GUIDANCE PACK FOR APPLICATION OF THE EMISSION LICENCE FOR STATIONARY SOURCES AS PER AIR QUALITY REGULATIONS, 2014.

Contents

- (1) Form IV: Application for Initial Emission License
- (2) Annex 1: Guidelines on Filling the Emission License Application Form
- (3) Annex 2: Stack Emission Report Format
- (4) Annex 3: Ambient Air Quality Monitoring Report Format
- (5) Annex 4: Guidelines for Construction of Particulate Matter Sampling and Testing Facilities
- (6) Annex 5: List of laboratories designated by NEMA to carry out air quality measurements
- (7) Annex 6: Emission License Fees as per the 13th Schedule
- (8) Annex 7: List of Controlled Areas as per 6th Schedule
- (9) Annex 8 : Emission Limits for Controlled and Non-Controlled Facilities

 3rd Schedule
- (9) Annex 9: Acceptable Emission Control Technologies 8 Schedule

NINTH SCHEDULE

EMISSION LICENSES

r41

Form IV: Application for Initial Emission License

2. Addre F T	e of Company
_	on LR No Street Area Division CownDistrict Province
4. Туре	of Industry
	e(s) of emitting Equipment
(a) I (b) F (c) N	Plan Layout, (attach sketch) Distance of the Stack to the nearest building
	rating Emission levels
, ,	
` '	
, ,	
8. Prope	osed Emission Control Mitigation Measures
` '	
, ,	
(viii)	

9. Additional	information required
10. Start-up, a) b) c) d) (ii) (iii) (iii) (iv)	and shut-down of the equipment Methods. Expected Frequency of Occurrence Duration of occurrence. Projected emitted Pollutants
ll. (a) Natur	e of emissions (gaseous, Particulates)
<i>(i)</i>	
(ii)	
(iii)	
(iv)	
(i) (ii)	entration of the emissions
	ture of ApplicantDate
FOR O	FFICIAL USE
	oved/Not approvedof 20
Signa (Seal)	ure

ANNEX 1: GUIDELINES ON FILLING THE EMISSION LICENCE APPLICATION FORM

Form IV: Application for Initial Emission License

r41

- 1. Name of Company: (As indicated in the Certificate of Registration)
- 2. Address:

P.O. Box: (As indicated in the Letter Head)

Tel: (As above); Fax: (As above); E-mail: (official email details)

Name of Contact Person: (Legal Entity of the company)

3. Location

LR No. (As indicated in the title deed); Street: (Name of road or street); Area: (Name of the location); Division: (Name of Sub-County); Town: (Name of urban center); District: (Name of County); Province: NA

- 4. Type of Industry: (Indicate production sector)
- 5. Name(s) of emitting Equipment:

e.g.:

- a) Generators;
- b) Boilers
- c) Incinerators
- d) Production equipment
- 6. Site Plan Layout, (attach sketch)
 - (a) Distance of the Stack to the nearest building (X meters)
 - (b) Height of the above referred building(Y meters)
 - (c)Nearest sensitive area or facility (As guided under 6th Schedule Annex 6)
 - (d) Immission (fall-out) point (Point determined from the dispersion model)
- 7. Operating Emission levels: (Allowable emission limits as indicated in the 3^{rd} Schedule –Annex 7)

For instance:

- (a) Parameter: $V \mu g/Nm^3$
- (b) Parameter: $X \mu g/Nm^3$
- (c) Parameter: Y µg/Nm³
- (d) Parameter: $Z \mu q/Nm^3$
- 8. Proposed Emission Control Mitigation Measures: (As guided under 7th Schedule-Annex 8)

For instance:

- (a) PM Mechanical Collectors (cyclones)
- (b) VOC Adsorption/absorption systems, flares, oxidizers and bio filters

- (c) SOX dry or wet scrubbers
- (d) NOX Catalytic reduction systems or EGR
- 9. Additional information required: Attach Stack Emission Report and Ambient Air Quality Monitoring Report as per the attached report format Annex 2 and Annex 3 respectively.

NB. Stack Emission and Ambient Air Quality Measurements shall be carried out by a laboratory designated by NEMA for the Air media. (List of designated laboratories indicated under Annex-4)

- 10. Start-up, and shut-down of the equipment
 - a) Methods. *Emission Reduction measures*
 - b) Expected Frequency of Occurrence: (Number of times in a day)
 - c) Duration of occurrence: (Time in minutes and start time of the occurrence)
 - d) Projected emitted Pollutants **For instance:**
 - (i) Particulate Matter
 - (ii) Volatile Organic Compounds
 - (iii) Sulphur Oxides
- 11. (a).Nature of emissions (gaseous, Particulates)
 - (i) **Particulates**
 - (ii) Gaseous
 - (iii) Odour

 (b) Concentration of the emissions Emission Results from the stack emission report. (a) Parameter: V₁ μg/Nm³ (b) Parameter: X₁ μg/Nm³ (c) Parameter: Y₁ μg/Nm³ (d) Parameter: Z₁ μg/Nm³
Signature of ApplicantDate
FOR OFFICIAL USE
Approved/Not approvedof 20 Signaturedayof

NB. Attach a copy of bank deposit slip for payment of Emission licence and its application fees as guided under Annex 5.

Annex 2: Stack Emission Measurement Report Format

The results summary report must be signed by the licensee.

Format for the Stack Emission Measurement Report

The stack emission measurements report must include:

1.0 General Source Information

- 1.1 Name of facility
- 1.2 Location of plant (Physical address)
- 1.3 GPS Reference
- 1.4 Legal Entity at the facility including their telephone numbers and Email.

2.0 Testing Firm Information

- 2.1 Name and address of the firm conducting the testing and Analysis.
- 2.2 Responsible person at the testing firm and telephone number.
- 2.3 Certification by team leader
- 2.4 Certification by reviewer

3.0 Introduction

- 3.1 Purpose of the test
- 3.2 Date and time of the test
- 3.3 Test location
- 3.4 Type of process
- 3.5 Air Pollutants tested
- 3.6 Aim of the Measurements
- 3.6 Observers' names if available
- 3.7 Any other important background information

4.0 Objective

- 4.1 Statement of objectives of the monitoring plan
- 4.2 Specific objectives

5.0 Production Process Information

- 5.1 Representative raw materials and operation
- 5.2 Types of fuel used in production
- 5.3 A description of the process equipment, including a process flow sheet.
- 5.4 Type and quantity of raw materials, catalysts, and products being used or produced in the process.
- 5.5 Maximum rated capacity of the process.
- 5.6 Actual maximum operating capacity of the process.
- 5.7 Operating capacity during the previous six-month period.
- 5.8 Process data being monitored and recorded to ensure representative operation throughout the test, including during the sampling interval.
- 5.9 Normal process operating schedule during a 24-hour period.
- 5.10 Normal maintenance schedule for this process.

6.0 Emission Control Equipment:

- 6.1 Description of emission control system, including the types of control equipment, manufacturer of control equipment, the rated capacity and efficiency of the control equipment.
- 6.2 Schematic diagram identifying gas flow from source point to exit point, including identification of all control devices, fans and bypasses
- 6.3 Data to be monitored and recorded to ensure representative operation of the control equipment during testing and the sampling period for recording.
- 6.4 Minimum acceptable values of all control devices parameters such as flow rates, pressure drops, temperature, and voltage of electrical input.
- 6.5 Description of any preconditioning of gases used with control equipment.
- 6.6 Normal maintenance schedule on control equipment for the previous year Description of source and control devises.
- 6.7 Process and control equipment diagram.
- 6.8 Process and control equipment data.
- 6.9 Demonstration of any specially-required operation

7.0 Operating Conditions

- 7.1 Type of process.
- 7.2 Plant operating conditions during the test
- 7.3 Brief description of the section of the plant under test relating it to the entire plant

8.0 Monitoring Plan Details:

- 8.1Name and model of equipment used,
- 8.2 Indicate the serial numbers of the equipment
- 8.3 A description of the emission sampling and analysis equipment including a schematic diagram of the sampling train.
- 8.4Calibration certificates of the equipment;
- 8.5 Methods of measurement, and
- 8.6 Quality assurance program; and
- 8.7 Relevant quality control steps for the monitoring

9.0 Stack Details:

- 9.1 Internal stack/duct diameters or stack/duct dimensions (metres) at the test plane;
- 9.2 Internal stack/duct diameter or stack/duct dimensions (metres) at the outlet of the stack;
- 9.3 Stack height in metres above ground level;
- 9.4 Location of sampling plane with respect to the nearest upstream and downstream flow disturbances
- 9.5 Stack/duct location (latitude and longitude with datum reference) and reference within the plant altitude above mean sea level of the base of the stack;
- 9.6 Attach photographs of the full stack showing measurements in progress.

10.0 Details of the Sampling Point:

- 10.1 A scale diagram showing each discharge point with each testing access port; and
- 10.2 Distances from upstream and downstream flow disturbances;
- 10.3 A scale diagram showing the sampling platform and access; the dimensions of all access ports.
- 10.4 Sampling port location and cross-section with dimensions.
- 10.5 Sampling port description, including labeling systems

11.0 Emission Point Information:

- 11.1 A diagram of the stack showing the dimensions, and the configuration of the sampling location, location of continuous emission monitor (CEM) probes, and the distances to the nearest upstream and downstream flow interferences.
- 11.2 A cross-sectional sketch of the stack at the sampling location, showing the locations of the sampling traverse points.
- 11.3 Estimated flue gas conditions at the sampling location, including temperature, moisture content, and velocity.

12.0 Test Methods

- 12.1Types of pollutants to be sampled and analyzed
- 12.2 A description of the sampling and analysis procedures
- 12.3 Brief description of test procedure(s) used and any variations from those procedure(s) and reasons for selecting non-recommended methods;
- 12.4 Description of analysis procedures, and any deviations from standard procedures
- 12.5 Documentation for any proposed variations from approved/standard procedures
- 12.6 Sample time if different than recommended
- 12.7 Table of results obtained
- 12.8 Provide sample calculations
- 12.9 Attach countersigned raw data

13.0 Data and Results

- 13.1 Gaseous Emissions
 - a) Legible copies of the measuring equipment data-Logger print-outs.
- b) a photograph of the whole stack presenting sampling in progress for gaseous emissions and of the stack sampling point.
 - c) Table of raw data
- d) Table of results referenced to STP and corrected for the relevant gas concentration

- 13.2 Particulate Matter Emissions
- a) Average stack/duct gas velocity (metres/second) at STP;
- b) Average ambient barometric pressure during the test (kilopascals);
- c) Average stack/duct gas pressure during the test (Pascals);
- d) Average stack/duct gas flow rate, (cubic metres/second) at STP;
- e) Average stack/duct gas temperature, K or degrees Celsius;
- f) Moisture content, % v/v or w/v;
- g) Composition of stack/duct gas where applicable, % volume, dry basis;
- h) Concentration of the pollutant under test such as total particulates, heavy metals, or any specific gaseous pollutant, mg/m3 at STP, dry basis, adjusted if necessary to reference conditions;
- i) Estimate of the measure of uncertainty for the test results;
- j) Comments on whether data exceed the Authority emission limits and any other factors which may have affected results.
- k) Indicate the last volume reading of the isokinetic particulate matter sampling equipment

14.0 Data Sheets:

Copies of all field data sheets used during the test, including data sheets used to record process and control equipment parameters.

15.0 Chain of Custody:

- a) A description of the procedures that were followed to maintain the integrity of the samples collected.
- b) Copies of chain of custody seals and data sheets.

16.0 Quality Control:

- a) Calibration sheets for the dry gas meter, pitot tube, nozzle, and other equipment requiring calibration.
- b) Quality assurance control charts for the analytical procedures used in the analysis of the test samples.
- c) A list of pre-weighed filters during particulate emissions testing.

17.0 Summary of Results

- a) Process data
- b) Emission results.
- c) Allowable emissions.
- d) Discussion of errors, both real and apparent.

18.0 Discussion and Recommendations

19.0 Annexes

- a) Complete results with example calculations.
- b) Copies of original field and laboratory data sheets.

- c) Copies of original production data signed by plant official.
- d) Test log- data loggers
- e) Project participants and titles.
- f) Related correspondence.

NB.

- (i) If an Air pollutant cannot be detected, results **must not** be quoted as zero but as less than the method's limit of detection.
- (ii) All volumes and concentrations are normally reported as dry at a temperature of 0°C and at an absolute pressure of 101.3 kilopascals (kPa).
- (iii) In case the sampled data deviates between consecutive runs beyond ±10%, you are required to carry out run 3

(Note: STP is 0° Celsius and 1 standard atmosphere pressure.)

Other Requirements:

- 1) Submission of the Stack Emission Measurement Report
 The Submission of the Stack Emission Measurement Report/Test results
 must be forwarded to the Authority as outlined in the Emission License
 conditions, and in any case within TEN (10) weeks after the test date.
 The report should be spiral bound.
- 2) Where testing for reporting to the Authority is of a continuous nature over an extended period of time (greater than that for a corresponding manual stack emission test), the results must be reported as indicated in the emission license or as agreed beforehand with the Authority.
- 3) Reports must be submitted to the Authority in the format required by the Authority
- 4) This is to be accompanied by a letter of certification from the licensee (facility).
- 5) The letter of certification is a statement by a person authorized to sign on behalf of the licensee that the provided report is a true and correct record of the testing conducted on the premises.
- 6) Testing and analytical laboratory information: All information concerning testing, monitoring and ancillary equipment, charts, raw data, calibration records and other relevant documentation must be attached to the report as indicated under **item 11 below.**

- 7) Monitoring equipment includes both ambient and stack monitoring equipment for both continuous and discrete testing. All data gathered from the monitoring equipment, along with a copy of the test report, must be stored on a computer data system available through the company premises for seven years or the operational period of the plant whichever is the lesser.
- 8) Information supplied may, at the discretion of the Authority, be published in full, in summary form, in annual or other reports with reference to the source.
- 9) The test and analytical laboratories must determine a person who is a director, or legally responsible person in the company to sign documents on behalf of the company, to verify the correctness of any reports or documents submitted to the Authority. All documents related to a test, submitted to the Authority or under emission licence requirement must be signed by the nominated person.

10. Adjustment to reference conditions

10.1 Under the Air Quality Regulations, it is a mandatory requirement that all stack emissions must be reported to 0°C, 1 standard atmosphere pressure and on a dry basis (STP dry).

10.2 In addition some emissions defined under the emission standards must be adjusted to a reference concentration of oxygen or carbon dioxide. This requires that oxygen and/or carbon dioxide concentration must also be measured. Adjustments are to be made as follows:

10.2.1 Adjustment to a reference concentration of carbon dioxide

$$Ca = Ci \times (CO_2r)/(CO_2m)$$

Where:

Ca = adjusted emission concentration at 0 °C and 1 standard atmosphere on a dry basis having a specified reference concentration of carbon dioxide volume % CO_{2r}

Ci = initial unadjusted pollutant emission concentration at 0 °C and 1 standard atmosphere on a dry basis and at known carbon dioxide concentration (CO₂m) before admixture with air or other gases

 $CO_2r = a$ specified reference concentration of carbon dioxide (volume %)

 CO_2m = actual measured carbon dioxide concentration (volume %)

10.2.2. Adjustment to a reference concentration of oxygen

Cb = Ci x
$$(20.9 - O_2r)/(20.9 - O_2m)$$

where:

- Cb = adjusted emission concentration at 0 °C and 1 standard atmosphere pressure having a specified reference concentration of oxygen volume % O₂r
- $O_2r = a$ specified reference concentration of oxygen (volume %)
- Ci = initial unadjusted emission concentration at 0 $^{\circ}$ C and 1 standard atmosphere on a dry basis and at a known oxygen concentration (O₂m) before admixture with air or other gases

 O_2m = actual measured oxygen concentration (volume %)

In both adjustments, the value reported to the Authority for purposes of compliance is the adjusted value (Ca or Cb) at its reference carbon monoxide or oxygen concentration along with the measured oxygen and/or carbon monoxide concentration.

11. Data Charts

USEPA Method 1a USEPA Method 1— Sample and Velocity Traverses for Stationary Sources

Table 1 Required Number of Traverse Point for Sampling Sites (Clause 5.3)

Stack/Chimney No.	Stack Diameter m	Total No. of Traverse	Number of Traverse		
			Plane 1	Plane 2	
	(2)	(3)	(4)	(5)	

Signed by: Labora	atory Ar	alyst	 Date	
Countersigned by	y Facility	Representative	 Date	

USEPA Method 2a USEPA Method 2: Velocity Traverse Determination $% \left\{ 1,2,\ldots ,n\right\} =0$

a)	Plant or Facility
b)	Date
c)	Run No
d)	Stack Dia. or Dimensions, (M.)
e)	Stack Height. or Dimensions, (M.)
f)	Barometric Press., (mm. Hg)
g)	Cross Sectional Area, (M ²)
h)	Operators
i)	Pitot Tube I.D. No
j)	Avg. Coefficient, Cp =
k)	Last Date Calibrated

Schematic Of Stack Cross Section

		Stack temperatur	re		
Traverse Pt. No.	Vel. Hd., Δp (mm.) H ₂ O	T _s , °C ((°deg;)	T _s , °K (°R)	P _g (mm. Hg)	(Δ p)1/2
	Average(1)				

Signed by: Labo	ratory Analyst	Date							
Countersigned b	ountersigned by Facility Representative								
USEPA Method USEPA Method	3a 3- Gas Analysis for the Detern	nination of Dry Molecula	r Weight						
Table 3: Samplin	g Rate Data								
			% Deviation ^a						

Time	Traverse point	Q (liter/min)	% Deviation ^a
Average			

^a % Dev.= $[(Q-Q_{avg})/Q_{avg}]\times 100$ (Must be $\leq \pm 10\%$)	
Signed by: Laboratory Analyst	Date
Countersigned by Facility Representative	Date
5 , , ,	

USEPA Method 4a

USEPA Method 4-1 Moisture Content Determination

Table 4: Moisture Content

			Pressure differential	Meter		Gas sample temperature at dry gas meter		Temperature of gas
	Sampling	Stack temperature	across orifice	reading gas sample		Inlet Tm _{in}	Outlet Tm _{out}	leaving condenser or last
Traverse Pt.	time	°C ((mm (in.)	volume	$\Delta V_m m^3$		°C ((impinger
No.	(Δ) , min	°deg;F)	H_2O	m^3 (ft ³)	(\mathbf{ft}^3)	°deg;F)	°deg;F)	°C ((°deg;F)
Average								

			yst Representativ							
004111011	g,	1 401111, 1	oprosoman							
USEPA N 1) P m) D n) R o) S p) S q) B r) C s) C t) P u) A	lant or Factorial lant or Factorial land land land land land land land la	Particula cility Or Dimer nt. Or Dir Press., (i onal Are L.D. No cient, Cr		I.)		-				
Traver se Point Numb er	Sampli ng Time (Min)	Vacuu m (mm Hg)	Stack Temperat ure (°C)	Velocity Head (ΔP)mm H ₂ O	Pressur e Differen tial across orifice meter (mmH ₂ O)	Gas Meter Readi ng	ure a Gas	perat at Dry Meter Outl et	Temperat ure	Temperat ure of Gas leaving condense r or last Impinger
							(°C	(°C)		

Signed by: Laboratory Analyst	Date
Countersigned by Facility Representative	Date

Annex 3: Ambient Air Quality Monitoring Report Format

The results summary report must be signed by the licensee

The report must contain at least the:

- a) Name and address of reporting facility
- b) Date of issue of the report
- c) The test method used and details of any deviation from that method
- d) Period of monitoring (start and end dates and percentage of time the instruments were online)
- e) Location of monitoring points (normal address and GPS reference, height above nominal ground level, and a description of the terrain features)
- f) The air pollutants measured, the monitoring instruments used, and a description of the air sampling system
- g) Maximum hourly average concentration, daily average concentration, and appropriate longer-term averages
- h) Appropriate statistical information to describe the variability and range of the concentrations
- i) Any factors that may have affected the monitoring results
- j) The precision of the results in accordance to ISO 5725
- k) Details of the most recent calibration of each instrument used to take measurements.

NB.

- (i) If an Air pollutant cannot be detected, results **must not** be quoted as zero but as less than the method's limit of detection.
- (ii) Attach a copy of the measuring equipment data-Logger print-out.

Annex 4: Guidelines for Construction of Particulate Matter Sampling and Testing Facilities

1 Introduction

Most often the installation of sampling/testing ports is an afterthought in the process of stack construction. However, sampling ports are essential to the stack testing process which is a requirement for demonstrating compliance. The Authority recommends that EPA method 1 should be taken into consideration before stack construction begins.

1.1. Selection of a sampling site

Selection of a proper sampling site is vital to assure representative measurement of pollutant emissions and accurate volumetric flow rate from a stationary source is obtained. Safety and accessibility are essential considerations of site selection. Clearance for sampling equipment, access to electrical power, exposure of personnel to weather, process heat, and presence of hazardous gases must also be considered in construction of sampling facilities.

2 Regulatory Basis

The Air Quality Regulations, 2014 states:

- 61 (2)An owner or operator of a controlled facility shall provide portholes, and platforms which shall be conveniently located for easy access and all other facilities required for taking samples of air or emission from any chimney, flue or duct, plant or vessel or any other outlets.
- 62. Where the Authority requires stack emission tests to be performed under these Regulations, an owner of a facility shall provide the following-
 - (a) sampling ports which are adequate for the test methods applicable to the facility;
 - (b) safe sampling platforms or other suitable and safe permanent or temporary structures or equipment; and
 - (c) safe access to sampling platforms.

3 Sampling Ports

The ideal sample location would have flow characteristics such that the gas flows in a linear pattern parallel to the axis of the stack or duct, have a uniform velocity profile and not flow in a cyclonic or swirling pattern. The following port location directions are provided to guide you in selecting a stack sampling location close to ideal.

1.1. Port Location.

Ports should be located at least eight stack or duct diameters downstream and two diameters upstream from any flow disturbance such as a bend, expansion, or contraction in the stack, or from a visible flame. For a rectangular stack or duct utilize an equivalent diameter to determine distances from flow disturbances. Calculate the equivalent diameter as follows:

Equivalent Diameter = De

$$D_{c} = \frac{2LW}{L+W}$$

where

D_{e}	Ш	equivalent diameter, in m;
		length of the rectangle, in m; and
W	=	width of the rectangle, in m.

3.2. Port Type

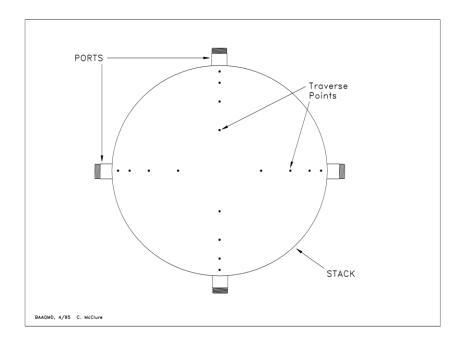
A sampling port should be a welded pipe nipple with screw cap or a pipe with flange. A gate valve should be used for stacks handling hazardous gases or gases over 94°C under positive pressure. Stacks or ducts having a thin wall may be provided with ports consisting of appropriately sized surface holes with plugs. For determination of Total Suspended Particulate (TSP) emissions, the sampling ports must be at least 87.5mm I. D. (Inside Diameter) pipe nipple. Particulate sampling to determine the PM10 or PM2.5 (Particulate Matter less than 10 or 2.5 microns) fraction of the TSP emissions requires sampling ports be 150mm I. D. pipe nipples.

3.3 Port Installation

Ports must be flush with the interior stack wall and extend outward from the exterior wall at least 50mm and not more than 150mm. Ports must be located at least 1.2m and not more than 1.5m above the work platform floor and meet clearance zone requirements described below. The ports must be located parallel and in a plane perpendicular to longitudinal axis of the stack.

3.4. Number of Ports

For circular stacks, if the sum of the stack inside diameter plus the port length is 1.625m or less, two ports must be installed 90 degrees apart. If this sum is over 1.625m, four ports must be installed 90 degrees apart. See Figure 1.



3.1 <u>Figure 1</u>

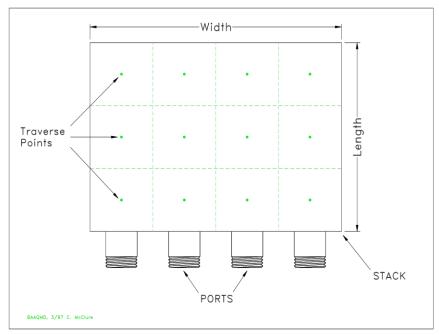
Example of port location for a round stack requiring four ports

In the case of rectangular stacks, an equivalent diameter will be calculated, and a cross section layout (matrix layout) applied. The cross section layout is specified in the Method 1 (EPA Method 1-Sample and Velocity Traverses for Stationary Sources). Locate the number of ports required by the cross section layout on the widest of the two stack sides. Contact the facility manager to provide assistance in determining the correct cross section layout (matrix layout) for rectangular port placement. See Figure 2.

4 Work Platform

1.2. Size

For a round stack with two-port installation the platform must serve that quarter of the stack and extend at least 900mm beyond each port. It must be at least 1200mm wide. For a circular stack with four-port installation the platform must serve the full circumference of the stack and be at least 1500mm wide. For a rectangular stack the platform must serve the side containing the ports. It must be at least 1200mm wide and extend across the full width of the stack.



4.1 <u>Figure 2</u> Example of port location for a rectangular stack with a 4X3 traverse point matrix

4.2. Construction

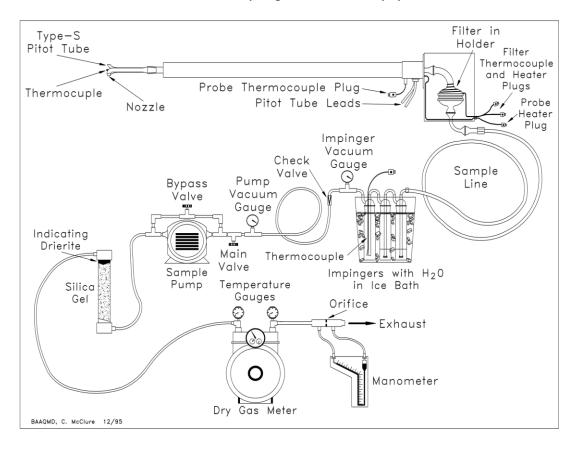
Safe and easy access to the work platform must be provided by stairway, caged ladder, or other suitable means. The platform must have a guardrail and toeboard. It must be able to support 500Kg live load. All construction must conform to the OSHA, 2007 requirements. Platforms more than 21m high require a swinging arm and pulley to assist the lifting of test apparatus.

2.0. Clearances

Insertion of a long stack sampling probe with filter assembly on its end is a difficult task. See Figure 3 for a diagram of the sampling probe and equipment. There must be a clearance zone, with no obstructions, around each port extending 300mm above, 600mm below, and 600mm on each side of the port. The zone must extend outward from the end of the port at least one stack diameter plus 900mm. Guardrails that enter the clearance zone must be equipped with removable sections or chains.

Conducting PM₁₀ or PM_{2.5} sampling requires site-specific clearances and probe support devices, i.e. a support rail system/bracket. See Figure 4. Contact the Facility Manager for site-specific guidance.

Particulate Sampling Probe and Equipment



4.2 Figure 3 Particulate Sampling Probe and Equipment

Typical Stack Sampling Access Requirements

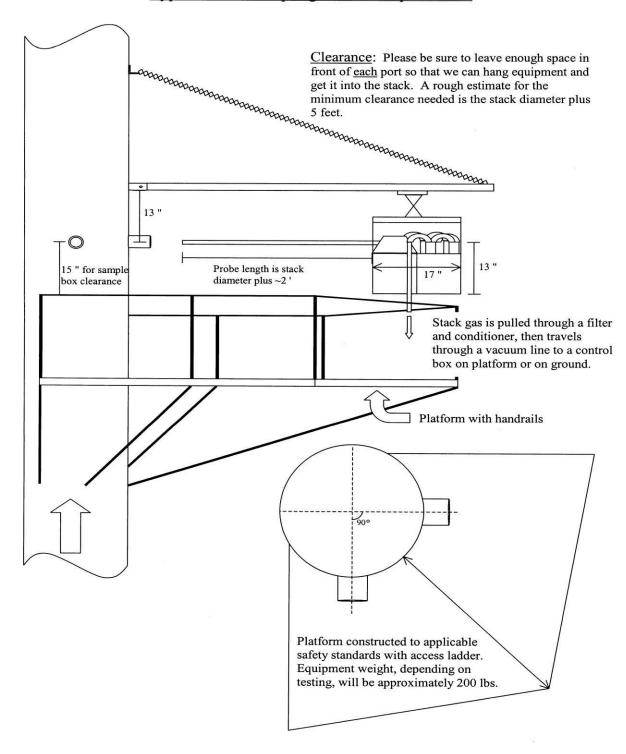


Figure 4: Typical Stack Sampling Access Requirements

5 Power Supply

Sampling platforms over 21m high require two electrical outlets, each capable of supplying 230/115-volt, 15-ampere, 60 cycle AC. Sampling platforms less than 21m high require that two electrical outlets, each capable of supplying 115-volt, 15-ampere, 60 cycle AC be located within 15m of the sampling site.

6 Additional Requirements

Additional requirements may be specified for situations with special needs such as flow pattern adjustment, hoist equipment, or variations in the number and location of sampling ports.

Mark the Pitot tube at the points to be sampled accordingly as explained above. Mark with tape, asbestos rope or other suitable material compatible with flue gas conditions the required length of Pitot for each of the points.

Table 1 Required Number of Traverse Point for Sampling Sites (Clause 5.3)

SI No.	Range of Stack Diameter m	Total No. of Traverse	Number o	f Traverse
			Plane 1	Plane 2
(1)	(2)	(3)	(4)	(5)
i)	ID up to 0.3	4	4	_
ii)	ID>0.3 up to 0.6	8	8	_
iii)	ID>0.6 up to 1.2	12	12	_
iv)	ID>1.2 up to 2.4	20	10	10
v)	ID>2.4 up to 5.0	32	16	16
vi)	ID > 5.0	52	26	26

3

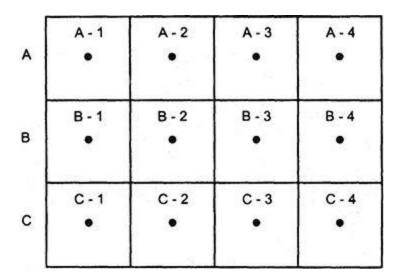
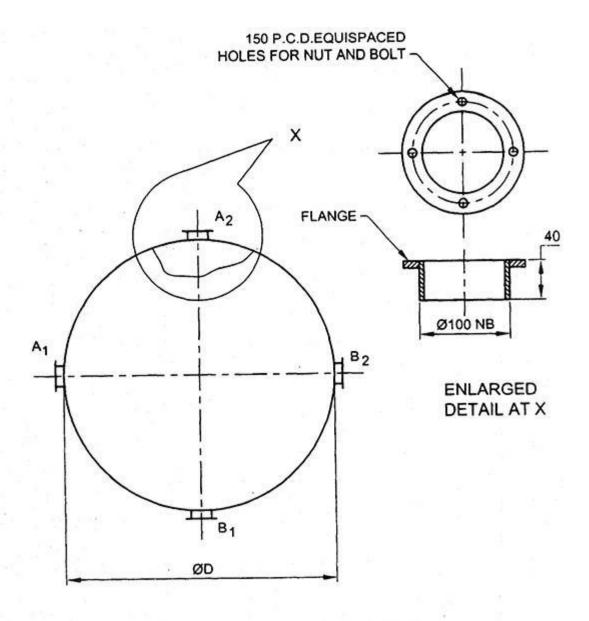


Fig. 3 Diagrammatic Representation of Equal Rectangular Sub-Areas for a Rectangular Duct



- 1) A_1 , A_2 When D < 2000
- 2) A_1 , A_2 , B_1 , B_2 When D < 2000

All dimensions in millimetres.

Fig. 4 Sampling Port

6

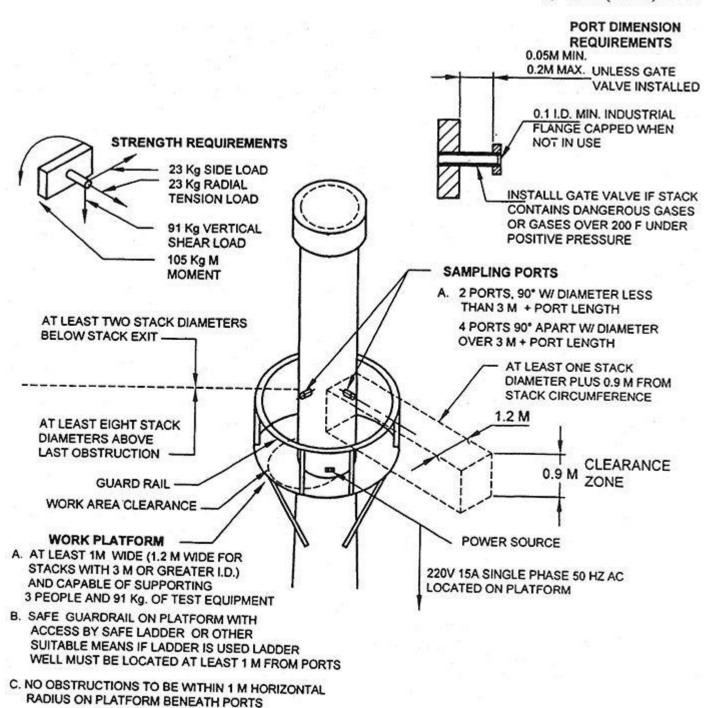


Fig. 5 Typical Sampling Provision

Annex 5: List of laboratories designated by NEMA to carry out air quality measurements: Available on the NEMA website- www.nema.go.ke; gazetted designated laboratories

	Laboratory			Sco	ре М	easurei	nent	
sn		Email & Mobile Contact.	Staci Emi:	k ssion	Amb Air Qua		Indo Qua	or Air lity
			PM	Gase s	PM	Gase s	PM	Gase s
1	SGS Kenya Limited P.O. Box 72118- 00200, NAIROBI	enquiries.kenya@sgs.com Landline +254 02 2733690/99 Tel: 0709633400, 0733637142	Yes	Yes	Yes	Yes	Yes	Yes
2	Ecoscience Engineering Services	0713566825/0711632342/ Ecoscience.engineering @gmail.com or info@ecoscience and engineering.co.ke	Yes	Yes	Yes	Yes	Yes	Yes
3	Eurolab Services Limited P.O Box 5050- 00200 NAIROBI	info@eurolabltd.com eurolablimited@gmail.c om 0722241766,0752435126	Yes	Yes	No	No	Yes	Yes
4	Labworks P.O.Box 9648 - 00100, NAIROBI	info@labworksea.com Tel: 0726209641	Yes	Yes	Yes	Yes	No	No
5	Polucon Services P.O. Box 99344 - 80100, MOMBASA	Polucon@polucon.com Tel: 0722229944	Yes	Yes	Yes	No	No	No
6	CSI International Ltd. P.O.Box 47846 - 00100, NAIROBI	info@csiinternationalke .co.ke Tel:0712603528/072347 9111/0705204610	Yes	Yes	Yes	No	No	No
7	Ecoserv Laboratory, P.O.Box 1303 - 00100, NAIROBI	gachanja@ecoservkeny a.com Tel: 0722882879	Yes	Yes	Yes	No	No	No
8	Instrumentation Engineers(EA) Limited P.O Box 18986- 00500, NAIROBI	info@instrumentation- engineers.com Tel: 0722357991/+254- 20-2108606	No	Yes	No	No	No	No

	Laboratory			Sco	pe M	easurei	ment	
sn		Email & Mobile Contact. Stack Emission No. Ves					Indo Qua	or Air lity
9	Kenya Bureau of Standards P.O.Box 54974- 00200, NAIROBI	info@kebs.org Tel: 0722203137	No	Yes	Yes	No	No	No
10	Institute of Nuclear Science & Technology P.O.Box 3097 - 00100, NAIROBI	inst@uonbi.ac.ke Tel: 020-318262	No	No	Yes	No	No	No

Key:

- PM: Particulate Matter/Dust (i)
- Yes: Capacity exists to measure the indicated parameters No: No capacity to measure the indicated parameters (ii)
- (iii)
- (iv) Equipment Inventory available for verification

THIRTEENTH SCHEDULE

FEES

The fees chargeable under these Regulations shall be as specified hereafter.

- (a) Application for:
 - (i) Emission License for listed emitting facility: KShs. 5, 000/=
 - (ii) Emission License for other emitting facility than (i) above: KShs. 5,000/=
 - (iii) Variation of emission license: KShs.3,000/=
 - (iv) Transfer of emission license: KShs.3,000/=
- (b) Annual License fee for Emission into the atmosphere
 - (i) Facility listed in 6th schedule under category I:- KShs. 50,000/=
 - (ii) Facility listed in 6th schedule under category II:- KShs. 30,000/=
 - (iii) Polluting facility not in 6th Schedule other than (i) and (ii) above:- KShs.20,000/=
- (c) Inspection of emission monitoring records/emission license register: KShs.200/=
- (d) Variation of emission License is 10% of the Annual License fee

The above fees are payable to the NEMA Revenue Account:

Details:

Account No. 1102298158

Bank Name: *KCB*, Branch: *KICC Branch*.

Annex 7: List of Controlled Areas

SIXTH SCHEDULE

LIST OF CONTROLLED AREAS

- a) Residential areas, Hospitals,
- b) National Parks,
- c) Reserves and Sanctuaries,
- d) Conservation areas,
- e) Central Business Districts
- f) Any other area declared by the Authority from time to time

Annex 8: Emission Limits for Controlled and Non-Controlled Facilities

7 r16,72

THIRD SCHEDULE

EMISSION LIMITS FOR CONTROLLED AND NON-CONTROLLED FACILITIES

Air Pollutant Industry	Opacity	Particulate (Dust) PM ₁₀ (mg/Nm ³)		Sulphur oxide	(SO _x) (mg/Nm ³)	Nitrogen oxides	(NO _x) (mg/Nm ³)	O ₂ %	Carbon monoxide	Carbon dioxide (mg/Nm ³)	Hydrocarbons	Hydrogen Sulphide	Hydrogen Chloride	Hydrogen Fluoride	Lead (mg/Nm ³)	Dioxins/Furans
		NDA	DA	NDA	DA	NDA	DA									
Aluminium recycling plants		10 – 30									20		*			*
Asphalt mixing batch plants		< 100 t: g/kg 100 to 300 t: 22g/k g 300 to 500 t: 31g/k g > 500 t: 33 g/kg		200		460			*	*	20					
Boilers	*	50		*		*			*	*	*	*				*
Cement plants		50		400		1500			*	500	30 0					0.5ng/N m ³
Ceramics manufacture		400				180- 250 ppm										

Air Pollutant Industry		Particulate (Dust)	I 17110 (1118/14111)	Sulphur oxide	$(SO_X) (mg/Nm^3)$	Nitrogen oxides	(NO _x) (mg/Nm ³)	O ₂ %	Carbon monoxide	Carbon dioxide (mg/Nm ³)	Hydrocarbons	Hydrogen Sulphide	Hydrogen Chloride	Hydrogen Fluoride	Lead (mg/Nm^3)	Dioxins/Furans
		NDA	DA	NDA	DA	NDA	DA									
Coke & coal plants		*		*		*			*	*	*	*	*			
Dairy		50														
Fertilizer plant		50		*		500					20	30		50		
Iron Foundry		50		560					*	*				5		
Brass bronze Foundry		50		20 - 50												
Glass Manufacture		20 - 50		Oil fire d: 1,8 00 Gas		1000 - 2000							5 0	5		
				fire d: 700												
Galvanizing operations	*	50														
Incinera tors	*	< 10 t: 4g/kg 10 to 30 t: 10g/k g		500		Exis ting: 130- 600 ppm			*		*					2.0 – ng TEQ/N
		30 to 50 t: 10g/k g														

Air Pollutant Industry		Opacity	Particulate (Dust)	1 1A10 (1118/1A111)	Sulphur oxide	(SO _x) (mg/Nm ³)	Nitrogen oxides	(NO _x) (mg/Nm ³)	O ₂ %	Carbon monoxide	Carbon dioxide	Hydrocarbons	Hydrogen Sulphide	Hydrogen Chloride	Hydrogen Fluoride	Lead (mg/Nm ³)	Dioxins/Furans
			NDA	DA	NDA	DA	NDA	DA									
			> 50 t: 17.5 g/kg				New : 60- 400 ppm										0.1 – 5 – TEQ/N
	Munic ipal waste		100				300			*	*						
	Medic al waste		20 (PM _{2.5}		500		300			*	*	*		*			*
	Indust rial waste	*	50		150		460			*	*	*	*	*			*
Kraft pulp	mills		100- 150		500		600			*	*	20	15	*	*		*
Lead Replants	ecycling		20 (PM _{2.5}		400											5 0 (B & R F u r n a c e	*

Air Pollutant Industry	Opacity	Particulate (Dust)	Particulate (Dust) PM ₁₀ (mg/Nm ³)		(SO _x) (mg/Nm ³)	Nitrogen oxides	(NO _x) (mg/Nm ³)	O ₂ %	Carbon monoxide	Carbon dioxide (mg/Nm ³)	Hydrocarbons	Hydrogen Sulphide	Hydrogen Chloride	Hydrogen Fluoride	Lead (mg/Nm ³)	Dioxins/Furans
		NDA	PA	NDA	DA	NDA	DA									
															1 0 P o t F u r n a c e)	
Mineral Processing Mining & Quarry	2	50 400														
	0 %															
Non-ferrous secondary smelters		50		20		*			*	*	*	*				
Non-ferrous secondary smelters	*	< 10 t: 7.5 g/kg (PM _{2.5}		800		*			*	*	20	15				*

Air Pollutant Industry	Opacity	Particulate (Dust)	Particulate (Dust) PM ₁₀ (mg/Nm ³)		(SO _x) (mg/Nm ³)	Nitrogen oxides	(NO _x) (mg/Nm ³)	02%	Carbon monoxide	$\begin{array}{ccc} \textbf{Carbon} & \textbf{dioxide} \\ (mg/Nm^3) & \end{array}$	Hydrocarbons	Hydrogen Sulphide	Hydrogen Chloride	Hydrogen Fluoride	Lead (mg/Nm ³)	Dioxins/Furans
		NDA	DA	NDA	DA	NDA	DA									
		10 to 30 t: 22.5 g/kg (PM _{2.5}														
		30 to 50 t: 37.5g/ kg (PM _{2.5}														
		> 50 t: 52.5 g/kg (PM _{2.5}														
Paint and varnish manufacturing		50 (PM _{2.5}									20	15	1 0			
Pesticides formulation		20 (PM _{2.5}									20		5			
Pesticide manufacturing		20									20					

Air Pollutant Industry	Opacity	Particulate (Dust) PM ₁₀ (mg/Nm ³)		Sulphur oxide	(SO _x) (mg/Nm ³)	Nitrogen oxides	(NO _x) (mg/Nm ³)	O ₂ %	Carbon monoxide	Carbon dioxide (mg/Nm ³)	Hydrocarbons	Hydrogen Sulphide	Hydrogen Chloride	Hydrogen Fluoride	Lead (mg/Nm ³)	Dioxins/Furans
		NDA	DA	NDA	DA	NDA	DA									
Petroleum Refineries		50		Sul phu r rec ove ry: 150 Co mb usti on unit s:5 00		460			*		20	15 2				*
Pharmaceuticals manufacturing plants		20									80		1 0			
Printing industry	*	Dwint		500		200			*		20		1 0			
Steel mills	*	Existi ng- 240 (PM _{2.5}) New- 120 (PM _{2.5})		500		180			. *							

Air Pollutant Industry	Opacity	Particulate (Dust)	T 1V10 (1118/1V111)	Sulphur oxide	(SO _x) (mg/Nm ³)	Nitrogen oxides	(NO _x) (mg/Nm ³)	O ₂ %	Carbon monoxide	Carbon dioxide (mg/Nm ³)	Hydrocarbons	Hydrogen Sulphide	Hydrogen Chloride	Hydrogen Fluoride	Lead (mg/Nm^3)	Dioxins/Furans
		NDA	DA	NDA	DA	NDA	DA									
Sulphuric acid Plants		50		SO 2: 2 kg/t aci d SO 3: 0.1 5 kg/t aci d												
				<pre>100 t: 3.7 5 g/k g 100 to 300 t: 10. 5 g/k g</pre>												

Air Pollutant Industry	Opacity	Particulate (Dust)	(III SIII) (III I	Sulphur oxide	$(\mathbf{SOx}) \; (\mathbf{mg/Nm}^3)$	Nitrogen oxides	(NO _x) (mg/Nm ³)	O ₂ %	Carbon monoxide	Carbon dioxide (mg/Nm ³)	Hydrocarbons	Hydrogen Sulphide	Hydrogen Chloride	Hydrogen Fluoride	Lead (mg/Nm ³)	Dioxins/Furans
Industry		NDA	DA	NDA	DA	NDA	DA									
				300 to 500 t: 34. 5g/ kg > 500 t: 48 g/k g												
Sugar Manufacture		(< 8.7 mw input boiler) : 150 (>8.7 mw input boiler) : 100		200		Liqu id fuels : 460 ppm Soli d fuels : 750 ppm										
Soda ash Manufacture		50											*			
Tanneries		50		100 0		1500					20	15	*			*
Textiles Geothermal Power plants		50		*		*					20	*				

Air Pollutant Industry	Opacity	Particulate (Dust)	(1118/1/8111) 0.1.41.1	Sulphur oxide	(SO _x) (mg/Nm ³)	Nitrogen oxides	(NO _x) (mg/Nm ³)	O ₂ %	Carbon monoxide	Carbon dioxide (mg/Nm³)	Hydrocarbons	Hydrogen Sulphide	Hydrogen Chloride	Hydrogen Fluoride	Lead (mg/Nm ³)	Dioxins/Furans
		NDA	DA	NDA	DA	NDA	DA									
Thermal Power Plants																
Small combustion facilities(3MWth – 50NMWth)Recip rocating internal Combustion Engine (RICE)																
Engine (Gas)		N/A		N/ A		200(SI) 1,60 0(CI										
Liquid	K S 1 5 1 5	50		Use 1.5 % sul phu r fuel (SF)*		Bore ø<4 00m m: 1460 Bore ø≥4 00m m: 1,85 0										
Turbine																. <u> </u>
Natural Gas																

Air Pollutant Industry	Opacity	Particulate (Dust)	()	Sulphur oxide	$(SO_X) (mg/Nm^3)$	Nitrogen oxides	(NO _x) (mg/Nm ³)	O ₂ %	Carbon monoxide	Carbon dioxide (mg/Nm³)	Hydrocarbons	Hydrogen Sulphide	Hydrogen Chloride	Hydrogen Fluoride	Lead (mg/Nm^3)	Dioxins/Furans
		NDA	DA	NDA	DA	NDA	DA									
3MWth to < 15MWth		N/A		N/ A		42pp m(El ectri c Gen erati on)(EG) 100p pm(Mec hani cal Driv e)(M D)										
15MWth to < 50MWth		N/A		N/ A		25pp m										
Other fuels		NT/A		0.5		06:-:-										
3MWth to < 15MWth		N/A		0.5 %S F		96pp m(E G) 150p pm(MD)										

Air Pollutant Industry	Opacity	Particulate (Dust)	1 1A10 (1118/1A111)	Sulphur oxide	(SO _x) (mg/Nm ³)	Nitrogen oxides	(NO _x) (mg/Nm ³)	O ₂ %	Carbon monoxide	Carbon dioxide (mg/Nm ³)	Hydrocarbons	Hydrogen Sulphide	Hydrogen Chloride	Hydrogen Fluoride	Lead (mg/Nm ³)	Dioxins/Furans
		NDA	DA	NDA	DA	NDA	DA									
15MWth to < 50MWth		N/A		0.5 % SF or low er		74pp m										
Boiler																
Gas		N/A		N/ A		320										
Liquid		50 or 150 *		200		460										
Solid		50 or 150 *		200		650										
Combustion Technology/Fuel RICE																
Natural Gas		N/A	N/ A	N/ A	N/ A	200(SI) 400(DF)	200 (SI) 400 (DF /CI)	1 5 %								

Air Pollutant Industry	Opacity	Particulate (Dust)	FIMIO (mg/nm²)	Sulphur oxide	(SO_X) (mg/Nm ³)	Nitrogen oxides	(NO _x) (mg/Nm ³)	O ₂ %	Carbon monoxide	Carbon dioxide (mg/Nm ³)	Hydrocarbons	Hydrogen Sulphide	Hydrogen Chloride	Hydrogen Fluoride	Lead (mg/Nm ³)	Dioxins/Furans
		NDA	DA	NDA	DA	NDA	DA									
Liquid fuels(>=50MWth to < 300MWth		50	30	1,1 70 or ≤2 %S F	0.5 % SF	1,46 0(CI ,Bor e ø<4 00m m) 1,85 0(CI , Bore ø≥4 00m m)		1 5 %								
Liquid fuels(plant≥ 300MWth		50	30	585 or ≤ 1% SF	0.2 % SF	740*	400	1 5 %								

Air Pollutant Industry	Opacity	Particulate (Dust)	(mig/min) 1 (mig/min)	Sulphur oxide	(SO _x) (mg/Nm ³)	Nitrogen oxides	(NO _x) (mg/Nm ³)	O ₂ %	Carbon monoxide	Carbon dioxide (mg/Nm ³)	Hydrocarbons	Hydrogen Sulphide	Hydrogen Chloride	Hydrogen Fluoride	Lead (mg/Nm ³)	Dioxins/Furans
		NDA	DA	NDA	DA	NDA	DA									
Biofuels/Gaseou s fuels other than Natural Gas		50	30	N/ A	N/ A	30% > Natu ral Gas & Liqu id Fuel s										
Combustion Turbine Natural Gas (All		N/A	N/	N/	N/	51	25									
turbine types of unit> 50MWth)			A	A	A	31	23									
Fuels other than Natural Gas(unit> 50MWth)		50	30	Use 1% SF	Us e 0.5 % SF											
Boiler Natural Gas		N/A	N/ A	N/ A	N/ A	240	240	3 %								
Other Gaseous fuels		50	30	400	40 0	240	240	3 %								

Air Pollutant Industry	Opacity	Particulate (Dust)	FM10 (mg/Nm ²)	Sulphur oxide	(SO _x) (mg/Nm ³)	Nitrogen oxides	(NO _x) (mg/Nm ³)	O ₂ %	Carbon monoxide	Carbon dioxide (mg/Nm³)	Hydrocarbons	Hydrogen Sulphide	Hydrogen Chloride	Hydrogen Fluoride	Lead (mg/Nm ³)	Dioxins/Furans
		NDA	DA	NDA	DA	NDA	DA									
Liquid Fuels(Plant >=50MWth to < 600MWth)		50	30	900 - 150 0	40 0	400	200	3 %								
Liquid Fuels(>=600MW th)		50	30	200 - 850	20 0	400	200	3 %								
Solid Fuels>=50MWth to < 600MWth)		50	30	900 - 150 0	40 0	510	200	6 %								
Solid Fuels(>=600MW th)		50	30	200 - 850	20 0	1,10 0 upto volat ile matt er of fuel < 10%	200	6 %								
Waste water treatment plants						NH ₃ (100 - 400)					40 0- 2,0 00	50 - 20 0				

And any other parameter as may be prescribed by the Authority from time to time

Legend SF: Suphur Fuel

- * 1.5-3.0% only justified by project specific considerations i.e. add secondary treatment to meet levels of 1.5% Sulphur
- ** dependent on water availability for injection

CI: Compression Ignition

SI: Spark Ignition DF: Dual Fuel

DA: Degraded Area

NDA: Non-degraded Area

The chimney or stack should have a minimum height of 10 metres above ground level and clear the highest of the building by not less than 3 metres for all roofs. The topography and height of adjacent buildings within 50 metres radius should be taken into account.

Toxic Equivalent (TEQ) is the sum of the toxic equivalent factors (TEF) of a mixture congeners contained in a compound. The compound 2,3,7,8-tetrachlorodibenzo-p-dioxin(TCDD) was assigned a TEF of 1 after being identified, by International Association of Radiology and Cancer (IARC) and World Health Organisation (WHO), as the most toxic of all compounds, and as carcinogenic to humans, based mainly on studies of cases involving accidental or occupational heavy exposure. Therefore the TEF is a weighting factor.

```
g - gram
µg- microgram
kg — kilogram (1,000g)
mg - milligram
µg- microgram
m³ — cubic metre
ppm — Parts per million
t — tonne
```

Annex 9: Acceptable Emission Control Technologies

SEVENTH SCHEDULE

Acceptable Emission Control Technologies

List of Acceptable Emission Control Technologies

	Air Pollutants	9 Emission Control technologies	Remarks
1.	Particulate Matter	Mechanical collectors (<u>dust cyclones</u> , multicyclones)	
		Electrostatic precipitators	
		Fabric filters (baghouses)	
		Particulate scrubbers	
2.	Nitrogen Oxides (Nox) *	Low NOx burners	
		Selective catalytic reduction (SCR)	
		Selective non-catalytic reduction (SNCR)	
		NOx scrubbers	
		Exhaust gas recirculation	
		Catalytic converter	
3.	Volatile Organic Compounds (VOC), hydrocarbons	Adsorption systems, such as activated carbon	
		<u>Flares</u>	
		Thermal oxidizers	
		Catalytic oxidizers	
		Biofilters	
		Absorption (scrubbing)	
		Cryogenic condensers	

	Air Pollutants	9 Emission Control technologies	Remarks
4.	Sulphur Oxides (SOx)	Wet scrubbers	
		<u>Dry scrubbers</u>	
		Flue gas desulphurization	
5.	Carbon Oxides	Thermal oxidizers	
6.	Hydrogen Sulphides	Absorption (scrubbing)	
7.	Hydrogen Chloride	Dry Scrubbers, Adsorption systems, such as activated carbon	
8.	Dioxins & Furans	Cyclone	
		Electrostatic precipitator	
		Bag filter	
		Wet scrubber	
		Quenching & subsequent wet scrubber	
		Catalytic oxidation (selective catalytic reaction)	
		Catalytic bag filter	
		Dry absorption in resins (carbon Particles dispersed in a polymer matrix)	
		Entrained flow reactor with added activated carbon or coke/lime or limestone solutions and subsequent fabric filter	

	Air Pollutants	9 Emission Control technologies	Remarks									
		Fixed bed or circulating fluidized bed reactor, adsorption with activated carbon or open hearth coke										
9.	Metals (Hg, Pb,)	Sorbent Injection Technology										
		Electro-Catalytic Oxidation (ECO)										
		K-Fuel										
10.	Any other technology approved by the Authority from time to time											

* Notes

Best Available Technology (BAT) for this category of equipment will consist of combustion modification technology including either:

- (a) low NO burner technology with low excess air
- (b) Air if technically feasible; or
- (c) flue gas re-circulation with low excess air.