





11-12 June & 30-31 July, 2013

Prepared for Alan Nuttalls Ltd

REC Report 71519p1r0

Issued: 30 August, 2013













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CONT	ENTS	Page No
EXEC	UTIVE SUMMARY	5
1. I	NTRODUCTION	7
1.1	Background	7
1.2	Scope of the Survey	7
1.3	Sampling Personnel	7
2. I	METHODOLOGY	8
2.1	Species & Techniques	8
2.2	Sampling & Analytical Methodology	8
2.3	Laboratory Analysis	10
3.	SAMPLING AND OPERATIONAL DETAILS	11
3.1	Process Description	11
3.2	Sampling Positions	11
3.3	Uncertainty	11
3.4	Emission Monitoring Survey Details	12
4. I	RESULTS AND DISCUSSION	13
4.1	Initial Velocity and Temperature Traverse	13
4.2	Particulate Matter	13
4.3	Formaldehyde	13
4.4	Hydrogen Chloride	13
4.5	HCN	13
4.6	Combustion Gases	14
4.7	Total VOC Emission Data	14

FIGURES (2 x Additional Pages)

- 1 Combustion Gas Emission Data Summary, Boiler stack, 31/08//13 (data expressed at 273K, 101.3kPa, Wet Gas with no correction for oxygen content).
- 2 VOC Emission Data Summary, Boiler stack, 31/08//13 (data expressed at 273K, 101.3kPa, Wet Gas with no correction for oxygen content).

TABLES (13 x Additional Pages)

- 1 Flow Data
- 2-9 Particulate Emission Data Summary
- 10 Particulate & Formaldehyde Emissions Data Summary
- 11 HCI Emissions Data Summary
- 12 HCN Emissions Data Summary
- 13 Combustion Gas & VOC Emissions Data Summary

APPENDICES (14 x Additional Pages)

- 1 Certificate of Analysis
- 2 Photographs of Sampling Locations
- 3 Calculations

EXECUTIVE SUMMARY

Resource & Environmental Consultants (REC) Ltd was commissioned by Alan Nuttalls Ltd to monitor to monitor emissions of pollutants released from various process stacks at their site in Dudley.

In accordance with the requirements of their site permit, monitoring has been undertaken for the following pollutants:-

- Combustion Gases including O₂ & CO
- Total Particulate Matter
- Hydrogen Chloride (HCl)
- Hydrogen Cyanide
- Formaldehyde
- Total Volatile Organic Compounds (VOCs) expressed as Carbon (C)

The following results were obtained from the emission monitoring survey and are compared with the current permit limit:-

Species	Accreditation Status	Particulate Emission Concentration (mg/Nm ³)	Permit Limit (mg/Nm ³)
Pre-treatment Drying Oven	В	1.1	10
Refrigeration Powder Coat	В	1.2	10
Camel Back Curing Oven	В	1.0	10
Batch Line Curing Oven	В	1.6	10
Automatic Line Curing Oven	В	0.5	50
Wood Spraying Booth No.1	В	0.5	50
Wood Spraying Booth No.2	В	<0.3	50

Oven and Spraybooths

NOTE 1: All data are expressed in mg/Nm³ at 273K, 101.3kPa, without correction for moisture and oxygen content unless otherwise stated.

NOTE: UKAS Status:- (B) REC Ltd accredited for sampling only, UKAS accredited analysis conducted by SAL Ltd .

Species	Accreditation Status	Emission Concentration (mg/Nm ³)	Permit Limit (mg/Nm ³)
Total VOCs (as C)	А	2179.9	20
Oxygen	А	15.2	n/a
Carbon Monoxide	А	599.6	250
Particulate Matter	В	63.1	200
Hydrogen Chloride	В	6.9	100
Hydrogen Cyanide	E	2.87	5
Formaldehyde	В	<0.1	5

Wood Burning Boiler

NOTE 1: All data are expressed in mg/Nm³ at 273K, 101.3kPa, without correction for moisture and oxygen

content unless otherwise stated. **NOTE: UKAS Status:-** (A) REC Ltd accredited for sampling and analysis. (B) REC Ltd accredited for sampling only, UKAS accredited analysis conducted by SAL Ltd or RPS Ltd. (E) REC Ltd not accredited for sampling, sub-contracted analysis not UKAS accredited.

1. INTRODUCTION

1.1 Background

Alan Nuttals Ltd commissioned REC Ltd to conduct an emission monitoring survey on various process stacks at their site in Dudley.

The mains processes at the site involve the manufacture and powder coating of shop furniture fittings such as shelving. The other part of their operations involves the manufacture and powder coating of refrigerated display and storage units.

1.2 <u>Scope of the Survey</u>

An emission monitoring survey was required to determine the release concentrations of various pollutants from the powder coat and wood spraying processes. Concentrations of the following pollutants were quantified during the survey:

- Combustion Gases including O₂ & CO
- Total Particulate Matter
- Hydrogen Chloride (HCl)
- Hydrogen Cyanide
- Formaldehyde
- Total Volatile Organic Compounds (VOCs) expressed as Carbon (C)

Ancillary measurements of stack dimensions, temperature and velocity were also made.

Sampling for combustion gases (CO & O_2) and VOCs was carried out on a continuous basis with measured concentrations being data-logged at 1 minute intervals over the sampling period.

All results were to be reported at 273K, 101.3kPa, wet gas without correction for oxygen content.

1.3 <u>Sampling Personnel</u>

Monitoring was conducted by the following REC Ltd permanent staff:-

11 June, 2013

 Michelle Edwards - Rachel Powis - 	Team Leader, MM05 659, MCERTS Level 2, TE1&2 Assistant, MM12 1203, MCERTS Level 1
12 June, 2013	
 Dave Burns - Michelle Edwards - 	Team Leader, MM05 579, MCERTS Level 2, TE1-4 Assistant, MM05 659, MCERTS Level 2, TE1&2
30-31 July, 2013	
Dave Burns Rachel Powis	Team Leader, MM05 579, MCERTS Level 2, TE1-4 Assistant, MM12 1203, MCERTS Level 1

2. METHODOLOGY

2.1 Species & Techniques

The following table shows the reference methods used for the emission monitoring survey:

Species	UKAS Status	Method	Uncertainty (±%)	Limit of Detection
Moisture	A	In house method MM0010 based on BS EN 14790	20	0.1%vol
Particulate Matter	В	In house method MM0004 based on BS EN 13284 & BS ISO 9096	10	1 mg/m ³
Hydrogen Chloride	В	In house method MM0006 based on BS EN 1911	15	0.1 mg/m ³
Hydrogen Cyanide	E	Based on US EPA CTM33	30	0.1 mg/m ³
Formaldehyde	В	In house method MM0015 based on US EPA M316A	20	0.1 mg/m ³
Carbon Monoxide	A	In house method MM0002 based on BS EN 15058	10	1 mg/m ³
Oxygen	А	In house method MM0002 based on BS EN 14789	10	0.1%vol
Total VOCs (as C)	A	In house method MM0002 based on BS EN 12619 or BS EN 13526	10	1 mg/m ³

NOTE: UKAS Status:- (A) REC Ltd accredited for sampling and analysis. (B) REC Ltd accredited for sampling only, UKAS accredited analysis conducted by SAL Ltd or RPS Ltd. (E) REC Ltd not accredited for sampling, sub-contracted analysis by not UKAS accredited.

2.2 Sampling & Analytical Methodology

Total Particulate Matter

To determine the concentration of particulate matter in emissions, isokinetic stack sampling equipment satisfying the requirements of BS EN 13284 was utilised and in-house method MM0004 followed.

The Standard describes the methodology for measuring particulate matter under defined conditions and at discrete locations in the duct. Sampling is carried out under isokinetic sampling conditions i.e. the flowrate through the sampling nozzle is adjusted to equal the flowrate in the duct at the sampling positions. Velocity pressures were recorded throughout the monitoring period by means of an 'S' type pitot integral to the sampling probe and nozzle assembly.

A sample of the exhaust stream was removed from the stack via a titanium nozzle and titanium lined heated probe. It was then passed through a quartz fibre filter contained in a heated oven compartment. The temperature of the probe and filter box were maintained at

 160° C i.e. above the dew point of the stack gases, to ensure moisture did not condense on the filter. Each filter used complied with the requirements of Section 6.2.7 of BS EN 13284-1:2001 in that the efficiency was better than 99.5% for particles of 0.3µm diameter (or 99.9% for particles of 0.6µm diameter).

The impinger train was seated in a water bath to cool the gas stream and condense out less volatile gases and water vapour.

The first two impingers encountered by the gas stream contained deionised water. The third impinger was left empty and the fourth contained anhydrous silica gel which was used to dry the gas stream before passing it through a dry gas meter (DGM) to measure the volume of gas sampled.

All the impingers were weighed before and after the sampling run in order to determine the mass of water condensed by the impinger train (in house Method MM0010).

The sample volume collected was in excess of the minimum requirement stated in MM0004. The minimum sample volume ensures the results would be representative of normal plant operating conditions.

Upon completion of sampling, the filter was removed to a clean petri dish, labelled and sealed. The probe and filter housing were rinsed with acetone and water. The washings were collected in a container and submitted for analysis along with the filter.

This methodology allows the determination of formaldehyde to be carried out in conjunction with the particulate sampling as per US EPA Method 316A.

HCI

To determine the concentration of HCl in emissions, non-isokinetic stack sampling equipment satisfying the requirements of BS EN 1911 was utilised and in-house method MM0006 followed.

A sample of the exhaust stream was removed from the stack via a probe. It was then passed through a quartz fibre filter. On leaving the filter, the sampled exhaust gas was passed into a series of impingers.

The first three impingers encountered by the gas stream contained deionised water to capture and absorb the volatile chloride (Cl⁻) ions. The fourth impinger was left empty and the fifth contained anhydrous silica gel which was used to dry the gas stream before passing it through a dry gas meter (DGM) to measure the volume of gas sampled.

Upon completion of sampling, the contents of impingers 1 & 2 were transferred to a sealed, labelled container. The contents of impinger 3 were transferred to a separate container. The absorbing solution was subsequently analysed for chloride (CI^{\circ}) via an ion chromatographic (IC) technique.

HCN

To determine the concentration of HCN in emissions, non-isokinetic stack sampling equipment satisfying the requirements of US EPA CTM33 was utilised.

A sample of stack gas was removed via a stainless steel sampling probe. The gas then passes into a series of impingers. The first Two impingers encountered by the gas stream contained 0.1M sodium hydroxide (NaOH) to capture and absorb Cyanide ions.

The third impinger was left empty and the fourth contained anhydrous silica gel.

Sampling was conducted using a pre-calibrated, low flow sampling pump and Dry gas meter.

On completion of sampling, the impinger solution was collected and analysed for HCN by a colorimetric procedure.

Stack Temperature and Velocity

To determine the stack temperature, a calibrated thermocouple and digital indicator were employed. The exhaust gas velocity was investigated using a pitot static probe (to MM0004) and digital manometer.

2.3 Laboratory Analysis

An approved UKAS accredited sub-contractor, SAL Ltd, would undertake the sample analysis for Particulate matter, Hydrogen Cyanide and Hydrogen Chloride. RPS Ltd would undertake the sample analysis for Formaldehyde.

A copy of the Certificate of Analysis is enclosed in Appendix 1.

3. SAMPLING AND OPERATIONAL DETAILS

3.1 <u>Process Description</u>

The operations at Alan Nuttalls Ltd are authorised under a Part B permit issued by the Local Authority under the Environmental Permitting Regulations, 2010.

The process is therefore under Local Authority regulation and must demonstrate compliance with the emission limits stipulated in the site permit.

The main process operations involve the automatic and manual coating of shop furniture fittings, with various powdered paints, spirit stains and/or solvent based lacquers.

Once products have been spray coated, they are fed to a gas fired oven where the coatings are cured at a temperature of 180° c

The Waste wood generated in the manufacturing process is used to fuel the Ranheat wood burning boiler. The heat produced is then utilised to heat the factory. During sampling the waste wood used to fire the boiler, comprised of sawdust, laminates, chipboards and fibreboard.

3.2 <u>Sampling Positions</u>

On stacks all stack a single 4" BSP sampling port was installed on a horizontal plane. The sampling points provided were less than 4 x hydraulic diameters from any flow disturbance both upstream and downstream from the sampling plane. Access was via small temporary scaffolding.

Both the port size and sample planes do not fully meet the requirements stated in Environment Agency Technical Guidance Note M1. TGN M1 of $2 \times 5^{\text{"}}$ BSP sockets located at least $5 \times$ hydraulic diameters from any flow disturbance both upstream and downstream of the sampling plane. However the initial temperature and velocity traverse conducted along the sample plane showed that the flow requirements of TGN M1 were however met.

Photographs detailing the sampling locations are provided in Appendix 2.

3.3 Uncertainty

Due to the absence of a second sampling port, only a single sampling plane could be utilised. However the number of sample points, were increased along this plane and therefore the standard uncertainties would still apply.

REC has calculated uncertainty budgets for all of the pollutants listed in the Method Details Table in Section 2.1 above in accordance with calculations and methodology supplied by the Source Testing Association (STA). These uncertainties are quoted in the Tables section of this report.

3.4 Emission Monitoring Survey Details

The emission monitoring survey was carried out on the powder coating and wood spraying processes over the period 11-12 June and 30-31 July, 2013. The table overleaf summarises the actual sampling periods.

Stack	Parameter	Sample Time (& Date)
Pre-treatment Drying Oven	Particulate Matter	13:02 – 13:50 (11/06/13)
Refrigeration Powder Coat	Particulate Matter	15:32 – 16:04 (11/06/13)
Camel Back Curing Oven	Particulate Matter	13:00 – 13:48 (12/06/13)
Automatic Line Curing Oven	Particulate Matter	14:44 – 15:18 (12/06/13)
Batch line Curing Oven	Particulate Matter	09:27 – 09:57 (30/07/13
Wood Spraybooth No.1	Particulate Matter	11:54 – 12:34 (30/07/13)
Wood Spraybooth No.2	Particulate Matter	13:49 – 14:29 (30/07/13)
	Particulate Matter & HCHO	09:36 - 10:06 (31/07/13)
Wood Purping Boilor	HCI	09:35 – 10:05 (31/07/13
Wood Burning Boiler	HCN	10:30 – 11:00 (31/07/13)
	Combustion Gas & VOC	10:01 – 11:01 (31/07/13

SAMPLING PERIODS

4. **RESULTS AND DISCUSSION**

4.1 Initial Velocity and Temperature Traverse

An initial pitot-static pressure and temperature traverse was carried out. From these data stack velocity, expressed in metres per second (m/s), and volumetric flowrates expressed in cubic metre per hour (m^3/hr) have been calculated.

The results are reported at actual stack conditions and the volumetric flowrate is further expressed at the standard reference conditions of 273K, 101.3kPa i.e. standard temperature and pressure (STP). The results are summarised in Table 1.

4.2 Particulate Matter

The results of the particulate sampling runs are summarised in Tables 2 to 10. From the mass of particulate matter on the filter and in the acetone/water wash residue and volume sampled an emission concentration was calculated.

The results are expressed in mg/m³ at 273K, 101.3kPa, without correction for water vapour content.

4.3 <u>Formaldehyde</u>

The results of the Formaldehyde sampling runs are also summarised in Table 10. From the mass of HCHO in the absorbing solution and measured sample volume an emission concentration was calculated.

The results are expressed in mg/m³ at 273K, 101.3kPa, without correction for water vapour content.

4.4 <u>Hydrogen Chloride</u>

The results of the volatile chloride sampling runs are summarised in Table 11. From the concentration of Cl⁻ and the measured volume of absorbing solution a total mass of HCl in microgram (μ g) was determined. From their respective molecular weights, equivalent weights of HCl were then calculated. From the measured sample volume, an emission concentration was calculated.

The results are expressed in mg/m³ at 273K, 101.3kPa, without correction for water vapour content

4.5 <u>HCN</u>

The results of the Hydrogen Cyanide sampling run are summarised in Table 12. From the concentration of HCN in the absorbing solution and measured sample volume an emission concentration was calculated.

The results are expressed in mg/m³ at 273K, 101.3kPa, without correction for water vapour content.

4.6 <u>Combustion Gases</u>

The results of the combustion gas monitoring tests are summarised in Table 13 and Figure 1. The table presents the average of concentrations measured throughout each of the sample period.

Concentrations are expressed in mg/m³ at the standard reference conditions of 273K, 101.3kPa without correction for water vapour content.

Measured concentrations on a dry gas basis have been converted to a wet gas basis using moisture measurements from the Particulate /Formaldehyde sampling runs.

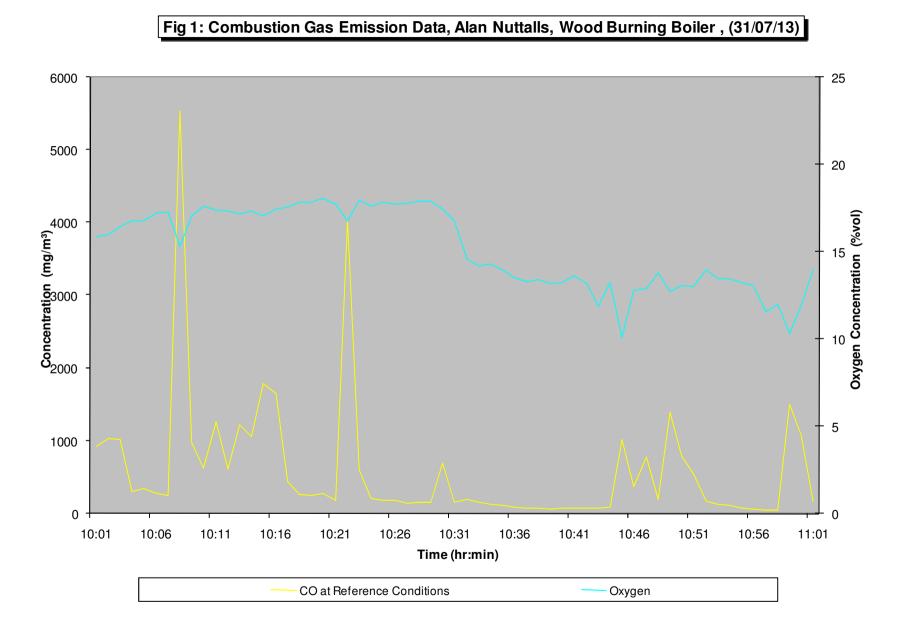
4.7 Total VOC Emission Data

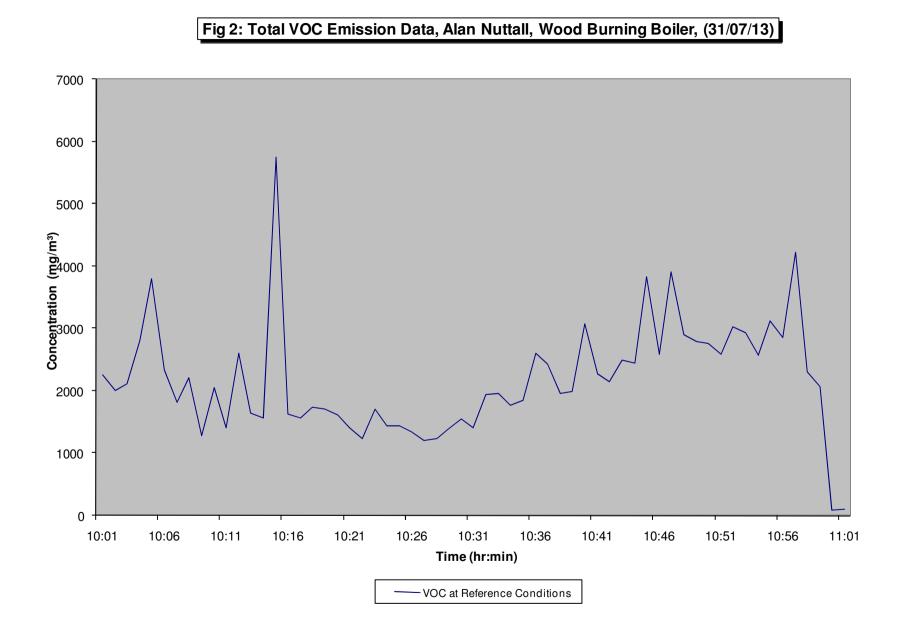
The results of the VOC monitoring tests are summarised in Table 13 and Figure 2. The table presents the average of concentrations measured throughout each of the sample periods.

Concentrations are expressed in mg/m³ as carbon (C) at the standard reference conditions of 273K, 101.3kPa without correction for water vapour content.

===== End of Report Text ====

FIGURES





FLOW DATA

Stack Ref.	Stack Temp	Av Pitot ∆P	Duct Diam	X-Sect. Area	Velocity (actual)		ne Flow ³/hr)
	(⁰ C)	(Pa)	(cm)	(m²)	(m/s)	(actual)	(@ ntp)
Batch Line Curing Oven Pre-treatment Drying Oven Refrigeration stack Camel Back Curing Oven Automatic Line Curing Oven Wet Spraybooth No.1 Wet Spraybooth No.2 Wood Burning Boiler	138 184 97 146 190 42 23 127	119 10 26 17 42 22 28 68	40 40 48x28 40 39 80 80 22	0.126 0.126 0.134 0.126 0.119 0.503 0.503 0.038	16.8 5.1 7.5 6.5 10.7 6.4 6.9 12.5	7,610 2,324 3,632 2,936 4,612 11,524 12,553 1,714	5,058 1,391 2,681 1,912 2,721 9,988 11,586 1,171

PARTICULATE EMISSION DATA SUMMARY – PRE-TREATMENT DRYING OVEN

13:02 - 13:50

Sampling Data 44 Run Time (min) 44 Total mass H ₂ O collected (g) 16. Pitot tube constant, Cp 0.8 Dry gas meter (DGM) volume (m ³) 0.7 Temperature DGM (°C) 25 Temperature stack (°C) 18 Mean pitot tube pressure drop, delta P (mm H ₂ O) 1.4 Orifice meter pressure drop, delta H (mm H ₂ O) 24. Sarometric Pressure (kPa) 98. X-sectional area of stack (m ²) 0.1 Nozzle size (mm) 10.0 Flow Data 5.4 Velocity, actual (m/s) 5.4 Velocity, ntp (m/s) 3.1 Vol. Flow, actual (m ³ /hr) 2.44 Vol. Flow, actual (m ³ /hr) 0.64 Volume sampled, ntp, dry gas (m ³) 0.64 Volume sampled, ntp, wet gas (m ³) 0.70 Analytical Data 5.4 Filter Weight Gain (mg) <0.4 Acetone Wash Residue Weight (mg) 0.4	
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Filter Weight Gain (mg) <0.	
Acetone Wash Residue Weight (mg) 01	
Total Particulates (mg) 0.4	
Partics Field Blank (mg) <0.	-
Blank % of ELV 13.	1
Emission Data	
H ₂ O (% vol) 2.9	2
Percentage Isokinetic 96.	
Particulates (mg/m ³) 1.	
Uncertainty $(\pm mg/m^3)$ 0.	
	1

DATE: 11/06/13

PARTICULATE EMISSION DATA SUMMARY – REFRIGERATION POWDER COAT

Sampling Data Run Time (min)	
Pup Time (min)	
	32
Total mass H_2O collected (g)	14.5
Pitot tube constant, Cp	0.84
Dry gas meter (DGM) volume (m ³)	0.896
Temperature DGM (°C)	29
Temperature stack (°C)	109
Mean pitot tube pressure drop, delta P (mm H ₂ O)	3.7
Orifice meter pressure drop, delta H (mm H ₂ O)	66.0
Barometric Pressure (kPa)	98.4
X-sectional area of stack (m ²)	0.134
Nozzle size (mm)	10.09
Flow Data	
Velocity, actual (m/s)	7.6
Velocity, ntp (m/s)	5.3
Vol. Flow, actual (m ³ /hr)	3,655
Vol. Flow, ntp (m ³ /hr)	2,577
Volume sampled, ntp, dry gas (m ³)	0.783
Volume sampled, ntp, wet gas (m ³)	0.801
Analytical Data	
Filter Weight Gain (mg)	<0.10
Acetone Wash Residue Weight (mg)	1.0
Total Particulates (mg)	1.0
Partics Field Blank (mg)	<0.4
Blank % of ELV	5.1
Emission Data	
H_2O (% vol)	2.3
Percentage Isokinetic	97.8
Particulates (mg/m ³)	1.2
Uncertainty (± mg/m³)	0.7

PARTICULATE EMISSION DATA SUMMARY – CAMEL BACK

DATE:	11/06/13
-------	----------

13:00 - 13:48

Sampling Data	
Run Time (min)	48
Total mass H ₂ O collected (g)	20.2
Pitot tube constant, Cp	0.84
Dry gas meter (DGM) volume (m ³)	0.899
Temperature DGM (°C)	29
Temperature stack (°C)	147
Mean pitot tube pressure drop, delta P (mm H ₂ O)	1.9
Orifice meter pressure drop, delta H (mm H ₂ O)	30.1
Barometric Pressure (kPa)	98.2
X-sectional area of stack (m ²)	0.126
Nozzle size (mm)	10.09
Flow Data	
Velocity, actual (m/s)	5.8
Velocity, ntp (m/s)	3.7
Vol. Flow, actual (m ³ /hr)	2,602
Vol. Flow, ntp (m³/hr)	1,664
Volume sampled, ntp, dry gas (m ³)	0.785
Volume sampled, ntp, wet gas (m³)	0.810
Analytical Data	
Filter Weight Gain (mg)	<0.10
Acetone Wash Residue Weight (mg)	0.8
Total Particulates (mg)	0.8
Partics Field Blank (mg)	<0.7
Blank % of ELV	1.8
Emission Data	
	0.4
H ₂ O (% vol) Percentage Isokinetic	3.1 95.7
Particulates (mg/m ³)	1.0

PARTICULATE EMISSION DATA SUMMARY – AUTOMATIC LINE

14:44 - 15:18

Sampling Data			
Run Time (min)	34		
Total mass H ₂ O collected (g)	13.1		
Pitot tube constant, Cp	0.84		
Dry gas meter (DGM) volume (m ³)	1.055		
Temperature DGM (°C)	27		
Temperature stack (°C)	193		
Mean pitot tube pressure drop, delta P (mm H ₂ O)	5.1		
Orifice meter pressure drop, delta H (mm H ₂ O)	74.5		
Barometric Pressure (kPa)	98.1		
X-sectional area of stack (m ²)	0.119		
Nozzle size (mm)	10.09		
Flow Data			
Velocity, actual (m/s)	9.9		
Velocity, ntp (m/s)	5.7		
Vol. Flow, actual (m ³ /hr)	4,270		
Vol. Flow, ntp (m³/hr)	2,460		
Volume sampled, ntp, dry gas (m ³)	0.926		
Volume sampled, ntp, wet gas (m ³)	0.942		
Analytical Data			
Filter Weight Gain (mg)	<0.10		
Acetone Wash Residue Weight (mg)	0.5		
Total Particulates (mg)	0.5		
Partics Field Blank (mg)	<0.4		
Blank % of ELV	4.3		
Emission Data			
H ₂ O (% vol) Berecentage lookingtig	1.7		
Percentage Isokinetic	101.0		
Particulates (mg/m ³) Uncertainty (± mg/m ³)	0.5		
Uncertainty (± mg/m)	0.6		

PARTICULATE EMISSION DATA SUMMARY – BATCH LINE

09:27 - 09:57

Sampling Data	
Dun Time (min)	30
Run Time (min) Total mass H ₂ O collected (g)	30 12.1
Pitot tube constant, Cp	0.84
Dry gas meter (DGM) volume (m ³)	0.671
Temperature DGM (°C)	22
Temperature stack (°C)	133
Mean pitot tube pressure drop, delta P (mm H ₂ O)	9.7
Orifice meter pressure drop, delta H (mm H ₂ O)	40.4
Barometric Pressure (kPa)	98.8
X-sectional area of stack (m ²)	0.126
Nozzle size (mm)	7.01
Flow Data	
Velocity, actual (m/s)	12.6
Velocity, ntp (m/s)	8.4
Vol. Flow, actual (m ³ /hr)	5,720
Vol. Flow, ntp (m ³ /hr)	3,800
Volume sampled, ntp, dry gas (m ³)	0.604
Volume sampled, ntp, wet gas (m ³)	0.619
Analytical Data	
Filter Weight Gain (mg)	<0.10
Acetone Wash Residue Weight (mg)	1.0
Total Particulates (mg)	1.0
Partics Field Blank (mg)	0.4
Blank % of ELV	6.6
Emission Data	
	0.4
H ₂ O (% vol) Percentage Isokinetic	2.4 105.9
Particulates (mg/m ³)	1.6
Uncertainty (± mg/m ³)	0.8
Uncertainty (± mg/m)	0.0

PARTICULATE EMISSION DATA SUMMARY - WOOD SPRAY BOOTH No.1

11:54 - 12:34

Sampling Data	
Run Time (min)	40
Total mass H ₂ O collected (g)	7.6
Pitot tube constant, Cp	0.85
Dry gas meter (DGM) volume (m ³)	1.059
Temperature DGM (°C)	22
Temperature stack (°C)	31
Mean pitot tube pressure drop, delta P (mm H_2O)	2.4
Orifice meter pressure drop, delta H (mm H ₂ O)	54.6
Barometric Pressure (kPa)	99.9
X-sectional area of stack (m ²)	0.503
Nozzle size (mm)	10.09
Flow Data	
Velocity, actual (m/s)	5.5
Velocity, ntp (m/s)	4.9
Vol. Flow, actual (m ³ /hr)	9,871
Vol. Flow, ntp (m³/hr)	8,803
Volume sampled, ntp, dry gas (m ³)	0.963
Volume sampled, ntp, wet gas (m ³)	0.973
Analytical Data	
Filter Weight Gain (mg)	<0.10
Acetone Wash Residue Weight (mg)	0.5
Total Particulates (mg)	0.5
Partics Field Blank (mg)	0.5
Blank % of ELV	1.0
Emission Data	
H₂O (% vol)	1.0
Percentage Isokinetic	104.3
Particulates (mg/m ³)	0.5
Uncertainty (± mg/m ³)	0.5

DATE: 30/07/13

PARTICULATE EMISSION DATA SUMMARY - WOOD SPRAY BOOTH No.2

13:49 - 14:29

Sampling Data	
Run Time (min)	40
Total mass H ₂ O collected (g)	11.9
Pitot tube constant, Cp	0.85
Dry gas meter (DGM) volume (m³)	1.292
Temperature DGM (°C)	26
Temperature stack (°C)	24
Mean pitot tube pressure drop, delta P (mm H ₂ O)	3.6
Orifice meter pressure drop, delta H (mm H ₂ O)	92.6
Barometric Pressure (kPa)	99.6
X-sectional area of stack (m ²)	0.503
Nozzle size (mm)	10.09
Flow Data	
Velocity, actual (m/s)	6.6
Velocity, ntp (m/s)	6.0
Vol. Flow, actual (m ³ /hr)	11,964
Vol. Flow, ntp (m³/hr)	10,923
Volume sampled, ntp, dry gas (m ³)	1.154
Volume sampled, ntp, wet gas (m ³)	1.169
Analytical Data	
Filter Weight Gain (mg)	<0.1
Acetone Wash Residue Weight (mg)	<0.3
Total Particulates (mg)	<0.4
Partics Field Blank (mg)	<0.4
Blank % of ELV	0.7
Emission Data	
H_2O (% vol)	1.3
Percentage Isokinetic	100.9
Particulates (mg/m ³)	<0.3
Uncertainty (± mg/m ³)	0.4

DATE: 30/07/13

PARTICULATE & HCHO EMISSION DATA SUMMARY – WOOD BURNING BOILER

DATE: 31/07/13	09:36 - 10:06
Sampling Data	
Run Time (min)	30
Total mass H ₂ O collected (g)	17.8
Pitot tube constant, Cp	0.85
Dry gas meter (DGM) volume (m ³)	0.763
Temperature DGM (°C)	20
Temperature stack (°C)	135
Mean pitot tube pressure drop, delta P (mm H ₂ O)	8.0
Orifice meter pressure drop, delta H (mm H_2O)	57.7
Barometric Pressure (kPa)	98.7
X-sectional area of stack (m ²)	0.126
Nozzle size (mm)	8.06
Flow Data	
Velocity, actual (m/s)	11.6
Velocity, ntp (m/s)	7.7
Vol. Flow, actual (m³/hr)	5,264
Vol. Flow, ntp (m³/hr)	3,474
Volume sampled, ntp, dry gas (m ³)	0.691
Volume sampled, ntp, wet gas (m³)	0.713
Analytical Data	
Filter Weight Coin (mg)	07.0
Filter Weight Gain (mg) Acetone Wash Residue Weight (mg)	27.0 18.0
Total Particulates (mg)	45.0
Partics Field Blank (mg)	0.4
Blank % of ELV	0.3
Mass of HCHO (µg)	<42.0
Emission Data	
H ₂ O (% vol)	3.1
Percentage Isokinetic	101.1
Particulates (mg/m ³)	63.1
Uncertainty (± mg/m ³)	2.1
HCHO (mg/m ³)	<0.1
Uncertainty (± mg/m ³)	0.01

HCL EMISSION DATA SUMMARY

Sampling Data	Boiler Stack
Start Time/Date End Time/Date Sampling Period (min) DGM start (dry m ³) DGM end (dry m ³) Volume Sampled (dry m ³) Ambient Temp (^o C) Ambient Press (kPa) Wt of Water (g) Volume Water (m ³) Volume Sampled, 273K, 101.3kPa (dry m ³) Volume Sampled, 273K, 101.3kPa (wet m ³) Volume H ₂ O Impingers 1+2 (ml) Volume H ₂ O Impinger 3 (ml)	09:35, 31/07/13 10:05, 31/07/13 30 44.773 45.093 0.320 20.5 98.7 80.9 0.101 0.290 0.391 330 170
Analytical Data	
HCI Blank (mg/l) HCI in H ₂ O Imps 1+2 (mg/l) HCI in Imps 1+2 (μ g) HCI in H ₂ O Imp 3 (mg/l) HCI in Imp 3 (μ g) % HCI in Imps 1+2	<0.05 7.5 2475 1.30 221 92
Emission Concentration Data	
Moisture (%vol) HCI (mg/m ³) HCI Measurement Uncertainty (± mg/m ³)	25.8 6.9 1.1

HCN EMISSION DATA SUMMARY

Sampling Data	Wood Burning Boiler			
Start Time/Date	10:30, 31/07/13			
End Time/Date	11:00, 31/07/13			
Sampling Period (min)	30			
DGM start (dry m ³)	45.101			
DGM end (dry m ³)	45.375			
Volume Sampled (dry m ³)	0.274			
Ambient Temp (°C)	21.5			
Ambient Press (kPa)	98.7			
Wt of Water (g)	13.3			
Volume Water (m ³)	0.017			
Volume Sampled, 273K, 101.3kPa (dry m ³)	0.247			
Volume Sampled, 273K, 101.3kPa (wet m ³)	0.264			
Volume NaOH Impinger 1 (ml)	330			
Analytical Data				
HCI Blank (mg/l)	<0.05			
HCN in NaOH Imp1 (mg/l)	2.30			
total HCN (μg)	759			
Emission Concentration Data				
Moisture (%vol)	6.3			
HCN (mg/m ³)	2.87			

TABLE 13

COMBUSTION GAS & VOC EMISSION DATA SUMMARY

	O ₂	C	:0	Total VOCs		
Stack Ref	(%vol) ppm mg/m ³		mg/m ³	ppm (as C ₃ H ₈)	mg/m ³ as C	
Wood Burning Boiler	15.2	130.4	599.6	1356.4	2179.9	
Uncertainty (±)	0.4	-	14.5	-	49.7	

APPENDIX 1

Certificate of Analysis



Scientific Analysis Laboratories Ltd

Certificate of Analysis

Hadfield House Hadfield Street Combrook Manchester M16 9FE Tel : 0161 874 2400 Fax : 0161 874 2404

Scientific Analysis Laboratories is a limited company registered in England and Wales (No 2514788) whose address is at Hadfield House, Hadfield Street, Manchester M16 9FE

Report Number: 335526-1

Date of Report: 24-Jun-2013

Customer: Resource Environmental Consultants Ltd Unit 19 Bordesley Trading Estate Bordesley Green Road Birmingham B8 1BZ

Customer Contact: Ms Michelle Edwards

Customer Job Reference: 71519 Customer Site Reference: 11&12 June, 2013 Date Job Received at SAL: 17-Jun-2013 Date Analysis Started: 19-Jun-2013 Date Analysis Completed: 24-Jun-2013

The results reported relate to samples received in the laboratory

Opinions and interpretations expressed herein are outside the scope of UKAS accreditation This report should not be reproduced except in full without the written approval of the laboratory Tests covered by this certificate were conducted in accordance with SAL SOPs All results have been reviewed in accordance with QP22





Report checked and authorised by : Mary Drury Project Manager Issued by : Mary Drury Project Manager

SAL Reference: 335526 Project Site: 11&12 June, 2013 Customer Reference: 71519

Analysed as Wash(Acetone)

Wash(Acetone)

Miscellaneous SAL Reference 335526 002 335526 004 335526 006 335526 008 335526 010 71519/8 71519/14 Customer Sample Reference 71519/2 71519/4 71519/6 Test Sample AR AR AR AR AR Determinand Method LOD Units Symbol <0.3 Particulates (Total) Grav 0.3 mg U 0.8 0.8 1.0 0.7

SAL Reference: 335526 Project Site: 11&12 June, 2013 Customer Reference: 71519

Wash(Acetone) Analysed as Wash(Acetone)

Miscellaneous

		SAL	Reference	335526 012	335526 014	335526 016				
0			SAL Reference							
Customer Sample Referenc										
	Test Sample				AR	AR				
Determinand Method LOD Units Symbol										
av 0	0.3	mg	U	0.8	<0.3	0.5				
			thod LOD Units	thod LOD Units Symbol	thod LOD Units Symbol	thod LOD Units Symbol				

SAL Reference: 335526

Project Site: 11&12 June, 2013 Customer Reference: 71519

Filter Quartz 110mm Analysed as Filter Quartz 110mm Miscellaneous

			335526 001	335526 003	335526 013	335526 015		
		Custor	71519/1	71519/3	71519/17	71519/19		
	Test Sample						AR	AR
	Filter Reference						477	459
Determinand	Method	LOD	Units	Symbol		126.00		1.5
Particulates (Total)	Grav (5 Dec)	0.10	mg	U	<0.10	<0.10	<0.10	<0.10

SAL Reference	: 335526							1
Project Site	: 11&12 June,	2013						
Customer Reference	: 71519							
Filter Quartz 85mm Miscellaneous	Analysed as	Filter Qua	artz 85mm					
			SA	L Reference	335526 005	335526 007	335526 009	335526 011
		Custo	mer Sampl	e Reference	71519/5	71519/7	71519/13	71519/15
			٦	Fest Sample	AR	AR	AR	AR
			Filte	r Reference	858	865	818	868
Determinand	Method	LOD	Units	Symbol				
Particulates (Total)	Grav (5 Dec)	0.10	mg	U	<0.10	<0.10	<0.10	<0.10

Index to symbols used in 335526-1

Value	Description	
AR	As Received	
U	Analysis is UKAS accredited	





		Test	Certificate		Date 19/08/2013
Client	REC Ltd Unit 19			Drder No.	71519
		na Estato	C	Certificate No.	WK13-4922
		Bordesley Trading Estate Bordesley Green Rd			1
	Birmingham				
	B8 1BZ				
Contact	Michelle Edw	ards	C	Date Received	09/08/2013
Description	3 solutions for fo	ormaldehyde		echnique	Wet Chemistry
Sample No.	755209	71519/22			Method
					Wellou
Formaldehyde		<0.1 µg/ml	420 ml		M103(U)
Sample No.	755210	71519/23			Method
Formaldehyde		<0.1 µg/ml	395 ml		M103(U)
Sample No.	755211	71519/24	·		Method
Formaldehyde		<0.1 µg/ml	228 ml		M103(U)



Test Certificate

		Test Certific			
Client	REC Ltd		Certificate No. Issue No.	WK13-4922 1	
Tested By	Ashley Lunt	Date	16/08/2013		
Approved By	g. Q	Date	19/08/2013		
For and on authority	Joanne Dewnurst Laboratory Manager of RPS Laboratories Ltd.				
Method Symbols	(U) Analysis is UKAS Accredited(N) Analysis is not UKAS Accredited				
accreditation.	/m 3 and ppm) are provided to assist with interpretation efering to the sample volume.	only, they are not cov	ered by the scope of UKAS		
RPS Laboratories terms and conditions apply - a copy is available on request. Analysis carried out on samples 'as received' This document may not be reproduced other than in full, except with the written approval of the issuing laboratory.					

APPENDIX 2

Photographs of Sampling Locations



Pre-treatment Drying Oven

Refrigeration Powder Coat





Automatic Line



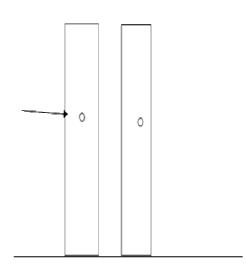




Wood Burning Boiler



Wood Spraybooth's No.1& 2



APPENDIX 3

Calculations

Conversion Factors

ppm ® mg/Nm³ (at 273K, 101.3kPa: STP)					
CO	х	1.25			
SO ₂	х	2.86			
VOC's	х	1.61	(ppm as C_3H_8 to mg/Nm ³ as C)		
NO _X	х	2.05	(ppm NO + NO ₂ to mg/m ³ as NO ₂)		

Oxygen Correction to Reference Value

Concentration at (STP) -> Concentration at 273K, 101.3kPa, reference O_2 and Dry Gas, i.e. Concentration X ((20.9- O_2 ref)/(20.9- O_2 measured)) = Concentration at ref Oxygen state.

Example Calculation

SO ₂ concentration at STP	=	170.7 mg/Nm ³			
Oxygen percentage in gas stream	=	13.8%			
Reference Oxygen	=	11%			
SO_2 concentration at reference O_2 conc	litions	=	170.7 ((20.9-11)/(20.9-13.8)) 238 mg/Nm³ at 273K, 101.3kPa, 11% O ₂ and Dry Gas		
Moisture Correction (Wet to Dry)					
Concentration of Gas Dry =	Concentration of x 100/100-Bws Gas Wet				
Concentration of Gas Wet =	Concer	ntration c	of x 100-Bws/100 Gas Dry		
Where Bws = moisture content of gas stream in percent (Vol/Vol).					
Example					

VOC concentration	=	25 mg/Nm ³ (Wet)
Moisture Content	=	27.1%
Concentration of VOC	=	25 (100/(100-27.1))

Carbon (C) to Trichloethylene (TCE)

ppm TCE = ppm C x 0.6715 TCE in mg/m³ = TCE ppm x 5.864 (Mol Wt/22.4)