

Liquid Solvents for Direct Air Capture and Utilization

Experimental Characterization, Optimization and Stability Testing

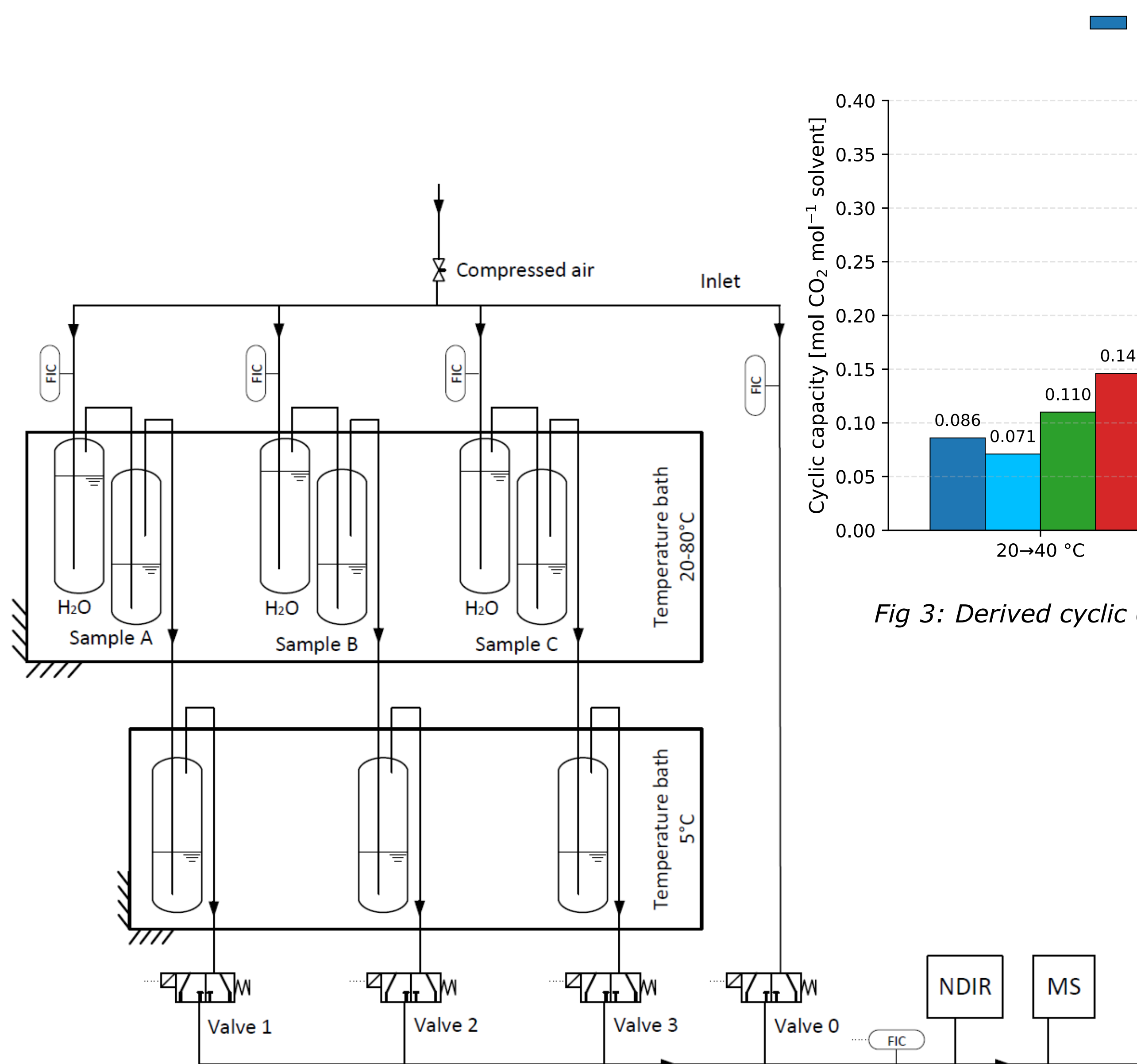


Fig 1: Schematic of experimental setup for parallel absorption and desorption experiments

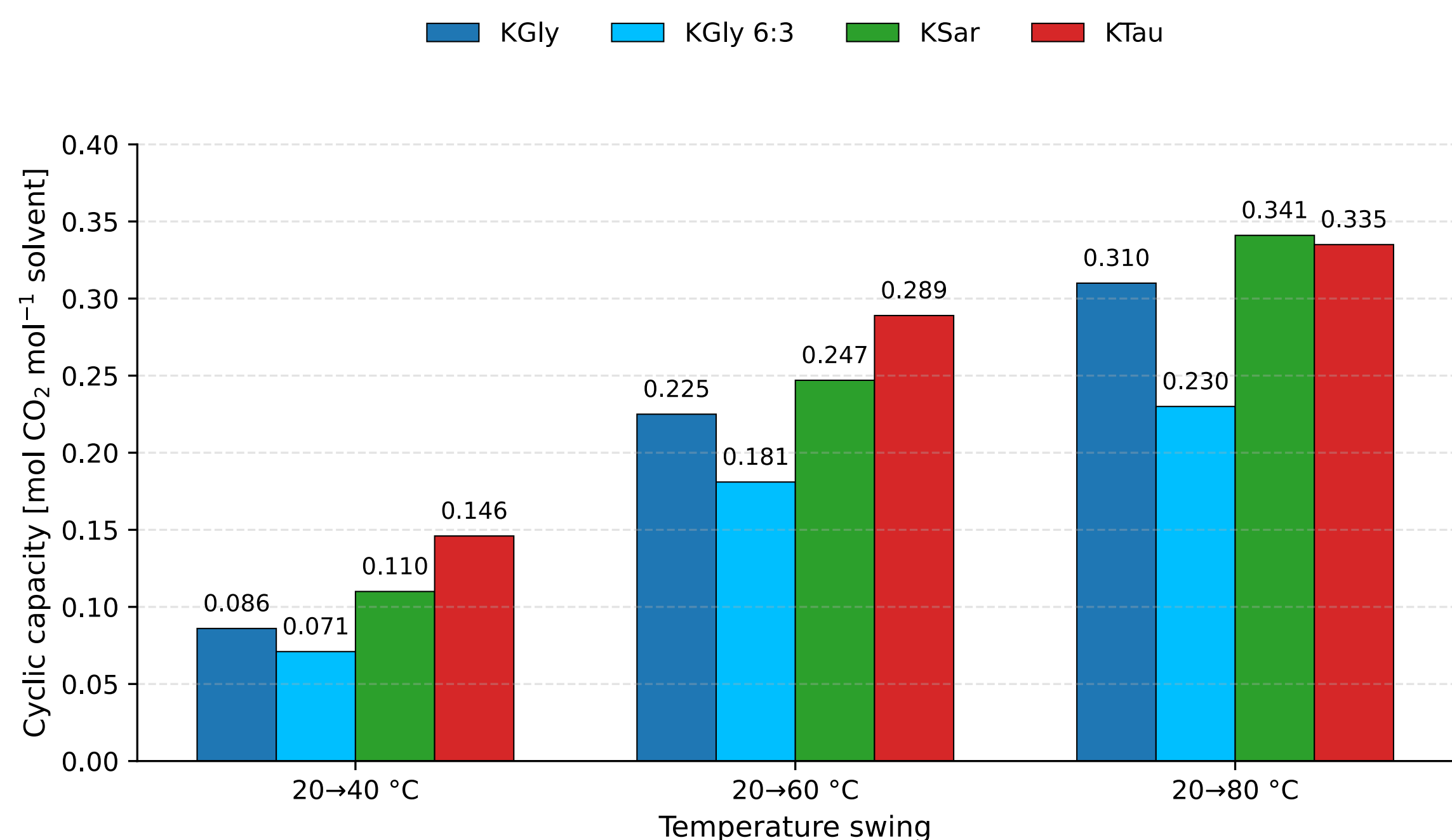


Fig 3: Derived cyclic capacities for tested solvents and temperature swings



Fig 2: Solvent samples after CO₂ absorption: thermally cycled >200 times left, <10 times right

Motivation

Direct air capture (DAC) can remove CO₂ from ambient air and provide a renewable CO₂ source, for example for greenhouse enrichment.

Because air contains only about 400 ppm CO₂, efficient solvents are needed. Amino acid salt solutions are promising because they chemically bind CO₂ and have low volatility.

Procedure

Three 3 mol/L aqueous amino acid salt solvents were tested:

- Potassium Glycinate (KGly)
- Potassium Sarcosinate (KSar)
- Potassium Taurinate (KTau)

The solvents were investigated in a parallel absorption-desorption setup using ambient air (Fig 1). CO₂ uptake and release were measured with an NDIR gas analyzer during temperature swings from 20 to 80 °C.

Additional tests evaluated solvent enhancement strategies using TiO₂ addition and excess KOH, as well as thermal stability by cycling loaded solvents 200 times between 20 and 90 °C.

Key Findings

Cyclic capacity is more relevant than absolute CO₂ loading for temperature-swing DAC operation, because it describes how much CO₂ can be repeatedly absorbed and released.

KSar showed the highest cyclic capacity for the 20 → 80 °C temperature swing, while thermal cycling over 200 cycles caused no measurable performance loss.

KTau showed the strongest response at moderate temperature swings (Fig 3), reaching 0.289 mol CO₂/mol solvent for 20 → 60 °C.

For the larger 20 → 80 °C swing, KSar reached the highest cyclic capacity with 0.341 mol CO₂/mol solvent, followed closely by KTau and KGly.

Excess KOH increased absolute CO₂ loading but did not improve cyclic capacity (KGly 6:3 in Fig 3).

TiO₂ addition showed no systematic performance improvement under the tested conditions.

Thermal cycling (Fig 2) caused no measurable loss in absorption/desorption performance, although further chemical analysis is needed to assess degradation

Céline Winkelhausen

Supervisor: Dr. Benjamin Fumey

Advisor: Prof. Dr. Mirko Kleingries

Expert: Dr. Andreas Borgschulte