

J R C T E C H N I C A L R E P O R T S

Evaluation of the Laboratory Comparison Exercise for SO₂, CO, O₃, NO and NO₂ 03rd - 06th October 2011

EC Harmonization Program for Air Quality Measurements



Maurizio Barbieri, Friedrich Lagler

2012

European Commission
Joint Research Centre
Institute for Environment and Sustainability

Contact information

Friedrich Lagler

Address: Joint Research Centre, Via Enrico Fermi 2749, TP 442, 21027 Ispra (VA), Italy

E-mail: friedrich.lagler@jrc.ec.europa.eu

Tel.: +39 0332 789990

Fax: +39 0332 789931

<http://www.jrc.ec.europa.eu/>

This publication is a Reference Report by the Joint Research Centre of the European Commission.

Legal Notice

Neither the European Commission nor any person acting on behalf of the Commission is responsible for the use which might be made of this publication.

Europe Direct is a service to help you find answers to your questions about the European Union Freephone number (*): 00 800 6 7 8 9 10 11

(*) Certain mobile telephone operators do not allow access to 00 800 numbers or these calls may be billed.

A great deal of additional information on the European Union is available on the Internet. It can be accessed through the Europa server <http://europa.eu/>.

JRC71998

EUR 25386EN

ISBN 978-92-79-25366-9

ISSN 1831-9424

doi:10.2788/33270

Luxembourg: Publications Office of the European Union, 2012

© European Union, 2012

Reproduction is authorised provided the source is acknowledged.

Printed in Italy

In collaboration with:

Thorsten Zang; Alfred Wagner; Pilar Morillo Gómez; David Martin Bermejo; Fabio Cadoni; Damiano Centioli; Natalija Ivanc; Ales Razpotnik; Premec Krunoslav; Petra Lepri; Tomasz Frączkowski; Andrzej Pindel; Jan Adams; Luk Van Camp; Christos Kizas; Christakis Papadopoulos; Leif Marsteen; Frank Dauge.



WHO Collaborating Centre for Air Quality
Management and Air Pollution Control
at the Federal Environmental Agency



	NAME	VERSION	DATE
AUTHOR	M. BARBIERE	DRAFT 1	15/05/2012
REVIEW	F. LAGLER	DRAFT 2	16/05/2012
REVIEW	EXTERNAL	DRAFT 3	04/06/2012
APPROVAL	N. JENSEN	1.0	11/06/2012

Executive Summary

From the 03rd to the 06th of October 2011 nine Laboratories of AQUILA (Network of European Air Quality Reference Laboratories) met at an laboratory comparison exercise in Ispra (IT) to evaluate their proficiency in the analysis of inorganic gaseous pollutants (SO₂, CO, NO, NO₂ and O₃) covered by the European Air Quality Directive 2008/50/EC.

The proficiency evaluation, where each participant's bias was compared to two criteria, provides information on compliance with Data Quality Objectives and measurement capabilities of the National Air Quality Laboratories to the European Commission and can be used by participants in their laboratory's quality system.

On the basis of criteria imposed by the European Directive, 79% of the results reported by AQUILA laboratories were good both in terms of measured values and reported uncertainties. Another 20% of the results had good measured values, but the reported uncertainties were either too high (12%) or too small (8%). A small number of values (1%) were questionable and the uncertainties "not ok".

The comparability of results among AQUILA participants at the highest generated concentration levels, excluding outliers, is acceptable for SO₂, CO and O₃ measurements while NO and NO₂ measurements showed less satisfactory results.

Contents

1. Introduction	9
1.1 Communication and time schedule	11
1.2 Participants	11
1.3 The preparation of test mixtures	13
2. The evaluation of laboratory's measurement proficiency.....	14
2.1 z' - score	14
2.2 E _n - number.....	18
3. Performance characteristics of individual laboratories	24
3.1 The efficiency of NO ₂ -to-NO converters of NO _x analyzers	24
4. Discussion	26
5. Conclusions	28
6. References	30
Annex A. Assigned values.....	32
Annex B. The results of the IE	35
Reported values for SO ₂	35
Reported values for CO	39
Reported values for O ₃	42
Reported values for NO	45
Reported values for NO ₂	51
Annex C. The precision of standardized measurement methods	54
Annex D. The scrutiny of results for consistency and outlier test	60

List of tables

Table 1: The list of participating organizations.	11
Table 2: The list of instruments used by participants.	12
Table 3: The sequence program of generated test gases	13
Table 4: The standard deviation for proficiency assessment (σ_p).	14
Table 5: The efficiency of NO ₂ -to-NO converters.	25
Table 6: The general assessment of proficiency results.	27
Table 7: Flags summary	28
Table 8: Z'-score summary	28
Table 9: The validation of assigned values (X)	33
Table 10: Reported values for SO ₂ run 0.	35
Table 11: Reported values for SO ₂ run 1.	36
Table 12: Reported values for SO ₂ run 2.	36
Table 13: Reported values for SO ₂ run 3.	37
Table 14: Reported values for SO ₂ run 4.	37
Table 15: Reported values for SO ₂ run 5.	38
Table 16: Reported values for CO run 0.	39
Table 17: Reported values for CO run 1.	39
Table 18: Reported values for CO run 2.	40
Table 19: Reported values for CO run 3.	40
Table 20: Reported values for CO run 4.	41
Table 21: Reported values for CO run 5.	41
Table 22: Reported values for O ₃ run 0.	42
Table 23: Reported values for O ₃ run 1	42
Table 24: Reported values for O ₃ run 2.	43
Table 25: Reported values for O ₃ run 3.	43
Table 26: Reported values for O ₃ run 4.	44
Table 27: Reported values for O ₃ run 5.	44
Table 28: Reported values for NO run 0.	45
Table 29: Reported values for NO run 1.	45
Table 30: Reported values for NO run 2.	46
Table 31: Reported values for NO run 3.	46
Table 32: Reported values for NO run 4.	47
Table 33: Reported values for NO run 5.	47
Table 34: Reported values for NO run 6.	48
Table 35: Reported values for NO run 7.	48
Table 36: Reported values for NO run 8.	49
Table 37: Reported values for NO run 9.	49
Table 38: Reported values for NO run 10.	50
Table 39: Reported values for NO ₂ run 0.	51
Table 40: Reported values for NO ₂ run 2.	51
Table 41: Reported values for NO ₂ run 4.	52
Table 42: Reported values for NO ₂ run 6.	52
Table 43: Reported values for NO ₂ run 8.	53
Table 44: Reported values for NO ₂ run 10.	53
Table 45: Critical values of t used in the repeatability (r) and reproducibility (R) evaluation.	54
Table 46: The R and r of SO ₂ standard measurement method.	55
Table 47: The R and r of CO standard measurement method.	56
Table 48: The R and r of O ₃ standard measurement method.	57
Table 49: The R and r of NO standard measurement method.	58
Table 50: The R and r of NO ₂ standard measurement method.	59
Table 51: "Genuine" statistical outliers according to Grubb's one outlying observation test.	60

List of figures

Figure 1: The z'-score evaluations of SO ₂ measurements	15
Figure 2: The z'-score evaluations of CO measurements	16
Figure 3: The z'-score evaluations of O ₃ measurements	16
Figure 4: The z'-score evaluations of NO measurements	17
Figure 5: The z'-score evaluations of NO ₂ measurements	17
Figure 6: Bias of participant's SO ₂ measurement results	19
Figure 7: Bias of participant's CO measurement results	20
Figure 8: Bias of participant's O ₃ measurement results	21
Figure 9: Bias of participant's NO measurement results	22
Figure 10: Bias of participant's NO ₂ measurement results	23
Figure 11: Bias of participant's NO ₂ measurements for run numbers 1, 3, 5, 7 and 9	24
Figure 12: The decision diagram for general assessment of proficiency results.	26
Figure 13: Reported values for SO ₂ run 0.	35
Figure 14: Reported values for SO ₂ run 1.	36
Figure 15: Reported values for SO ₂ run 2.	36
Figure 16: Reported values for SO ₂ run 3.	37
Figure 17: Reported values for SO ₂ run 4.	37
Figure 18: Reported values for SO ₂ run 5.	38
Figure 19: Reported values for CO run 0.	39
Figure 20: Reported values for CO run 1.	39
Figure 21: Reported values for CO run 2.	40
Figure 22: Reported values for CO run 3.	40
Figure 23: Reported values for CO run 4.	41
Figure 24: Reported values for CO run 5.	41
Figure 25: Reported values for O ₃ run 0.	42
Figure 26: Reported values for O ₃ run 1.	42
Figure 27: Reported values for O ₃ run 2.	43
Figure 28: Reported values for O ₃ run 3.	43
Figure 29: Reported values for O ₃ run 4.	44
Figure 30: Reported values for O ₃ run 5.	44
Figure 31: Reported values for NO run 0.	45
Figure 32: Reported values for NO run 1.	45
Figure 33: Reported values for NO run 2.	46
Figure 34: Reported values for NO run 3.	46
Figure 35: Reported values for NO run 4.	47
Figure 36: Reported values for NO run 5.	47
Figure 37: Reported values for NO run 6.	48
Figure 38: Reported values for NO run 7.	48
Figure 39: Reported values for NO run 8.	49
Figure 40: Reported values for NO run 9.	49
Figure 41: Reported values for NO run 10.	50
Figure 42: Reported values for NO ₂ run 0.	51
Figure 43: Reported values for NO ₂ run 2.	51
Figure 44: Reported values for NO ₂ run 4.	52
Figure 45: Reported values for NO ₂ run 6.	52
Figure 46: Reported values for NO ₂ run 8.	53
Figure 47: Reported values for NO ₂ run 10.	53
Figure 48: The R and r of SO ₂ standard measurement method as a function of concentration.	55
Figure 49: The R and r of CO standard measurement method as a function of concentration.	56
Figure 50: The R and r of O ₃ standard measurement method as a function of concentration.	57
Figure 51: The R and r of NO standard measurement method as a function of concentration.	58
Figure 52: The R and r of NO ₂ standard measurement method as a function of concentration.	59

Abbreviations

AQUILA	Network of National Reference Laboratories for Air Quality
CO	Carbon monoxide
DQO	Data Quality Objective
ERLAP	European Reference Laboratory of Air Pollution
EC	European Commission
GPT	Gas Phase Titration
IE	Inter-laboratory Comparison Exercise
IES	Institute for Environment and Sustainability
ISO	International Organization for Standardization
JRC	Joint Research Centre
NO	Nitrogen monoxide
NO ₂	Nitrogen dioxide
NO _x	the oxides of nitrogen, the sum of NO and NO ₂
NRL	National Reference Laboratory
O ₃	Ozone
SO ₂	Sulphur dioxide
WHO	World Health Organization Collaborating Centre for Air Quality
CC-EURO	Management and Air Pollution Control, Berlin

Mathematical Symbols

symbol explanation

α	converter efficiency (EN 14211; [4])
E_n	E_n – number statistic (ISO 13528; [13])
r	repeatability limit (ISO 5725; [14])
R	reproducibility limit (ISO 5725; [14])
σ_p	standard deviation for proficiency assessment (ISO 13528; [13])
x^*	robust average (Annex C ISO 13528; [13])
s^*	robust standard deviation (Annex C ISO 13528; [13])
s_r	repeatability standard deviation (ISO 5725; [14])
s_R	reproducibility standard deviation (ISO 5725; [14])
U_X	expanded uncertainty of the assigned/reference value (ISO 13528; [13])
U_{xi}	expanded uncertainty of the participant's value
u_X	standard uncertainty of the assigned/reference value (ISO 13528; [13])
X	assigned/reference value (ISO 13528; [13])
x_i	average of three values reported by the participant i (for particular parameter and concentration level) (ISO 5725; [14])
$x_{i,j}$	j -the reported value of participant i (for particular parameter and concentration level) (ISO 5725; [14])
z'	z' -score statistic (ISO 13528; [13])

1. Introduction

As a result of the revision of the legislation framework on air quality in the CAFÉ (Clean Air for Europe) thematic strategy, former mother and most daughter directives were integrated into a single rule. With the adoption of Directive 2008/50/EC [1] on ambient air quality and cleaner air for Europe, a framework for a harmonized air quality assessment in Europe was set. One important objective of the Directive is that the ambient air quality shall be assessed on the basis of common methods and criteria. It deals with the air pollutants sulphur dioxide (SO₂), nitrogen dioxide (NO₂) and monoxide (NO), particulate matter, lead, benzene, carbon monoxide (CO) and ozone (O₃). Among others it specifies the reference methods for measurements and Data Quality Objectives (DQOs) for the accuracy of measurements.

The European Commission (EC) has supported the development and publication of reference measurement methods for CO [2], SO₂ [3], NO-NO₂ [4] and O₃ [5] as European standards. Appropriate calibration methods [6], [7] and [8] have been standardized by the International Organization for Standardization (ISO).

As foreseen in the Air Quality Directive, the European Reference Laboratory of Air Pollution (ERLAP) of the Institute for Environment and Sustainability (IES) at the Joint Research Centre (JRC) organizes inter-laboratory comparison exercises (IE) to assess and improve the status of comparability of measurements of National Reference Laboratories (NRL) of each Member State of the European Union.

The World Health Organization Collaborating Centre for Air Quality Management and Air Pollution Control, Berlin (WHO CC) is carrying out similar activities since 1994 [9] [10], but with a view to obtaining harmonized air quality data for health related studies. Their program integrates within the WHO EURO region, which includes public health institutes and other national institutes - especially from the Central Eastern Europe, Caucasus and countries from Central Asia.

Starting in 2004, it has been decided to bring together the efforts of both the JRC-ERLAP and WHO CC and to coordinate activities as far as possible, with a view to optimize resources and have better international harmonization. The following report deals with the IE that took place from 03rd to the 06th of October 2011 in Ispra (IT) in joint cooperation of EC/ JRC/IES/ERLAP and WHO CC.

Since 1990 ERLAP organizes IE aiming at evaluating the comparability of measurements carried out by NRLs and promoting information exchange among the expert laboratories. Currently, a more systematic approach has been adopted, in accordance with the Network of National Reference Laboratories for Air Quality (AQUILA) [11], aiming both at providing an alert mechanism for the purposes of the EC legislation and at supporting the implementation of quality schemes by NRLs. The methodology for the organization of IE was developed by ERLAP in collaboration with AQUILA and is described in a paper on the organization of laboratory comparison exercises for gaseous air pollutants [12].

This evaluation scheme was adopted in December 2008 and is applied to all IE since then. It contains common criteria to alert the EC on possible performance failures which do not rely solely on the uncertainty claimed by participants. The evaluation scheme implements the z'-score method [13] with the uncertainty requirements for calibration gases stated in the European standards [2], [3], [4] and [5], which are consistent with the DQOs of European Directives.

According to the said document, NRLs with an overall unsatisfactory performance in the z'-score evaluation (one unsatisfactory or two questionable results per parameter) ought to repeat their participation in the following IE in order to demonstrate remediation measures [12]. In addition, considering that the evaluation scheme should be useful to participants for accreditation according to ISO 17025, they are requested to include their measurement

uncertainty. Hence, participants' results (measurement values and uncertainties) are compared to the assigned values applying the E_n – number method [13]. Beside the proficiency of participating laboratories, the repeatability and reproducibility of standardized measurement methods [14], [15] and [16] are evaluated as well. These group evaluations are useful indicators of trends in measurement quality over different IE.

1.1 Communication and time schedule

The IE was announced in March 2011 to the members of the AQUILA network and the WHO CC representative. Registration was opened on March 2011 and due to the number of request ERLAP decided to organize two consecutive sessions of IE exercises.

The participants were required to bring their own measurement instruments, data acquisition equipment and travelling standards (to be used for calibrations or checks during the IE).

The participants were invited to arrive on Monday, 03rd October 2011, for the installation of their equipment. The calibration of NO_x and O₃ analysers was carried out on Tuesday morning and the generation of NO_x and O₃ gas mixtures started at 11:00. The calibration of SO₂ and CO analysers was carried out on Wednesday 18:00 and the generation of CO and SO₂ gas mixtures started at 20:00. The test gases generation finished on Thursday at 9:30.

1.2 Participants

All participants were organizations dealing with the routine ambient air monitoring or institutions involved in public health protection. The national representatives came from EU member states: Germany, Spain, Italy, Slovenia, Croatia, Poland, Belgium, Cyprus and Norway.

Country	Laboratory	Code
Germany	Landesamt für Natur, Umwelt und Verbraucherschutz (LANUV)	A
Spain	Instituto De Salud Carlos III (ISCIII)	B
Italy	Italian National Institute for Environmental Protection and Research (ISPRA)	C
Slovenia	Slovenian Environment Agency (SEA)	D
Croatia	Meteorological and Hydrological Service (DHMZ)	E
Poland	Chief Inspectorate of Environmental Protection (GIOS)	F
European Commission	European Reference Laboratory of Air Pollution (ERLAP)	G
Belgium	Flemish Environment Agency (VMM)	H
Cyprus	Dept. of Labour Inspection (DLI)	I
Norway	Norwegian Institute for Air Research (NILU)	L

Table 1: The list of participating organizations.

In Table 2 are reported the manufacturer and model of the instrumentation used by every participant during the inter-laboratory comparison exercise included those used in the calculation of the assigned values.

As a whole, the instrumentation belongs to 5 different manufacturers for NO_x, 4 for CO and SO₂, and 3 brands are present for O₃. The list contains the information reported by participants and by no means can be considered as an implicit or explicit endorsement of the organizers to any specific type of instrumentation.

Gas	Lab Code	Instrument
SO ₂	A	Teledyne API 200E and Ansyco AF 21 M
	B	API100E, 2010
	C	
	D	Horiba, 2002, APSA 360 A
	E	Horiba, 2010, APSA-370
	F	Thermo 43C, 2004
	G	Thermo Electrom Corporation, 2009, 43i
	H	Thermo Scientific 43i 2010
	I	Ecotech Australia / 2005 / EC 9850B
	L	API, 2005, 100E
NO _x	A	ECO Physics CLD 700 AL and Ansyco AC 32 M
	B	API 200E, 2010
	C	Thermo Electron Corporation 42i
	D	Horiba, 2010, APNA 370
	E	
	F	Thermo 42C, 2004
	G	Thermo Electrom Corporation, 2010, 42i
	H	Thermo Scientific 42i , 2011
	I	Ecotech Australia / 2005 / EC 9841B
	L	API, 2005, 200E
CO	A	TE 49 i and Ansyco CO 11 M
	B	API300E, 2006
	C	Thermo Electron Corporation 48i
	D	Horiba, 2002, APMA 360 CE
	E	Horiba, 2010, APMA-370
	F	Thermo 48C, 2004
	G	Thermo Electronic Corporation, 2000, 48C
	H	API 300, 2001
	I	Ecotech Australia / 2005 / EC 9830B
	L	API, 2005, 300E
O ₃	A	Horiba APOA 370 and Ansyco O3 41 M
	B	API 400E, 2008
	C	Thermo Electron Corporation 49i
	D	TEI, 2002, O3 analyser 49C
	E	
	F	Thermo 49C , 2004
	G	Thermo Electronic Corporation, 1996, 49C
	H	Thermo Scientific 49i 2010
	I	Thermo / 2005 / 49i
	L	API, 2000, 400

Table 2: The list of instruments used by participants.

1.3 The preparation of test mixtures

The ERLAP IE facility has been described in several reports [17] and [18]. During this IE, gas mixtures were prepared for SO₂, CO, O₃, NO and NO₂ at concentration levels around limit values, critical levels and assessment thresholds set by European Air Quality Directive [1].

The test mixtures were prepared by the dilution of gases from cylinders containing high concentration of NO, SO₂ or CO using thermal mass flow controllers [8]. O₃ was added using an ozone generator and NO₂ was produced applying the gas phase titration method [19] in a condition of NO excess.

The participants were required to report three half-hour-mean measurements for each concentration level (run) in order to evaluate the repeatability of standardized measurement methods. Zero concentration levels were generated for one hour and one half-hour-mean measurement was reported. The sequence program of generated test gases is given in Table 3.

day	start time	duration	parameter	installation	calibration	Zero Air	NO	NO2	O3	CO	SO2
		h				nmol/mol	nmol/mol	nmol/mol	nmol/mol	mmol/mol	nmol/mol
3-Oct	12:00	5	/	X							
4-Oct	8:00	3	/		X						
4-Oct	11:00	1	NO-NO2-O3			0					
4-Oct	12:00	2	NO-NO2				520				
4-Oct	14:00	2	NO-NO2				390	130			
4-Oct	16:00	2	O3						130		
4-Oct	18:00	2	NO-NO2				60				
4-Oct	20:00	2	NO-NO2				35	25			
4-Oct	22:00	2	O3						25		
5-Oct	0:00	2	NO-NO2				175				
5-Oct	2:00	2	NO-NO2				120	55			
5-Oct	4:00	2	O3						55		
5-Oct	6:00	2	NO-NO2				260				
5-Oct	8:00	2	NO-NO2				165	95			
5-Oct	10:00	2	O3						95		
5-Oct	12:00	2	NO-NO2				20				
5-Oct	14:00	2	NO-NO2				6	14			
5-Oct	16:00	2	O3						14		
5-Oct	< 18:00	2	calibration		X						
5-Oct	20:00	1	CO-SO2			0					
5-Oct	21:00	2:30	CO-SO2							8	8
5-Oct	23:30	2	CO-SO2							4,5	50
6-Oct	1:30	1	CO-SO2	Zero Air not reported						0	0
6-Oct	2:30	2	CO-SO2							6	20
6-Oct	4:30	2	CO-SO2							3	120
6-Oct	6:30	2	CO-SO2							1	3
6-Oct	8:30	1				0					
6-Oct	9:30		END								

Table 3: The sequence program of generated test gases

2. The evaluation of laboratory's measurement proficiency

To evaluate the participants measurement proficiency the methodology described in ISO 13528 [13] was applied. It has been agreed among the AQUILA members to take the measurement results of ERLAP as the assigned/reference values for the whole IE [12]. The traceability of ERLAP's measurement results and the method applied to validate them are presented in Annex A. In the following proficiency evaluations, the uncertainty of test gas homogeneity (Annex A) was added to the uncertainties of ERLAP's measurement results.

All data reported by participating laboratories are presented in Annex B.

As it is described in the said position paper [12], the proficiency of the participants was assessed by calculating two performance indicators. The first performance indicator (z'-score) tests whether the difference between the participants measured value and the assigned/reference value remains within the limits of a common criterion. The second performance indicator (E_n-number) tests if the difference between the participants measured values and assigned/reference value remains within the limits of a criterion, that is calculated individually for each participant, from the uncertainty of the participants measurement result and the uncertainty of the assigned/reference value.

2.1 z' - score

The z'- score statistic is calculated according to ISO 13528 [13] as:

$$z' = \frac{x_i - X}{\sqrt{\sigma_p^2 + u_x^2}} = \frac{x_i - X}{\sqrt{(a \cdot X + b)^2 + u_x^2}} \quad \text{Equation 1}$$

where 'x_i' is a participant's run average value, 'X' is the assigned/reference value, 'σ_p' is the 'standard deviation for proficiency assessment' and 'u_x' is the standard uncertainty of assigned value. For 'a' and 'b' see Table 4.

In the European standards [2], [3], [4] and [5] the uncertainties for calibration gases used in ongoing quality control are prescribed. In fact, it is stated that the maximum permitted expanded uncertainty for calibration gases is 5% and that 'zero gas' shall not give instrument reading higher than the detection limit. As one of the tasks of NRLs is to supply calibration gas mixtures, the 'standard deviation for proficiency assessment' (σ_p) [13] is calculated in fitness-for-purpose manner from requirements given in European standards. Over the whole measurement range σ_p is calculated by linear interpolation between 2.5% at the calibration point (75% of calibration range) and the limit of detection at zero concentration level. The limits of detection of studied measurement methods were evaluated from the data of previous IE. The linear function parameters of σ_p are given in Table 4:

Gas	σ _p =a·c+b	
	a	b
		nmol/mol
SO ₂	0.022	1
CO	0.024	100
O ₃	0.020	1
NO	0.024	1
NO ₂	0.020	1

Table 4: The standard deviation for proficiency assessment (σ_p).

σ_p is a linear function of concentration (c) with parameters: slope (a) and intercept (b).

The assessment of results in the z'-score evaluation is made according to the following criteria:

- $|z'| \leq 2$ are considered satisfactory.
- $2 < |z'| \leq 3$ are considered questionable.
- $|z'| > 3$ are considered unsatisfactory. Scores falling in this range are very unusual and are taken as evidence that an anomaly has occurred that should be investigated and corrected.

The results of z'-score evaluation are presented in bar plots (Figure 1 to Figure 5) in which the z'-scores of each participant are grouped together, and assessment criteria are presented as $z'=\pm 2$ and $z'=\pm 3$ lines.

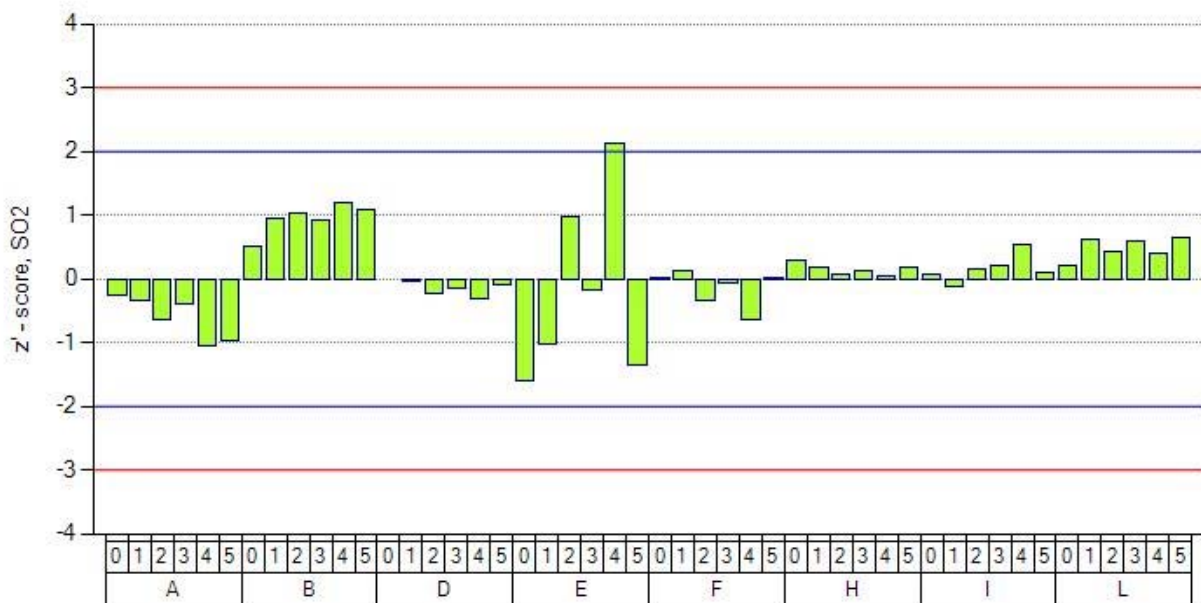


Figure 1: The z'-score evaluations of SO₂ measurements

Scores are given for each participant and each tested concentration level (run). Run number order (with nominal concentration) is: 0 (0 nmol/mol), 1 (8 nmol/mol), 2 (50 nmol/mol), 3 (20 nmol/mol), 4 (120 nmol/mol), 5 (3 nmol/mol). The assessment criteria are presented as $z'=\pm 2$ (blue line) and $z'=\pm 3$ (red line). They represent the limits for the questionable and unsatisfactory results.



Figure 2: The z'-score evaluations of CO measurements

Scores are given for each participant and each tested concentration level (run). Run number order (with nominal concentration) is: 0 (0 µmol/mol), 1 (8 µmol/mol), 2 (4.5 µmol/mol), 3 (6 µmol/mol), 4 (3 µmol/mol), 5 (1 µmol/mol). The assessment criteria are presented as $z'=\pm 2$ (blue line) and $z'=\pm 3$ (red line). They represent the limits for the questionable and unsatisfactory results.

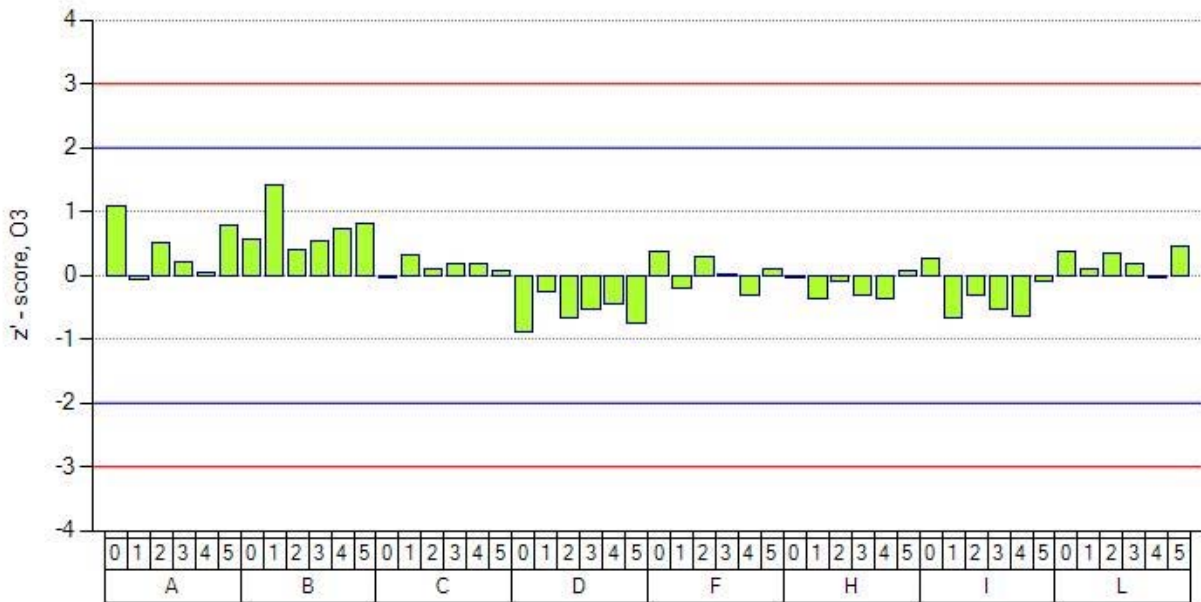


Figure 3: The z'-score evaluations of O₃ measurements

Scores are given for each participant and each concentration level (run). Run number order (with nominal concentration) is: 0 (0 nmol/mol), 1 (130 nmol/mol), 2 (25 nmol/mol), 3 (55 nmol/mol), 4 (95 nmol/mol), 5 (14 nmol/mol). The assessment criteria are presented as $z'=\pm 2$ (blue line) and $z'=\pm 3$ (red line). They represent the limits for the questionable and unsatisfactory results.

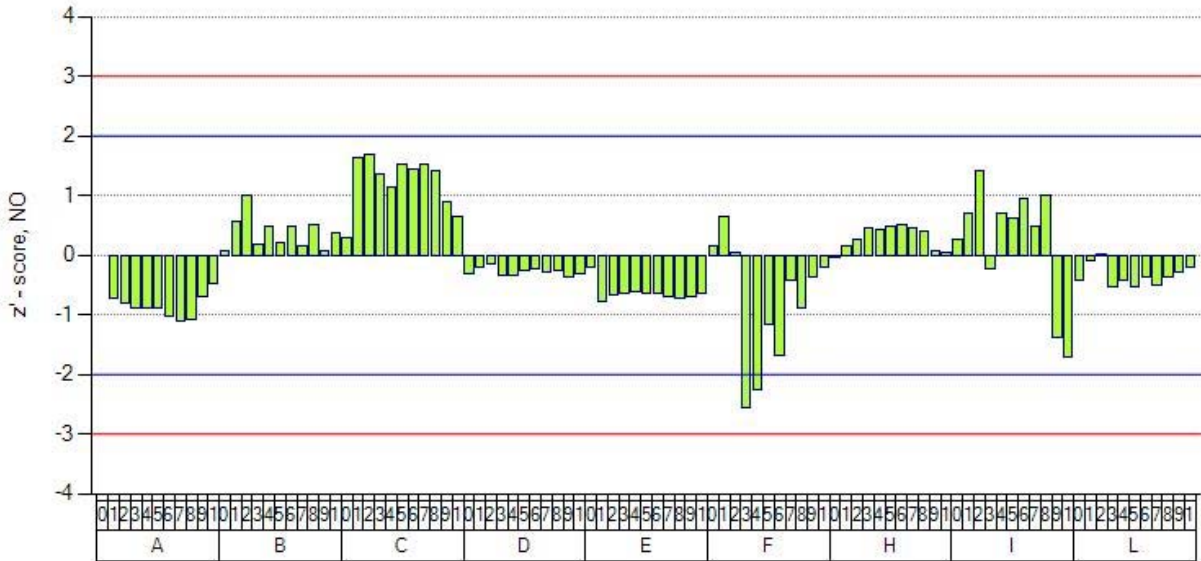


Figure 4: The z'-score evaluations of NO measurements

Scores are given for each participant and each tested concentration level (run). Run number order (with nominal concentration) is: 0 (0 nmol/mol), 1 (520 nmol/mol), 2 (390 nmol/mol), 3 (60 nmol/mol), 4 (35 nmol/mol), 5 (175 nmol/mol), 6 (120 nmol/mol), 7 (260 nmol/mol), 8 (165 nmol/mol), 9 (20 nmol/mol), 10 (6 nmol/mol). The assessment criteria are presented as $z'=\pm 2$ (blue line) and $z'=\pm 3$ (red line). They represent the limits for the questionable and unsatisfactory results.

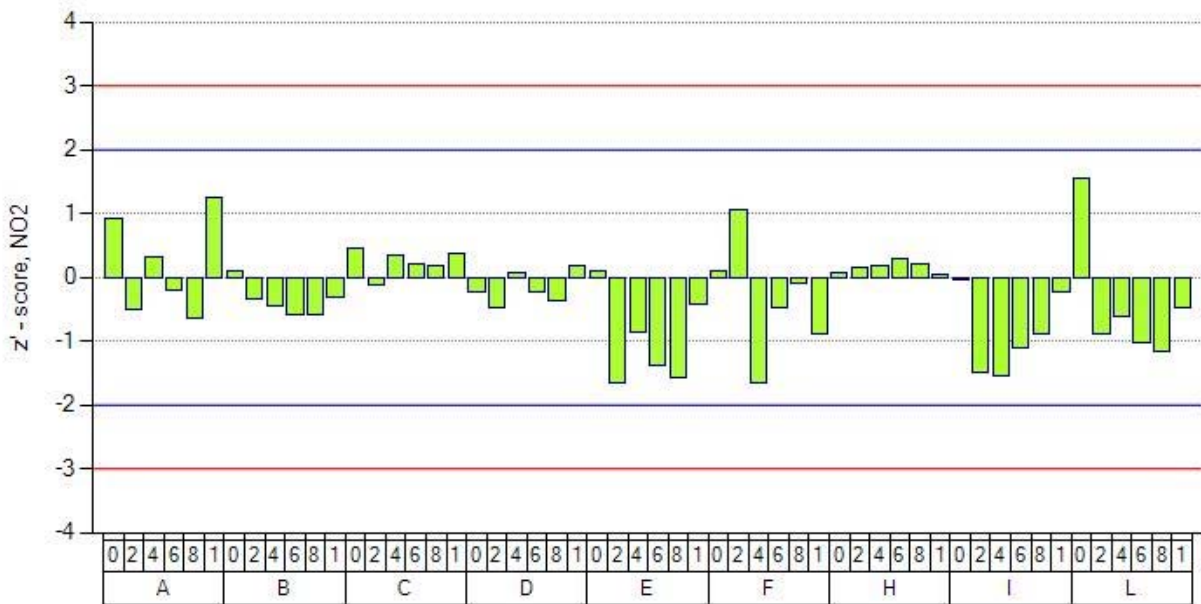


Figure 5: The z'-score evaluations of NO₂ measurements

Scores are given for each participant and each concentration level (run). Run number order (with nominal concentration) is: 0 (0 nmol/mol), 1 (130 nmol/mol), 2 (25 nmol/mol), 3 (55 nmol/mol), 4 (95 nmol/mol), 5 (14 nmol/mol). The assessment criteria are presented as $z'=\pm 2$ (blue line) and $z'=\pm 3$ (red line). They represent the limits for the questionable and unsatisfactory results.

2.2 E_n - number

The normalized deviations [13] (E_n) were calculated according to:

$$E_n = \frac{x_i - X}{\sqrt{U_{x_i}^2 + U_X^2}} \quad \text{Equation 2}$$

where 'X' is the assigned/reference value with an expanded uncertainty 'U_X' and 'x_i' is the participant's average value with an expanded uncertainty 'U_{x_i}'. Satisfactory results are the ones for which $|E_n| \leq 1$.

In Figure 6 to Figure 10 the bias of each participant (x_i-X) are plotted and error bars are used to show the value of denominator of equation 2 ($\sqrt{U_{x_i}^2 + U_X^2}$). These plots represent also the E_n-number evaluations where, considering the E_n criteria ($|E_n| \leq 1$), all results with error bars touching or crossing x-axis are satisfactory. Reported standard uncertainties (Annex B) that are bigger than "standard deviation for proficiency assessments" (σ_p, Table 4) are considered not fit-for-purpose and are denoted with "*" in the x-axis of each figure.

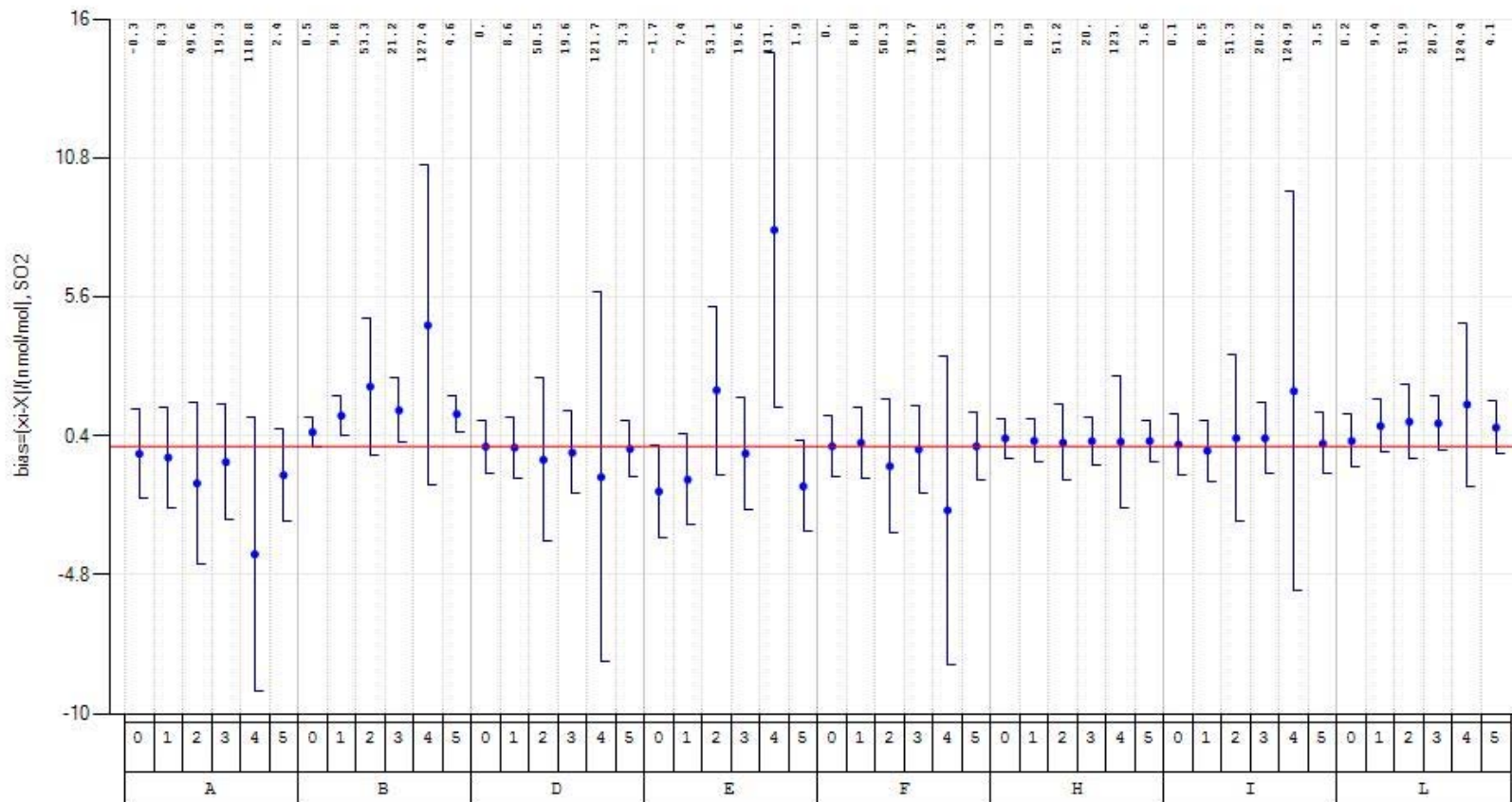


Figure 6: Bias of participant's SO₂ measurement results

Expanded uncertainty of bias for each run is presented as error bar. The results with error bars touching or crossing the x-axis are satisfactory. For each evaluation the run number (numbers 0 to 5) together with the participants rounded run average (nmol/mol) is given. The '*' mark indicates reported standard uncertainties bigger than σ_p .

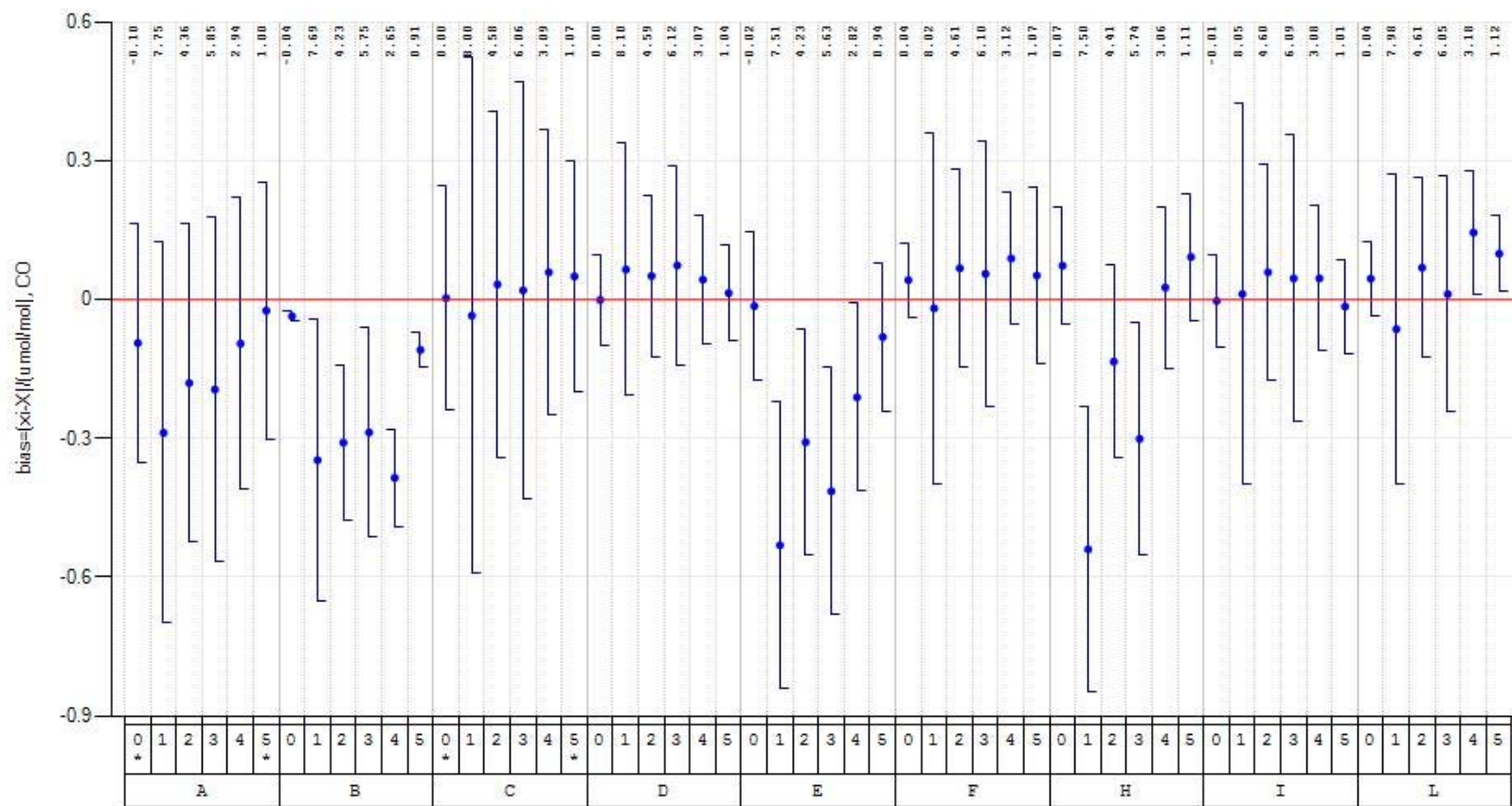


Figure 7: Bias of participant's CO measurement results

Expanded uncertainty of bias for each run is presented as error bar. Results with error bars touching or crossing the x-axis are satisfactory. For each evaluation the run number (numbers 0 to 5) together with the participants rounded run average (µmol/mol) is given. The '*' mark indicates reported standard uncertainties bigger than σ_p .

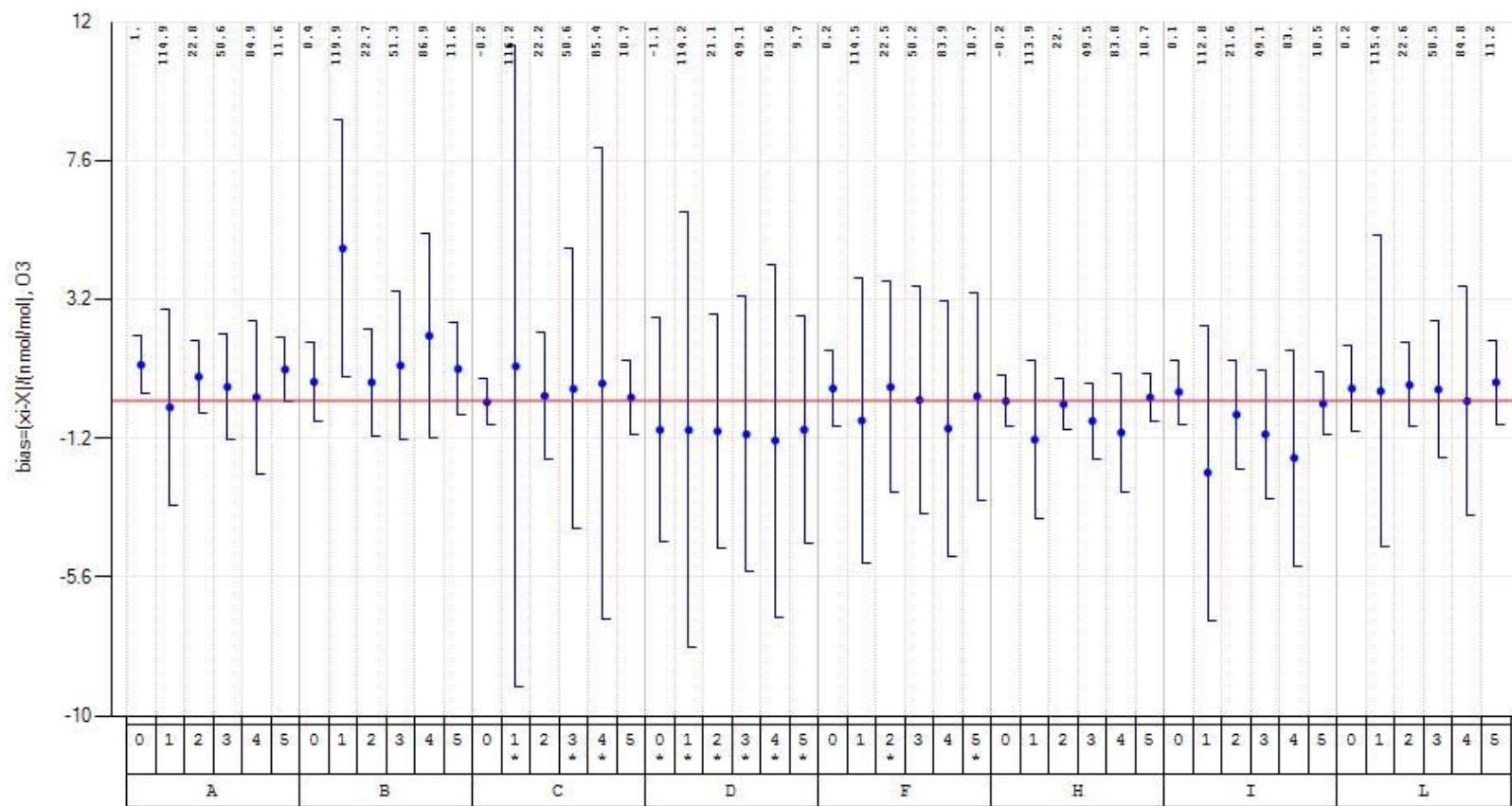


Figure 8: Bias of participant's O₃ measurement results

Expanded uncertainty of bias for each run is presented as error bar. Results with error bars touching or crossing the x-axis are satisfactory. For each evaluation the run number (numbers 0 to 5) together with the participants rounded run average (nmol/mol) is given. The '*' mark indicates reported standard uncertainties bigger than σ_p .

EC harmonization program for Air Quality Measurement
 Evaluation of the Laboratory Comparison Exercise for SO₂, CO, O₃, NO and NO₂, 03rd -06th October 2011

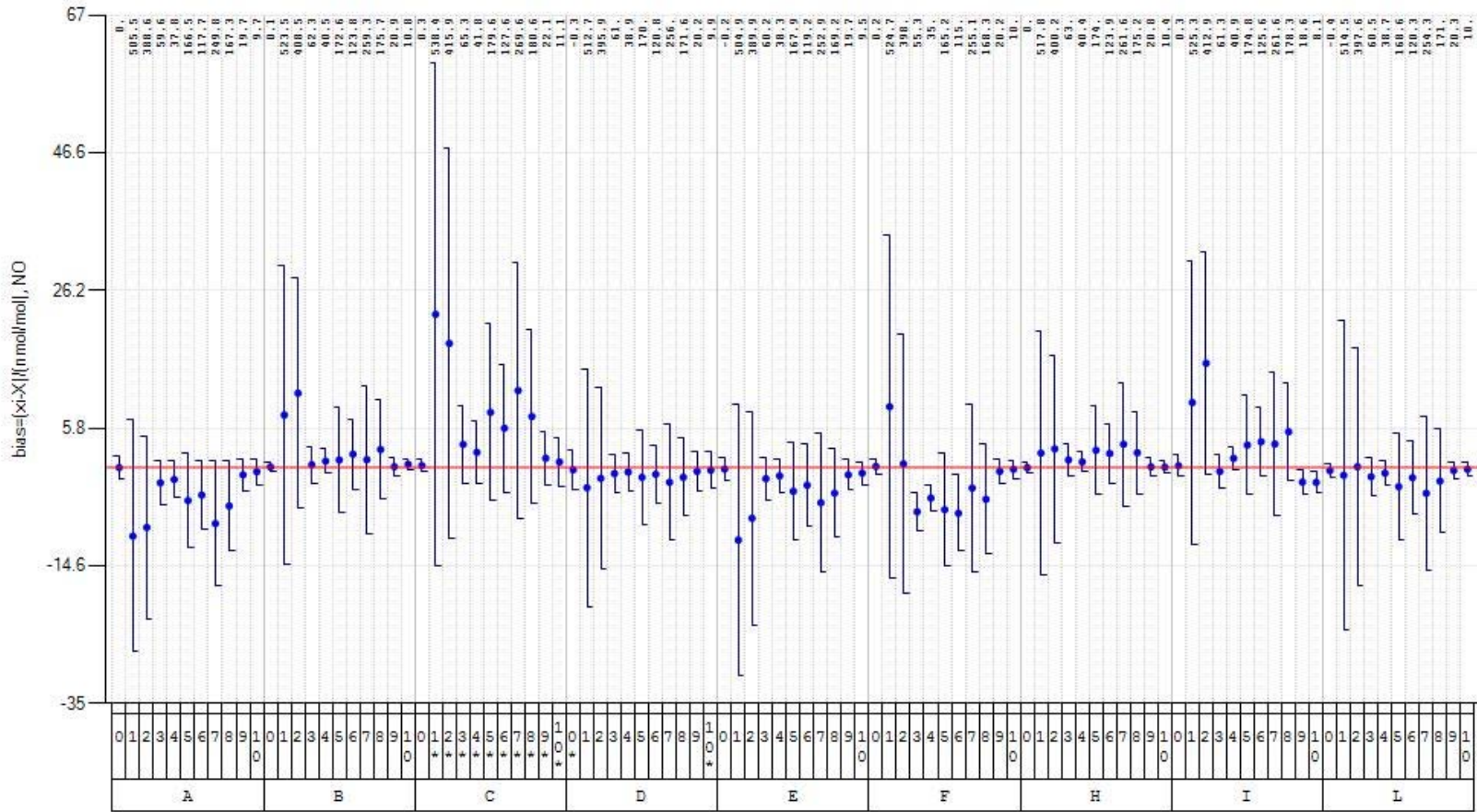


Figure 9: Bias of participant's NO measurement results

Expanded uncertainty of bias for each run is presented as error bar. Results with error bars touching or crossing the x-axis are satisfactory. For each evaluation the run number (numbers 0 to 10) together with the participants rounded run average (nmol/mol) is given. The '*' mark indicates reported standard uncertainties bigger than σ .

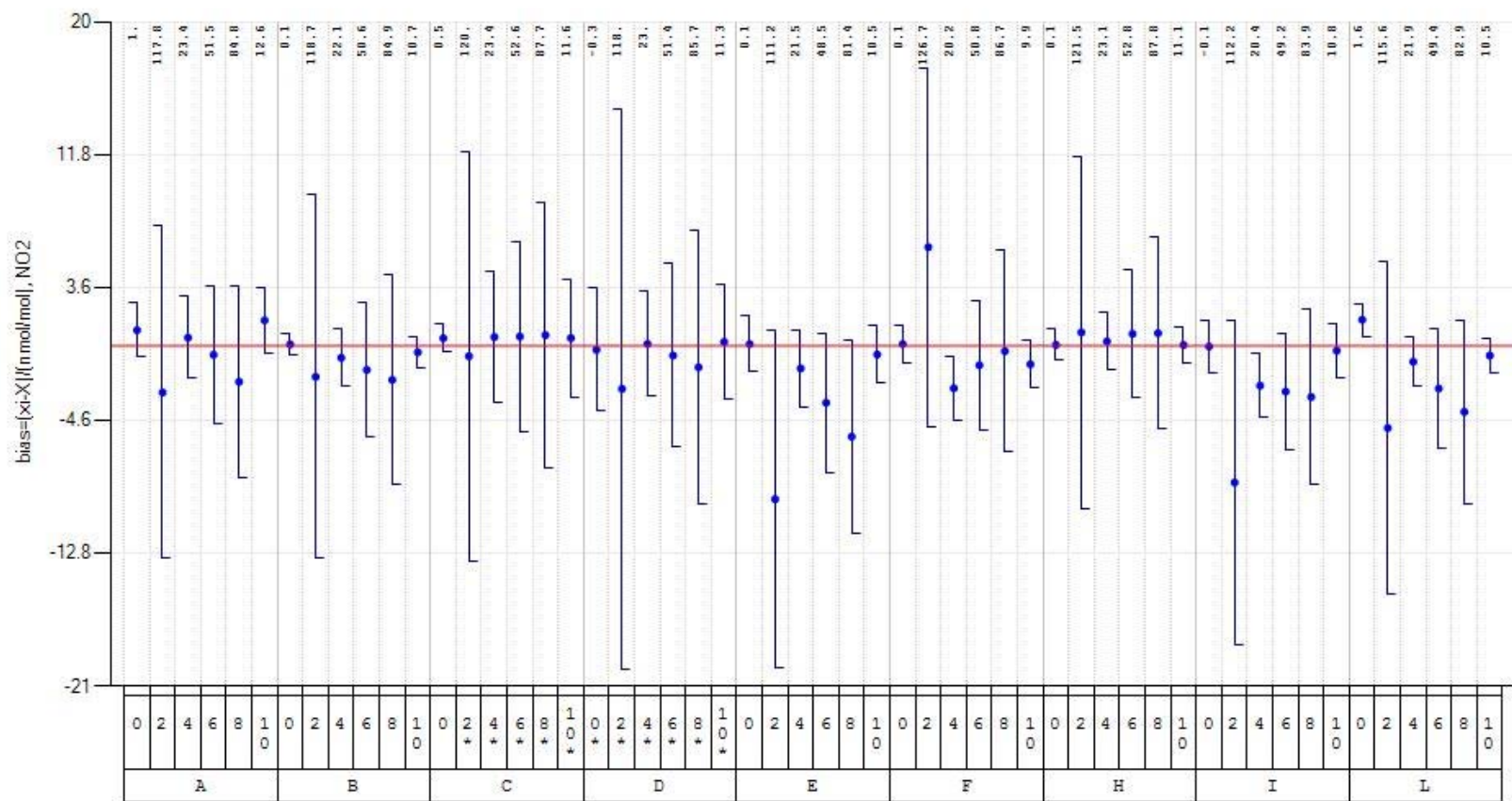


Figure 10: Bias of participant's NO₂ measurement results

Expanded uncertainty of bias is presented as error bar for NO₂ run numbers 0, 2, 4, 6, 8 and 10 (see Table 3). Results with error bars touching or crossing the x-axis are satisfactory. For each evaluation the run number together with the participants rounded run average (nmol/mol) is given. The '*' mark indicates reported standard uncertainties bigger than σ_p .

3. Performance characteristics of individual laboratories

Individual participants' bias were evaluated and are presented in chapter 2 (Figure 6-Figure 10). Since the results of NO₂ runs 1, 3, 5, 7 and 9 were not treated in proficiency evaluation the bias of these runs are presented in Figure 11.

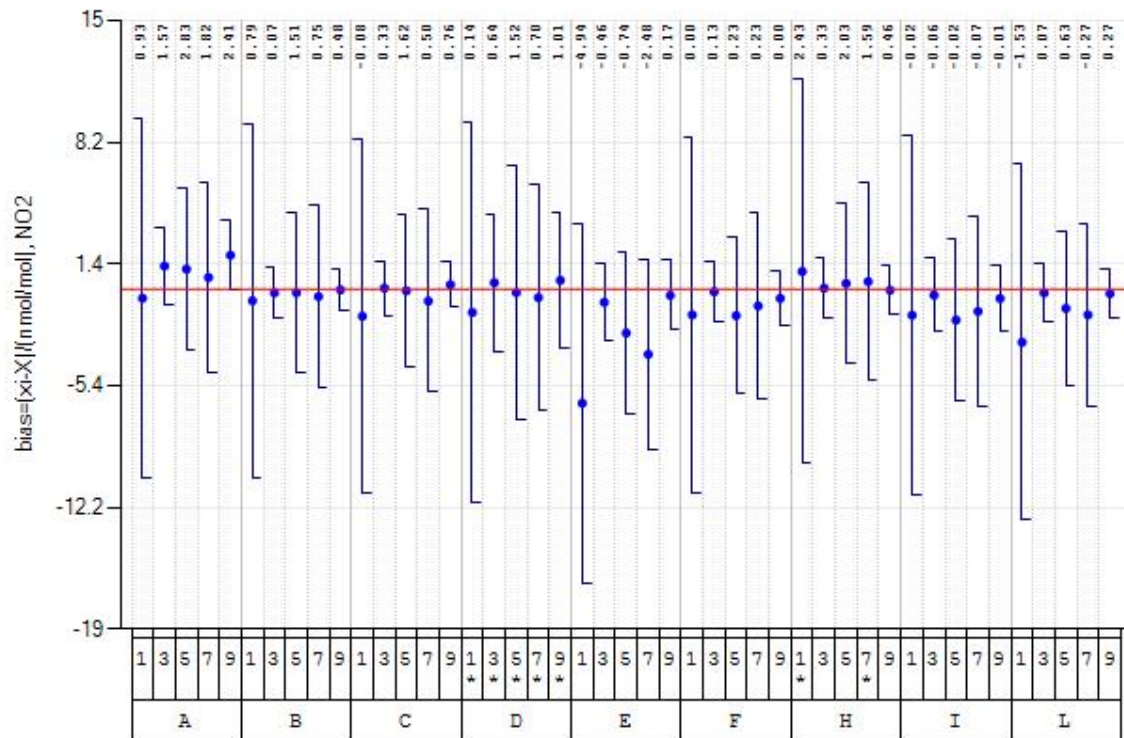


Figure 11: Bias of participant's NO₂ measurements for run numbers 1, 3, 5, 7 and 9
 At these test gas mixtures the concentration levels of NO₂ were zero and the concentration levels of NO were not zero (see Table 3). In that perspective the figure shows the effect of NO concentration on NO₂ measurements. For each evaluation the run number together with the participants rounded run average (nmol/mol) is given.

3.1 The efficiency of NO₂-to-NO converters of NO_x analyzers

Since NO and NO₂ test gases were produced by gas phase titration it is possible to evaluate the efficiency of NO₂-to-NO converter of each participant's NO_x analyser. The evaluation takes each participant's NO and NO₂ measurements before and after oxidation by O₃. The converter efficiency (α) is calculated using Equation 3 [4]:

$$\alpha = \frac{[NO_2]_i - [NO_2]_{i-1}}{[NO]_{i-1} - [NO]_i} \cdot 100\% \quad \text{Equation 3}$$

The O₃ measurements of each participant can also be compared to either NO or NO₂ change by calculating Δ^{NO} or Δ^{NO_2} using Equation 4 and Equation 5 respectively:

$$\Delta^{NO} = [O_3]_{i+1} - ([NO]_{i-1} - [NO]_i) \quad \text{Equation 4}$$

$$\Delta^{NO_2} = [O_3]_{i+1} - ([NO_2]_i - [NO_2]_{i-1}) \quad \text{Equation 5}$$

Ideal value for α is 100% while for Δ^{NO} and Δ^{NO_2} it is 0 nmol/mol.

IE	NO ₂	α	Δ^{NO}	Δ^{NO_2}
code	nmol/mol	%	nmol/mol	nmol/mol
A	14	102.5		
A	95	100.6	2.4	2
A	55	99.6	1.8	2
A	25	99.6	0.9	1
A	130	99.9	-2.1	-1.9
B	14	100.9		
B	95	100.7	3.3	2.7
B	55	100.5	2.5	2.3
B	25	100.9	0.8	0.6
B	130	102.6	4.9	2
C	14	98.0		
C	95	98.0	-3.6	-1.8
C	55	98.0	-1.4	-0.4
C	25	98.0	-1.3	-0.8
C	130	98.0	-6.3	-3.9
D	14	100.1		
D	95	100.7	-0.8	-1.4
D	55	101.5	-0.1	-0.8
D	25	100.9	-1	-1.2
D	130	100.9	-2.7	-3.7
F	14	98		
F	95	99.6	-2.8	-2.5
F	55	100.9	0	-0.4
F	25	98.9	2.2	2.4
F	130	100	-12.2	-12.3

IE	NO ₂	α	Δ^{NO}	Δ^{NO_2}
code	nmol/mol	%	nmol/mol	nmol/mol
G	14	101		
G	95	100.9	-0.3	-1
G	55	101.4	0.5	-0.1
G	25	100.9	-0.3	-0.5
G	130	100.8	-3.1	-4.1
H	14	101.7		
H	95	99.8	-2.6	-2.4
H	55	101.2	-0.6	-1.2
H	25	100.5	-0.7	-0.8
H	130	101.2	-3.7	-5.2
I	14	102.9		
I	95	100.8	-0.3	-0.9
I	55	100.1	0	-0.1
I	25	100.1	1.2	1.2
I	130	99.8	0.4	0.6
L	14	99.3	0	0
L	95	99.9	1.5	1.6
L	55	100.9	2.2	1.8
L	25	99.8	0.7	0.8
L	130	100.1	-1.6	-1.7

Table 5: The efficiency of NO₂-to-NO converters.

The evaluation of Equation 4 and Equation 5 cannot be made at the lowest NO₂ level (14 ppb) because, due to the low concentration of NO, O₃ and NO₂ are not detectable with the necessary accuracy. The evaluation of equations 3, 4 and 5 for each participant at different concentration levels are given in Table 5.

4. Discussion

For a general assessment of the quality of each result a decision diagram was developed (Figure 12) that results in seven categories (1 to 7). The general comments for each category are:

- **1**: measurement result is completely satisfactory
- **2**: measurement result is satisfactory (z'-score satisfactory and En-number ok) but the reported uncertainty is too high
- **3**: measured value is satisfactory (z'-score satisfactory) but the reported uncertainty is underestimated (En-number not ok)
- **4**: measurement result is questionable (z'-score questionable) but due to a high reported uncertainty can be considered valid (En-number ok)
- **5**: measurement result is questionable (z'-score questionable and En-number not ok)
- **6**: measurement result is unsatisfactory (z'-score unsatisfactory) but due to a high reported uncertainty can be considered valid (En-number ok)
- **7**: measurement result is unsatisfactory (z'-score unsatisfactory and En-number not ok)

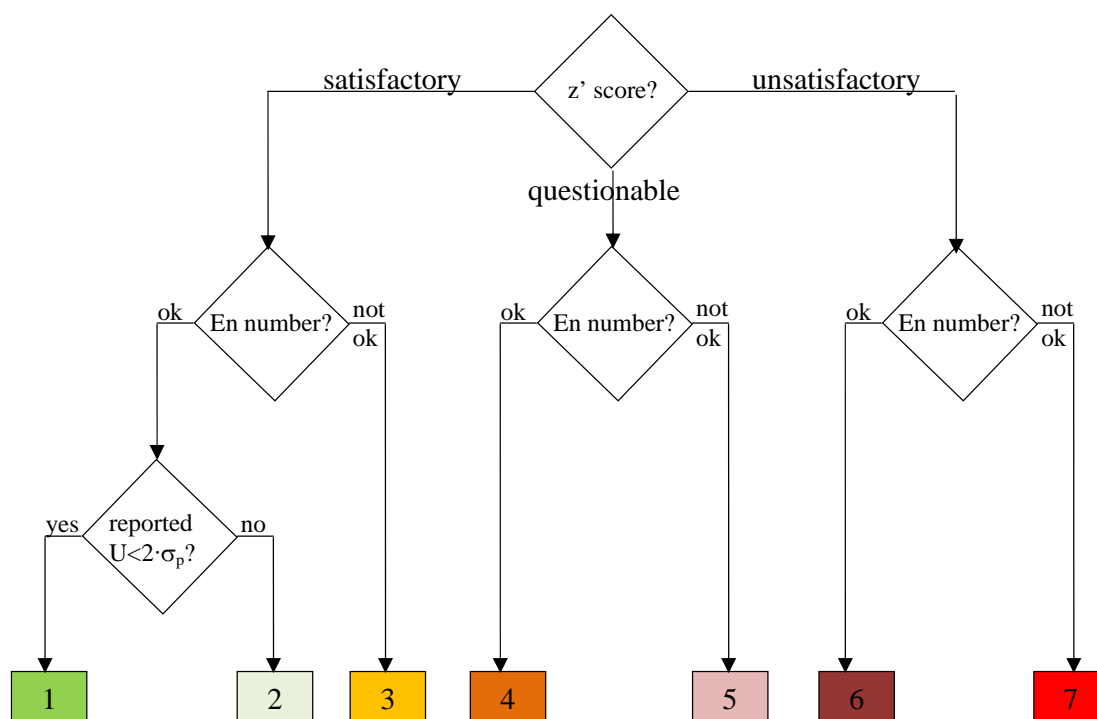


Figure 12: The decision diagram for general assessment of proficiency results.

The results of the IE were assigned to categories according to the diagram given in Figure 12 and are presented in Table 6.

	run number	Ref. conc. level	IE code								
			A	B	C	D	E	F	H	I	L
CO (µmol/mol)	0	-0.002	2	3	2	1	1	1	1	1	1
	1	8.039	1	3	1	1	3	1	3	1	1
	2	4.542	1	3	1	1	3	1	1	1	1
	3	6.041	1	3	1	1	3	1	3	1	1
	4	3.031	1	5	1	1	1	1	1	1	3
	5	1.021	2	3	2	1	1	1	1	1	3
NO (nmol/mol)	0	0.0	1	1	1	2	1	1	1	1	1
	1	515.7	1	1	2	1	1	1	1	1	1
	2	397.5	1	1	2	1	1	1	1	1	1
	3	61.9	1	1	2	1	1	5	1	1	1
	4	39.5	1	1	2	1	1	5	1	1	1
	5	171.4	1	1	2	1	1	1	1	1	1
	6	121.8	1	1	2	1	1	3	1	1	1
	7	258.1	1	1	2	1	1	1	1	1	1
	8	173.0	1	1	2	1	1	1	1	1	1
	9	20.7	1	1	2	1	1	1	1	3	1
	10	10.3	1	1	2	2	1	1	1	3	1
NO ₂ (nmol/mol)	0	0.0	1	1	1	2	1	1	1	1	3
	2	120.6	1	1	2	2	1	1	1	1	1
	4	22.8	1	1	2	2	1	3	1	3	1
	6	52.0	1	1	2	2	1	1	1	1	1
	8	87.0	1	1	2	2	1	1	1	1	1
	10	11.1	1	1	2	2	1	1	1	1	1
O ₃ (nmol/mol)	0	-0.2	3	1	1	2	nd	1	1	1	1
	1	115.1	1	3	2	2	nd	1	1	1	1
	2	22.1	1	1	1	2	nd	2	1	1	1
	3	50.2	1	1	2	2	nd	1	1	1	1
	4	84.8	1	1	2	2	nd	1	1	1	1
	5	10.6	1	1	1	2	nd	2	1	1	1
SO ₂ (nmol/mol)	0	0.0	1	1	nd	1	1	1	1	1	1
	1	8.7	1	3	nd	1	1	1	1	1	1
	2	51.0	1	1	nd	1	1	1	1	1	1
	3	19.8	1	3	nd	1	1	1	1	1	1
	4	122.9	1	1	nd	1	5	1	1	1	1
	5	3.4	1	3	nd	1	1	1	1	1	1

Table 6: The general assessment of proficiency results. "nd" is referring to values not reported.

5. Conclusions

The proficiency evaluation scheme has provided an assessment of the participants measured values and their evaluated uncertainties. In terms of the criteria imposed by the European Directive (σ_p) 78.5% of the results reported (Table 7) by AQUILA laboratories fall into category '1' and are good both in terms of measured values and evaluated uncertainties. Among the remaining results the majority presented good measured values, but the evaluated uncertainties were either too high, category '2' (12.5%), or too small, category '3' (7.6%) and 1.3% of results, category '5', are questionable compared to z-score and Not OK for the En-number.

IE	Categories %						
	1	2	3	4	5	6	7
Apr-08	68.4	18.1	7.3	1.0	1.0	2.6	1.6
Oct-08 (I)	37.9	40.8	14.2	0.6	3.6	1.0	1.9
Oct-08 (II)	34.3	38.9	23.7	1.0	2.0	0.0	0.0
Sep-09	60.8	29.9	3.1	4.1	1.0	1.0	0.0
Oct-09	85.0	5.7	7.5	0.4	1.4	0.0	0.0
Jun-10	84.6	8.1	4.4	0.7	2.3	0.0	0.0
Sep-11	86.0	7.9	5.4	0.0	0.3	0.0	0.3
Oct-11 (I)	78.5	12.5	7.6	0.0	1.3	0.0	0.0

Table 7: Flags summary

As in previous IE, the adopted criteria for high concentrations were the standard deviations for proficiency assessment, deriving from the European Standards' uncertainty requirements. The reproducibility standard deviation obtained at this (Annex C) and previous IE [20], [21], [22], [23], [24], [25] are comparable to the mentioned criteria. On the other hand, the uncertainty criteria for zero levels were those set in AQUILA's position paper [12]. In the present IE a high share of '1' results can be observed confirming the trend of the most recent IEs.

In this exercise there were no unsatisfactory results in the z'-score evaluations (Table 8). Laboratory B obtained 1 questionable result for CO, laboratory E obtained 1 questionable result for SO₂ and laboratory F obtained 2 questionable results for NO.

Note: One unsatisfactory or two questionable results per parameter require participation at next IE.

Inter-comparison	Site	Questionable	Unsatisfactory	Satisfactory
June-05	Ispra (IT)	2.3%	2.3%	95.5%
June-07	Ispra (IT)	1.9%	0.3%	97.8%
October-07	Essen (DE)	4.6%	2.2%	93.2%
April-08	Ispra (IT)	2.1%	4.1%	93.8%
October 2008_1	Ispra (IT)	4.2%	2.9%	92.9%
October 2008_2	Ispra (IT)	3.0%	0.0%	97.0%
September-09	Langen (DE)	4.7%	0.9%	94.3%
October-09	Ispra (IT)	1.8%	0.0%	98.2%
June-10	Ispra (IT)	3.0%	0.0%	97.0%
September-11	Ispra (IT)	0.3%	0.3%	99.7%
October-11	Ispra (IT)	1.3%	0.0%	98.7%

Table 8: Z'-score summary

Comparability of results among AQUILA participants at the highest concentration level, excluding outliers, is acceptable for SO₂, CO and O₃ measurements while NO and NO₂ measurements showed less satisfactory results.

The relative reproducibility limits, at the highest studied concentration levels, are 9.7% for SO₂, 9.4% for CO, 5.7% for O₃ and for NO 6.3% all within the objective derived from criteria imposed by the European Commission (σ_p). As shown by the Figure 51 there is a slightly poor reproducibility around 10 nmol/mol for NO. The poor reproducibility for NO₂ is more relevant and the relative reproducibility limit 10.9% is beyond the target 9.02% (see Table 4).

During this IE the performance of all NRL has been quite good. Only one outlier has been identified at zero level for SO₂ (Table 51).

6. References

- [1] Directive 2008/50/EC of the European Parliament and of the Council of 21 May 2008 on ambient air quality and cleaner air for Europe, L 152, 11.06.2008
- [2] EN 14626:2005, Ambient air quality - Standard method for the measurement of the concentration of carbon monoxide by non-dispersive infrared spectroscopy
- [3] EN 14212:2005, Ambient air quality - Standard method for the measurement of the concentration of sulphur dioxide by ultraviolet fluorescence
- [4] EN 14211:2005, Ambient air quality - Standard method for the measurement of the concentration of nitrogen dioxide and nitrogen monoxide by chemiluminescence
- [5] EN 14625:2005, Ambient air quality - Standard method for the measurement of the concentration of ozone by ultraviolet photometry
- [6] ISO 6143:2001, Gas analysis - Comparison methods for determining and checking the composition of calibration gas mixtures
- [7] ISO 6144:2003, Gas analysis - Preparation of calibration gas mixtures - Static volumetric method
- [8] ISO 6145-7:2001, Gas analysis - Preparation of calibration gas mixtures using dynamic volumetric methods - Part 7: Thermal mass-flow controllers
- [9] Mücke H.-G., (2008), Air quality management in the WHO European Region – Results of a quality assurance and control programme on air quality monitoring (1994-2004), Environment International, EI-01718
- [10] Mücke H.-G., et al. (2000), European Intercomparison workshop on air quality monitoring vol.4 – Measuring NO, NO₂, O₃ and SO₂ – Air Hygiene Report 13, WHO Collaboration Centre for Air Quality Management and Air Pollution Control, ISSN 0938 - 9822
- [11] <http://ies.jrc.ec.europa.eu/aquila-project/aquila-homepage.html>
- [12] AQUILA POSITION PAPER N. 37, (2008) Protocol for intercomparison exercise. Organisation of intercomparison exercises for gaseous air pollution for EU national air quality reference laboratories and laboratories of the WHO EURO region http://ies.jrc.ec.europa.eu/uploads/fileadmin/H04/Air_Quality/N%2037%20final%20version%20IE%20organisation%20and%20evaluation.pdf
- [13] ISO 13528:2005, Statistical methods for use in proficiency testing by interlaboratory comparisons
- [14] ISO 5725-1:1994, Accuracy (trueness and precision) of measurement methods and results – Part 1: General principles and definitions
- [15] ISO 5725-2:1994, Accuracy (trueness and precision) of measurement methods and results – Part 2: Basic method for the determination of repeatability and reproducibility of a standard measurement method
- [16] ISO 5725-6:1994, Accuracy (trueness and precision) of measurement methods and results - Part 6: Use in practice of accuracy values
- [17] Harmonisation of Directive 92/72/EEC on air pollution by ozone, E. De Saeger et al., EUR 17662, 1997
- [18] De Saeger E. et al., (1997) European comparison of Nitrogen Dioxide calibration methods, EUR 17661
- [19] ISO 15337:2009, Ambient air - Gas phase titration - Calibration of analysers for ozone
- [20] Kapus M. et al. (2009) The evaluation of the Intercomparison Exercise for SO₂, CO, O₃, NO and NO₂ carried out in June 2007 in Ispra . JRC scientific and technical reports. EUR 23804.
- [21] Kapus M. et al. (2009) The evaluation of the Intercomparison Exercise for SO₂, CO, O₃, NO and NO₂ - April 2008. JRC scientific and technical reports. EUR 23805.

- [22] Kapus M. et al. (2009) The evaluation of the Intercomparison Exercise for SO₂, CO, O₃, NO and NO₂ 6-9 October 2008. JRC scientific and technical reports. EUR 23806.
- [23] Kapus M. et al. (2009) The evaluation of the Intercomparison Exercise for SO₂, CO, O₃, NO and NO₂ 13-16 October 2008.. JRC scientific and technical reports. EUR 23807.
- [24] Belis C. A. et al. (2010) The evaluation of the Interlaboratory comparison Exercise for SO₂, CO, O₃, NO and NO₂ Langen 20-25 September 2009.
- [25] Belis C. A. et al. (2010) The evaluation of the Interlaboratory comparison Exercise for SO₂, CO, O₃, NO and NO₂ 19-22 October 2009.
- [26] Viallon J. et al 2009 Metrologia 46 08017. Final report, on-going key comparison BIPM.QM-K1: Ozone at ambient level, comparison with JRC, 2008. doi: 10.1088/0026-1394/46/1A/08017
- [27] Viallon, J., et al. (2006), International comparison CCQM-P28: Ozone at ambient level, Metrologia, 43, Tech. Suppl., 08010, doi:10.1088/0026-1394/43/1A/08010
- [28] Tanimoto, H., et al. (2006), Intercomparison of ultraviolet photometry and gas-phase titration techniques for ozone reference standards at ambient levels, Journal of Geophysical Research, vol. 111, D16313, doi:10.1029/2005JD006983
- [29] GUM Workbench, The Tool for Expression of Uncertainty of Measurements
- [30] VDI 2449 Part3: 2001, Measurement methods test criteria- General method for the determination of the uncertainty of calibratable measurement methods.

Annex A. Assigned values

The assigned values of tested concentration levels (run) were derived from ERLAPs measurements which are calibrated against the certified reference values of CRMs and are traceable to international standards. In this perspective the assigned values are reference values as defined in the ISO 13528 [13].

ERLAP's SO₂, CO and NO analysers were calibrated according to the methodology described in the ISO 6143 [6]. Reference gas mixtures were produced from the primary reference materials (produced and certified by NMI Van Swinden Laboratorium) by dynamic dilution method using mass flow controllers [8]. All flows were measured with a certified molbloc/molbox1 system. For O₃ measurements, the analyzers were calibrated using the JRC SRP42 primary standard (constructed by NIST) which has been compared to BIPM primary standard [26]. The photometer absorption cross section uncertainty (1.06%) was included in the uncertainty budget [27] [28].

The reference gas mixture and the calibration experiment evaluation were carried out using two computer applications, the "GUM WORKBENCH" [29] and "B-least" [30] respectively. For extending calibration from the NO to NO₂ channel of NO_x analyser the GPT test was performed to establish the efficiency of NO₂-converter.

ERLAP's measurement results were validated by comparison to the group statistics (x* and s*) for every parameter and concentration level of the IE. These statistics are calculated from participants, applying the robust method described in the Annex C of the ISO 13528 [13]. The validation is taking into account ERLAP's measurement result (X) and its standard uncertainty (u_X) as given in Equation 6 [13]:

$$\frac{|x^* - X|}{\sqrt{\frac{(1,25 \cdot s^*)^2}{p} + u_X^2}} < 2 \quad \text{Equation 6}$$

Where 'x*' and 's*' represent robust average and robust standard deviation respectively and 'p' is the number of participants.

In Table 9 all inputs for expression 6 are given and all ERLAP's measurement results are confirmed to be valid.

As a group evaluation robust average (x*) and robust standard deviation (s*) were calculated (applying the procedure described in Annex C of ISO 13528) for each run, and are presented in the following table.

run	unit	X	uX'	x*	s*	p	val.
NO_0	nmol/mol	0.04	0.3	0.021	0.273	10	OK
NO_1	nmol/mol	515.683	3.97	517.773	10.19	10	OK
NO_2	nmol/mol	397.453	3.05	399.457	8.494	10	OK
NO_3	nmol/mol	61.887	0.58	61.222	1.674	10	OK
NO_4	nmol/mol	171.42	1.77	170.836	4.435	10	OK
NO_5	nmol/mol	39.52	0.44	39.326	1.778	10	OK
NO_6	nmol/mol	121.77	0.99	121.541	4.014	10	OK
NO_7	nmol/mol	258.14	2.04	257.348	5.16	10	OK
NO_8	nmol/mol	173.037	1.37	172.939	4.738	10	OK
NO_9	nmol/mol	20.74	0.49	20.302	0.82	10	OK
NO_10	nmol/mol	10.287	0.31	10.056	0.579	10	OK
NO2_0	nmol/mol	-0.01	0.31	0.116	0.21	10	OK
NO2_1	nmol/mol	1.43	4.96	0.179	1.297	10	OK
NO2_2	nmol/mol	120.637	4.54	118.161	4.157	10	OK
NO2_3	nmol/mol	0.267	0.7	0.204	0.242	10	OK
NO2_4	nmol/mol	22.843	0.63	22.268	1.147	10	OK
NO2_5	nmol/mol	1.7	2.11	1.202	1.084	10	OK
NO2_6	nmol/mol	52.027	1.54	50.883	1.654	10	OK
NO2_7	nmol/mol	1.16	2.54	0.565	0.927	10	OK
NO2_8	nmol/mol	87	2.24	85.318	2.28	10	OK
NO2_9	nmol/mol	0.507	0.56	0.477	0.445	10	OK
NO2_10	nmol/mol	11.063	0.37	10.932	0.604	10	OK

run	unit	X	uX'	x*	s*	p	val.
CO_0	µmol/mol	-0.002	0.005	0.001	0.04	10	OK
CO_1	µmol/mol	8.039	0.041	7.937	0.144	10	OK
CO_2	µmol/mol	4.5423	0.024	4.531	0.088	10	OK
CO_3	µmol/mol	6.0413	0.031	6.012	0.096	10	OK
CO_4	µmol/mol	3.0313	0.016	3.052	0.078	10	OK
CO_5	µmol/mol	1.021	0.007	1.031	0.074	10	OK
O3_0	nmol/mol	-0.18	0.32	0.048	0.43	9	OK
O3_1	nmol/mol	115.107	0.93	114.876	1.197	9	OK
O3_2	nmol/mol	22.077	0.29	22.21	0.534	9	OK
O3_3	nmol/mol	50.18	0.42	50.143	0.735	9	OK
O3_4	nmol/mol	84.82	0.68	84.484	1.123	9	OK
O3_5	nmol/mol	10.597	0.3	10.759	0.341	9	OK
SO2_0	nmol/mol	-0.01	0.29	0.032	0.307	9	OK
SO2_1	nmol/mol	8.657	0.31	8.684	0.39	9	OK
SO2_2	nmol/mol	51.003	0.47	51.289	1.27	9	OK
SO2_3	nmol/mol	19.837	0.33	19.892	0.447	9	OK
SO2_4	nmol/mol	122.853	0.9	123.416	3.257	9	OK
SO2_5	nmol/mol	3.42	0.3	3.453	0.364	9	OK

Table 9: The validation of assigned values (X)

by comparison to the robust averages (x*) with taking into account the standard uncertainties of assigned values (uX'), and robust standard deviations (s*) as denoted by Equation 6.

The homogeneity of test gas was evaluated from measurements at the beginning and end of the distribution line. From the relative differences between beginning and end measurements,

average and standard deviation were calculated, and the uncertainty of test gas due to lack of homogeneity was calculated as the sum of squares of these average and standard deviation. The upper and lower limits of bias due to homogeneity was evaluated to be smaller than 0.5% which constitutes the relative standard uncertainty of 0,3% of each concentration level. The standard uncertainties of assigned/reference values (u_X) were calculated with Equation 7 and used in the proficiency evaluations of chapter 2.

$$u_{X'}^2 = u_X^2 + (X \cdot u_{\text{homogeneity}})^2$$

Equation 7

Annex B. The results of the IE

In this annex are reported participant's results, presented both in tables and graphs. For each run, participants were asked to report 3 results representing 30 minutes measurement each (x_{ij}). In this annex are presented the reported data and their uncertainty $u(x_i)$ and $U(x_i)$ expressed in mol/mol units.

For all the runs except concentration levels 0, also average (\bar{x}_i) and standard deviation (s_i) of each participant are presented.

The assigned value is indicated on the graphs with the red line and the individual laboratories expanded uncertainties (Ux_i) are indicated with error bars.

Reported values for SO₂

values	laboratories									
	A	B	D	E	F	G	H	I	L	
$x_{i,1}$ (nmol/mol)	-0.28	0.53	-0.01	-1.69	0.00	-0.01	0.30	0.07	0.20	
$u(x_i)$ (nmol/mol)	0.78	0.03	0.40	0.80	0.50	0.29	0.24	0.49	0.40	
$U(x_i)$ (nmol/mol)	1.55	0.06	0.81	1.60	1.00	0.58	0.48	0.98	0.80	

Table 10: Reported values for SO₂ run 0.

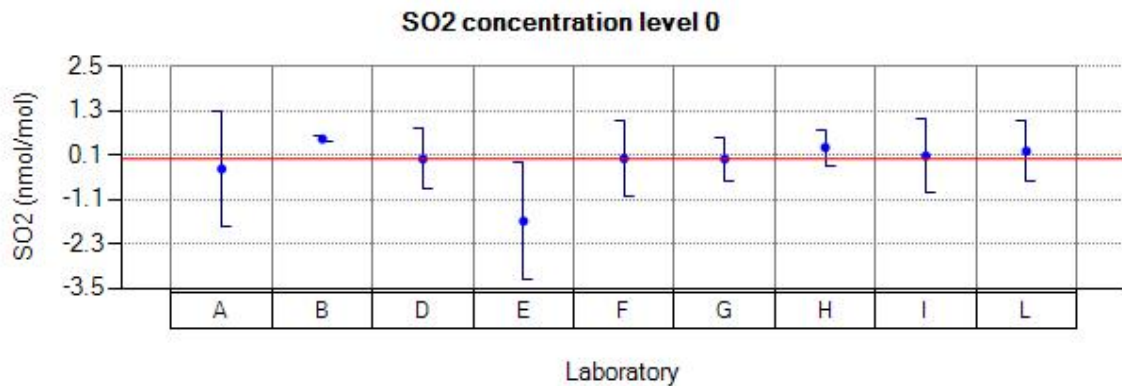


Figure 13: Reported values for SO₂ run 0.

values	laboratories									
	A	B	D	E	F	G	H	I	L	
xi,1 (nmol/mol)	8.17	9.89	8.58	7.42	8.8	8.65	8.86	8.43	9.3	
xi,2 (nmol/mol)	8.51	9.9	8.59	7.37	8.8	8.63	8.87	8.51	9.5	
xi,3 (nmol/mol)	8.08	9.65	8.7	7.48	8.8	8.69	8.89	8.56	9.5	
Xi (nmol/mol)	8.25	9.81	8.62	7.42	8.80	8.65	8.87	8.50	9.43	
Si (nmol/mol)	0.22	0.14	0.06	0.05	0.00	0.03	0.01	0.06	0.11	
u(xi) (nmol/mol)	0.89	0.23	0.48	0.80	0.60	0.30	0.25	0.49	0.40	
U(xi) (nmol/mol)	1.78	0.46	0.96	1.60	1.20	0.61	0.51	0.98	0.80	

Table 11: Reported values for SO₂ run 1.

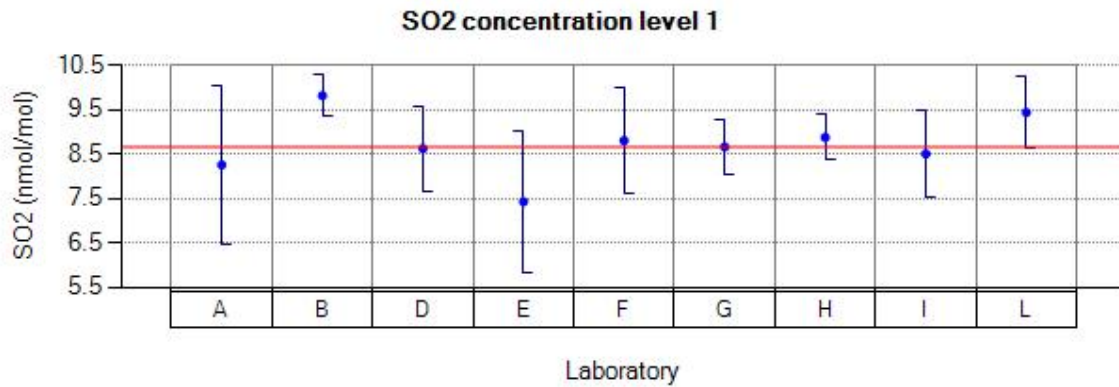


Figure 14: Reported values for SO₂ run 1.

values	laboratories									
	A	B	D	E	F	G	H	I	L	
xi,1 (nmol/mol)	49.69	53.17	50.48	52.79	50.20	50.94	51.00	50.96	51.80	
xi,2 (nmol/mol)	49.44	53.42	50.53	53.21	50.30	51.05	51.30	51.59	51.90	
xi,3 (nmol/mol)	49.77	53.16	50.52	53.33	50.30	51.02	51.16	51.42	52.10	
Xi (nmol/mol)	49.63	53.25	50.51	53.11	50.26	51.00	51.15	51.32	51.93	
Si (nmol/mol)	0.17	0.14	0.02	0.28	0.05	0.05	0.15	0.32	0.15	
u(xi) (nmol/mol)	1.45	1.19	1.45	1.50	1.20	0.47	0.53	1.49	0.51	
U(xi) (nmol/mol)	2.90	2.38	2.91	3.00	2.30	0.94	1.06	2.98	1.02	

Table 12: Reported values for SO₂ run 2.

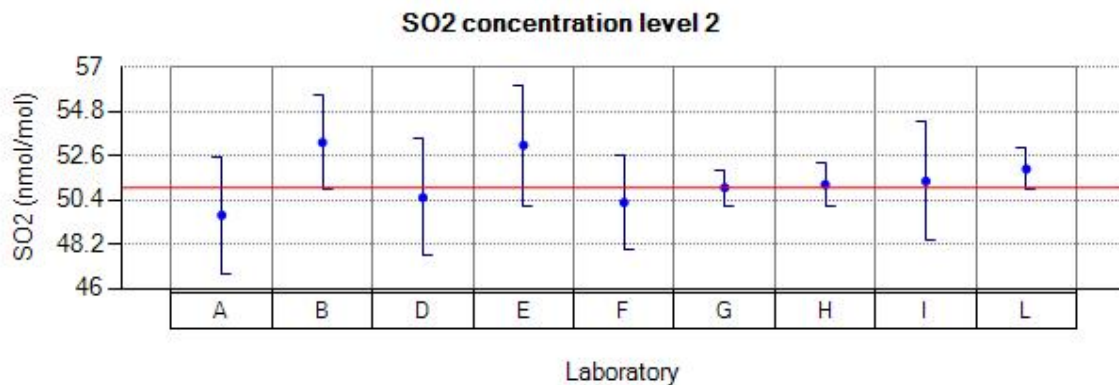


Figure 15: Reported values for SO₂ run 2.

values	laboratories									
	A	B	D	E	F	G	H	I	L	
xi,1 (nmol/mol)	18.97	21.56	19.69	19.57	19.70	19.85	20.01	20.11	20.60	
xi,2 (nmol/mol)	19.35	21.14	19.56	19.56	19.70	19.85	20.07	20.10	20.70	
xi,3 (nmol/mol)	19.45	20.86	19.58	19.60	19.80	19.81	20.03	20.23	20.80	
Xi (nmol/mol)	19.25	21.18	19.61	19.57	19.73	19.83	20.03	20.14	20.70	
Si (nmol/mol)	0.25	0.35	0.07	0.02	0.05	0.02	0.03	0.07	0.10	
u(xi) (nmol/mol)	1.04	0.51	0.70	1.00	0.70	0.33	0.30	0.58	0.40	
U(xi) (nmol/mol)	2.08	1.02	1.40	2.00	1.50	0.65	0.61	1.17	0.80	

Table 13: Reported values for SO₂ run 3.

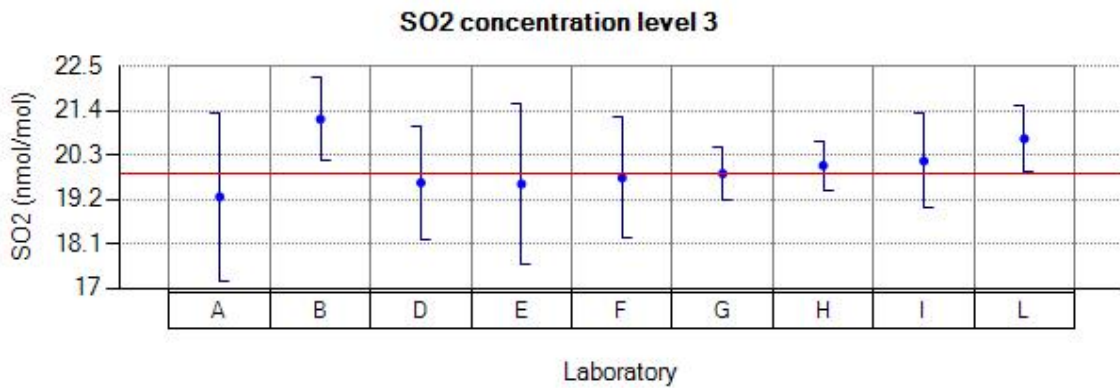


Figure 16: Reported values for SO₂ run 3.

values	laboratories									
	A	B	D	E	F	G	H	I	L	
xi,1 (nmol/mol)	118.66	127.03	121.52	130.80	120.50	122.81	122.90	124.84	124.70	
xi,2 (nmol/mol)	118.87	127.27	121.94	130.91	120.60	123.03	122.92	125.16	124.30	
xi,3 (nmol/mol)	118.92	127.86	121.66	131.17	120.30	122.72	123.28	124.79	124.30	
Xi (nmol/mol)	118.81	127.38	121.70	130.96	120.46	122.85	123.03	124.93	124.43	
Si (nmol/mol)	0.13	0.42	0.21	0.19	0.15	0.15	0.21	0.20	0.23	
u(xi) (nmol/mol)	2.38	2.85	3.33	3.20	2.80	0.90	0.85	3.62	1.23	
U(xi) (nmol/mol)	4.77	5.70	6.66	6.40	5.50	1.81	1.71	7.25	2.46	

Table 14: Reported values for SO₂ run 4.

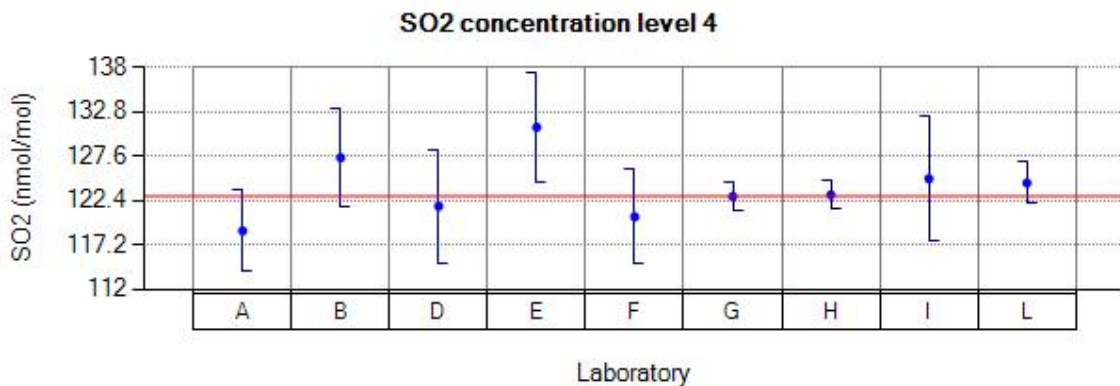


Figure 17: Reported values for SO₂ run 4.

values	laboratories									
	A	B	D	E	F	G	H	I	L	
xi,1 (nmol/mol)	2.40	4.49	3.41	1.99	3.50	3.46	3.69	3.57	4.30	
xi,2 (nmol/mol)	2.35	4.81	3.30	1.93	3.40	3.40	3.65	3.50	3.90	
xi,3 (nmol/mol)	2.30	4.61	3.29	1.88	3.40	3.40	3.55	3.53	4.20	
Xi (nmol/mol)	2.35	4.63	3.33	1.93	3.43	3.42	3.63	3.53	4.13	
Si (nmol/mol)	0.05	0.16	0.06	0.05	0.05	0.03	0.07	0.03	0.20	
u(xi) (nmol/mol)	0.81	0.14	0.42	0.80	0.60	0.30	0.24	0.49	0.40	
U(xi) (nmol/mol)	1.62	0.28	0.85	1.60	1.10	0.59	0.48	0.98	0.80	

Table 15: Reported values for SO₂ run 5.

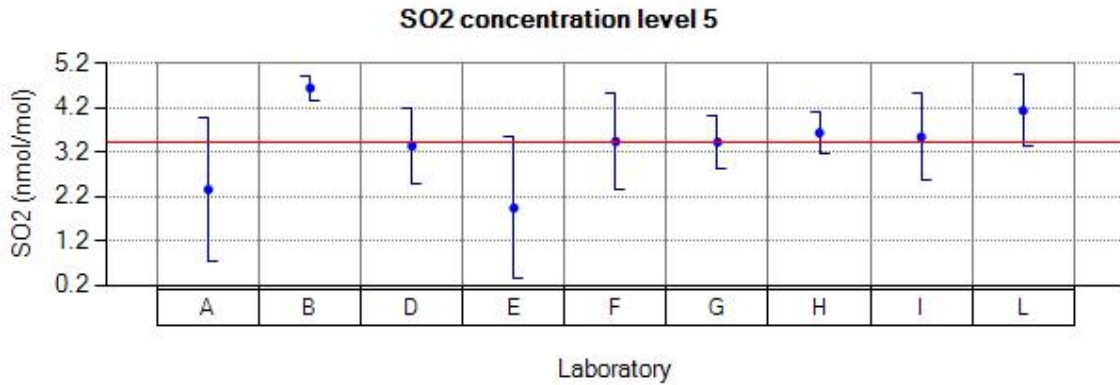


Figure 18: Reported values for SO₂ run 5.

Reported values for CO

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
$x_{i,1}$ ($\mu\text{mol/mol}$)	-0.096	-0.038	0.001	-0.003	-0.016	0.040	-0.002	0.071	-0.005	0.043
$u(x_i)$ ($\mu\text{mol/mol}$)	0.129	0.001	0.121	0.049	0.080	0.040	0.005	0.063	0.050	0.040
$U(x_i)$ ($\mu\text{mol/mol}$)	0.258	0.002	0.241	0.097	0.160	0.080	0.010	0.126	0.100	0.080

Table 16: Reported values for CO run 0.

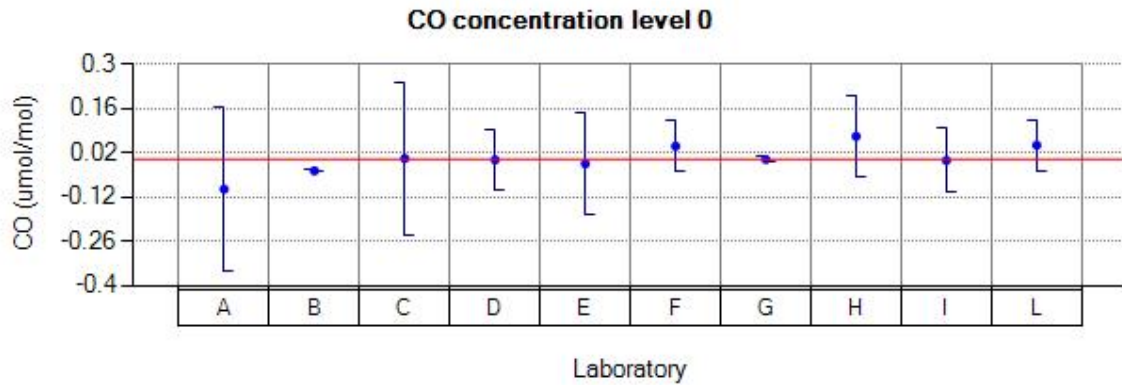


Figure 19: Reported values for CO run 0.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
$x_{i,1}$ ($\mu\text{mol/mol}$)	7.744	7.695	8.001	8.096	7.510	8.020	8.038	7.491	8.046	7.975
$x_{i,2}$ ($\mu\text{mol/mol}$)	7.759	7.692	8.004	8.104	7.509	8.020	8.040	7.504	8.054	7.975
$x_{i,3}$ ($\mu\text{mol/mol}$)	7.750	7.690	8.008	8.111	7.506	8.020	8.039	7.501	8.054	7.974
\bar{X}_i ($\mu\text{mol/mol}$)	7.751	7.692	8.004	8.104	7.508	8.020	8.039	7.499	8.051	7.975
S_i ($\mu\text{mol/mol}$)	0.008	0.003	0.004	0.008	0.002	0.000	0.001	0.007	0.005	0.001
$u(x_i)$ ($\mu\text{mol/mol}$)	0.202	0.146	0.276	0.130	0.150	0.180	0.041	0.149	0.201	0.163
$U(x_i)$ ($\mu\text{mol/mol}$)	0.403	0.292	0.551	0.260	0.300	0.370	0.082	0.297	0.403	0.325

Table 17: Reported values for CO run 1.

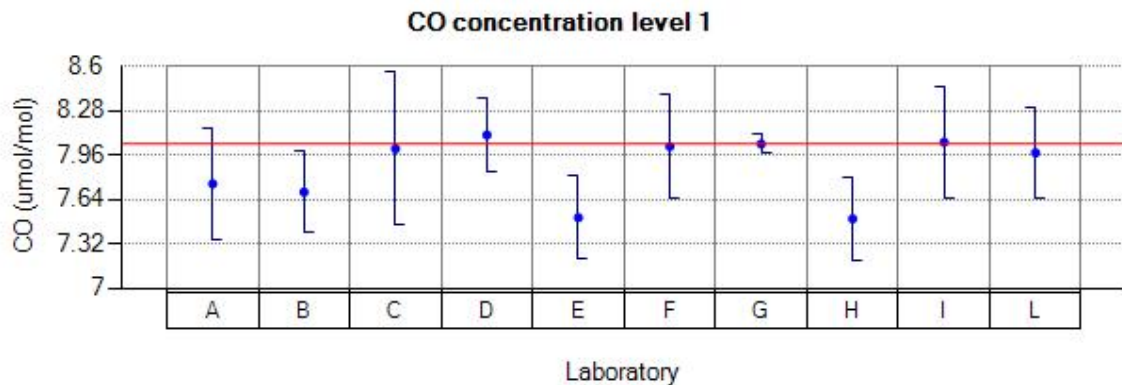


Figure 20: Reported values for CO run 1.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
x _{i,1} (μmol/mol)	4.353	4.230	4.574	4.592	4.235	4.610	4.542	4.409	4.586	4.616
x _{i,2} (μmol/mol)	4.364	4.233	4.575	4.592	4.234	4.610	4.543	4.401	4.592	4.612
x _{i,3} (μmol/mol)	4.368	4.235	4.577	4.594	4.233	4.610	4.542	4.414	4.626	4.606
X _i (μmol/mol)	4.362	4.233	4.575	4.593	4.234	4.610	4.542	4.408	4.601	4.611
S _i (μmol/mol)	0.008	0.003	0.002	0.001	0.001	0.000	0.001	0.007	0.022	0.005
u(x _i) (μmol/mol)	0.170	0.080	0.186	0.084	0.120	0.110	0.024	0.102	0.115	0.094
U(x _i) (μmol/mol)	0.341	0.160	0.372	0.168	0.240	0.210	0.047	0.204	0.230	0.188

Table 18: Reported values for CO run 2.

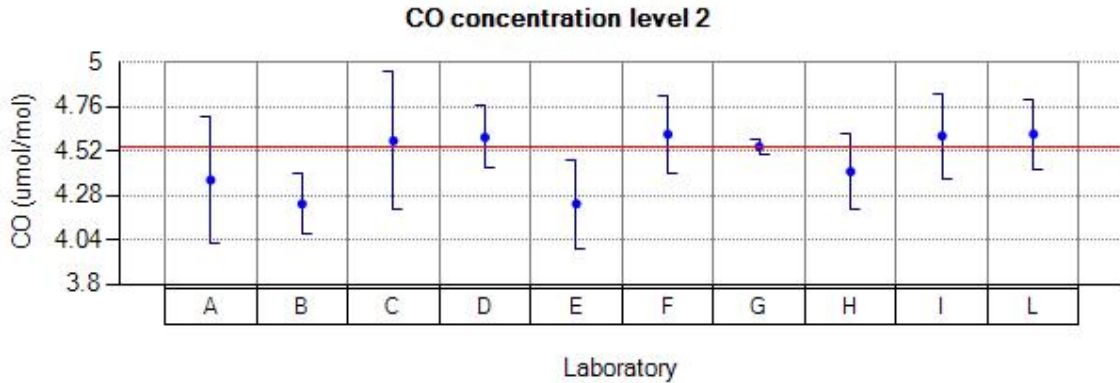


Figure 21: Reported values for CO run 2.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
x _{i,1} (μmol/mol)	5.849	5.759	6.058	6.115	5.626	6.100	6.040	5.736	6.079	6.051
x _{i,2} (μmol/mol)	5.847	5.750	6.061	6.113	5.628	6.100	6.042	5.741	6.089	6.056
x _{i,3} (μmol/mol)	5.846	5.753	6.063	6.116	5.626	6.090	6.042	5.742	6.094	6.053
X _i (μmol/mol)	5.847	5.754	6.061	6.115	5.627	6.097	6.041	5.740	6.087	6.053
S _i (μmol/mol)	0.002	0.005	0.003	0.002	0.001	0.006	0.001	0.003	0.008	0.003
u(x _i) (μmol/mol)	0.184	0.109	0.223	0.103	0.130	0.140	0.031	0.121	0.152	0.123
U(x _i) (μmol/mol)	0.368	0.218	0.446	0.206	0.260	0.280	0.062	0.242	0.304	0.247

Table 19: Reported values for CO run 3.

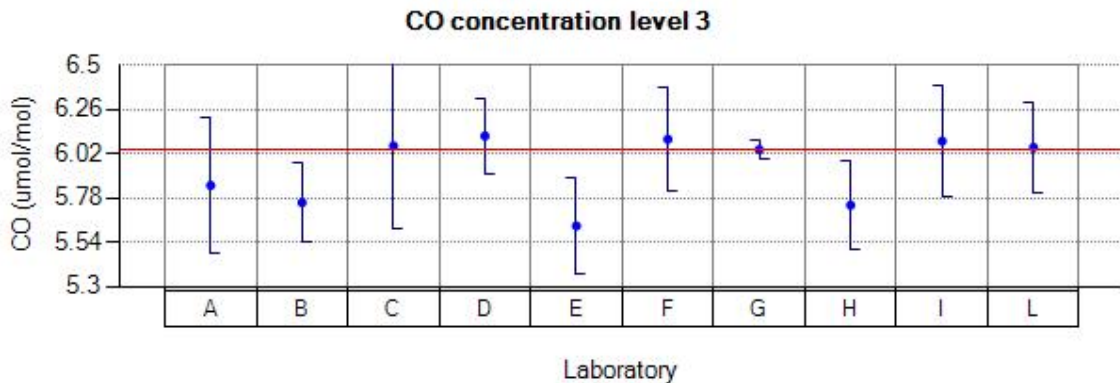


Figure 22: Reported values for CO run 3.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
$x_{i,1}$ ($\mu\text{mol/mol}$)	2.926	2.650	3.089	3.078	2.820	3.120	3.031	3.057	3.079	3.178
$x_{i,2}$ ($\mu\text{mol/mol}$)	2.942	2.646	3.089	3.071	2.819	3.120	3.031	3.054	3.076	3.177
$x_{i,3}$ ($\mu\text{mol/mol}$)	2.940	2.641	3.091	3.074	2.820	3.120	3.032	3.061	3.075	3.174
X_i ($\mu\text{mol/mol}$)	2.936	2.646	3.090	3.074	2.820	3.120	3.031	3.057	3.077	3.176
S_i ($\mu\text{mol/mol}$)	0.009	0.005	0.001	0.004	0.001	0.000	0.001	0.004	0.002	0.002
$u(x_i)$ ($\mu\text{mol/mol}$)	0.157	0.050	0.154	0.067	0.100	0.070	0.016	0.085	0.077	0.065
$U(x_i)$ ($\mu\text{mol/mol}$)	0.314	0.100	0.308	0.134	0.200	0.140	0.032	0.171	0.154	0.130

Table 20: Reported values for CO run 4.

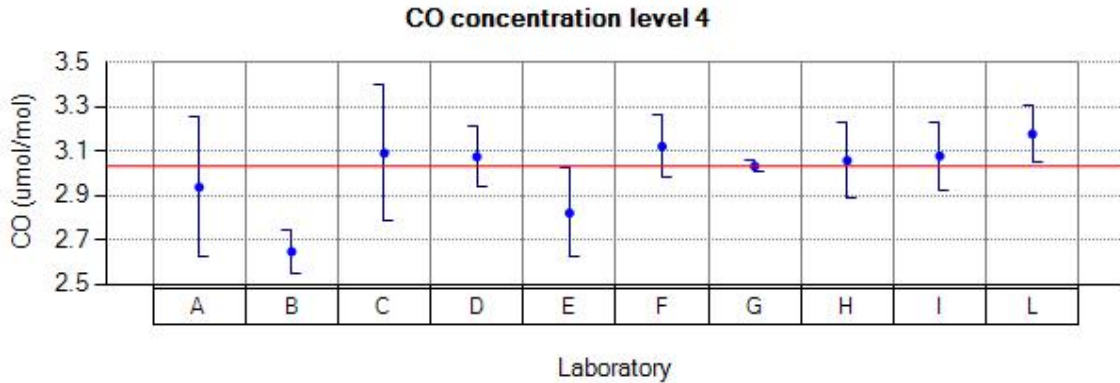


Figure 23: Reported values for CO run 4.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
$x_{i,1}$ ($\mu\text{mol/mol}$)	0.999	0.909	1.071	1.038	0.941	1.080	1.022	1.114	1.003	1.121
$x_{i,2}$ ($\mu\text{mol/mol}$)	0.995	0.912	1.070	1.034	0.940	1.070	1.021	1.116	1.008	1.120
$x_{i,3}$ ($\mu\text{mol/mol}$)	0.998	0.915	1.071	1.034	0.940	1.070	1.020	1.110	1.008	1.119
X_i ($\mu\text{mol/mol}$)	0.997	0.912	1.071	1.035	0.940	1.073	1.021	1.113	1.006	1.120
S_i ($\mu\text{mol/mol}$)	0.002	0.003	0.001	0.002	0.001	0.006	0.001	0.003	0.003	0.001
$u(x_i)$ ($\mu\text{mol/mol}$)	0.139	0.017	0.125	0.051	0.080	0.100	0.007	0.068	0.050	0.040
$U(x_i)$ ($\mu\text{mol/mol}$)	0.278	0.034	0.250	0.102	0.160	0.190	0.014	0.136	0.100	0.080

Table 21: Reported values for CO run 5.

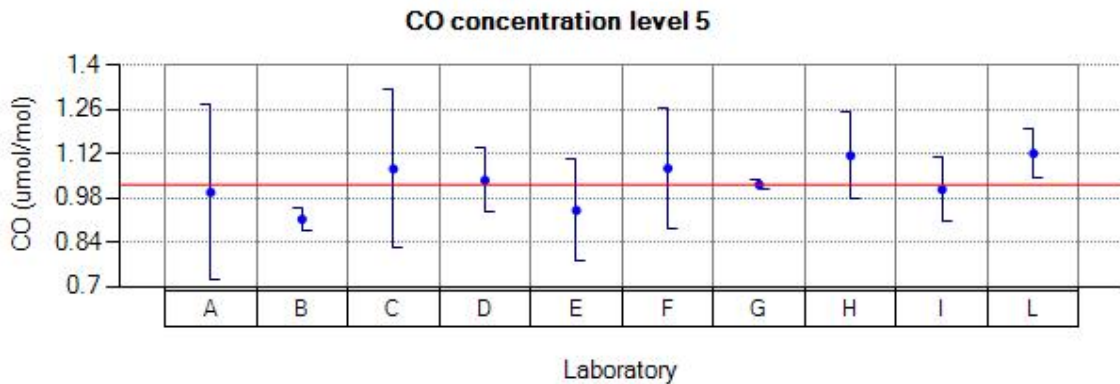


Figure 24: Reported values for CO run 5.

Reported values for O₃

values	laboratories									
	A	B	C	D	F	G	H	I	L	
xi,1 (nmol/mol)	0.95	0.41	-0.23	-1.11	0.20	-0.18	-0.20	0.09	0.20	
u(xi) (nmol/mol)	0.32	0.55	0.20	1.76	0.50	0.32	0.24	0.40	0.60	
U(xi) (nmol/mol)	0.65	1.10	0.39	3.51	1.00	0.63	0.48	0.80	1.20	

Table 22: Reported values for O₃ run 0.

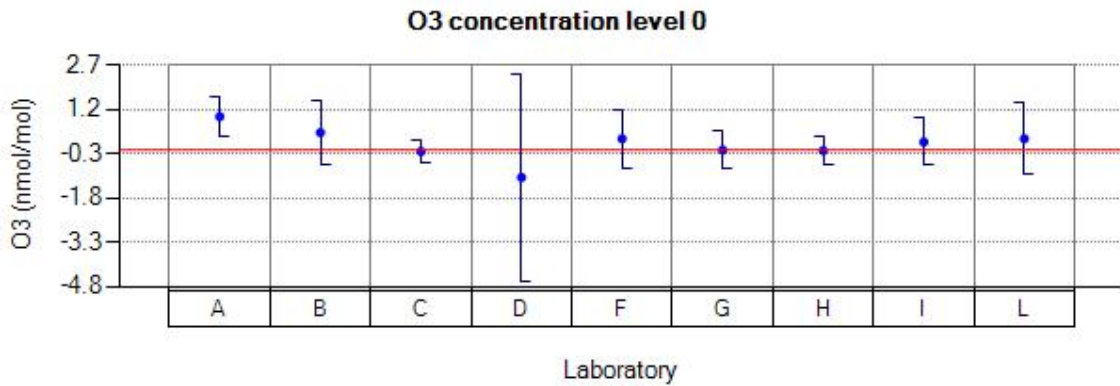


Figure 25: Reported values for O₃ run 0.

values	laboratories									
	A	B	C	D	F	G	H	I	L	
xi,1 (nmol/mol)	114.78	120.30	116.03	114.09	114.40	114.70	113.68	112.56	115.20	
xi,2 (nmol/mol)	114.84	119.89	116.30	114.20	114.50	115.21	113.88	112.90	115.30	
xi,3 (nmol/mol)	115.06	119.60	116.23	114.22	114.50	115.41	114.05	113.01	115.70	
Xi (nmol/mol)	114.89	119.93	116.18	114.17	114.46	115.10	113.87	112.82	115.40	
Si (nmol/mol)	0.14	0.35	0.14	0.07	0.05	0.36	0.18	0.23	0.26	
u(xi) (nmol/mol)	1.25	1.82	5.00	3.32	2.06	0.93	0.85	2.14	2.29	
U(xi) (nmol/mol)	2.50	3.64	10.00	6.64	4.12	1.86	1.71	4.29	4.59	

Table 23: Reported values for O₃ run 1

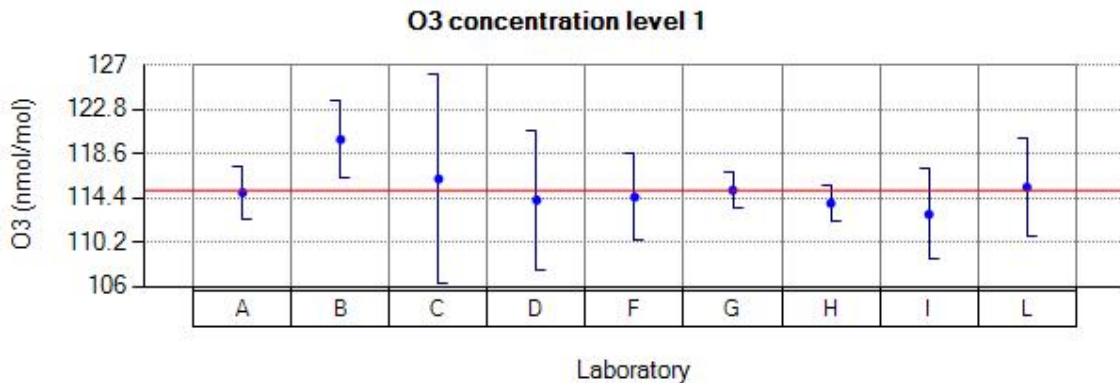


Figure 26: Reported values for O₃ run 1.

values	laboratories									
	A	B	C	D	F	G	H	I	L	
xi,1 (nmol/mol)	22.82	22.82	22.17	21.09	22.5	22.04	21.93	21.58	22.6	
xi,2 (nmol/mol)	22.83	22.51	22.23	21.13	22.5	22.07	21.98	21.63	22.6	
xi,3 (nmol/mol)	22.83	22.62	22.26	21.09	22.5	22.12	21.98	21.69	22.5	
Xi (nmol/mol)	22.82	22.65	22.22	21.10	22.50	22.07	21.96	21.63	22.56	
Si (nmol/mol)	0.00	0.15	0.04	0.02	0.00	0.04	0.02	0.05	0.05	
u(xi) (nmol/mol)	0.50	0.79	0.98	1.83	1.65	0.29	0.29	0.82	0.60	
U(xi) (nmol/mol)	1.00	1.58	1.95	3.67	3.29	0.57	0.57	1.64	1.20	

Table 24: Reported values for O₃ run 2.

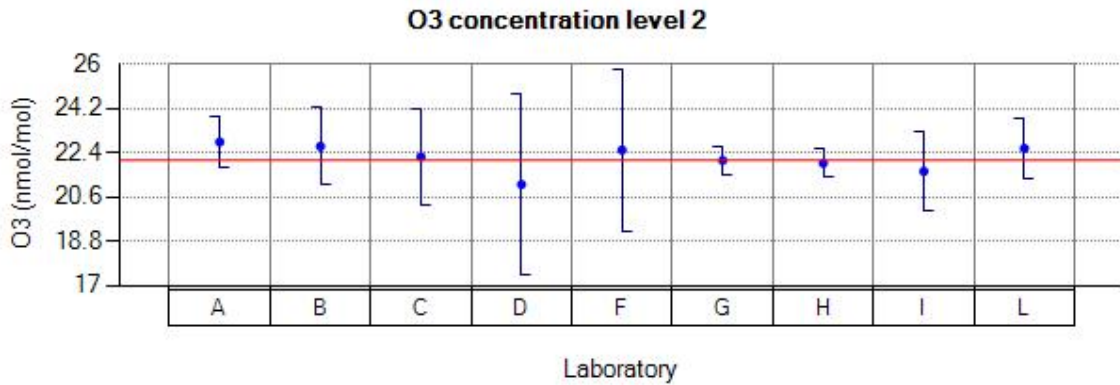


Figure 27: Reported values for O₃ run 2.

values	laboratories									
	A	B	C	D	F	G	H	I	L	
xi,1 (nmol/mol)	50.47	51.12	50.45	49.02	50.20	50.12	49.41	48.99	50.40	
xi,2 (nmol/mol)	50.69	51.34	50.58	49.14	50.20	50.20	49.53	49.15	50.60	
xi,3 (nmol/mol)	50.68	51.41	50.63	49.17	50.20	50.22	49.65	49.18	50.60	
Xi (nmol/mol)	50.61	51.29	50.55	49.11	50.20	50.18	49.53	49.10	50.53	
Si (nmol/mol)	0.12	0.15	0.09	0.07	0.00	0.05	0.12	0.10	0.11	
u(xi) (nmol/mol)	0.73	1.09	2.18	2.13	1.75	0.42	0.43	0.93	1.00	
U(xi) (nmol/mol)	1.45	2.18	4.37	4.27	3.50	0.84	0.86	1.87	2.01	

Table 25: Reported values for O₃ run 3.

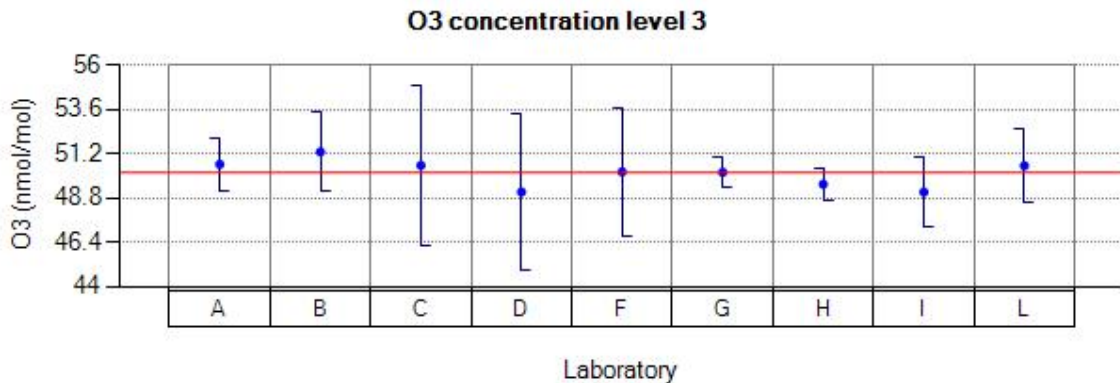


Figure 28: Reported values for O₃ run 3.

values	laboratories									
	A	B	C	D	F	G	H	I	L	
xi,1 (nmol/mol)	84.78	86.74	85.29	83.49	83.90	84.70	83.68	82.79	84.60	
xi,2 (nmol/mol)	85.01	86.93	85.38	83.53	83.90	84.86	83.85	83.07	84.90	
xi,3 (nmol/mol)	84.97	86.94	85.41	83.64	84.00	84.90	83.88	83.14	84.90	
\bar{X}_i (nmol/mol)	84.92	86.87	85.36	83.55	83.93	84.82	83.80	83.00	84.80	
S_i (nmol/mol)	0.12	0.11	0.06	0.07	0.05	0.10	0.10	0.18	0.17	
$u(x_i)$ (nmol/mol)	1.01	1.46	3.68	2.71	1.91	0.68	0.65	1.57	1.69	
$U(x_i)$ (nmol/mol)	2.02	2.92	7.35	5.42	3.82	1.35	1.30	3.15	3.37	

Table 26: Reported values for O₃ run 4.

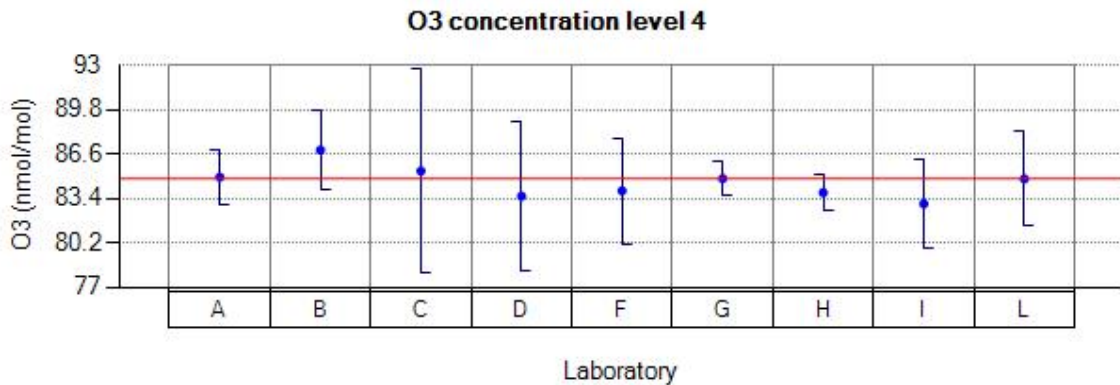


Figure 29: Reported values for O₃ run 4.

values	laboratories									
	A	B	C	D	F	G	H	I	L	
xi,1 (nmol/mol)	11.6	11.58	10.66	9.67	10.7	10.54	10.68	10.47	11.1	
xi,2 (nmol/mol)	11.53	11.57	10.72	9.68	10.8	10.62	10.69	10.51	11.2	
xi,3 (nmol/mol)	11.62	11.65	10.7	9.65	10.7	10.63	10.7	10.53	11.2	
\bar{X}_i (nmol/mol)	11.58	11.60	10.69	9.66	10.73	10.59	10.69	10.50	11.16	
S_i (nmol/mol)	0.04	0.04	0.03	0.01	0.05	0.04	0.01	0.03	0.05	
$u(x_i)$ (nmol/mol)	0.41	0.67	0.50	1.77	1.62	0.30	0.25	0.40	0.60	
$U(x_i)$ (nmol/mol)	0.82	1.34	1.00	3.55	3.24	0.59	0.50	0.80	1.20	

Table 27: Reported values for O₃ run 5.

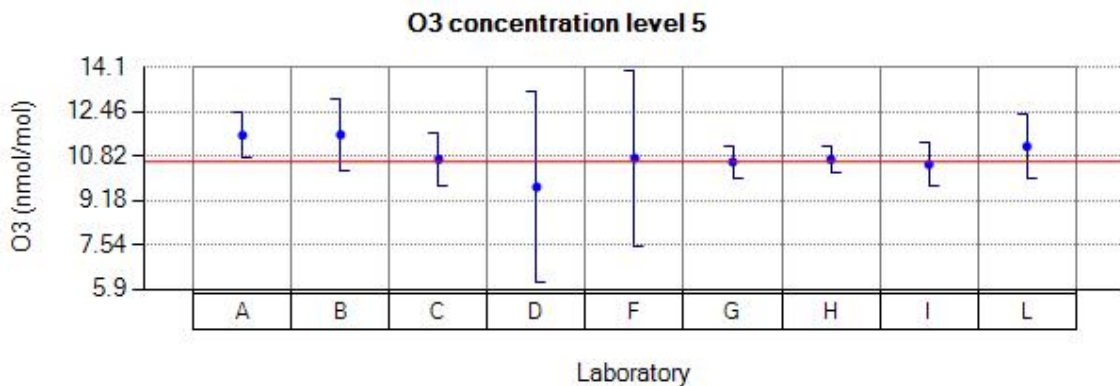


Figure 30: Reported values for O₃ run 5.

Reported values for NO

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
xi,1 (nmol/mol)	0.04	0.12	0.34	-0.29	-0.18	0.20	0.04	0.02	0.30	-0.40
u(xi) (nmol/mol)	0.78	0.01	0.30	1.43	0.80	0.50	0.29	0.24	0.75	0.40
U(xi) (nmol/mol)	1.56	0.02	0.59	2.85	1.60	1.00	0.59	0.48	1.50	0.80

Table 28: Reported values for NO run 0.

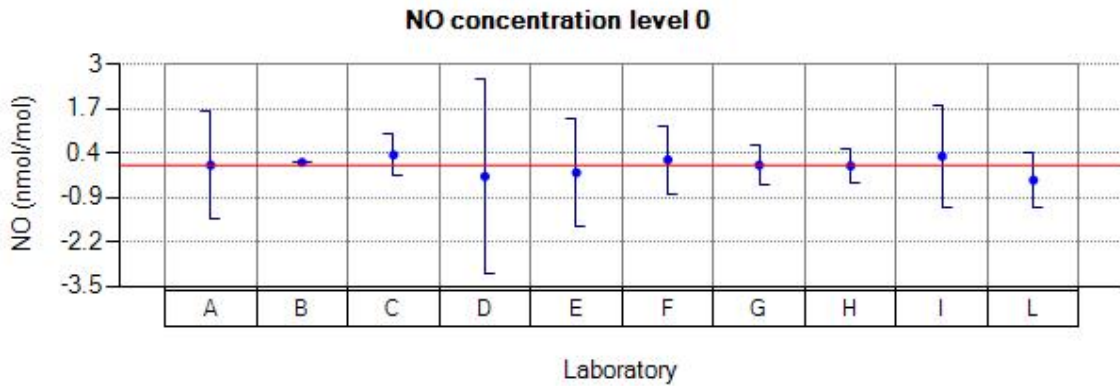


Figure 31: Reported values for NO run 0.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
xi,1 (nmol/mol)	504.52	525.03	537.72	512.08	504.07	524.80	515.14	516.68	525.00	513.60
xi,2 (nmol/mol)	506.24	524.78	538.37	512.68	505.11	524.70	515.75	517.73	525.77	514.80
xi,3 (nmol/mol)	505.81	520.60	539.07	513.34	505.60	524.60	516.16	519.05	525.17	515.20
Xi (nmol/mol)	505.52	523.47	538.38	512.70	504.92	524.70	515.68	517.82	525.31	514.53
Si (nmol/mol)	0.89	2.48	0.67	0.63	0.78	0.10	0.51	1.18	0.40	0.83
u(xi) (nmol/mol)	7.59	10.31	18.23	7.86	9.20	12.10	3.97	8.14	9.72	10.75
U(xi) (nmol/mol)	15.22	20.62	36.46	15.72	18.40	24.10	7.94	16.28	19.44	21.50

Table 29: Reported values for NO run 1.

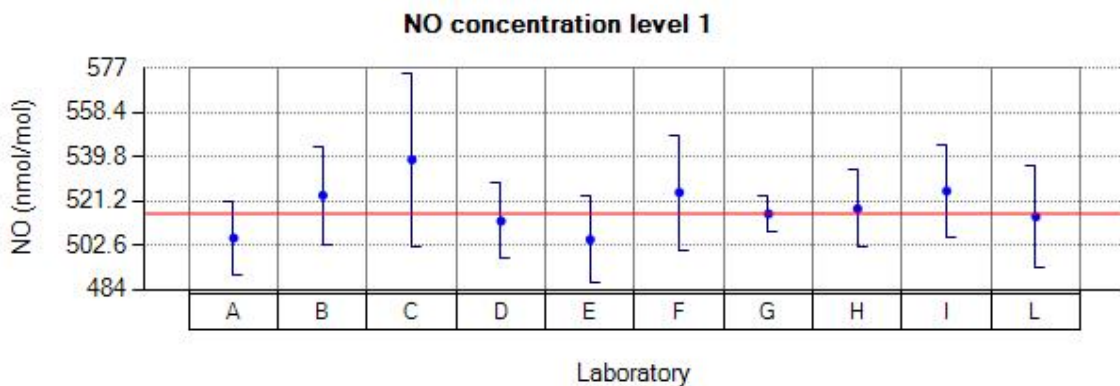


Figure 32: Reported values for NO run 1.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
xi,1 (nmol/mol)	387.85	408.48	415.66	395.91	389.88	398.00	397.43	400.69	412.96	397.00
xi,2 (nmol/mol)	388.76	408.59	416.08	396.03	389.89	398.00	397.51	399.83	413.03	397.80
xi,3 (nmol/mol)	389.09	408.35	415.82	395.65	390.04	398.00	397.42	400.18	412.75	397.90
Xi (nmol/mol)	388.56	408.47	415.85	395.86	389.93	398.00	397.45	400.23	412.91	397.56
Si (nmol/mol)	0.64	0.12	0.21	0.19	0.09	0.00	0.04	0.43	0.14	0.49
u(xi) (nmol/mol)	6.02	7.97	14.11	5.99	7.30	9.20	3.05	6.26	7.64	8.30
U(xi) (nmol/mol)	12.06	15.94	28.23	11.98	14.60	18.30	6.10	12.52	15.28	16.61

Table 30: Reported values for NO run 2.

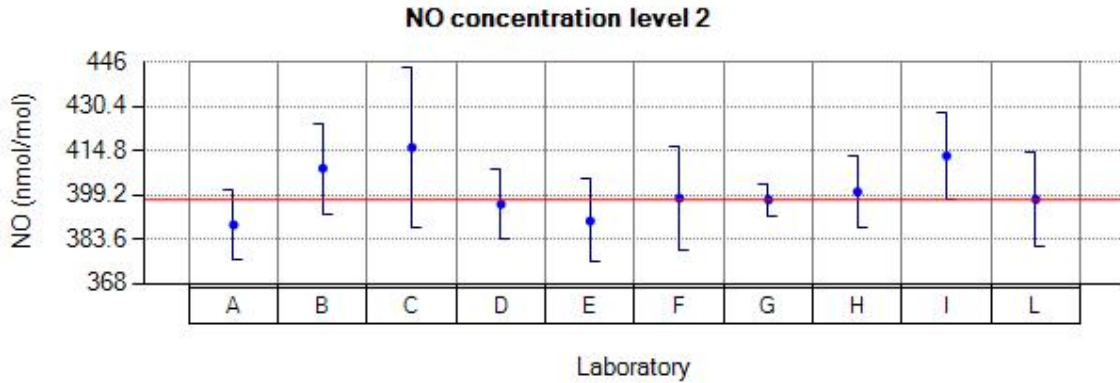


Figure 33: Reported values for NO run 2.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
xi,1 (nmol/mol)	59.31	62.43	65.25	60.89	60.17	55.30	61.75	62.95	61.03	60.50
xi,2 (nmol/mol)	59.95	62.37	65.26	61.05	60.30	55.30	61.96	63.03	61.36	60.50
xi,3 (nmol/mol)	59.63	62.12	65.45	61.04	60.22	55.40	61.95	63.11	61.54	60.60
Xi (nmol/mol)	59.63	62.30	65.32	60.99	60.23	55.33	61.88	63.03	61.31	60.53
Si (nmol/mol)	0.32	0.16	0.11	0.09	0.06	0.05	0.11	0.08	0.25	0.05
u(xi) (nmol/mol)	1.58	1.22	2.81	1.34	1.50	1.30	0.58	1.04	1.13	1.26
U(xi) (nmol/mol)	3.17	2.44	5.61	2.68	3.00	2.50	1.16	2.09	2.27	2.53

Table 31: Reported values for NO run 3.

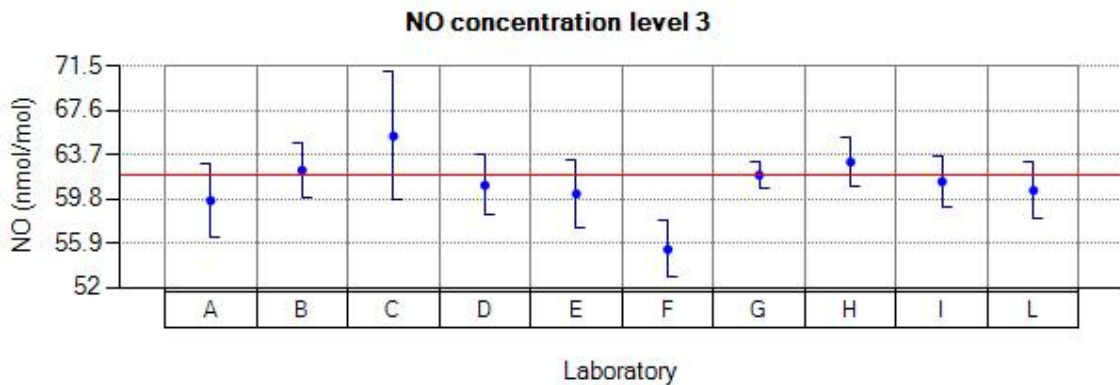


Figure 34: Reported values for NO run 3.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
xi,1 (nmol/mol)	37.70	40.41	41.75	38.90	38.19	35.00	39.49	40.30	40.86	38.60
xi,2 (nmol/mol)	37.77	40.50	41.81	38.84	38.26	35.00	39.53	40.38	40.76	38.80
xi,3 (nmol/mol)	37.79	40.46	41.80	38.83	38.43	35.00	39.54	40.36	41.05	38.70
Xi (nmol/mol)	37.75	40.45	41.78	38.85	38.29	35.00	39.52	40.34	40.89	38.70
Si (nmol/mol)	0.04	0.04	0.03	0.03	0.12	0.00	0.02	0.04	0.14	0.10
u(xi) (nmol/mol)	1.29	0.79	2.24	1.31	1.20	0.80	0.44	0.57	0.75	0.81
U(xi) (nmol/mol)	2.58	1.58	4.47	2.62	2.40	1.60	0.87	1.14	1.51	1.62

Table 32: Reported values for NO run 4.

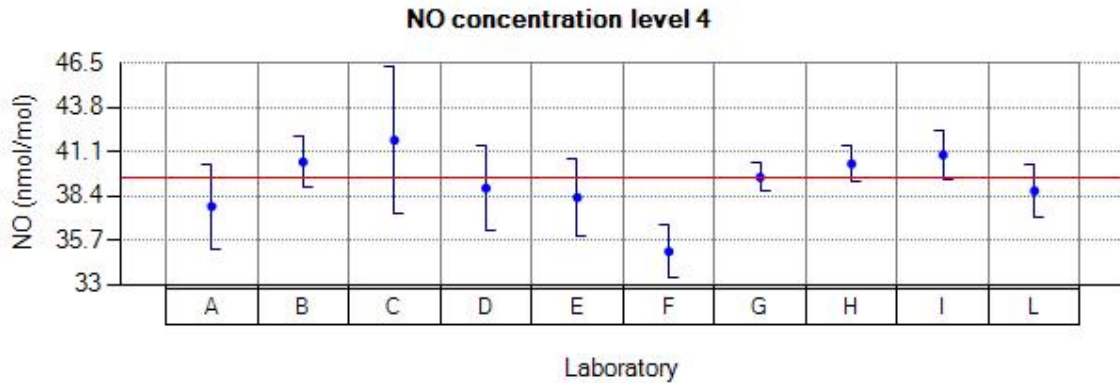


Figure 35: Reported values for NO run 4.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
xi,1 (nmol/mol)	164.96	170.48	177.89	168.51	166.36	165.20	169.71	172.34	174.85	166.30
xi,2 (nmol/mol)	167.01	173.27	180.33	170.58	168.62	165.00	172.15	174.68	174.89	169.40
xi,3 (nmol/mol)	167.62	173.89	180.58	170.86	168.71	165.30	172.40	174.95	174.50	170.10
Xi (nmol/mol)	166.53	172.54	179.60	169.98	167.89	165.16	171.42	173.99	174.74	168.60
Si (nmol/mol)	1.39	1.81	1.48	1.28	1.33	0.15	1.48	1.43	0.21	2.02
u(xi) (nmol/mol)	3.03	3.52	6.32	3.05	3.10	3.80	1.77	2.75	3.23	3.52
U(xi) (nmol/mol)	6.07	7.04	12.64	6.10	6.20	7.60	3.54	5.50	6.47	7.04

Table 33: Reported values for NO run 5.

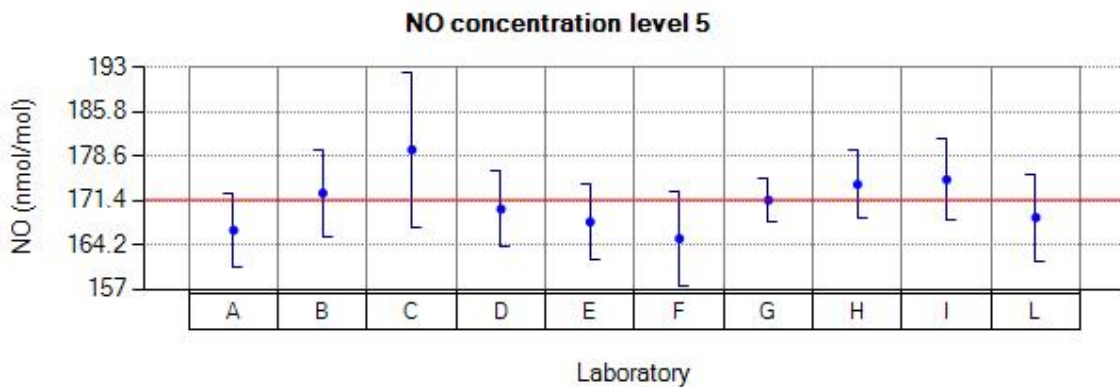


Figure 36: Reported values for NO run 5.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
xi,1 (nmol/mol)	117.82	123.90	127.72	120.83	119.20	115.00	121.79	123.89	125.82	120.00
xi,2 (nmol/mol)	117.55	123.64	127.56	120.69	119.05	115.00	121.74	123.82	125.64	120.50
xi,3 (nmol/mol)	117.66	123.71	127.52	120.83	119.23	115.00	121.78	123.85	125.32	120.30
Xi (nmol/mol)	117.67	123.75	127.60	120.78	119.16	115.00	121.77	123.85	125.59	120.26
Si (nmol/mol)	0.13	0.13	0.10	0.08	0.09	0.00	0.02	0.03	0.25	0.25
u(xi) (nmol/mol)	2.37	2.41	4.64	1.87	2.90	2.60	0.99	1.97	2.32	2.51
U(xi) (nmol/mol)	4.74	4.82	9.27	3.73	5.80	5.30	1.98	3.94	4.65	5.02

Table 34: Reported values for NO run 6.

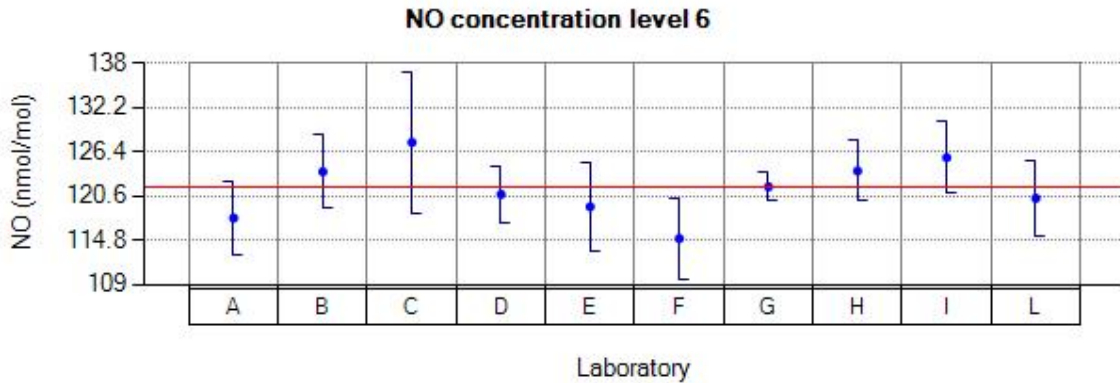


Figure 37: Reported values for NO run 6.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
xi,1 (nmol/mol)	248.66	258.27	269.00	255.59	252.45	255.20	257.37	260.94	261.38	253.60
xi,2 (nmol/mol)	250.07	259.77	269.84	256.13	253.24	255.00	258.45	261.83	261.41	254.70
xi,3 (nmol/mol)	250.75	259.90	269.82	256.17	253.11	255.10	258.60	261.99	262.10	254.70
Xi (nmol/mol)	249.82	259.31	269.55	255.96	252.93	255.10	258.14	261.58	261.63	254.33
Si (nmol/mol)	1.06	0.90	0.47	0.32	0.42	0.10	0.67	0.56	0.40	0.63
u(xi) (nmol/mol)	4.16	5.08	9.25	3.81	4.70	5.90	2.04	4.07	4.84	5.31
U(xi) (nmol/mol)	8.31	10.16	18.50	7.63	9.40	11.70	4.09	8.15	9.68	10.63

Table 35: Reported values for NO run 7.

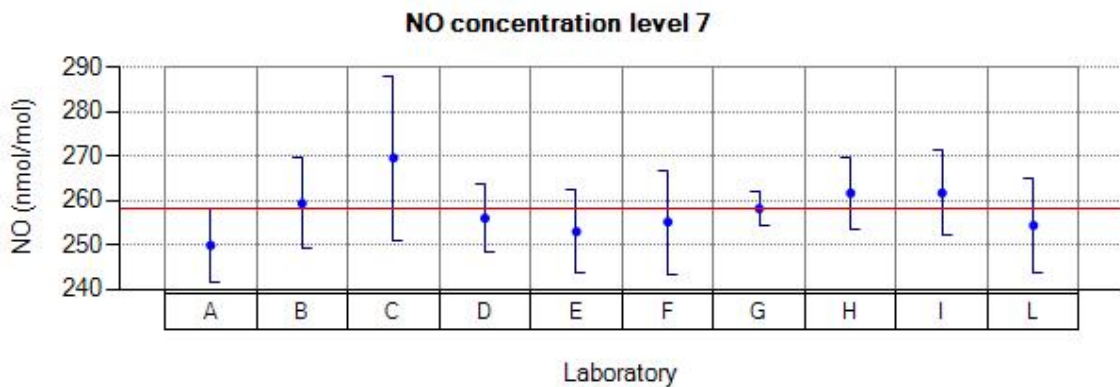


Figure 38: Reported values for NO run 7.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
xi,1 (nmol/mol)	167.43	175.78	180.83	171.74	169.40	168.30	173.23	175.57	178.55	171.10
xi,2 (nmol/mol)	167.12	175.58	180.62	171.77	169.35	168.30	173.13	175.28	178.45	171.00
xi,3 (nmol/mol)	167.48	175.78	180.35	171.29	168.97	168.40	172.75	174.85	178.03	171.00
Xi (nmol/mol)	167.34	175.71	180.60	171.60	169.24	168.33	173.03	175.23	178.34	171.03
Si (nmol/mol)	0.19	0.11	0.24	0.26	0.23	0.05	0.25	0.36	0.27	0.05
u(xi) (nmol/mol)	3.04	3.43	6.33	2.55	3.00	3.90	1.37	2.75	3.30	3.57
U(xi) (nmol/mol)	6.07	6.86	12.66	5.09	6.00	7.70	2.74	5.50	6.60	7.15

Table 36: Reported values for NO run 8.

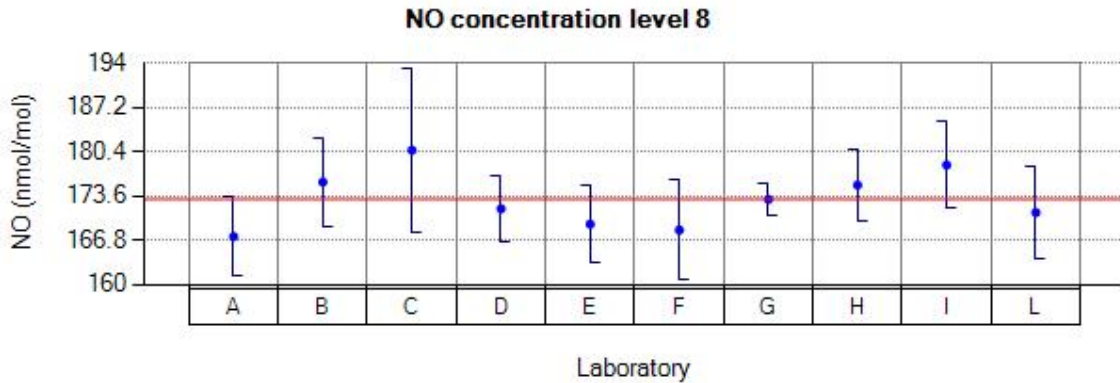


Figure 39: Reported values for NO run 8.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
xi,1 (nmol/mol)	18.92	20.29	21.48	19.51	19.09	20.10	20.08	20.14	18.30	20.30
xi,2 (nmol/mol)	19.92	21.23	22.38	20.56	19.92	20.20	21.02	21.10	18.60	20.30
xi,3 (nmol/mol)	20.11	21.10	22.49	20.48	19.97	20.20	21.12	21.28	18.83	20.30
Xi (nmol/mol)	19.65	20.87	22.11	20.18	19.66	20.16	20.74	20.84	18.57	20.30
Si (nmol/mol)	0.63	0.50	0.55	0.58	0.49	0.05	0.57	0.61	0.26	0.00
u(xi) (nmol/mol)	1.04	0.50	1.89	1.41	1.00	0.70	0.49	0.39	0.75	0.42
U(xi) (nmol/mol)	2.10	1.00	3.79	2.82	2.00	1.50	0.98	0.79	1.50	0.85

Table 37: Reported values for NO run 9.

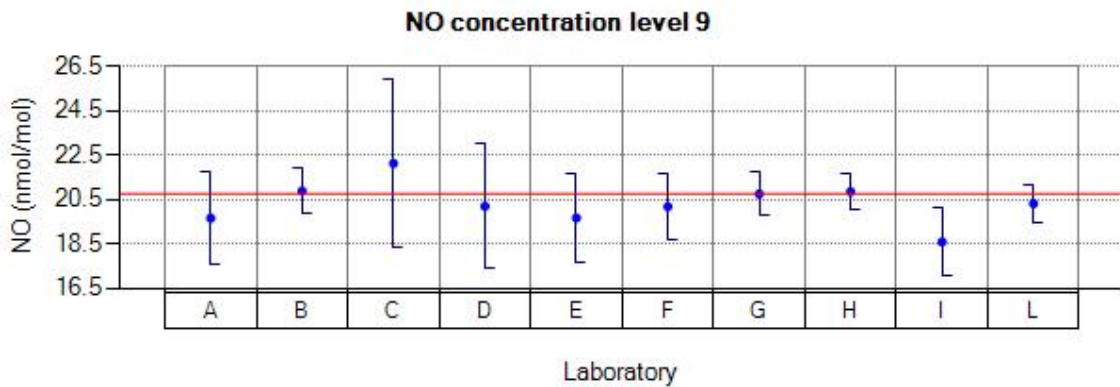


Figure 40: Reported values for NO run 9.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
xi,1 (nmol/mol)	9.65	10.80	11.16	9.95	9.46	10.00	10.34	10.44	8.04	10.20
xi,2 (nmol/mol)	9.77	10.74	11.08	9.82	9.42	10.10	10.28	10.35	8.13	10.10
xi,3 (nmol/mol)	9.59	10.77	11.10	9.90	9.49	10.00	10.24	10.27	8.13	9.80
Xi (nmol/mol)	9.67	10.77	11.11	9.89	9.45	10.03	10.28	10.35	8.10	10.03
Si (nmol/mol)	0.09	0.03	0.04	0.06	0.03	0.05	0.05	0.08	0.05	0.20
u(xi) (nmol/mol)	0.91	0.21	1.78	1.32	0.80	0.60	0.31	0.29	0.75	0.40
U(xi) (nmol/mol)	1.82	0.42	3.55	2.64	1.60	1.20	0.62	0.57	1.50	0.80

Table 38: Reported values for NO run 10.

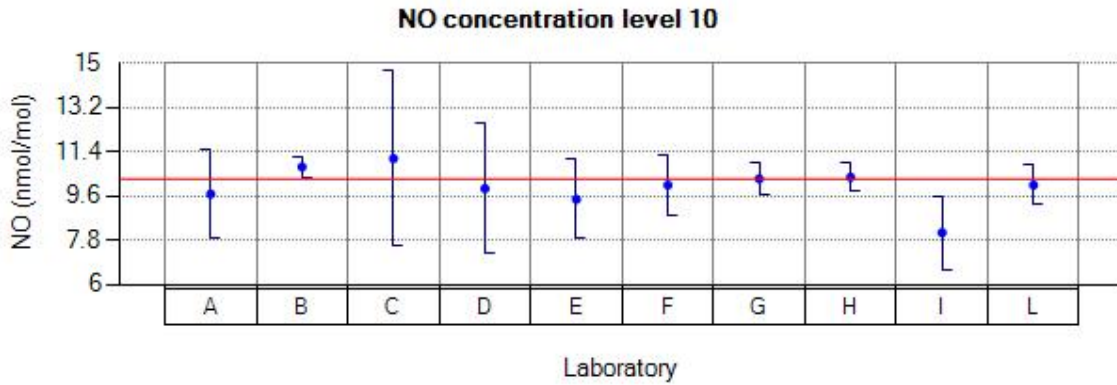


Figure 41: Reported values for NO run 10.

Reported values for NO₂

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
xi,1 (nmol/mol)	0.96	0.08	0.46	-0.25	0.10	0.10	-0.01	0.05	-0.05	1.60
u(xi) (nmol/mol)	0.78	0.01	0.30	1.87	0.80	0.50	0.31	0.37	0.75	0.40
U(xi) (nmol/mol)	1.56	0.02	0.59	3.74	1.60	1.00	0.62	0.73	1.50	0.80

Table 39: Reported values for NO₂ run 0.

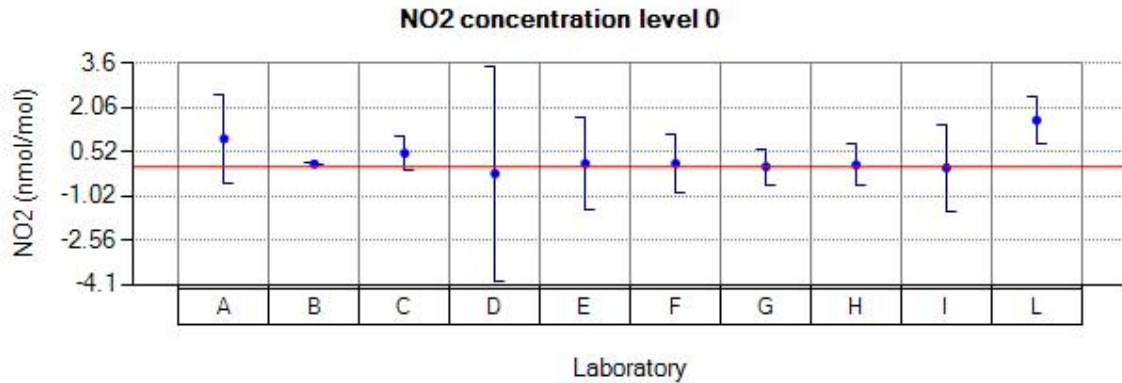


Figure 42: Reported values for NO₂ run 0.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
xi,1 (nmol/mol)	117.99	119.10	120.19	118.04	111.18	126.80	120.98	121.80	112.20	115.50
xi,2 (nmol/mol)	118.17	118.69	119.77	117.83	111.29	126.70	120.60	121.30	112.10	115.50
xi,3 (nmol/mol)	117.13	118.40	120.03	118.11	111.06	126.70	120.33	121.32	112.32	115.70
Xi (nmol/mol)	117.76	118.73	119.99	117.99	111.17	126.73	120.63	121.47	112.20	115.56
Si (nmol/mol)	0.55	0.35	0.21	0.14	0.11	0.05	0.32	0.28	0.11	0.11
u(xi) (nmol/mol)	2.37	3.28	4.40	7.35	2.50	3.20	4.54	2.98	2.07	2.41
U(xi) (nmol/mol)	4.75	6.57	8.81	14.70	5.00	6.30	9.09	5.96	4.15	4.83

Table 40: Reported values for NO₂ run 2.

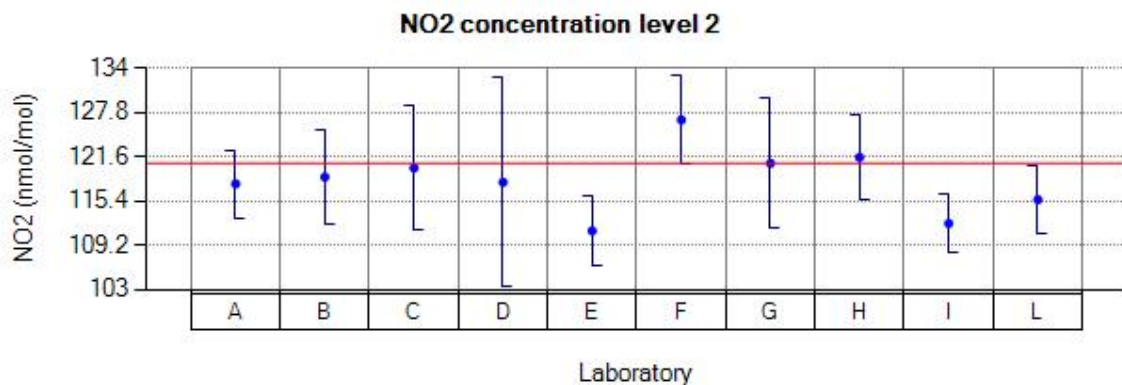


Figure 43: Reported values for NO₂ run 2.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
xi,1 (nmol/mol)	23.24	22.28	23.42	22.96	21.54	20.20	22.84	23.18	20.27	21.90
xi,2 (nmol/mol)	23.46	21.97	23.37	23.01	21.41	20.30	22.87	23.05	20.42	21.70
xi,3 (nmol/mol)	23.35	22.08	23.38	22.93	21.43	20.20	22.82	23.14	20.48	22.00
Xi (nmol/mol)	23.35	22.11	23.39	22.96	21.46	20.23	22.84	23.12	20.39	21.86
Si (nmol/mol)	0.11	0.15	0.02	0.04	0.07	0.05	0.02	0.06	0.10	0.15
u(xi) (nmol/mol)	1.10	0.62	1.91	1.48	1.00	0.80	0.63	0.61	0.75	0.46
U(xi) (nmol/mol)	2.19	1.24	3.81	2.95	2.00	1.50	1.25	1.22	1.50	0.91

Table 41: Reported values for NO₂ run 4.

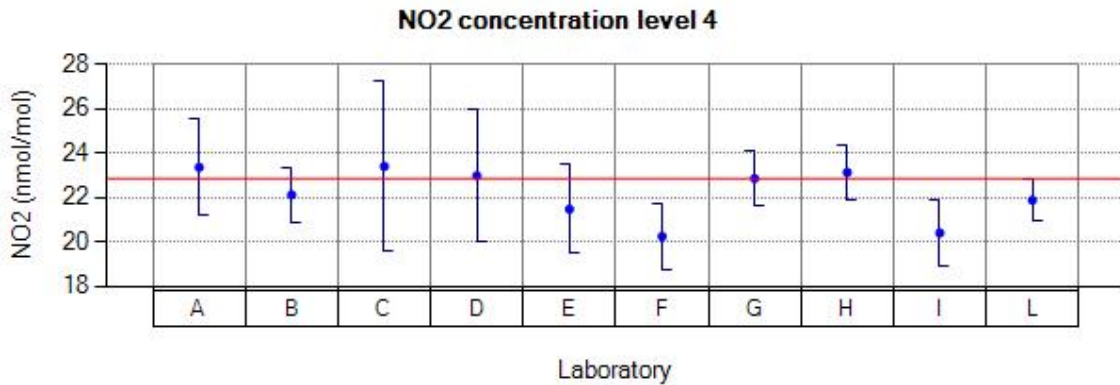


Figure 44: Reported values for NO₂ run 4.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
xi,1 (nmol/mol)	51.55	50.38	52.48	51.38	48.46	50.80	52.07	52.77	49.34	49.40
xi,2 (nmol/mol)	51.68	50.60	52.63	51.55	48.57	50.80	51.96	52.80	49.00	49.40
xi,3 (nmol/mol)	51.20	50.67	52.68	51.39	48.54	50.90	52.05	52.74	49.29	49.40
Xi (nmol/mol)	51.47	50.55	52.59	51.44	48.52	50.83	52.02	52.77	49.21	49.40
Si (nmol/mol)	0.24	0.15	0.10	0.09	0.05	0.05	0.05	0.03	0.18	0.00
u(xi) (nmol/mol)	1.48	1.40	2.48	2.37	1.50	1.30	1.54	1.23	0.91	1.03
U(xi) (nmol/mol)	2.95	2.80	4.96	4.75	3.00	2.50	3.08	2.46	1.82	2.06

Table 42: Reported values for NO₂ run 6.

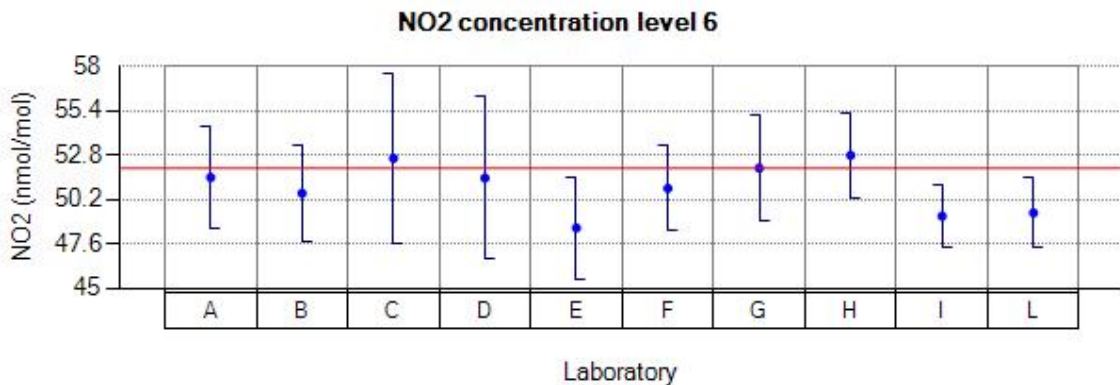


Figure 45: Reported values for NO₂ run 6.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
xi,1 (nmol/mol)	84.58	84.40	87.44	85.49	81.30	86.70	86.80	87.88	83.81	82.90
xi,2 (nmol/mol)	84.44	85.06	87.65	85.72	81.22	86.70	87.00	87.71	83.54	82.80
xi,3 (nmol/mol)	85.35	85.28	87.91	85.82	81.67	86.60	87.20	87.77	84.20	83.10
\bar{X}_i (nmol/mol)	84.79	84.91	87.66	85.67	81.39	86.66	87.00	87.78	83.85	82.93
S_i (nmol/mol)	0.49	0.45	0.23	0.16	0.24	0.05	0.20	0.08	0.33	0.15
$u(x_i)$ (nmol/mol)	1.92	2.36	3.43	3.57	2.00	2.20	2.24	1.93	1.55	1.73
$U(x_i)$ (nmol/mol)	3.84	4.72	6.86	7.15	4.00	4.30	4.47	3.86	3.10	3.46

Table 43: Reported values for NO₂ run 8.

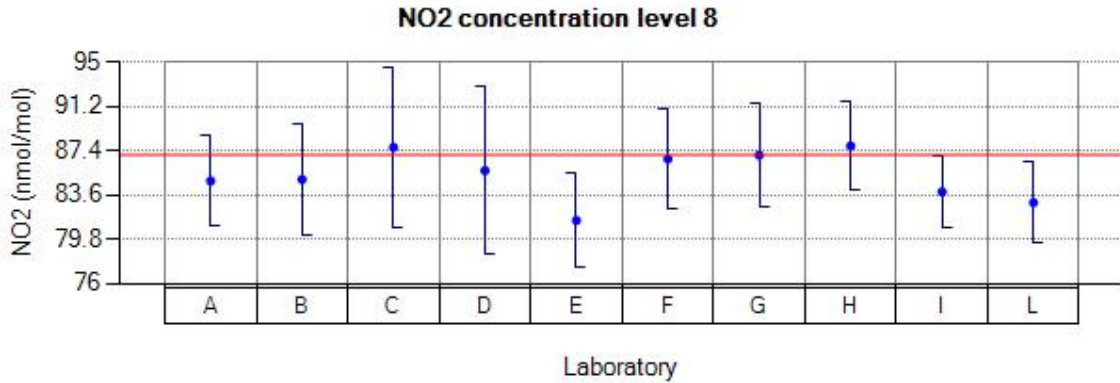


Figure 46: Reported values for NO₂ run 8.

values	laboratories									
	A	B	C	D	E	F	G	H	I	L
xi,1 (nmol/mol)	12.57	10.66	11.50	11.27	10.47	10.00	11.06	11.10	10.87	10.40
xi,2 (nmol/mol)	12.59	10.66	11.58	11.35	10.66	10.00	11.09	11.10	10.74	10.40
xi,3 (nmol/mol)	12.77	10.69	11.56	11.32	10.47	9.80	11.04	11.16	10.71	10.60
\bar{X}_i (nmol/mol)	12.64	10.67	11.54	11.31	10.53	9.93	11.06	11.12	10.77	10.46
S_i (nmol/mol)	0.11	0.01	0.04	0.04	0.11	0.11	0.02	0.03	0.08	0.11
$u(x_i)$ (nmol/mol)	0.95	0.29	1.78	1.73	0.80	0.60	0.36	0.43	0.75	0.40
$U(x_i)$ (nmol/mol)	1.90	0.58	3.56	3.46	1.60	1.30	0.73	0.85	1.50	0.80

Table 44: Reported values for NO₂ run 10.

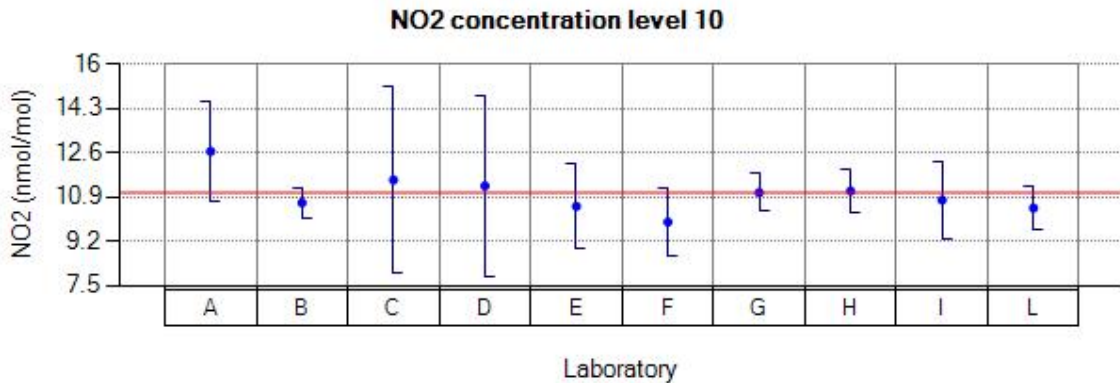


Figure 47: Reported values for NO₂ run 10.

Annex C. The precision of standardized measurement methods

For the main purpose of monitoring trends between different IE undertaken by ERLAP the precision of standardized SO₂, CO, O₃ and NO_x measurement methods [2], [3], [4] and [5] as implemented by NRLs was evaluated. Applied methodology is described in ISO 5725-1, -2 and -6 [14], [15] and [16]. The precision experiment has involved a total of nine laboratories the actual number of labs (p_j) varying from run to run (Table 45). Six concentration levels (for run 0 is requested only one value so repeatability cannot be evaluated) were tested for O₃, CO, SO₂ and NO₂, and eleven for NO. Outlier tests were performed and results are reported in Annex D.

The repeatability standard deviation (s_r) was calculated in accordance with ISO 5725-2 as the square root of average within laboratory variance. The repeatability limit (r) is calculated using Equation 8 [16]. It represents the biggest difference between two test results found on an identical test gas by one laboratory using the same apparatus within the shortest feasible time interval, that should not be exceeded on average more than once in 20 cases in the normal and correct operation of method.

$$r = t_{95\%,v} \cdot \sqrt{2} \cdot s_r \quad \text{Equation 8}$$

The reproducibility standard deviation (s_R) was calculated in accordance with ISO 5725-2 as the square root of sum of repeatability and between laboratory variance. The reproducibility limit (R) is calculated using Equation 9 [16]. It represents the biggest difference between two measurements on an identical test gas reported by two laboratories, which should not occur on average more than once in 20 cases in the normal and correct operation of method.

$$R = t_{95\%,v} \cdot \sqrt{2} \cdot s_R \quad \text{Equation 9}$$

The repeatability standard deviation was evaluated with (p_j*(3-1)) degrees of freedom (v) and reproducibility standard deviation with (p_j-1) degrees of freedom. The critical range student factors (t_{α,v}) are reported in Table 45.

parameter	run	p _j	t critical value 95% for r	t critical value 95% for R
CO	1,2,3,4,5	10	2.086	2.228
NO	1,2,3,4,5,6,7,8,9,10	10	2.086	2.228
NO ₂	1,2,3,4,5,6,7,8,9,10	10	2.086	2.228
O ₃	1,2,3,4,5	9	2.101	2.262
SO ₂	1,2,3,4,5	9	2.101	2.262

Table 45: Critical values of t used in the repeatability (r) and reproducibility (R) evaluation.

The repeatability (r) and reproducibility (R) limits of measurement methods are presented from Table 46 to Table 50 and from Figure 48 to Figure 52. It is also reported the 'reproducibility from common criteria (R (from σ_p))' calculated by substituting s_R in Equation 9 with a 'standard deviation for proficiency assessment' (Table 4). Comparison between R and R (from σ_p) serves to indicate that σ_p is realistic ([13] 6.3.1) or from the other point of view, that the general methodology implemented by NRLs is appropriate for σ_p .

SO ₂ data (nmol/mol) without outliers			
group average	repeatability limit : r	reproducibility limit : R	reproducibility limit (relative)
-0.1		0.0	
3.4	0.3	2.7	
8.7	0.3	2.2	
20.0	0.5	2.0	
51.4	0.5	4.0	
123.8	0.7	12.0	9.7%

Table 46: The R and r of SO₂ standard measurement method.

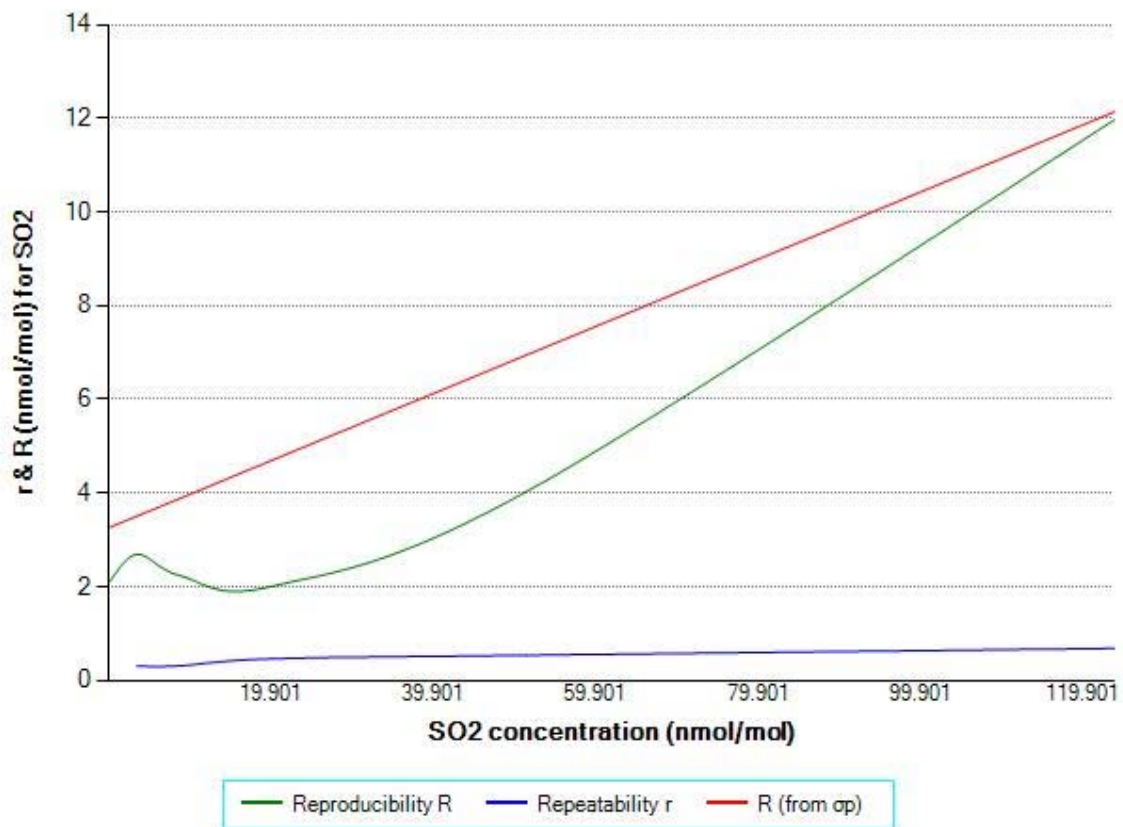


Figure 48: The R and r of SO₂ standard measurement method as a function of concentration.

CO data (µmol/mol) without outliers			
group average	repeatability limit : r	reproducibility limit : R	reproducibility limit (relative)
-0.001		0.149	
1.029	0.008	0.22	
3.003	0.011	0.513	
4.477	0.023	0.496	
5.942	0.012	0.58	
7.864	0.014	0.741	9.4%

Table 47: The R and r of CO standard measurement method.

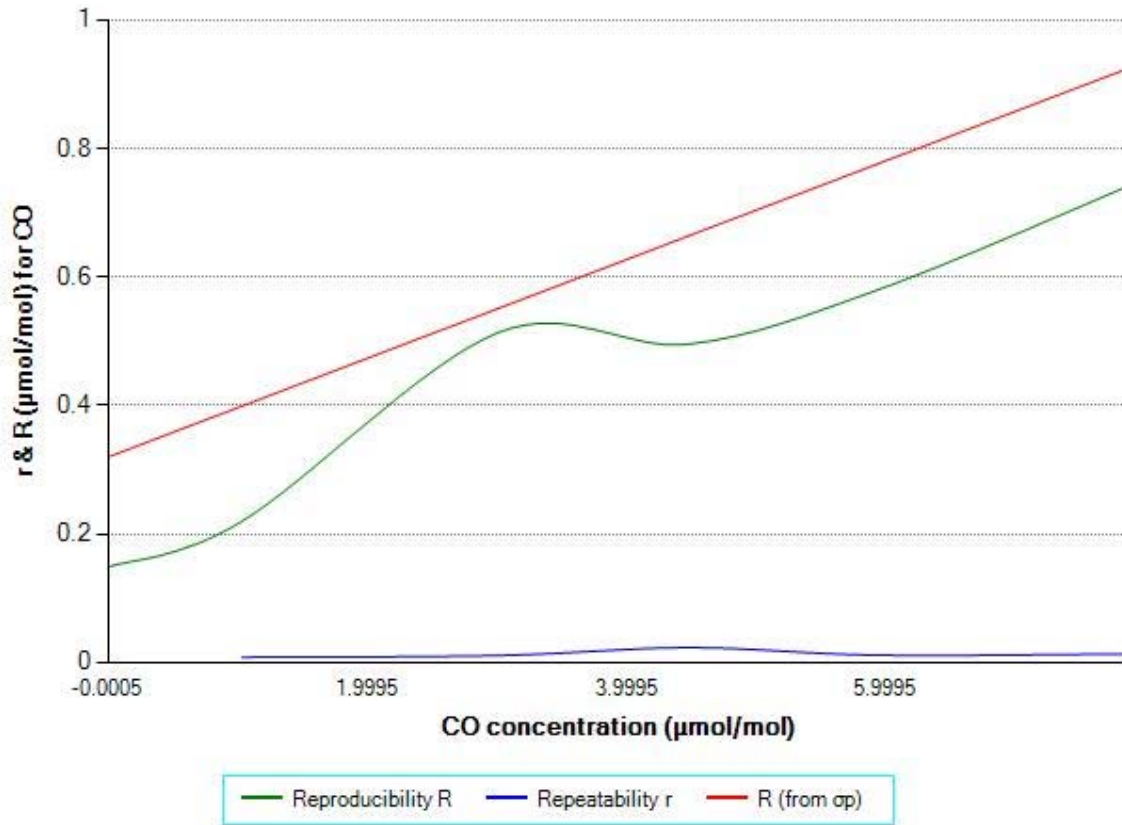


Figure 49: The R and r of CO standard measurement method as a function of concentration.

O ₃ data (nmol/mol) without outliers			
group average	repeatability limit : r	reproducibility limit : R	reproducibility limit (relative)
0.0		1.8	
10.8	0.1	1.9	
22.2	0.2	1.8	
50.1	0.3	2.4	
84.6	0.4	3.8	
115.2	0.7	6.6	5.7%

Table 48: The R and r of O₃ standard measurement method.

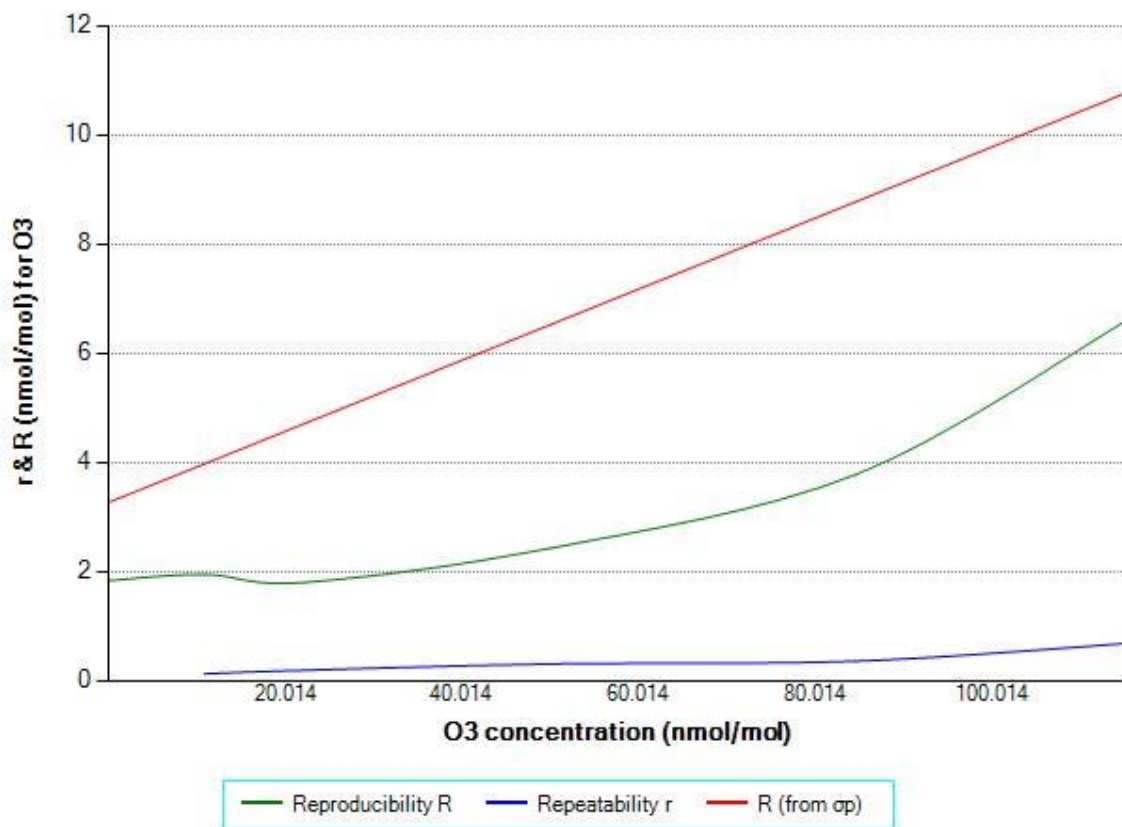


Figure 50: The R and r of O₃ standard measurement method as a function of concentration.

NO data (nmol/mol) without outliers			
group average	repeatability limit : r	reproducibility limit : R	reproducibility limit (relative)
0.0		0.8	
10.0	0.3	2.6	
20.3	1.4	3.3	
39.2	0.2	6.2	
61.1	0.5	8.3	
121.5	0.4	12.1	
171.0	4.1	14.4	
173.0	0.7	14.0	
257.8	1.8	17.9	
400.5	0.9	29.3	
518.3	3.1	32.4	6.3%

Table 49: The R and r of NO standard measurement method.

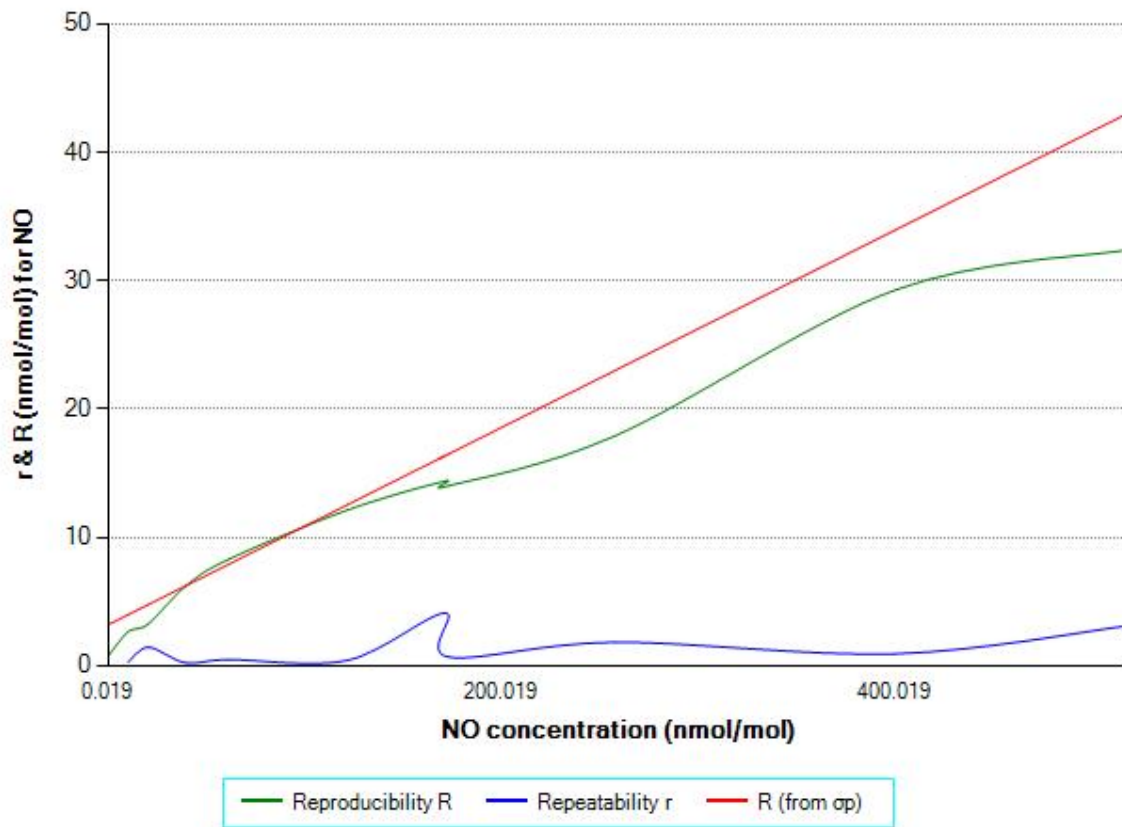


Figure 51: The R and r of NO standard measurement method as a function of concentration.

NO ₂			
group average	repeatability limit : r	reproducibility limit : R	reproducibility limit (relative)
0.0		1.0	
13.4	0.2	2.3	
20.2	0.1	2.7	
58.8	0.3	6.9	
99.6	0.6	12.6	
119.1	0.8	13.0	10.92%

Table 50: The R and r of NO₂ standard measurement method.

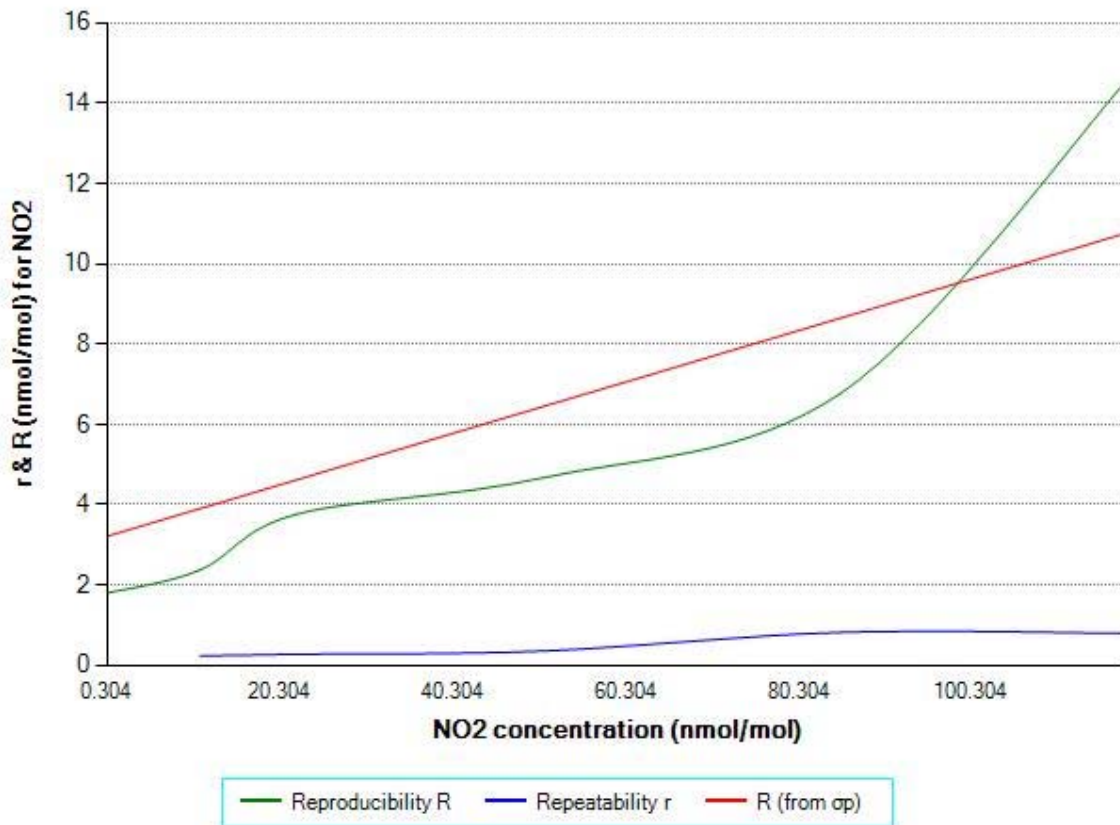


Figure 52: The R and r of NO₂ standard measurement method as a function of concentration.

Annex D. The scrutiny of results for consistency and outlier test

The precision evaluation (Annex C) focuses on data that are as much as possible the reflection of every day work of NRLs and thus represents the comparability of participant's standard operating procedures.

For that reason a procedure for the detection of exceptional errors (error during typing, slip in performing the measurement or the calculation, wrong averaging interval, malfunction of instrumentation, etc.) was applied. In this procedure were carried out tests for data consistency and statistical outliers as described in ISO 5725-2.

Laboratories showing some form of statistical inconsistency were requested to investigate the cause of discrepancies.

Laboratories were allowed to correct their results in case of identification of exceptional errors. Subsequently, data were considered definitive and "Grubb's one outlying observation test" was performed.

For runs where outliers were detected outliers were removed and "Grubb's one outlying observation test" was repeated until no more outliers were observed. Statistical outliers obtained at this stage are not considered as due to extraordinary errors but due to significant difference in participant's standard operating procedure.

During this IE the statistical outliers presented in the table below are related only to a zero level:

parameter	run	laboratory	measured value	failing test	confidence level
SO ₂	0	E	-1.69	G1 minimum	1%, 5%

Table 51: "Genuine" statistical outliers according to Grubb's one outlying observation test.

The precision of standardized measurement methods reported in Annex C are calculated using the database without outliers.

European Commission
EUR – Joint Research Centre – Institute for Environment and Sustainability

Title: Evaluation of the Laboratory Comparison Exercise for SO₂, CO, O₃, NO and NO₂, 3rd-6th October 2011

Author(s): Maurizio Barbieri, Friedrich Lagler

Luxembourg: Publications Office of the European Union

2012 – 63 pp. – 21.0 x 29.7 cm

EUR – Scientific and Technical Research series – ISSN 1831-9424

ISBN 978-92-79-25366-9

doi:10.2788/33270

Abstract

From the 03rd to the 06th of October 2011 in Ispra (IT), 9 Laboratories of AQUILA (Network of European Air Quality Reference Laboratories) met for a laboratory comparison exercise to evaluate their proficiency in the analysis of inorganic gaseous pollutants covered by European Directive about air quality (SO₂, CO, NO, NO₂ and O₃).

The proficiency evaluation, where each participant's bias was compared to two criteria, provides information on the current situation and capabilities to the European Commission and can be used by participants in their quality control system.

On the basis of criteria imposed by the European Commission, 78.5% of the results reported by AQUILA laboratories were good both in terms of measured values and reported uncertainties. Another 20.1% of the results had good measured values, but the reported uncertainties were either too high (12.5%) or too small (7.6%). Four values have been classified in category '5' (1.3%).

Comparability of results among AQUILA participants at the highest concentration level, excluding outliers, is acceptable in SO₂, CO and O₃ measurements while NO and NO₂ measurement methods showed less satisfactory results.

As the Commission's in-house science service, the Joint Research Centre's mission is to provide EU policies with independent, evidence-based scientific and technical support throughout the whole policy cycle.

Working in close cooperation with policy Directorates-General, the JRC addresses key societal challenges while stimulating innovation through developing new standards, methods and tools, and sharing and transferring its know-how to the Member States and international community.

Key policy areas include: environment and climate change; energy and transport; agriculture and food security; health and consumer protection; information society and digital agenda; safety and security including nuclear; all supported through a cross-cutting and multi-disciplinary approach.



Publications Office

ISBN 978-92-79-25366-9



9 789279 253669