



## **A direct and simultaneous method for detecting atmospheric Ar, O<sub>2</sub>, and N<sub>2</sub> by a gas chromatograph equipped with a thermal conductivity detector (GC-TCD)**

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Carbon dioxide (CO<sub>2</sub>) and oxygen (O<sub>2</sub>) fluxes are inversely linked through the processes of fossil fuel combustion, terrestrial photosynthesis, and respiration, while with no correlation during air-sea exchange. This behavior of CO<sub>2</sub> and O<sub>2</sub> fluxes provides an insight into distinguishing the terrestrial and oceanic sinks for anthropogenic CO<sub>2</sub> by measuring atmospheric O<sub>2</sub> and concurrent CO<sub>2</sub> concentrations. In addition, atmospheric Ar/N<sub>2</sub> ratio is expected to undergo very slight variations due to exchanges of Ar and N<sub>2</sub> across the air-sea interface, driven by ocean solubility changes and thus observing such minute variations may provide useful constraints on large-scale fluxes of heat across the air-sea interface. In our study, we described a direct and simultaneous method for detecting atmospheric Ar, O<sub>2</sub>, and N<sub>2</sub> by a gas chromatograph equipped with a thermal conductivity detector (GC-TCD) with the aid of a HP-PLOT molecule sieve capillary column and semiconductor cryogenic technology. Preliminary results showed this method could provide a good separation between Ar and O<sub>2</sub> in the atmosphere, but a little difficulty to measuring such minute changes in the atmospheric Ar/N<sub>2</sub> and O<sub>2</sub>/N<sub>2</sub> ratios.