



ACTRIS 2 (ECAC-ACMCC) Intercomparison of Aerosol Chemical Speciation Monitors, November 2018

Project No.: ACSM-2018-1-11

Basic information

Location of the quality assurance: SIRTA/ACMCC, LSCE, Lab. 705

Delivery date: Before November 13th, 2018

Setup in the laboratory: November 14th 2018

Workshop period: November 19th to November 23th, 2018

Principal investigator	Institute	Participant	Instrument
Axel Eriksson	Lund University	Axel Eriksson Erik Ahlberg	ToF-ACSM-#19 equipped with PM1 lens and standard vaporizer

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Summary

This report summarizes results obtained for the University of Lund (LUND) Time-of-Flight Aerosol Chemical Speciation Monitor (ToF-ACSM) during the 2018 ACMCC ACTRIS-2 workshop. It is divided into four main sections. The first section describes the instrumental set-up and data treatment methodologies used for the workshop. The second section presents results obtained during the initial intercomparison, where instruments were operated as delivered by participants. The third section describes calibrations and maintenance (if any) performed during the workshop. The last section presents results obtained during a second intercomparison campaign, where instruments were tuned and calibrated at ACMCC.

Performance evaluation includes:

- 1) The instrument performance is within the acceptable limits evaluated using a Z-score analysis, as described by ISO 5725-2.
- 2) The instrument is in overall good agreement - i.e., within $\pm 30\%$ - with a reference dataset, the latter corresponding to a reference instrument and/or the robust median of intercomparison participants datasets.

The reference dataset is that measured by the SIRTa instrument. This instrument successfully participated in the previous ACSM intercomparisons and didn't travel prior to the workshop (unlike other instruments). It is regularly maintained and calibrated by the ACMCC staff. The robust median refers to the median value of all participating PM1 QACSM and excludes the PM2.5 QACSM instruments.

The LUND instrument appeared to underestimate concentrations of every chemical species during both intercomparison campaigns. This was notably due to a leak in the three-way valve system, that could be solved only after the workshop. It cannot be excluded that this technical issue was due to damage during transport and/or installation. Calibrations performed during the workshop seemed however to have improved the determination of the RIE SO₄ value.

1. Overview of the instrumental set-up and data treatment methodologies

Instruments were asked to be delivered at ACMCC no later than the 13th of November. They were installed and switched on by ACMCC personnel during the week from the 12th to the 16th of November. There were four different tables, each containing four instruments. Each table had its own inlet, fitted with a common sampling head. All sampling lines were composed of ½ inch copper tubes and were the same length for each instrument. Each instrument sampled 3 liters from the main inlet line, this flow was controlled by external sample line flow pumps. Instruments were fitted with or without an individual dryer, as used at their usual sampling site and/or as requested by participants. For the ToF-ACSM, data analysis was performed using the Igor Pro (Wavemetrics, v6.3.7) procedure Tofware (v2.5.13). A composition dependent collection efficiency was applied to all data sets following the guidelines available in Middlebrook et al., (2012).

Instruments were compared with the SIRTa reference instrument and also with the robust median of all instruments through Z-score analysis. For comparison to the reference datasets, it is expected that good instrument performance is met when the participant's results fall within a well characterized uncertainty for aerosol mass spectrometers, taken here as $\pm 30\%$ (Bahreini et al., 2009, Middlebrook et al., 2012; Crenn et al., 2015).

The Z-score analysis was applied following the standards defined by the international standard organization (ISO). This method has been evaluated according to ISO 5725-2 and provides a means to evaluate instrument performance relative to other instruments participating in the intercomparison.

Such a method has been applied within other European intercomparison exercises (JRC technical intercomparison reports), and was validated during the first ACTRIS1 ACSM intercomparison (Crenn et al., 2015). The Z-score test evaluates if the variations in the different instruments from the reference value fall within a defined criterion, following Eq.1:

$$Z_i = \frac{X_I - X^*}{\sigma_*} \quad (\text{Eq. 1})$$

Where X_I is the robust median of the selected instruments, X^* is the value of the instrument being compared and σ^* is the standard deviation of this latter instrument. The robust median target values were determined based on datasets obtained by PM₁ ACSM only (i.e., for consistency, the few PM_{2.5} ACSMs were not considered for this target value calculation). According to this test, instrument performance is considered acceptable when values fall between 2 and -2 (indicated by the green lines in Figures 2.1 and 4.4).

2. Results from initial intercomparison campaign

Once installed by ACMCC personnel, instruments were turned on with tunings indicated by participants, and sampled ambient air at the station from the 16th to the 19th of November 2018 for the initial intercomparison. Fifteen instruments were compared all together during a limited period of 18th Nov 18hrs to 19th Nov 08 hrs. The Z-score test illustrates the instrument performances during this period, (Figure 2.1) where each instrument was compared to the robust median of the participating instruments.

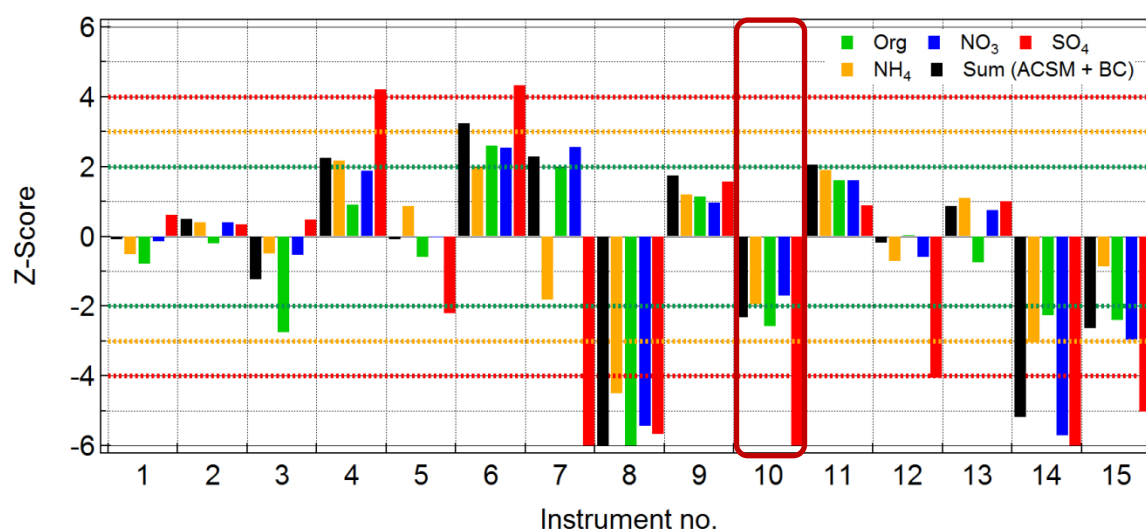


Figure 2.1. Z-score calculation for each species measured by the ACSM for each instrument that participated in the initial intercomparison campaign (18th Nov 18hrs to 19th Nov 08 hrs). The red rectangle highlights the LUND instrument.

The LUND instrument underestimated concentrations from all species when compared with the robust mean, failing to reach acceptable performance (especially for sulfate.). It should also be noted that, given the large variability in instrument performance prior to calibration, the median value can be considered as a weak reference for comparison. This is illustrated in Figure 2.2 below where the median NR-PM₁ + BC mass concentration is compared with the NR-PM₁ mass concentration measured from the Fine Dust Aerosol Spectrometer (FIDAS, Palas ®) and the Tapering Elemental Oscillating Microbalance with Filter Dynamics Measurement System (TEOM-FDMS) instruments,

giving a slope of 0.72 and 0.37 respectively. In comparison the NR-PM1 + BC mass concentration measured from the SIRTa instrument compares better with slope values of 1.08 and 0.56 respectively for the FIDAS and TEOM-FDMS instruments (Figure 2.2b).

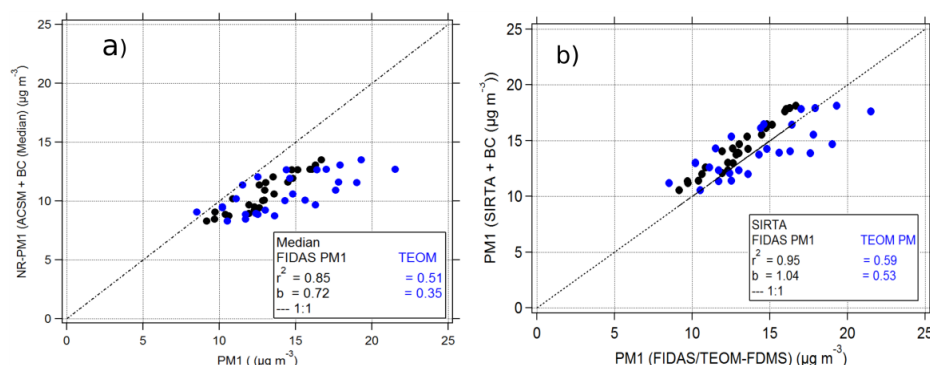


Figure 2.2. Comparison of a) Robust Median NR-PM1 + BC and b) SIRTa NR-PM1 + BC mass concentrations with those of the FIDAS and TEOM PM1 mass concentration.

For this reason, comparison with the SIRTa instrument seems to be more representative of the NR-PM1 than the median of all ACSM instruments. Similar to the observations from the z-score calculation, the LUND measured mass concentrations were much lower than those from both SIRTa and the FIDAS instrument.

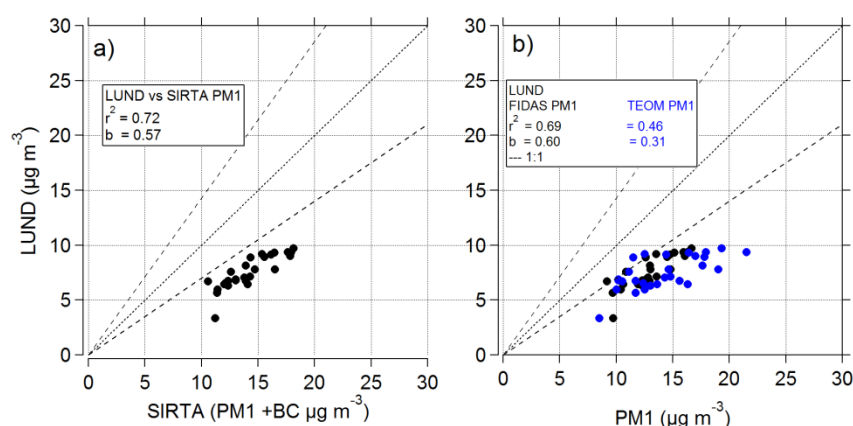
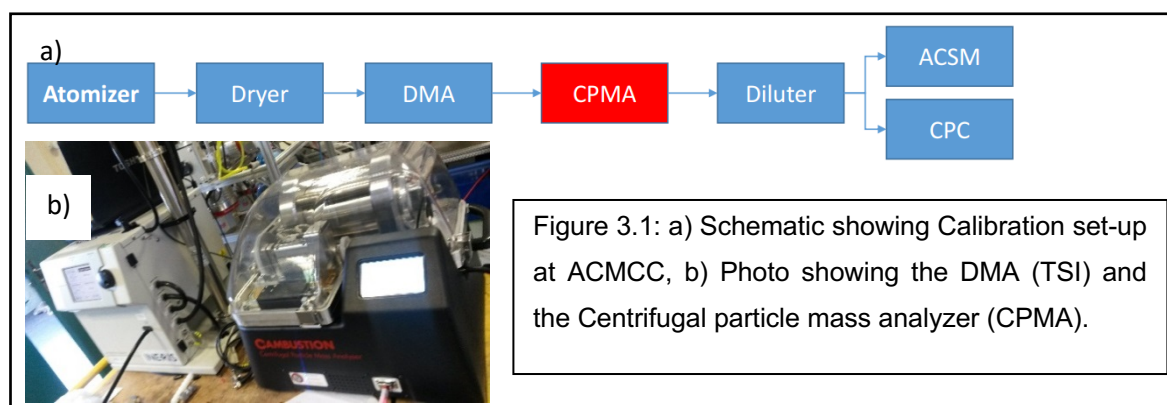


Figure 2.3. Comparison of LUND NR-PM1 + BC with the a) SIRTa NR-PM1 + BC mass concentrations and b) with PM1 mass concentration from FIDAS and TEOM-FDMS PM1 mass concentration.

3. Optimization of (relative) ionization efficiency values

Following initial intercomparison, instruments were checked, maintained and/or tuned (when needed) during the following week (19th to 23rd Nov). A series of calibrations were performed on each individual instrument at ACMCC during and after the workshop, using monodisperse (300 nm) ammonium nitrate and ammonium sulfate nebulized particles. The calibration set up is illustrated in Fig. 3.1.



The calibration set up included a differential mobility analyzer (DMA, TSI®) to select particles of diameter 300 nm. This DMA was calibrated prior to use, using 300 nm polystyrene latex spheres. The aerosol particles were then passed into a Centrifugal Particle Mass Analyser (CPMA) that separates particles by their actual mass and removes doubly charged aerosol particles. These particles are then passed simultaneously into the ACSM and into a condensation particle counter (CPC). This CPC was calibrated prior to the workshop, based on comparison with an electrometer.

The nitrate response factor (RF) was calculated for each instrument and applied to the raw ACSM signal to obtain quantitative information. This value is determined from a known quantity of a known chemical species that enters into the instrument. The relative ionization efficiency (RIE) is a chemical dependent value that is commonly determined experimentally for the ammonium and sulfate species.

Calibration values provided by LUND for initial intercomparison as well as the ones recommended by the ACMCC for the workshop period (based on results from the full scan calibration mode) are listed in Table 3.1. The NH_4 measured vs NH_4 predicted values are illustrated in Figure 3.2, showing poor agreement between the two values with a slope of 0.15 with the new RIE NH_4 value.

LUNDToF-ACSM-#19	RF NO_3	RIE NH_4	RIE SO_4	Ref Airbeam	Vap Temp (°C)
Initial	70	4	1.2	172000	XXX
Calibrated	26.97	2.96	0.88	149578	575

Table 3.1: Calibration values for the LUND instrument for each calibration method.

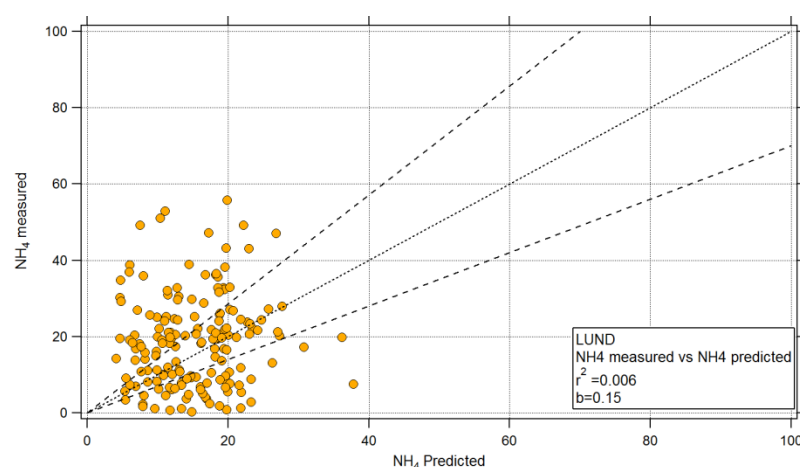


Figure 3.2 NH_4 measured vs NH_4 predicted for the LUND instrument using the calibrated RIE NH_4 values listed in Table 3.2.

It should be noticed here that the LUND instrument was actually influenced by a leak in the three-way valve system, which could only be detected and fully solved after the workshop. It cannot be excluded that this technical issue was due to damage during transport and/or installation.

4. Results from the second intercomparison campaign

A second intercomparison campaign took place from the 23th to the 26th of November, considering calibration results and tuning performed at the ACMCC. The performance of the LUND instrument was again evaluated through comparison with the SIRTA instrument, and later with the robust median of the participating PM1 QACSM instruments. As mentioned earlier, the instrument performance can be considered as acceptable if data points were within $\pm 30\%$ of these reference values. The LUND instrument shows poor temporal agreement with both the median and that of the SIRTA instrument (Fig 4.1), and does not fall within $\pm 30\%$ (Fig 4.2). This is attributed to a leak in the sampling system, as previously mentioned.

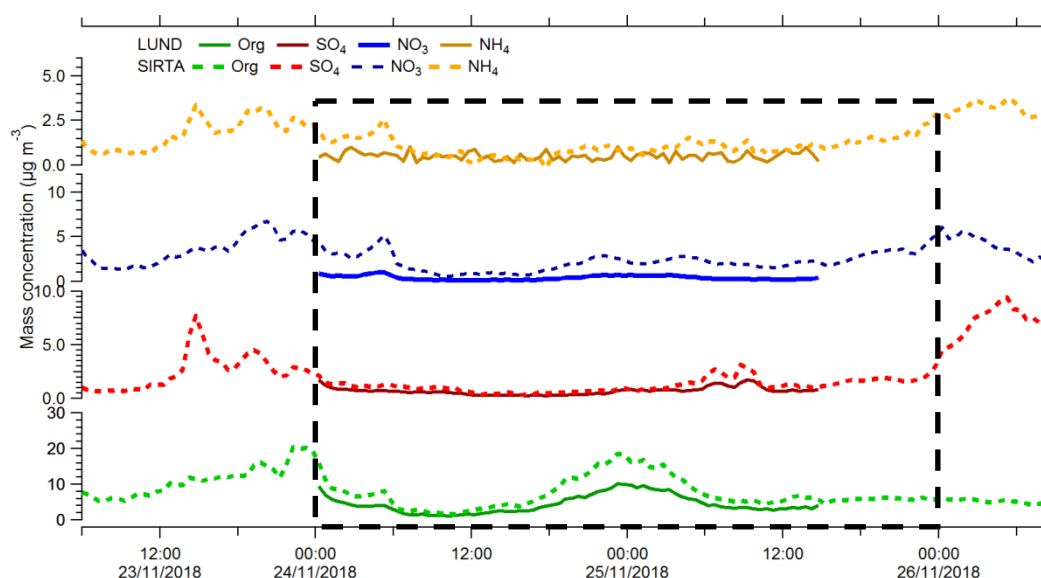


Figure 4.1. Comparison of LUND with SIRTA instrument for Ammonia, Nitrate, sulphate, and organic, during the second intercomparison campaign. The highlighted box indicates the intercomparison period (24/11 00hrs to 26/11 00hrs).

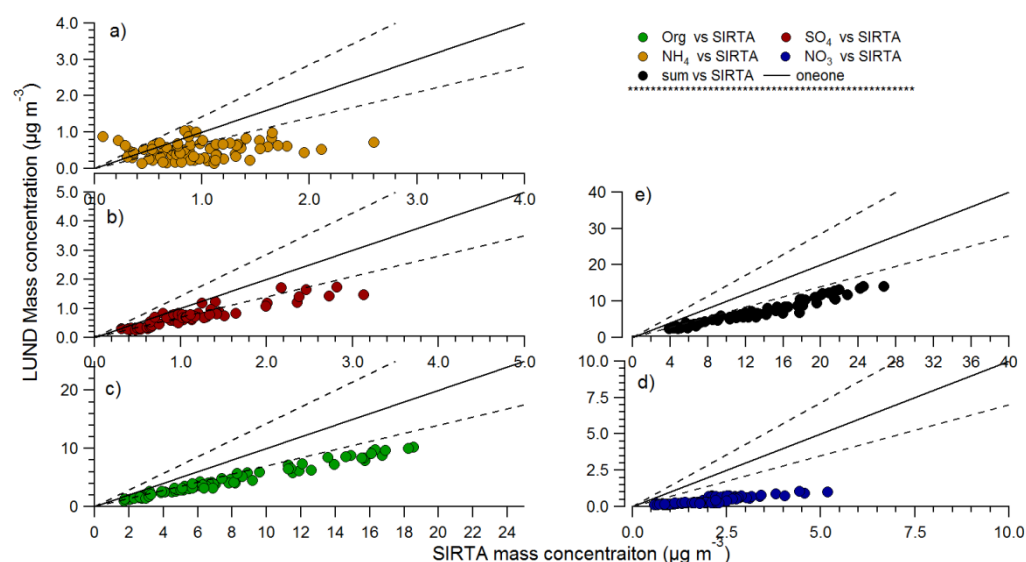


Figure 4.2. Correlations between different chemical species measured by the LUND and the SIRTa instrument during the second intercomparison campaign (24th Nov 00hrs to 26th Nov 00hrs).

The second method of evaluation was made using the Z-score comparison. As described in section 1, instrument performance is considered acceptable when Z-score values fall between 2 and -2 (indicated by the green lines in Figures 4.3). Figure 4.3 shows the z-score calculated for each instrument using the robust median as a reference. The robust median was calculated as the median of all instruments but outliers (#10) and PM_{2.5}-inlet ACSMs (# 14, 15, and 16). The LUND instrument is no. 10 (red rectangle), showing that this instrument was still outside of the acceptable limits of the z-score test for NH₄, NO₃, and therefore also the total PM₁, due to temporary technical issues.

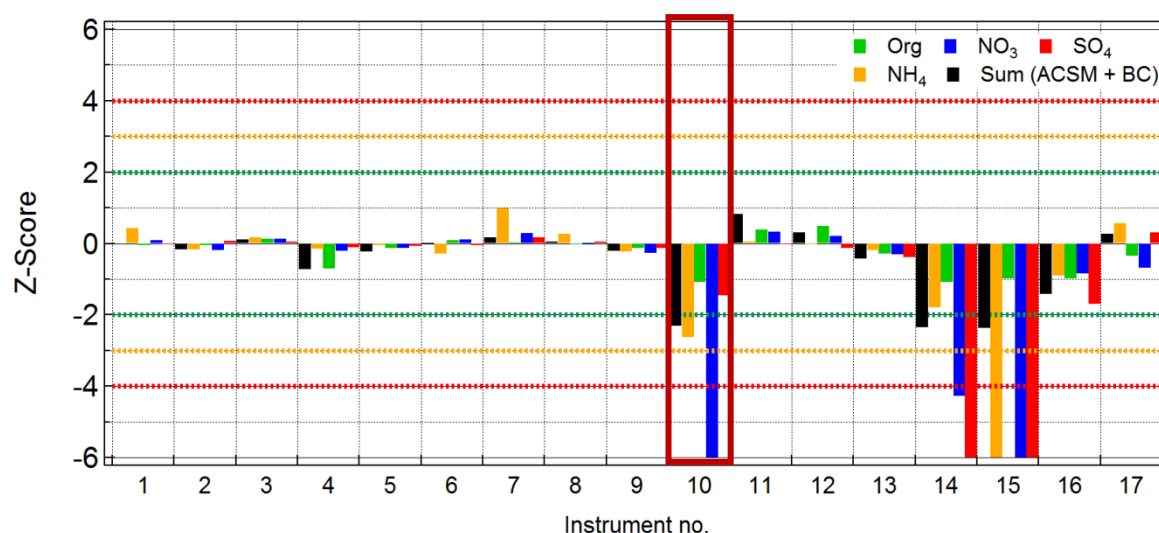


Figure 4.3. Z-score calculation for each species measured by the ACSM for each instrument that participated in the second intercomparison campaign (period 24th Nov 00hrs to 26th Nov 00hrs). The red rectangle highlights the LUND (no.10) instrument.

5. Conclusion

Mass concentrations measured by the LUND instrument during both intercomparison campaigns were lower than those of the reference datasets. This was expected to be due to a leak in the three-way valve system, that could be changed only after the second intercomparison campaign. However, calibrations performed during the workshop seemed to improve at least the RIE SO₄ value.

QA/QC checks and calibrations will be needed before the instrument is re-installed at the monitoring station.

6. References

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- ISO, 5725-5, 1998. Accuracy (trueness and precision) of measurement methods and results-Part 5: Alternative Methods for the Determination of the Precision of a standard measurement method. International Organization for Standardization.
- Middlebrook, A. N. R. Bahreini, J. L. Jimenez, and M. R. Canagaratna (2012) *Aerosol Sci. Tech.*, 46:258–271.