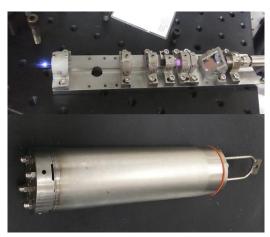


Figure 7. Picture of a LIBS probe

Summary

In-situ monitoring with NETL's sensor capabilities can be developed for deployment into the postcombustion carbon capture streams. These sensors will provide feedback on the carbon capture efficiency, solvent health and reduce operational costs.

In EY2022:

- Identified key indicators for amine degradation as sensing targets.
- Surveyed and selected low-cost existing sensor technologies for these targeted indicators, instead of expensive full-on laboratory chemical analysis Interviewed industry stakeholders such as The National Carbon Capture Center (NCCC) and Ion
- Clean Energy to learn the monitoring needs for post-combustion carbon capture process. 4. A report is prepared on monitoring needs, sensor technology survey, and recommendation for cost-
- effective online monitoring of amine degradation

Next Step:

In EY2023, pilot-scale testing of NETL-developed optical fiber sensors for amine degradation and CO₂ monitoring at NCCC.

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