## Global Flow Measurement Workshop 22 - 24 October 2024

## **Extended Abstract**

# Online measurement of impurities in CO<sub>2</sub> by using electrical permittivity

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## 1 INTRODUCTION

Carbon Capture and Storage (CCS) is expected to be an important part of the effort of reducing the emission of greenhouse gases in the years to come. Depending on the capture process, there will be different impurities in the captured  $CO_2$ . These impurities can cause problems for the transport of  $CO_2$  from the place of capture to the reservoir where it will be stored. To comply with  $CO_2$  specifications, monitoring of impurities is crucial. Common impurities expected in the CCS streams are  $O_2$ ,  $N_2$ ,  $A_7$ ,  $H_2O$ ,  $SO_2$ ,  $H_2$  and  $CH_4$ . This work explores the use of electrical permittivity for measurement of impurities.

## 2 THE MEASUREMENT CONCEPT

The concept focuses on identifying changes in the stream concentration from the measured stream permittivity. The measured permittivity can then be compared to the dielectric constant of pure  $CO_2$ , or to a predetermined stream permittivity range. The hypothesis is that the measurement system can report a level of impurity if the difference between the measured and the reference permittivities is above a certain level. The method requires a contrast in permittivity between the  $CO_2$  and the impurities. Impurities like water, where the difference in permittivity is larger, will be possible to detect in smaller quantities compared to impurities with permittivity closer to that of  $CO_2$  [1].

In this work two metering principles were used, which allows technology intercomparison. The low-frequency electrical permittivity was measured using (i) contact electrodes and (ii) microwave resonance measurements. The impurity used for the tests was nitrogen since  $CO_2$ - $N_2$  mixtures are well documented and agree well with predictions from GERG-2008 EOS [2].

#### 3 THE EXPERIMENTAL WORK

The experimental work was done at the thermal engineering laboratories of Sintef and NTNU in Trondheim, Norway. The equipment under test were installed in a temperature-controlled room. The fluids,  $CO_2$  and  $N_2$ , were injected from feeding cylinders into the rig and permittivity was measured by electrode measurements using an Emerson's Roxar Multiphase Flow Meter and by microwave resonance measurements using a Roxar Watercut Meter. The two meters were installed horizontally in series. The rig has 8mm stainless steel tubing and includes temperature and pressure sensors and a pump for circulating the fluid. The tests encompassed liquid  $CO_2$ , at temperatures ranging from -6 °C to 3 °C and pressure between 40 bara and 80 bara. The mass fraction of nitrogen ranges from 0 to 1.1%. The measured permittivities were compared with theoretical values by Harvey and Lemmon [3] as implemented in REFPROP v10.

An approach to identify the presence of impurities is to measure the permittivity and compare this with the calculated permittivity of  $CO_2$  at the actual temperature and pressure. This was investigated by looking at a series of test points with mass fraction  $N_2$  of 0%, 0.6% and 0.8%. Results are plotted in figure 3. This deviation appears as expected to be proportional to the mass fraction of  $N_2$  for both technologies.

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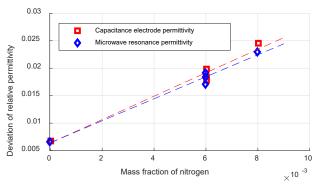


Fig. 1 - Deviation between measured relative permittivity of mix and calculated relative permittivity of  $CO_2$  vs mass fraction of  $N_2$ .

The effect of  $N_2$  on the measured permittivity is also seen in Figure 2, where measured and calculated permittivities are plotted for two test points. The temperature (-3 °C) and pressure (45 bar) are the same for the test points. Test point 1 is pure  $CO_2$  and for test point 2 there is 1.1% mass  $N_2$ . The uncertainty of the measurements is indicated in the figure, the error bars are showing one standard deviation.

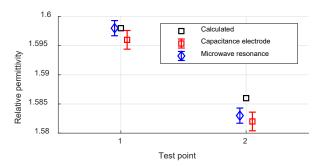


Fig. 2 – Theoretical and measured relative permittivity for two test points, without impurity (test point 1) and with 1.1% mass  $N_2$  (test point 2).

## 4 CONCLUSIONS

This work demonstrates the feasibility of using electrical permittivity measurement to find impurities in  $CO_2$  flow. The results indicate that nitrogen levels of 0.2%mass fraction can be feasible to detect using this concept for both technologies tested. The concept can be used for determining the presence of contamination but does not identify which impurity is present. The detectable level of impurity depends on the relative permittivity of the impurity.

#### **5 REFERENCES**

- [1] Y. Arellano, S. H. Stavland, E. C. Panduro, B. Hamre and B. T. Hjertaker, "Imaging measurement technologies for CCS," 2022 IEEE 12th Sensor Array and Multichannel Signal Processing Workshop (SAM), Trondheim, Norway, 2022, pp. 56-60, doi: 10.1109/SAM53842.2022.9827823.
- [2] O. Kunz, W. Wagner, The GERG-2008 wide-range equation of state for natural gases and other mixtures: an expansion of GERG-2004, J. Chem. Eng. Data 57 (11) (2012) 3032e3091, http://dx.doi.org/10.1021/je300655b.

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[3] A. H. Harvey and E. W. Lemmon, "Method for Estimating the Dielectric Constant of Natural Gas Mixtures," International Journal of Thermophysics, vol. 26, no. 1, 2005, doi: 10.1007/s10765-005-2351-5.