



Preliminary results from the Lauder site of the Total Carbon Column Observing Network TCCON



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Abstract

In this poster we give a brief description of the Total Carbon Column Observing Network (TCCON), a new network of ground-based Fourier Transform spectrometers (FTS) dedicated to measurement of greenhouse gas absorption (CO₂, CO, CH₄, N₂O) in the near infrared.

We present preliminary retrievals of CO₂ total column densities from the TCCON site in Lauder, New Zealand between July 2004 – when routine measurements began – and April 2005, and compare these retrievals with in-situ CO₂ measurements from the surface monitoring network station at Baring Head, New Zealand.

A discussion of how we plan to compare the ground-based FTS retrievals with in-situ surface observations and greenhouse gas retrievals from satellite radiances follows.

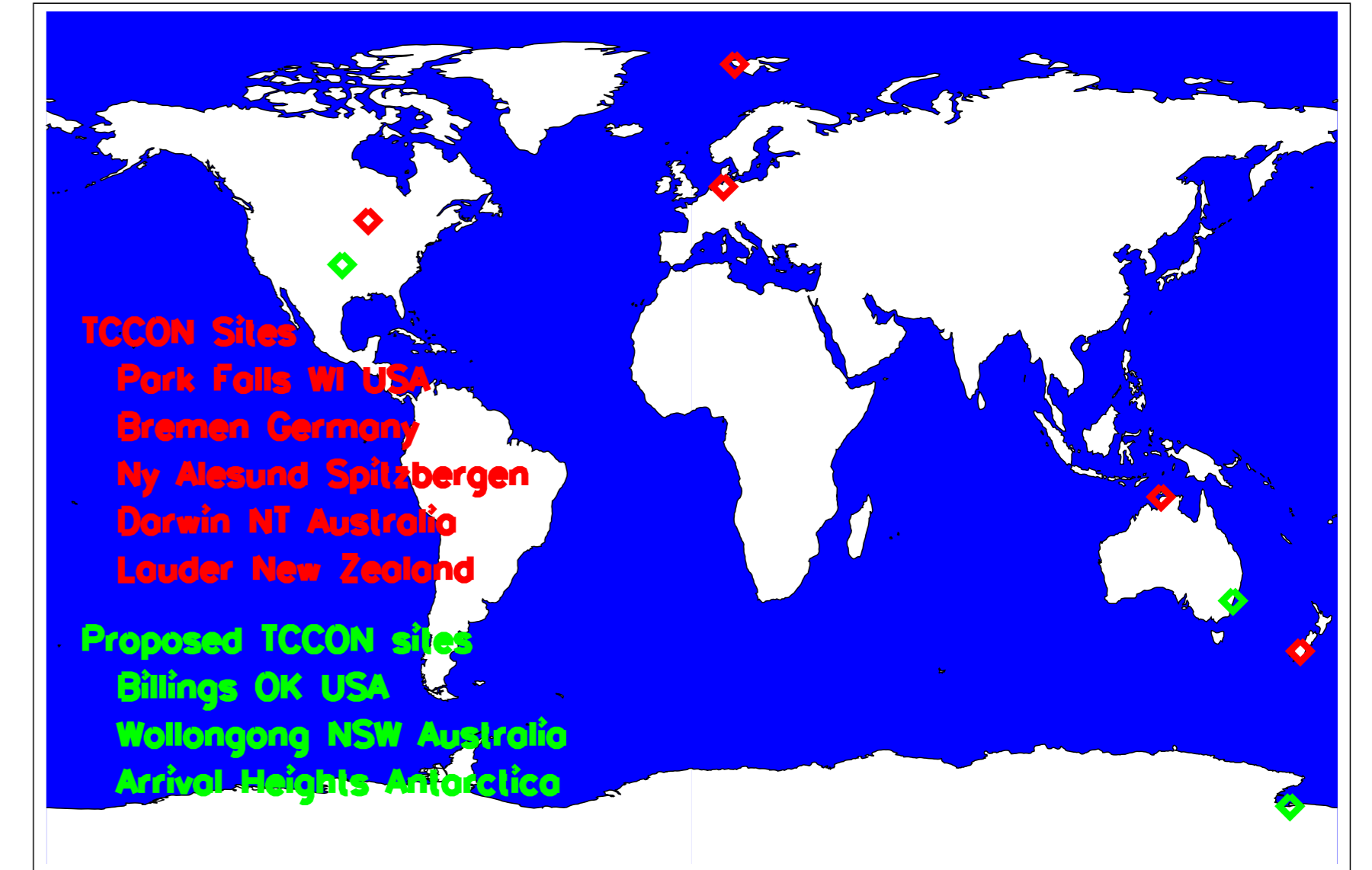
Overview of the TCCON network

TCCON is a global network of ground-based Fourier Transform spectrometers. Near infrared solar absorption spectra are acquired in the 4000–14000 cm⁻¹ interval at 0.02 cm⁻¹ resolution and analysed to retrieve CO₂, CO, CH₄, N₂O column densities and column average mixing ratios (denoted X_Y).

Column-integrated trace gas measurements are less sensitive to local sources and sinks and seasonal and diurnal rectifier effects than in-situ surface measurements.

However, if the TCCON network measurements are to provide useful additional constraints for the global carbon budget – both directly, and through validation of satellite column measurements – they must achieve a precision of 0.1% and an accuracy of 0.3%.

Accurate solar tracking, surface pressure measurements, spectroscopy and retrieval algorithms are all key to achieving



these error targets.

At each TCCON site the FTS measurement is complemented with measurements of surface pressure and in-situ surface trace gas concentrations. The FTS instrument line shape (ILS) is routinely monitored using a HCI cell.

Preliminary results from Lauder using a profile scaling retrieval algorithm

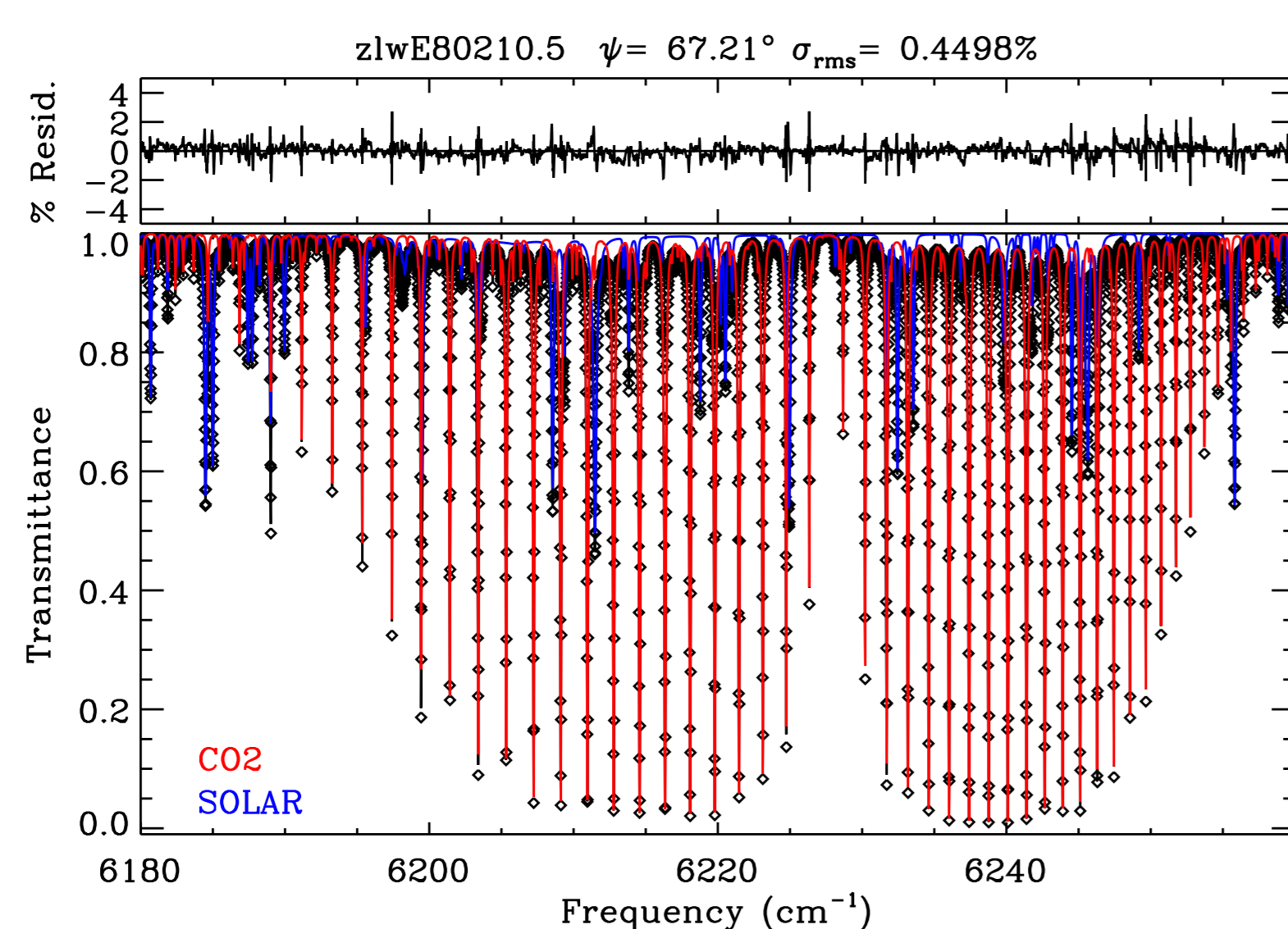


FIGURE 1: An example of simulated (solid black lines) and observed (diamonds) transmittance spectra in the 1.6 micron CO₂ band, identifying CO₂ and solar absorption lines. Fitting residuals are illustrated in the upper panel.

Timeseries of 1.6 micron band CO₂ retrievals

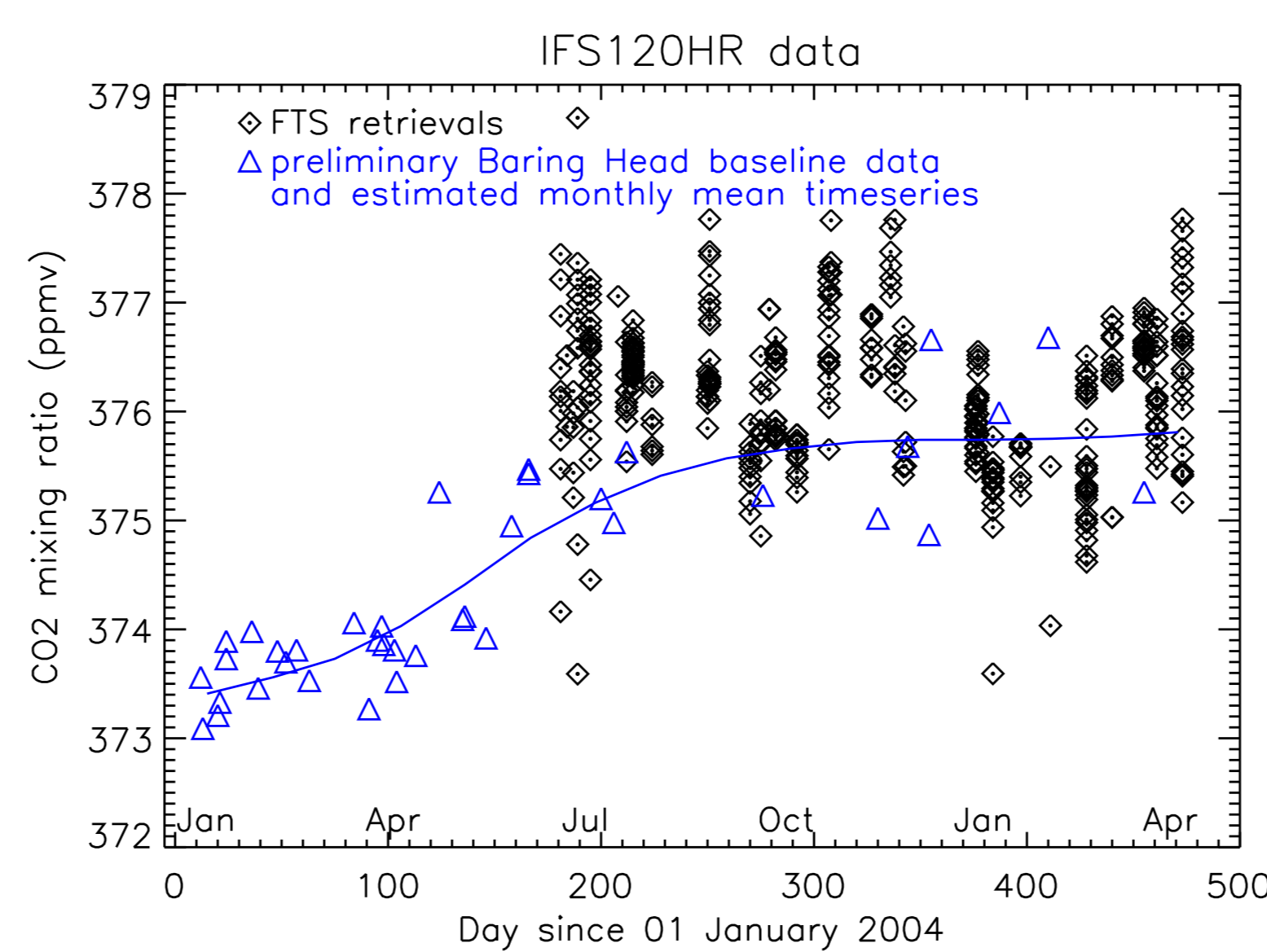


FIGURE 2: CO₂ column average volume mixing ratios derived from 1.6 micron band CO₂ retrievals. The mean mixing ratio for the time-series is 376.2 ± 0.7 ppmv (0.2%). The standard deviation of diurnal variations range from 0.1 to 1.0 ppmv and the day-to-day variation is of the order of 0.5 ppmv. In-situ CO₂ measurements from Baring Head are illustrated for comparison. Modelling studies predict diurnal and seasonal variations in the CO₂ column of 0–0.6 ppmv and 1 ppmv respectively in the Southern Hemisphere [Olsen and Randerson, 2004].

Use of O₂ column densities as an internal network standard

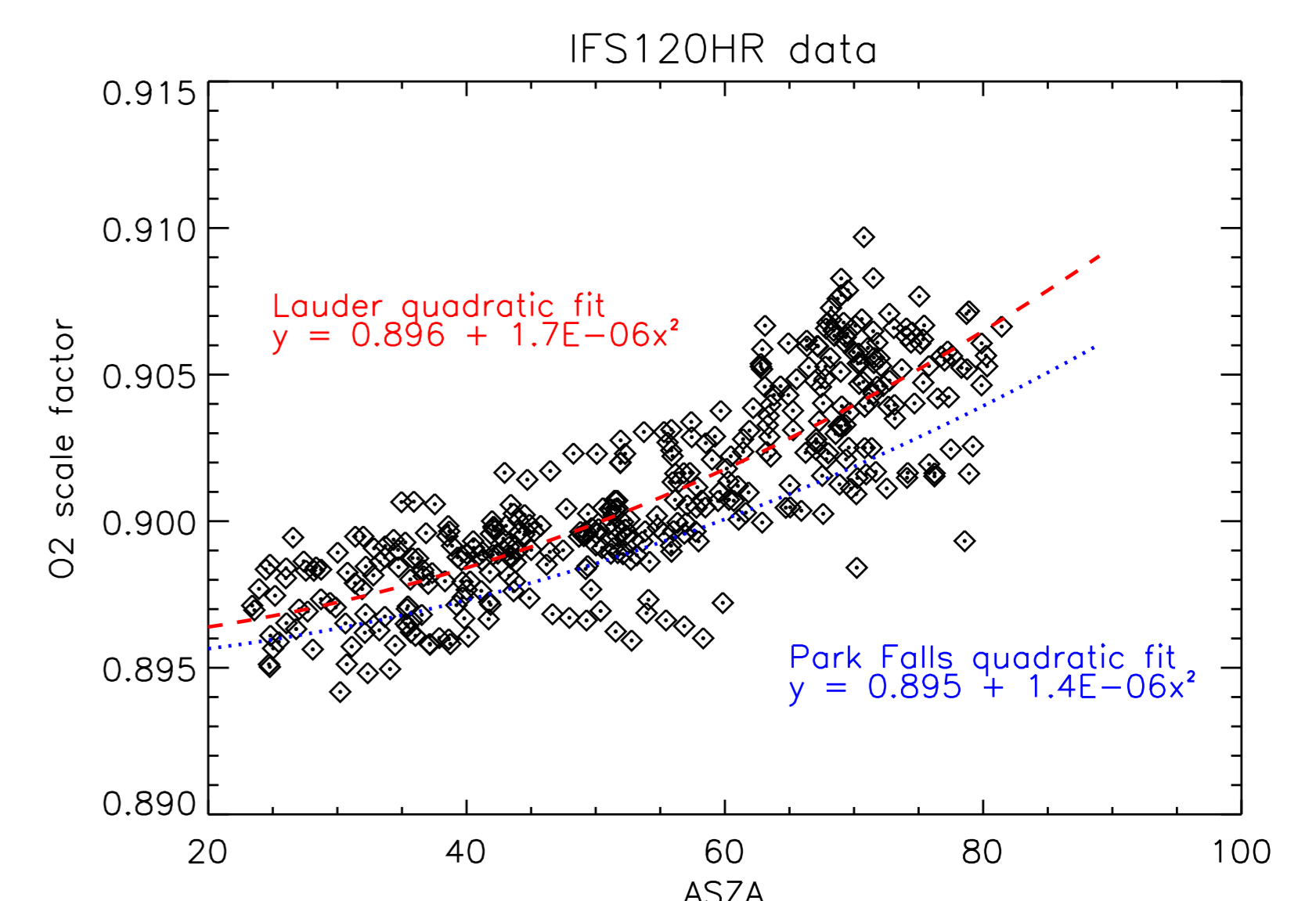


FIGURE 3: O₂ profile scale factors retrieved from the 1.27 micron band as a function of solar zenith angle. Scale factors should be unity at all zenith angles. The non-unity, zenith angle dependent character of retrievals should be corrected with improved band strength and spectroscopic parameter estimates. At a given solar zenith angle retrieval precision is of the order of 0.002 (0.2%). Target precision is 0.001 (0.1%). Lauder and Park Falls retrievals agree to within 0.1–0.3%.

Profile retrieval and applications

Profile retrieval has advantages over profile scaling algorithms

- improves precision for column retrievals due to reduced smoothing error (nearly ideal SZA-independent total column averaging kernel)
- provides some vertical resolution (typically 3–4 degrees of freedom for signal)

but it is potentially more sensitive to forward model errors e.g. specification of ILS, spectroscopy, atmospheric temperature.

Tabulated retrieval error budgets assume a signal to noise ratio of 500:1 and a perfect forward model. The a priori standard deviation of X_{CO₂} is 12 ppmv¹.

Planned comparison of ground based FTS and satellite measurements and retrievals will take retrieval errors and averaging kernels into account explicitly, following the work of Rodgers and Connor [2003]. One way of doing this is to apply satellite retrieval averaging kernels to ground based FTS profile retrievals.

Boundary layer and free troposphere partial column profile retrievals should improve comparisons with in-situ surface measurements and provide a potential link between surface and satellite measurements.

Ground based FTS CO₂ total column retrievals

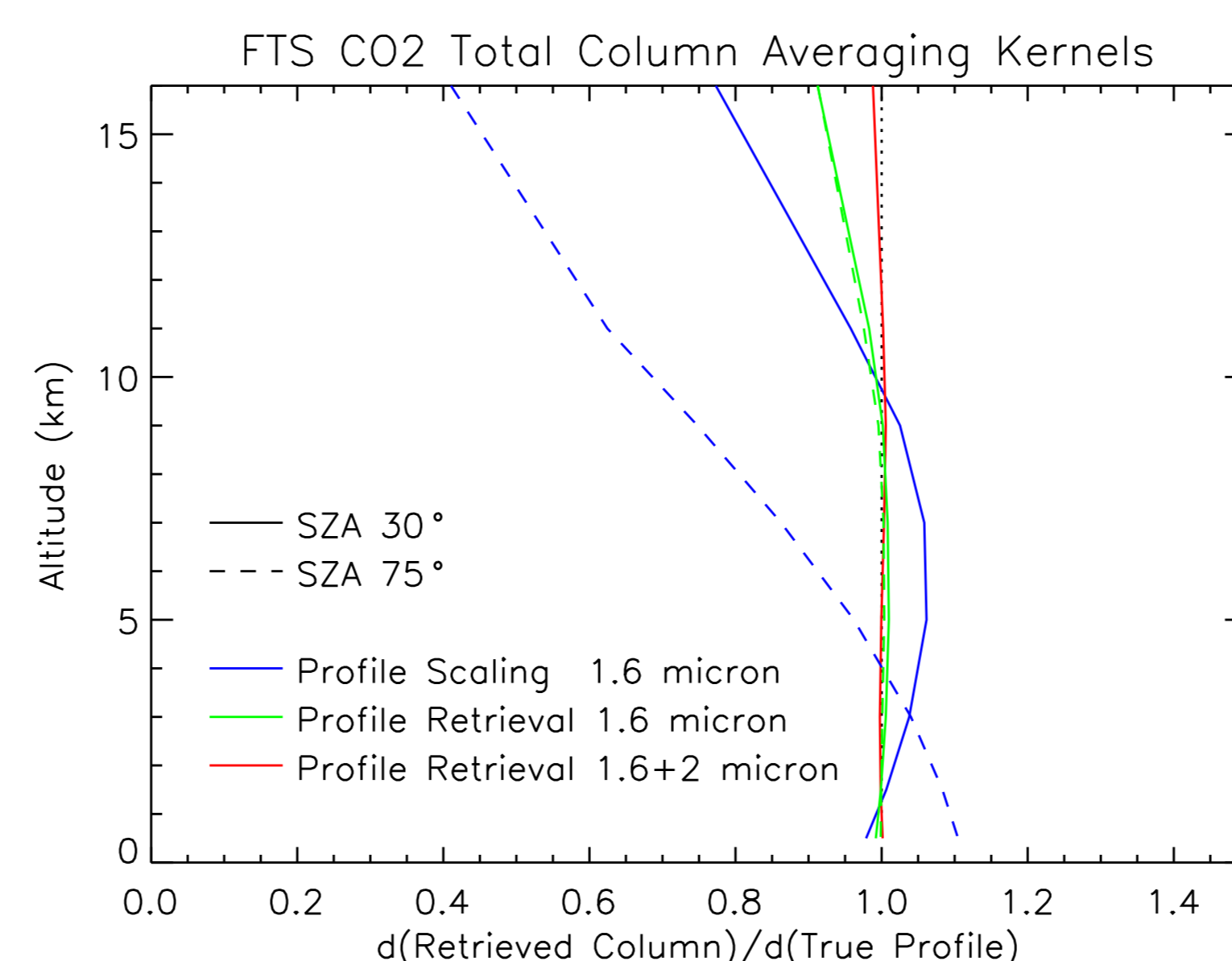


FIGURE 4: FTS CO₂ total column averaging kernels for profile retrieval and profile scaling algorithms and solar zenith angles of 30 and 75 degrees. The ideal column averaging kernel is unity at all levels.

Retrieval	SZA	Noise	Smoothing	Total	Units
Profile scaling	30°	0.10	0.24	0.26 (0.07%)	ppmv
	75°	0.05	0.67	0.67 (0.18%)	ppmv
Profile retrieval		0.13	0.06	0.14 (0.04%)	ppmv

Ground based FTS CO₂ partial column retrievals

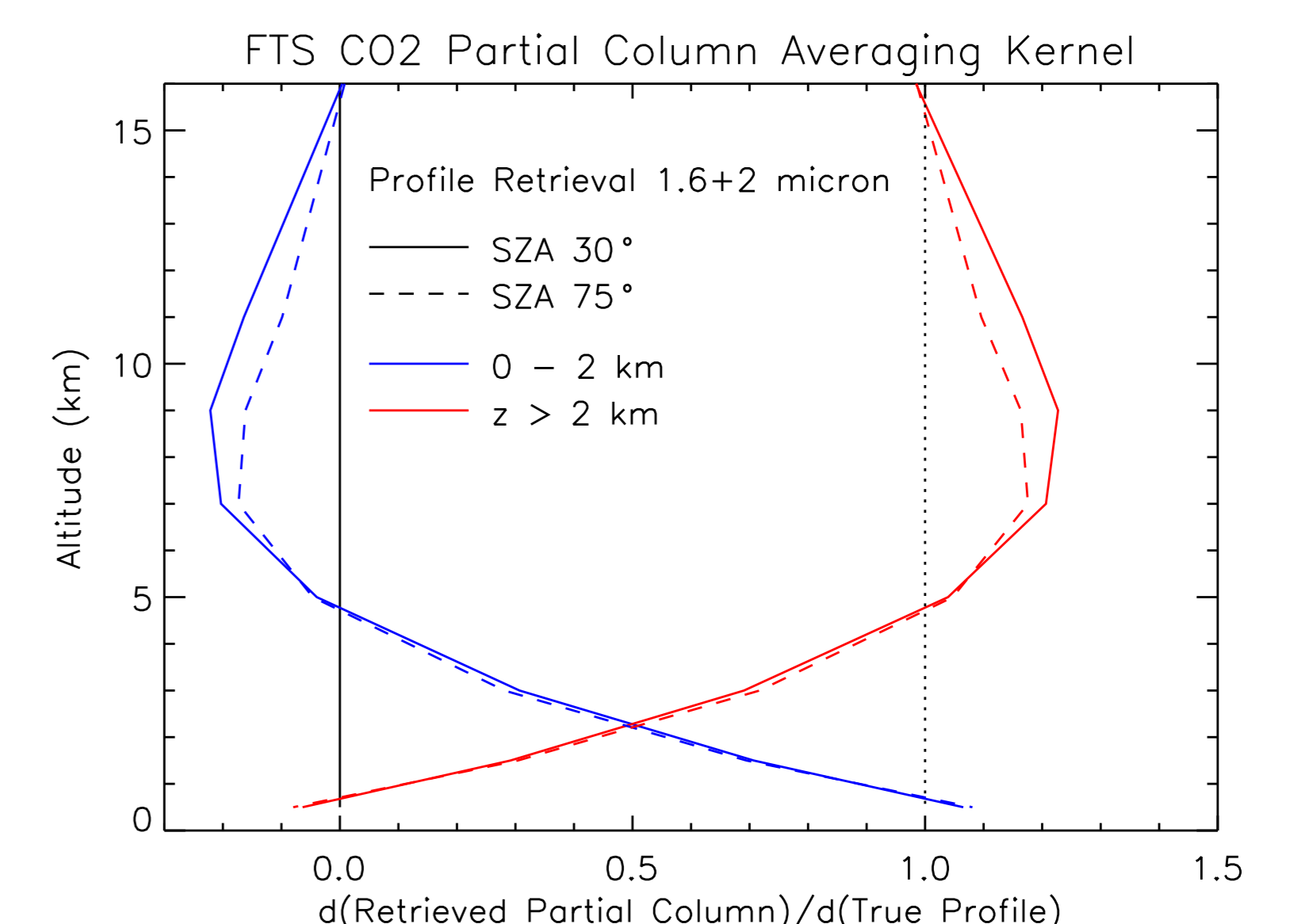


FIGURE 5: FTS CO₂ partial column averaging kernels for profile retrieval integration from 0–2 km (boundary layer) and for z > 2 km (free troposphere) and solar zenith angles of 30 and 75 degrees.

Partial column	SZA	Noise	Smoothing	Total	Units
0–2 km	30°	0.7	0.3	0.7	ppmv
	75°	0.2	0.1	0.2	ppmv
Total column		0.06	0.02	0.06 (0.02%)	ppmv

Conclusions

Preliminary results are encouraging: Lauder O₂ and CO₂ retrievals have a precision of 0.2% and O₂ retrievals are in good agreement with data from the Park Falls site.

Work is ongoing to identify the sources of observed variability and improve the precision and accuracy of retrievals to attain the TCCON error targets (0.1% precision, 0.3% accuracy).

This work will include monitoring and improving solar tracker accuracy, improving near infrared spectroscopic parameterisations (particularly O₂ collision induced absorption), the development and comparison of profile scaling and profile retrieval algorithms, with due consideration of the impact of forward model errors, and preliminary comparisons with in-situ CO₂ measurements at the Lauder site.

References

- S. C. Olsen and J. T. Randerson. Differences between surface and column atmospheric CO₂ and implications for carbon cycle research. *J. Geophys. Res.*, 109:D02301, 10.1029/2003JD003968, 2004.
- C. D. Rodgers and B. J. Connor. Intercomparison of remote sounding instruments. *J. Geophys. Res.*, 108(D3):4116, 10.1029/2002JD002299, 2003.

Acknowledgements

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¹ Supplementary material is available which details the assumed CO₂ vertical error covariance.

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