



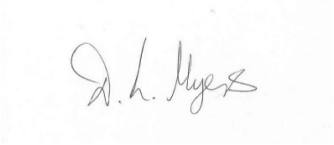
Client Linx Printing Technologies Limited
 Linx House
 33 Edison Road
 St Ives
 Cambridgeshire
 PE27 3LF

Part 1: Executive Summary

Report for the Periodic Monitoring of Emissions to Air.

Site	St Ives
Plant	Ink Manufacturing Plant
Sampling Date	3rd November 2015
Report Date	7th December 2015
Job Number	EM-2112
Permit Number	B04/94

Report Prepared by:	Print	Harpreet Badwal
	MCERTS No.	MM03 149 Level 2 TE: 1,2,3,4

Report Approved by:	Sign	
	Print	Derek Myers
	MCERTS No.	MM02 115 Level 2 TE: 1,2,3,4



1783

REC Ltd Environmental Monitoring
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 Bordesley Green Road
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Contents

Page 1	Part 1: Executive Summary
Page 2	Contents
Page 3	Monitoring Objectives
Page 3	Special Monitoring Requirements
Page 3	Summary Of Methods
Page 4	Summary Of Results
Page 5	Summary Of Results, Exhaust Gases
Page 6	Operating Information
Page 6	Comments On Monitoring Procedures
Page 7	Part 2: Supporting Information
Page 8	Appendix 1
Page 9	Emission Monitoring Procedures and Instrumentation
Page 10	Sampling Personnel
Page 10	Equipment References
Page 10	Subcontracted Analysis
Page 11	Appendix 2
Page 12	Preliminary Velocity Traverse & Moisture Data
Page 13	Exhaust Gases - Continuous Analysis Data
Page 14	Exhaust Gases - Continuous Analysis Data, Continued
Page 15	Chart 1 - VOC Emissions Data
Page 16	Total Particulate Matter - Sampling Data
Page 17	Total Particulate Matter - Emission Results
Page 18	Appendix 3
Page 19	Diagram of Sampling Location
Page 20	Generic Calculations
Page 21	Flow Calculations
Page 22	Concentration Calculation
Page 23	Uncertainty Estimate Calculations - Manual Techniques
Page 24	Uncertainty Estimate Calculations - Instrumental Techniques
Page 25	Appendix 4
Page 26	Uncertainty Estimates:- TOC
Page 27	Uncertainty Estimates:- Total Particulate Matter

Monitoring Objectives

The monitoring was undertaken to check compliance with authorised emission limits.

All monitoring procedures were carried out to the MCERTS requirements under the REC Environmental Monitoring quality system to ISO 17025: 2005.

Monitoring was undertaken for the listed emissions from the following sampling positions:

Sampling Location	Emission
Ink Manufacturing Plant	Particulates Total organic carbon

Special Monitoring Requirements

There were no special requirements for this monitoring campaign.

Summary of Methods

Emission	Method number	Method standard
Particulate	TPM/04	BS EN 13284-1 : 2002
Gas velocity and volume flow	TPM/01A	BS EN ISO 16911-1:2013
Water vapour	TPM/37	BS EN 14790 : 2005
TOC	TPM/13	BS EN 12619 : 2013

Summary Of Results

The table presents the atmospheric emissions from the tests undertaken on behalf of **Linx Printing Technologies Limited**
The results were measured from the sample positions downstream of the arrestment plant.

Emission at St Ives Ink Manufacturing Plant	Sampling Time			Emission Result	Authorised Limit	Uncertainty +/-	Detection Limit	Mass Emission
	Date	Start	End	mg/m ³ *	mg/m ³ *	mg/m ³ *	mg/m ³ *	g/h
TOC	03/11/15	12:45	16:45	309.4	150	6.3	0.2	822.3
s Particulate	03/11/15	11:49	19:49	ND	20	-	0.05	0.15

* at reference conditions	Stack Gas Weight 29.00 Kg/kmol	0 °C 101.3 kPa	Without correction for moisture Oxygen No Correction %
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Where applicable

Oxides of nitrogen results are expressed as nitrogen dioxide
TOC results are expressed as total carbon

Throughout Report:

* Reference conditions (see above)
** Analysis not required
ND Non detectable
s - Subcontracted laboratory analysis

Nm³ 273 K, 101.3 kPa
- UKAS accredited only
- Not Accredited
N/A Not applicable

The reported expanded uncertainty is based on a standard uncertainty multiplied by a coverage factor k=2, providing a 95% confidence level. The uncertainty evaluation has been carried out in accordance with UKAS requirements.



All tests included in this report are accredited under UKAS and MCERTS accreditation schemes unless otherwise stated.
Opinions and interpretations expressed herein are outside the scope of MCERTS and UKAS accreditation.



Summary Of Results, Exhaust Gases

The table presents the atmospheric emissions from the tests undertaken on behalf of **Linx Printing Technologies Limited**. The results were measured from the sample positions downstream of the arrestment plant.

Emission at St Ives Ink Manufacturing Plant		Sampling Time			Emission Result	Authorised Limit	Uncertainty +/-	Detection Limit	Mass Emission
		Date	Start	End	mg/m ³ *	mg/m ³ *	mg/m ³ *	mg/m ³ *	g/h
TOC	Test 1	03/11/15	12:45	13:15	355.4	150	7.2	0.2	944.7
TOC	Test 2	03/11/15	13:15	13:45	359.4	150	7.3	0.2	955.2
TOC	Test 3	03/11/15	13:45	14:15	275.8	150	5.6	0.2	733.2
TOC	Test 4	03/11/15	14:15	14:45	233.0	150	4.8	0.2	619.5
TOC	Test 5	03/11/15	14:45	15:15	189.3	150	4.0	0.2	503.2
TOC	Test 6	03/11/15	15:15	15:45	205.1	150	4.3	0.2	545.1
TOC	Test 7	03/11/15	15:45	16:15	551.2	150	11.1	0.2	1465.1
TOC	Test 8	03/11/15	16:15	16:45	305.8	150	6.2	0.2	812.7

* at ref	Stack Gas Weight	0 °C	Without correction for moisture
Conditions	29.00 Kg/kmol	101.3 kPa	Oxygen No Correction %

Where applicable

Oxides of nitrogen results are expressed as nitrogen dioxide
TOC results are expressed as total carbon

Throughout Report:

* Reference conditions (see above)
** Analysis not required
ND Non detectable
s - Subcontracted laboratory analysis

Nm³ 273 K, 101.3 kPa
- UKAS accredited only
- Not Accredited
N/A Not applicable

The reported expanded uncertainty is based on a standard uncertainty multiplied by a coverage factor k=2, providing a 95% confidence level. The uncertainty evaluation has been carried out in accordance with UKAS requirements.



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Operating Information

The table below shows details of the operating information on each sampling date for:

Ink Manufacturing Plant

Date	Process type	Process duration	Fuel	Feedstock	Abatement	Load
03/11/2015	Ink Manufacturing Plant	Continuous	N/A	Methyl Ethyl Ketone Acetone Ethyl Lactate Diethyl Ketone	None	Normal Operation

There are no CEM's available on this process.

Comments & Monitoring Deviations

A waste gas homogeneity test to BS EN 15259:2007 (MID) is not required:-

The homogeneity test is not applicable to non-combustion processes.

The homogeneity test is not applicable to duct areas less than 1m².

All monitoring was performed in accordance with the relevant procedures.

The sampling location is a vertical duct.

Due to the positioning of the scaffolding erected, sampling can only be undertaken from one sample port.

The particulate monitoring was within the required 95 to 115 % isokinetic rate as stated in BS EN 13284-1: 2002

The velocity and temperature profile at the sampling location met the requirements of BS EN 13284-1: 2002.

The calculated particulate emission level was greater than the control blank.

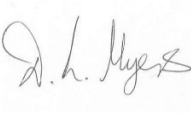
When the results are expressed as non-detected the mass emissions are calculated from the detection limit and therefore they are worst case results.



Part 2: Supporting Information

Report for the Periodic Monitoring of Emissions to Air.

Client	Linx Printing Technologies Limited
Site	St Ives
Plant	Ink Manufacturing Plant
Sampling Date	3rd November 2015
Report Date	7th December 2015
Job Number	EM-2112
Permit Number	B04/94

Report Prepared by:	Print	Harpreet Badwal	
	MCERTS No.	MM03 149	Level 2 TE: 1,2,3,4
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	MCERTS No.	MM02 115	Level 2 TE: 1,2,3,4



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APPENDIX 1

Emission Monitoring Procedures And Instrumentation

Gas velocity and temperature

Documented in-house procedure TPM01/A to the main procedural requirements of BS EN ISO 16911-1:2013. Velocity and temperature measurements are performed using a calibrated Pitot tube, a calibrated pressure differential reading device and a calibrated thermocouple. Velocity and possible flow deviation measurements are carried out at selected, representative points in the gas stream.

Water vapour

Documented in-house procedure TPM/37 to the main procedural requirements of BS EN 14790:2005. A measured volume of gas is extracted from the gas stream through a moisture trapping unit. The mass gain of moisture trapped is divided by the volume of gas sampled to determine the mass concentration of water vapour. For water saturated stacks the temperature of the gas stream is measured and the water vapour concentration is determined using liquid-gas equilibrium tables. Sampling points are selected in accordance with the findings of any BS EN 15259 assessment.

Total particulate matter

Documented in-house procedure TPM/04 to the main procedural requirements of BS EN 13284-1:2002. Stack gases are extracted from representative sampling points at isokinetic flow rates through a sharp-edged nozzle. Particulate matter is collected on a pre-weighed filter conditioned at 180°C. Deposits upstream of the filter are also recovered and weighed. The increase of mass of the filter and mass collected upstream of the filter is divided by the volume sampled to determine the mass concentration.

Total organic carbon

Documented in-house procedure TPM/13 to the main procedural requirements of BS EN 12619:2013. Continuous analysis using probe, sample line and multi range Flame Ionisation Detector (FID) analyser. The analyser is calibrated before and during the tests using certified gas mixtures of nitrogen, oxygen and propane. Sampling points are selected in accordance with the findings of any BS EN 15259 assessment.

Sampling Project Personnel Competency And Expiry Dates

		MCERTS No	Level 1	Level 2	TE1	TE2	TE3	TE4
Report prepared by:	Harpreet Badwal	MM03 149	-	30/09/2018	30/09/2018	30/04/2019	31/10/2019	31/10/2019
Report authorised by:	Derek Myers	MM02 115	-	30/09/2017	31/05/2018	30/11/2018	30/11/2018	30/11/2018
Team leader:	Harpreet Badwal	MM03 149	-	30/09/2018	30/09/2018	30/04/2019	31/10/2019	31/10/2019
Team leader:	Aidan Wrynn	MM08 921	-	31/05/2017	31/05/2017	31/11/2018	30/11/2017	30/06/2018

Equipment References

Equipment	Reference Number
FID	AQ271
Heated Line	HL40
Heated Filter	Sintered
Probe	AQ187
Control Box	CU16
Timer / Stopwatch	CU16/7 & ST41
Barometer	WS03
Pitot	PT116 & PT97
Callipers	CV15
Hot Box	HB19
Impinger Thermocouple	TV24
Sample Rate Calculation Equipment	Laptop
Thermometer	CU16/3 & TK28
Manometer	CU16/1 & PI03
Balance	BL21
Weights	W17 & W18

Subcontracted Analysis

REC Environmental Monitoring has, with your approval, used the following sub-contracted laboratories for the laboratory analyses referenced below:

Laboratory:	Scientific Analysis Laboratories
Parameters:	Particulates - Internal method, Gravimetric
Accreditation:	UKAS Accredited testing laboratory number 1549

APPENDIX 2

Date	03/11/2015
Time	11:30
Pitot Cp	0.79

Barometric pressure	101.3	kPa
Duct static pressure	-0.01	kPa
Stack Area	0.196	m ²

Stack Diameter (circular)	0.50	m
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Traverse	Traverse	Depth	ΔP	T	Angle	velocity	Traverse	Depth	ΔP	T	Angle	velocity
Point	Line	cm	mmH ₂ O	°C	°	m/s	Line	cm	mmH ₂ O	°C	°	m/s
1	A	5.0	3.7	20	<15	6.1	B	5.0	2.0	20	<15	4.6
2	A	5.0	3.4	20	<15	5.8	B	5.0	1.5	20	<15	3.9
3	A	5.9	3.1	20	<15	5.6	B	5.9	1.4	20	<15	3.8
4	A	8.9	2.8	20	<15	5.3	B	8.9	1.2	20	<15	3.5
5	A	12.5	2.7	20	<15	5.2	B	12.5	1.1	20	<15	3.4
6	A	17.8	1.7	20	<15	4.2	B	17.8	1.3	20	<15	3.7
7	A	32.2	0.8	20	<15	2.9	B	32.2	1.5	20	<15	3.9
8	A	37.5	0.4	20	<15	2.0	B	37.5	1.7	20	<15	4.2
9	A	41.2	0.8	20	<15	2.9	B	41.2	1.5	20	<15	3.9
10	A	44.1	0.8	20	<15	2.9	B	44.1	1.9	20	<15	4.4
11	A	45.0	0.4	20	<15	2.0	B	45.0	2.4	20	<15	5.0
12	A	45.0	0.7	20	<15	2.7	B	45.0	2.3	20	<15	4.9

Average Pitot DP	1.60	mmH ₂ O
Average Temperature	293.2	K
Average Velocity	4.0	m/s
Average volumetric flow rate	0.79	m ³ /s at stack conditions
Average volumetric flow rate	0.74	m ³ /s (wet STP)

Sampling plane requirements Re: BS EN 13284-1:2001 5.2

a	Angle of gas flow less than 15° with regard to duct axis	YES
b	No local negative flow	YES
c	Minimum pitot greater than 5Pa	NO
d	Ratio of highest to lowest local gas velocity less than 3:1	YES
	Minimum local gas velocity	2.0
	Maximum local gas velocity	6.1
	Ratio of highest to lowest local gas velocity	3.00

Moisture Determination BS EN 14790:2005

	Volume	Temp	Pressure
	m ³	°C	mbar
Meter start	3.095	13	1013
Meter end	13.668	25	1013
Meter Yd	1.052		
Gas volume	10.399		

Impinger	1	2	3	4
Mass start (g)	763.5	716.6	607.4	897.2
Mass End (g)	773.2	722.2	613.8	926.5
Total Mass collected (g)	51.0			

Water vapour content (% v/v)	0.6
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If water droplets are present in the gas, the water vapour content is calculated using BS EN 14790 Annex A

N/A % v/v

Preliminary Velocity Traverse & Moisture Data

Date 03/11/2015

From 12:45 to 13:15 30 minute mean

Volatile organic compounds	vppm, wet	221.13	mg/m ³ *	355.39
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From 13:15 to 13:45 30 minute mean

Volatile organic compounds	vppm, wet	223.60	mg/m ³ *	359.35
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From 13:45 to 14:15 30 minute mean

Volatile organic compounds	vppm, wet	171.64	mg/m ³ *	275.84
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From 14:15 to 14:45 30 minute mean

Volatile organic compounds	vppm, wet	145.00	mg/m ³ *	233.04
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Sampling Detection Limits

Volatile organic compounds	vppm	0.10	mg/m ³ *	0.16
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Reference Gas Details

Species	Units	Value	Cylinder Reference	Analyser Range	Uncertainty <i>k</i> = 2
Nitrogen	%	99.999	VCK01959	-	± 2
Volatile organic compounds	vppm	74.7	VC59841	10	± 2

Zero And Span Gas Details

Species	units	Initial Time	12:30	Final Time	17:00
		Initial Zero	Initial Span	Final Zero	Final Span
Volatile organic compounds	vppm	0.00	74.70	0.31	75.02

Exhaust Gases Continuous Analysis Data

Date 03/11/2015

From 14:45 to 15:15 30 minute mean

Volatile organic compounds	vppm, wet	117.79	mg/m ³ *	189.30
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From 15:15 to 15:45 30 minute mean

Volatile organic compounds	vppm, wet	127.61	mg/m ³ *	205.08
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From 15:45 to 16:15 30 minute mean

Volatile organic compounds	vppm, wet	342.95	mg/m ³ *	551.17
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From 16:15 to 16:45 30 minute mean

Volatile organic compounds	vppm, wet	190.25	mg/m ³ *	305.76
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Sampling Detection Limits

Volatile organic compounds	vppm	0.10	mg/m ³ *	0.16
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Reference Gas Details

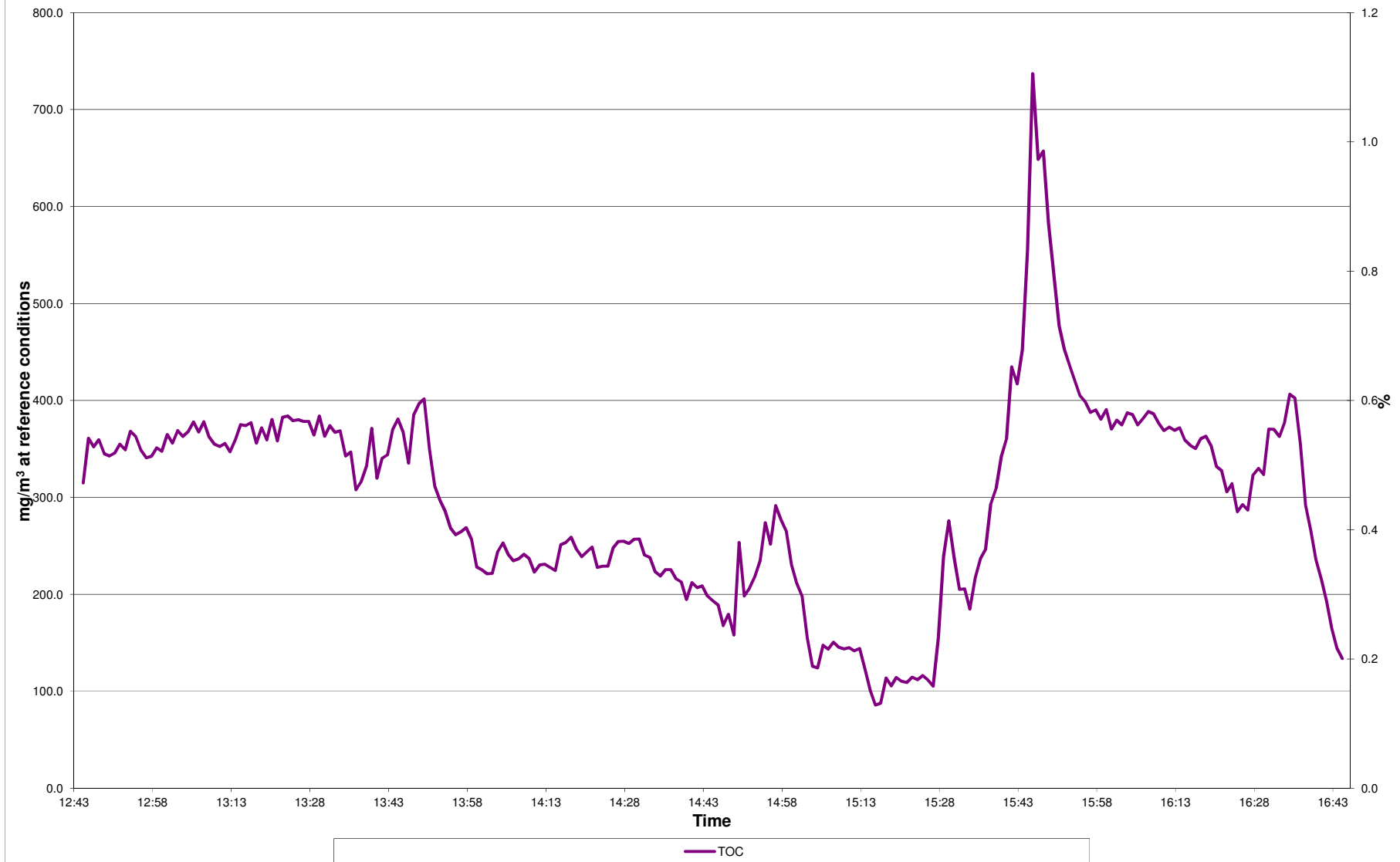
Species	Units	Value	Cylinder Reference	Analyser Range	Uncertainty <i>k</i> = 2
Nitrogen	%	99.999	VCK01959	-	± 2
Volatile organic compounds	vppm	74.7	VC59841	10	± 2

Zero And Span Gas Details

Species	units	Initial Time	12:30	Final Time	17:00
		Initial Zero	Initial Span	Final Zero	Final Span
Volatile organic compounds	vppm	0.00	74.70	0.31	75.02

Exhaust Gases - Continuous Analysis Data, Continued

Chart 1 - VOC Emissions Data - Ink Manufacturing Plant, 03/11/2015



Probe/Pitot ID No	AQ187/PT116	Nozzle Diameter (d)	mm	9.99
Pitot Calibration Factor (Cp)	0.84	In-stack Filtration	Y/N	N
Gas Calibration Factor (Y)	1.052	Leak check OK	Y/N	Y
Barometric Pressure	kPa 101.3	Meter ID No		CU16/2
Duct static Pressure	kPa -0.01	Initial Gas Meter Reading	m³	3.095
Date	03/11/2015	Final Gas Meter Reading	m³	13.668
Start Time	hh:mm 11:49	Sampled Vol, Dry at Meter	m³	10.573
End Time	hh:mm 19:49	Sampled Vol, Dry STP	Nm³	10.298
Sampling Duration	hh:mm 08:00	Moisture content of stack gas	% v/v	0.6
		Isokinetic	%	102.2

Impinger	1	2	3	4
Initial mass (g)	763.5	716.6	607.4	897.2
Final mass (g)	773.2	722.2	613.8	926.5

Sample Point Number	Line	Sampling Time hh:mm	Pitot Reading (h) mm w.g	Temperatures			Orifice Pressure (Dh) mm w.g
				Sample Gas Temperature (Ts) °C	Filter Temperature °C	Meter Temperature (Tm) °C	
1	A	00:00	2.6	20	160	13	65.3
		00:05	2.6	20	160	14	65.3
		00:10	2.6	20	160	15	65.3
		00:15	2.6	20	160	16	65.3
		00:20	2.6	20	160	17	65.3
		00:25	2.6	20	160	18	65.3
		00:30	2.6	20	160	19	65.3
		00:35	2.6	20	160	20	65.3
		00:40	2.6	20	160	20	65.3
		00:45	2.6	20	160	21	65.3
		00:50	2.6	20	160	22	65.3
		00:55	2.6	20	160	22	65.3
		01:00	2.6	20	160	23	65.3
		01:05	2.6	20	160	23	65.3
		01:10	2.6	20	160	23	65.3
		01:15	2.6	20	160	23	65.3
		01:20	2.6	20	160	24	65.3
		01:25	2.6	20	160	24	65.3
		01:30	2.6	20	160	24	65.3
		01:35	2.6	20	160	24	65.3
		01:40	2.6	20	160	24	65.3
		01:45	2.6	20	160	24	65.3
		01:50	2.6	20	160	24	65.3
		01:55	2.6	20	160	25	65.3
		02:00	2.6	20	160	25	65.3
		02:05	2.6	20	160	25	65.3
		02:10	2.6	20	160	25	65.3
		02:15	2.6	20	160	25	65.3
		02:20	2.6	20	160	25	65.3
		02:25	2.6	20	160	25	65.3
		02:30	2.6	20	160	25	65.3
		02:35	2.6	20	160	25	65.3
		02:40	2.6	20	160	25	65.3
		02:45	2.6	21	160	25	65.3
		02:50	2.6	21	160	25	65.3
		02:55	2.6	20	160	25	65.3
		03:00	2.6	20	160	25	65.3
		03:05	2.6	20	160	25	65.3
		03:10	2.6	20	160	25	65.3
		03:15	2.6	20	160	25	65.3
		03:20	2.6	20	160	25	65.3
		03:25	2.6	20	160	25	65.3
		03:30	2.6	20	160	25	65.3
		03:35	2.6	20	160	25	65.3
		03:40	2.6	20	160	25	65.3
		03:45	2.6	20	160	25	65.3
		03:50	2.6	20	160	25	65.3
		03:55	2.6	20	160	25	65.3
2	A	04:00	1.4	20	160	25	35.2
		04:05	1.4	20	160	25	35.2
		04:10	1.4	20	160	24	35.2
		04:15	1.4	20	160	24	35.2
		04:20	1.4	20	160	24	35.2
		04:25	1.4	21	160	24	35.2
		04:30	1.4	21	160	24	35.2
		04:35	1.4	21	160	24	35.2
		04:40	1.4	21	160	24	35.2
		04:45	1.4	21	160	24	35.2
		04:50	1.6	21	160	24	40.2
		04:55	1.6	21	160	24	40.2
		05:00	1.6	21	160	24	40.2
		05:05	1.6	21	160	24	40.2
		05:10	1.4	21	160	24	35.2
		05:15	1.4	21	160	24	35.2
		05:20	1.4	20	160	24	35.2
		05:25	1.4	21	160	24	35.2
		05:30	1.4	21	160	24	35.2
		05:35	1.4	21	160	24	35.2
		05:40	1.6	21	160	24	40.2
		05:45	1.6	21	160	24	40.2
		05:50	1.6	20	160	24	40.2
		05:55	1.6	20	160	24	40.2
		06:00	1.6	19	160	24	40.2
		06:05	1.6	19	160	24	40.2
		06:10	1.6	19	160	24	40.2
		06:15	1.6	19	160	24	40.2
		06:20	1.4	19	160	24	35.2
		06:25	1.4	19	160	24	35.2
		06:30	1.4	19	160	24	35.2
		06:35	1.4	22	160	24	35.2
		06:40	1.4	22	160	23	35.2
		06:45	1.2	19	160	23	30.1
		06:50	1.2	19	160	23	30.1
		06:55	1.2	18	160	23	30.1
		07:00	1.2	21	160	23	30.1
		07:05	1.2	20	160	23	30.1
		07:10	1.4	18	160	23	35.2
		07:15	1.4	18	160	23	35.2
		07:20	1.4	18	160	23	35.2
		07:25	1.4	18	160	23	35.2
		07:30	1.4	18	160	23	35.2
		07:35	1.4	18	160	23	35.2
		07:40	1.4	18	160	23	35.2
		07:45	1.6	18	160	23	40.2
		07:50	1.6	18	160	23	40.2
		07:55	1.6	18	160	23	40.2
	End	08:00					
		Averages	2.0	19.9	160.0	23.3	50.7

Average velocity	4.82	m/s
Average flow rate	0.95	m³/s
Average flow rate	0.88	m³/s*

Blank Filter Weight		Weight Collected	
Filter No.	EM - 2112/2	0.0001	g
Washings	EM - 2112/1	0.0003	g
Total		0.0004	g

Filter Weight Test 1		Weight Collected	
Filter No.	EM - 2112/4	0.0001	g
Washings	EM - 2112/3	0.0004	g
Total		0.0005	g

Particulate Emission Results		Blank	
Detection Limit		0.05	mg/m ^{3*}
Particulate Emission		0.04	mg/m ^{3*}

Particulate Emission Results		Test 1	
Detection Limit		0.05	mg/m ^{3*}
Particulate Emission		0.05	mg/m ^{3*}

Laboratory	SAL	
Method N° & Accreditation Status	Gravimetric	UKAS
Date of analysis	13/11/2015	

Total Particulate Matter - Emission Results

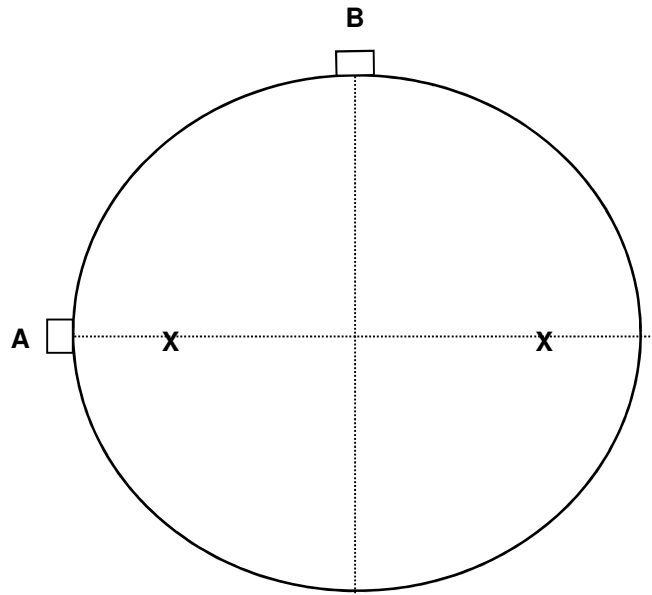
APPENDIX 3

Diagram Of The Sampling Location

Diagram of sampling points across the cross section of the duct (not to scale).

Traverse length = 0.50 m

Point	% of D	Location cm
1	14.6	7.3
2	85.4	42.7



General Calculations

Stack area:

$$\text{Area of a circle} = \frac{\pi \cdot D^2}{4}$$

D = Diameter (m)

π = 3.142

Pressure conversion:

1mmH₂O = 0.00980665 kPa

1mmH₂O = 9.80665 Pa

1 mar = 0.1 kPa

Water vapour concentration:

From reference calculations (taken from BS EN 14790):

$$V_{WC(\%)} = \frac{\frac{m_{WC} \cdot V_{mol(std)}}{M_w}}{\frac{m_{WC} \cdot V_{mol(std)}}{M_w} + V_{m(std)}} \times 100$$

WVC (%) = Water vapour content on wet basis, in volume % (m³ of water vapour in m³ of wet gas)

V_{m(std)} = Dry gas volume measured, corrected to standard conditions (m³)

m_{WC} = Mass of water collected in the impingers (g)

M_w = Molecular weight of water, 18.01534 rounded to 18 (g/mol)

V_{mol(std)} = Molar volume of water at standard conditions = 0.0224 (m³/mol)

Gas meter volume at standard conditions (STP)

From reference calculations (taken from BS EN 14790):

$$V_{m(std)} = y_d \times (V_2 - V_1) \times \frac{T_{std}}{T_m} \times \frac{p_m}{p_{std}}$$

V_{m(std)} = Dry gas meter volume at standard conditions (m³)

y_d = Gas meter calibration coefficient

(V₂-V₁) = Dry gas meter volume at actual conditions (m³)

T_m = Actual Temperature (K)

T_{std} = Standard temperature (273 K)

p_m = Absolute pressure at the gas meter (kPa)

p_{std} = Standard gas pressure (101.3 kPa)

Isokenetic Ratio (%):

From reference calculations (taken from EA TGN M2):

$$\text{IsokeneticRatio}(\%) = \frac{\text{Velocity at the sampling nozzle}}{\text{Velocity of the stack gas}} \times 100$$

Estimating Measurement Uncertainty

Uncertainty estimates are calculated using the general rule of uncertainty propagation. Guidance is taken from publications including UKAS document M3003 and ISO 20988:2007.

Flow Calculations

Velocity:

From reference calculations (taken from ISO 10780):

$$\bar{v} = KC \sqrt{\frac{T_s \Delta p}{p_e M_s}}$$

\bar{v} = Average velocity (m/s)

C = velocity calculation constant = 129

T_s = Average stack temperature (K)

M_s = Molar mass of gas; assume 29 kg/kmol unless the molar mass is < 27 kg/kmol or > 31 g/kmol

K = Pitot calibration coefficient

p_e = Absolute gas pressure (kPa)

Δp = Average pitot tube pressure differential (kPa)

Volume flow rate

From reference calculations (taken from ISO 10780):

$$q_{va} = \bar{v}A$$

q_{va} = Average flow rate (m³/s)

\bar{v} = Average velocity (m/s)

A = Stack cross-sectional area (m²)

Volume flow rate corrected for moisture

From reference calculations (taken from BS ISO 9096):

$$q_m = q_{va} \frac{(100 - H_a)}{(100 - H_m)}$$

q_m = Corrected volume flowrate (m³/s)

q_{va} = Volume flow rate at actual conditions (m³/s)

H_a = Moisture at actual conditions (%volume)

H_m = Reference moisture (%volume)

Volume flow rate corrected for temperature and pressure

From reference calculations (taken from BS ISO 9096):

$$q_m = q_{va} \frac{(T_m p_a)}{(T_a p_m)}$$

q_m = Corrected volume flowrate (m³/s)

q_{va} = Volume flow rate at actual conditions (m³/s)

T_a = Temperature at actual conditions (K)

T_m = Reference Temperature (K)

p_a = Absolute gas pressure at actual conditions (kPa)

p_m = Reference pressure (kPa)

Volume flow rate corrected for oxygen

From reference calculations (taken from BS ISO 9096):

$$q_m = q_{va} \frac{(20.9 - O_{2,ref})}{(20.9 - O_{2,m})}$$

q_m = Corrected volume flowrate (m³/s)

q_{va} = Volume flow rate at actual conditions (m³/s)

$O_{2,m}$ = Actual oxygen concentration (%)

$O_{2,ref}$ = Reference oxygen concentration (%)

Concentration Calculations

Concentration:

From reference calculations (taken from BS EN 13284-1):

$$c = \frac{m}{V}$$

c = Concentration
 m = Mass of substance
 V = Volume sampled

Mass Emission

$$\text{Mass emission} = c \times q_m$$

c = Concentration
 q = Volume flow rate

Concentration corrected for oxygen:

From reference calculations (taken from BS ISO 9096):

$$c_m = c_a \times \frac{20.9 - O_{2,ref}}{20.9 - O_{2,a}}$$

c_m = Concentration at reference conditions
 c_a = Actual concentration
 $O_{2, ref}$ = Reference oxygen (%)
 $O_{2, a}$ = Actual Oxygen (%)

Concentration corrected for moisture:

From reference calculations (taken from BS ISO 9096):

Convert wet gas to dry gas

$$c_{dry} = c_{wet} \times \frac{100}{100 - H_a}$$

Convert dry gas to wet gas

$$c_{wet} = c_{dry} \times \frac{100 - H_a}{100}$$

c_{wet} = Concentration wet gas
 c_{dry} = Concentration dry gas
 H_a = Water vapour content (%vol)

Conversion of parts per million (ppm) to mg/m³

From reference calculations (taken from EA TGN M2):

$$\text{Concentration}(\text{mg} / \text{m}^3) = \frac{\text{Concentration}(\text{ppm}) \times \text{molecular weight}(\text{g})}{\text{molar volume}(\text{l}) \text{ at a given temperature}}$$

molar volume at 273K = 22.4 litres

When Converting TOC

$$\text{Concentration}(\text{mg} / \text{m}^3) = \frac{\text{Concentration}(\text{ppm}) \times \text{molecular weight of carbon in span gas}(\text{g})}{\text{molar volume}(\text{l}) \text{ at a given temperature}}$$

molar volume at 273K = 22.4 litres

Calculation of Uncertainty Estimates - Manual Monitoring Techniques

$$C_m = \frac{Q_m}{V_{std}}$$

$$V_{std} = V_{T.p} \times \frac{T_{std}}{T} \times \frac{p}{P_{std}}$$

Where:

C_m	=	the measured concentration	in	mg/m ³
Q_m	=	mass concentration collected in absorber solution and filter	in	mg
v_s	=	the volume of the sample solution	in	l
V_{std}	=	volume sampled at standard conditions	in	m ³
T_m	=	mean temperature of gas meter	in	K
T_{std}	=	standard temperature	=	273K
$p = p_{rel} + p_{atm}$	=	absolute pressure = gas meter pressure + atmospheric pressure	in	kPa
P_{std}	=	standard pressure	=	101.325 kPa
$V_{T.p}$	=	volume sampled as indicated by the gas meter	in	m ³

Expression for calculation of the combined uncertainty of the measured concentration

$$\frac{u^2(C_m)}{(C_m)^2} = \frac{u^2(Q_m)}{(Q_m)^2} + \frac{u^2(v_s)}{(v_s)^2} + \frac{u^2(V_{std})}{(V_{std})^2}$$

Expression for calculation of the combined uncertainty of the gas volume

$$V_{std} = V_{T.p} \times \frac{T_{std}}{T_m} \times \frac{p}{P_{std}}$$

Assuming that uncertainties associated with the standard quantities T_{std} and P_{std} are negligible

Expression for calculation of the combined uncertainty of the measured concentration

$$\frac{u^2(C_m)}{(C_m)^2} = \frac{u^2(Q_m)}{(Q_m)^2} + \frac{u^2(v_s)}{(v_s)^2} + \frac{u^2(V_{T.p})}{(V_{T.p})^2} + \frac{u^2(T)}{(T)^2} + \frac{u^2(p_{rel})}{(p)^2} + \frac{u^2(p_{atm})}{(p)^2}$$

$$u(C_m) = \sqrt{\left(\frac{u^2(Q_m)}{(Q_m)^2} + \frac{u^2(v_s)}{(v_s)^2} + \frac{u^2(V_{T.p})}{(V_{T.p})^2} + \frac{u^2(T)}{(T)^2} + \frac{u^2(p_{rel})}{(p)^2} + \frac{u^2(p_{atm})}{(p)^2} \right)} \times (C_m)$$

Overall expanded uncertainty ($k = 2$)

$$U(C_m) = u(C_m) \times k$$

Uncertainty of the measured concentration at oxygen reference concentration

$$u(C_{m,O_2,ref}) = \sqrt{\left(\frac{u^2(Q_m)}{(Q_m)^2} + \frac{u^2(v_s)}{(v_s)^2} + \frac{u^2(V_{T.p})}{(V_{T.p})^2} + \frac{u^2(T)}{(T)^2} + \frac{u^2(p_{rel})}{(p)^2} + \frac{u^2(p_{atm})}{(p)^2} + \frac{u^2(O_{2,meas,dry})}{(21 - O_{2,meas,dry})^2} \right)} \times (C_{m,O_2,ref})$$

Where:

$u(C_{m,O_2,ref})$	=	uncertainty associated with the mass concentration at O ₂ reference concentration	in	mg/m ³
$C_{m,O_2,ref}$	=	mass concentration at O ₂ reference concentration	in	mg/m ³
$O_{2,meas}$	=	O ₂ measured concentration	in	% volume
$u((O_{2,meas})_{dry})$	=	uncertainty associated to the measured O ₂ concentration	in	% (relative to O _{2,meas})

Calculation of Uncertainty Estimates - Instrumental Monitoring Techniques

Model equation

$$C_{ppm} = C_{reading} + Corr_{fit} + Corr_{0,dr} + Corr_{s,dr} + Corr_{rep} + Corr_{adj} + \sum_{i=1}^p Corr_{inf} + Corr_{int}$$

Where:

C_{ppm}	=	concentration in ppm	$Corr_{rep}$	=	correction of repeatability of measurement
$C_{NO,reading}$	=	concentration given by analyser	$Corr_{adj}$	=	correction of adjustment
$Corr_{fit}$	=	correction of lack of fit	$Corr_{inf}$	=	correction of influence quantities
$Corr_{0,dr}$	=	correction of zero drift	$Corr_{int}$	=	correction of interferences
$Corr_{s,dr}$	=	correction of span drift			

Calculation of partial uncertainties

$$u(Corr_{fit}) = \frac{\left(\frac{X_{fit,max}}{100 \times range} \right)}{\sqrt{3}} \quad \text{Where: } X_{fit,max} \text{ is the maximum allowable deviation from linearity}$$

Expressed as % of the range and calculated by applying a rectangular probability distribution

$$u(Corr_{0,dr}) = \frac{X_{0,dr}}{\sqrt{3}} \quad u(Corr_{s,dr}) = \frac{X_{s,dr}}{\sqrt{3}}$$

$$u(Corr_{rep}) = \max(S_{0,rep}; S_{s,rep}) = S_{rep} \quad \text{Where: } S_{r,0} \text{ is the standard uncertainty at zero level}$$

$$S_{r,s} \text{ is the standard uncertainty at span level}$$

$$u(Corr_{adj}) = u(Corr_{loss}) + u(Corr_{cal}) \quad \text{Where: } u(Corr_{loss}) \text{ is the uncertainty due to losses in sample line}$$

$$u(Corr_{loss}) = \frac{C_{j,loss}}{\sqrt{3}} \quad u(Corr_{cal}) \text{ is the uncertainty due to losses in sample line}$$

$$u(Corr_{cal}) = \frac{U_{cal}}{2} \quad c_{j,loss} \text{ is the concentration of sample loss at span level}$$

$$U_{cal} \text{ is the expanded uncertainty of the calibration gas}$$

$$u(Corr_{int}) = c_j \sqrt{\frac{(x_{j,max} - x_{j,adj})^2 + (x_{j,min} - x_{j,adj})^2 + (x_{j,max} - x_{j,adj})(x_{j,min} - x_{j,adj})}{3}}$$

Where:

- c_j is the sensitivity coefficient of the influence quantity
- $x_{j,min}$ is the minimum value of the influence quantity during monitoring
- $x_{j,max}$ is the maximum value of the influence quantity during monitoring
- $x_{j,adj}$ is the value of the influence quantity during adjustment

$$u(Corr_{int}) = \frac{c_j}{Int_{j,test}} \sqrt{\frac{Int_{j,max}^2 + Int_{j,min} \times Int_{j,max} + Int_{j,min}^2}{3}}$$

and

$$u(\Sigma Corr_{int}) = \max[S_{Int,p}; S_{Int,n}] \quad \text{Where: } c_j \text{ is the sensitivity coefficient of the interferent j}$$

- $Int_{j,test}$ is the concentration of the interferent j used to determine c_j
- $Int_{j,min}$ is the minimum value of the interferent j quantity during monitoring
- $Int_{j,max}$ is the maximum value of the interferent j quantity during monitoring
- $Int_{j,adj}$ is the concentration of the interferent j in the cal gas used to adjust the analyser
- $S_{Int,p}$ is the sum of interferences with positive impact
- $S_{Int,n}$ is the sum of interferences with negative impact

Combined uncertainty

$$u(C_{ppm}) = \sqrt{u^2(Corr_{fit}) + u^2(Corr_{0,dr}) + u^2(Corr_{s,dr}) + u^2(Corr_{rep}) + u^2(Corr_{s,vf}) + u^2(Corr_{a,press}) + u^2(Corr_{temp}) + u^2(Corr_{volt}) + u^2(Corr_{adj}) + S_{Int}^2}$$

$$\text{Overall expanded uncertainty (k = 2)} \quad U(C_m) = u(C_m) \times k$$

Uncertainty of NOx measurements

$$u(C_{NOx,conv}) = \frac{C_{NOx} \times R \times \eta}{\sqrt{3}} \quad \text{Where: } C_{NOx} \text{ is the concentration of NOx measured by the analyser}$$

- R is the ratio of NO:Nox in the stack gas
- η is the NOx converter efficiency

Combined uncertainty NOx measurements

$$u(C_{NOx,stack}) = \sqrt{u^2(Corr_{fit}) + u^2(Corr_{0,dr}) + u^2(Corr_{s,dr}) + u^2(Corr_{rep}) + u^2(Corr_{s,vf}) + u^2(Corr_{a,press}) + u^2(Corr_{temp}) + u^2(Corr_{volt}) + u^2(Corr_{adj}) + S_{Int}^2 + u^2(Corr_{NOx,conv})}$$

Uncertainty of mass concentration at oxygen reference concentration

$$u(C_{O_2,ref}) = \sqrt{u^2(Corr_{fit}) + u^2(Corr_{0,dr}) + u^2(Corr_{s,dr}) + u^2(Corr_{rep}) + u^2(Corr_{s,vf}) + u^2(Corr_{a,press}) + u^2(Corr_{temp}) + u^2(Corr_{volt}) + u^2(Corr_{adj}) + S_{Int}^2 + \left(\frac{u^2(O_{2,meas,dry})}{(21 - O_{2,meas,dry})^2} \right)}$$

Where:

$u(C_{O_2,ref})$	=	uncertainty associated with the mass concentration at O ₂ ref. concentration	in	mg/m ³
$C_{O_2,ref}$	=	mass concentration at O ₂ reference concentration	in	mg/m ³
$O_{2,meas}$	=	O ₂ measured concentration	in	% volume
$u((O_{2,meas})_{dry})$	=	uncertainty associated to the measured O ₂ concentration	in	% (relative to O ₂ meas)

APPENDIX 4

Uncertainty Estimate For The Measurement Of Total Organic Carbon

Analysers Type/Model
Reference Oxygen %

Sick Maihak
0 (0 = No correction)

		Test 1	Test 2	Test 3	Test 4	Test 5	Test 6	Test 7	Test 8
Limit value	mg/m ³	150	150	150	150	150	150	150	150
Limit value	ppm	93.3	93.3	93.3	93.3	93.3	93.3	93.3	93.3
Measured concentration	ppm	221.1	223.6	171.6	145.0	117.8	127.6	343.0	190.2
Measured concentration	mg/m ³	355.4	359.4	275.8	233.0	189.3	205.1	551.2	305.8

Calibration gas	ppm	74.7	74.7	74.7	74.7	74.7	74.7	74.7	74.7
Calibration gas	mg/m ³	120.1	120.1	120.1	120.1	120.1	120.1	120.1	120.1
Analysers range	ppm	9.3	9.3	9.3	9.3	9.3	9.3	9.3	9.3
Analysers range	mg/m ³	15.0	15.0	15.0	15.0	15.0	15.0	15.0	15.0

Correction of Lack of Fit

Lack of fit	% range	2.0	2.0	2.0	2.0	2.0	2.0	2.0	2.0
	u(Corr _{lof})	0.11	0.11	0.11	0.11	0.11	0.11	0.11	0.11

Corrections of Zero and Span Drift

(*All drift is calculated for ∴ the residual is assumed to be < 5% u(max))

Zero Drift	% range	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
	u(Corr _{zdr})	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Span Drift	% range	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
	u(Corr _{spd})	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00

Correction of Repeatability of Measurement

Repeatability SD at span level (Not reported)	% range	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
	u(Corr _{rep})	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00

Correction of adjustment

losses in the line	% range	0.04	0.04	0.04	0.04	0.04	0.04	0.04	0.04
	u(Corr _{loss})	0.05	0.05	0.04	0.03	0.03	0.03	0.08	0.04
Uncertainty of calibration gas	% range	2.0	2.0	2.0	2.0	2.0	2.0	2.0	2.0
	u(Corr _{cga})	2.21	2.24	1.72	1.45	1.18	1.28	3.43	1.90

Correction of Influence of Interferents

N ₂ O	% range								
	u(Corr _{N2O})	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
CO ₂	% range								
	u(Corr _{CO2})	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
CH ₄	% range								
	u(Corr _{CH4})	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Total of interferent influences $u(\Sigma Corr_{int}) = \max[S_{int,1}; S_{int,n}]$	% range	2.50	2.50	2.50	2.50	2.50	2.50	2.50	2.50
	u(ΣCorr _{int})	0.16	0.16	0.16	0.16	0.16	0.16	0.16	0.16

Correction of Influence Quantities

Sensitivity to sample volume flow	% range	1.60	1.60	1.60	1.60	1.60	1.60	1.60	1.60
	u(Corr _{svf})	0.09	0.09	0.09	0.09	0.09	0.09	0.09	0.09
Sensitivity to atmospheric pressure (Not reported)	% range								
	u(Corr _{press})	0.00	0.00	0.00	0.00	0.00	0.00	0.00	0.00
Sensitivity to ambient temperature	% range	-2.40	-2.40	-2.40	-2.40	-2.40	-2.40	-2.40	-2.40
	u(Corr _{temp})	-0.28	-0.28	-0.28	-0.28	-0.28	-0.28	-0.28	-0.28
Sensitivity to electrical voltage (Not reported)	% range	0.50	0.50	0.50	0.50	0.50	0.50	0.50	0.50
	u(Corr _{volt})	0.10	0.10	0.10	0.10	0.10	0.10	0.10	0.10

Maximum standard uncertainty	u(Corr _{max})	2.21	2.24	1.72	1.45	1.18	1.28	3.43	1.90
5% of maximum standard uncertainty	u(Corr _{5%})	0.11	0.11	0.09	0.07	0.06	0.06	0.17	0.10

Interferent Concentration Variations	Minimum	Maximum	Value at cal	Performance	Units
CH ₄ range	0	10	0	50	mg/m ³
N ₂ O range	0	0	0	20	mg/m ³
CO ₂ range	8	12	0	15	mg/m ³
Oxygen effect variations	Minimum	Maximum	Value at cal	Performance	Units
Oxygen effect	0	20	0	2	mg/m ³

Influence Quantity Variations

	Minimum	Maximum	Value at cal	Performance	Units
Sensitivity to sample volume flow	55	65	60	5	l/h
Sensitivity to atmospheric pressure	99	100	99	1	kPa
Sensitivity to ambient temperature	278	313	288	10	K
Sensitivity to electrical voltage	187	250	230	5	V

Measurement uncertainty		Test 1	Test 2	Test 3	Test 4	Test 5	Test 6	Test 7	Test 8
Combined uncertainty	ppm	2.23	2.26	1.75	1.50	1.23	1.33	3.44	1.94
Combined uncertainty	mg/m ³	3.59	3.63	2.82	2.40	1.98	2.13	5.53	3.11
Combined uncertainty at oxygen reference	mg/m ³	3.59	3.63	2.82	2.40	1.98	2.13	5.53	3.11

Expanded uncertainty expressed with a level of confidence of 95%, k=2

Overall uncertainty	ppm	4.5	4.5	3.5	3.0	2.5	2.7	6.9	3.9
Overall uncertainty	mg/m ³	7.2	7.3	5.6	4.8	4.0	4.3	11.1	6.2
Overall uncertainty relative to measured value	%	2.0	2.0	2.0	2.1	2.1	2.1	2.0	2.0
Overall uncertainty relative to range	%	47.9	48.4	37.6	32.0	26.4	28.4	73.7	41.5
Overall uncertainty relative to ELV	%	6.0	6.0	4.7	4.0	3.3	3.6	9.2	5.2

The uncertainty evaluation has been carried out in accordance with UKAS requirements.

Uncertainty Estimate For The Measurement Of Total Particulate Matter

Total Particulate Matter

Symbol	Mass concentration of TPM	Units
C_m	0.05	mg/m ³
C_m at O_2 ref	N/A	mg/m ³

Filter	0.10	mg
Washings	0.40	mg

Symbol	Reference Oxygen	Units
O_2 ,ref (0=No Correction)	0	%

Symbol	Daily emission limit value	Units
DELV	20.0	mg/m ³

Mass of Particulate

Symbol	Source of Uncertainty	Value	Units	Probability Distribution	Divisor	c_i	U_i
$u(\text{calBal})$	Calibration	0.1180	mg	Normal	2	1	0.0590
$u(\text{repBal})$	Repeatability Filter	0.0110	mg	Normal	1	2	0.0220
$u(\text{repBal})$	Repeatability Washings	0.0300	mg	Normal	1	2	0.0600
$u(\text{driftBal})$	Drift	0.0312	mg	Rectangular	$\sqrt{3}$	1	0.0180
$u(\text{resBal})$	Resolution	0.0050	mg	Rectangular	$\sqrt{3}$	1	0.0029
$u(\text{aceBal})$	Residual acetone	0.0400	mg	Normal	2	1	0.0200
$u(\text{buoBal})$	Air buoyancy	0.0015	mg	Normal	1	1	0.0015
$u(\text{mass})$	Combined Uncertainty	-	-	Normal	-	-	0.0911
$u(\text{mass})/\text{mass}$	Relative	0.500	mg	-	-	-	1.8763
$u(\text{mass})^2/\text{mass}^2$							3.520556

$U(W_m)$	Expanded Combined Weighing Uncertainty As Percentage of DELV $k=2$	\pm	0.91	%
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Volume of sampled gas

Symbol	Source of Uncertainty	Value	Units	Probability Distribution	Divisor	c_i	U_i
$u(\text{calV}_{T,p})$	Calibration	0.0505	m ³	Normal	2	1	0.0252
$u(\text{repV}_{T,p})$	Repeatability	0.0051	m ³	Normal	1	1	0.0051
$u(\text{driftV}_{T,p})$	Drift	0.0278	m ³	Rectangular	$\sqrt{3}$	1	0.0161
$u(\text{resV}_{T,p})$	Resolution	0.0002	m ³	Rectangular	$\sqrt{3}$	1	0.0001
$u(V_{T,p})$	Combined Uncertainty	-	-	Normal	-	-	0.0303
$u(V_{T,p})/V_{T,p}$	Relative	10.298	m ³	-	-	-	0.0029
$u(V_m)^2/V_m^2$							0.000009

Temperature of the DGM

Symbol	Source of Uncertainty	Value	Units	Probability Distribution	Divisor	c_i	U_i
$u(\text{calT}_m)$	Calibration	1.0	K	Normal	2	1	0.5000
$u(\text{calT}_m)$	Calibration	1.0	K	Normal	2	1	0.5000
$u(\text{resT}_m)$	Resolution	0.1	K	Rectangular	$\sqrt{3}$	1	0.0577
$u(\text{driftT}_m)$	Drift	1.0	K	Rectangular	$\sqrt{3}$	1	0.5774
$u(T_m)$	Combined Uncertainty	-	-	Normal	-	-	0.9147
$u(T_m)/T_m$	Relative	296.3	K	Normal	-	-	0.0031
$u(T_m)^2/T_m^2$							0.000010

Atmospheric Pressure

Symbol	Source of Uncertainty	Value	Units	Probability Distribution	Divisor	c_i	U_i
$u(P_{\text{atm}})$	Metrological Office	300	Pa	Normal	$\sqrt{3}$	1	173
$u(P_{\text{atm}})$	Combined Uncertainty	-	-	Normal	-	-	173
$u(P_{\text{atm}})/P$	Relative	101300	-	Normal	-	-	0.0017
$u(P_{\text{atm}})^2/P^2$							0.000003

Relative DGM Pressure

Symbol	Source of Uncertainty	Value	Units	Probability Distribution	Divisor	c_i	U_i
$u(P_{\text{rel}})$	DGM Pressure	40	Pa	Normal	1	1	40
$u(P_{\text{rel}})$	Combined Uncertainty	-	-	Normal	-	-	40
$u(P_{\text{rel}})/P$	Measured pressure	101798	-	-	-	-	0.0004
$u(P_{\text{rel}})^2/P^2$							0.000002

Oxygen reference concentration

Symbol	Source of Uncertainty	Value	Units	Probability Distribution	Divisor	c_i	U_i
$u((O_2, \text{meas})_{\text{dry}})_{\text{rel}}$	Measurement of oxygen	0.20	%	Normal	1	1	0.2000
	Uncertainty of oxygen correction	-	-	-	-	-	N/A
	Measured Oxygen	N/A	%	-	-	-	$u(O_{\text{rel}})^2/O^2$

Measurement uncertainty

$u(C_m)$	Combined Standard Uncertainty			\pm	0.13	mg/m ³
$u(C_m)$ at ref O_2	Combined Standard Uncertainty			\pm	N/A	mg/m ³

Expanded uncertainty expressed with a level of confidence of 95%, $k=2$

$U(C_m)$	Expanded Combined Uncertainty	$k = 2$		\pm	0.26	mg/m ³
$U(C_m, \text{rel})$	Expanded Combined Uncertainty	$k = 2$		\pm	530.7	%
$U(C_m, \text{rel}, \text{ELV})$	Expanded Combined Uncertainty	$k = 2$		\pm	1.3	%

The reported expanded uncertainty is based on a standard uncertainty multiplied by a coverage factor ($k = 2$), providing a level of confidence of approximately 95%. The uncertainty evaluation has been carried out in accordance with UKAS requirements.