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EXECUTIVE BODY FOR THE CONVENTION ON LONG-RANGE TRANSBOUNDARY AIR POLLUTION

Working Group on Abatement Techniques

CONTROL TECHNIQUES FOR EMISSIONS OF NQ FROM STATIONARY SOURCES

Prepared by the Task Force on the Assessment of Control Options/Techniques for NQ, led by Germany $^{\star\prime}$

<u>Introduction</u>

1. This document covers the stationary sources of NQ emissions listed in table 1.

2. The aim of this document is to provide Parties to the Convention with guidance on identifying best available techniques to enable them to meet the obligations of the Protocol. The presented options for best available techniques (BAT) can be applied in general.

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 \star / At the fifth meeting of the Task Force held in Karlsruhe (Germany) from 21-22 October 1998.

Table 1:	Considered	stationary	source	categories	for 1	QN	emissions
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	Source categories
1.	Combustion installations
	(a) Boilers
	(b) Gas turbines
	(c) Stationary engines
2.	Mineral oil refineries
3.	Coke oven furnaces
4.	Production and processing of metals:
	- Metal ore roasting or sintering installations
	- Installations for the production of pig iron or steel (primary $lpha$
	secondary fusion) including continuous casting
	- Installations for the processing of ferrous metals (hot rolling)
5.	Installations for the production of cement clinker in rotary kilns or
	in other furnaces
6.	Installations for the manufacture of glass including glass fibre
7.	Installations for the production of nitric acid
8.	Installations for the incineration of municipal waste, hazardous waste,
	medical waste and sludges from waste-water treatment

3. This document is based on options and techniques for NQ emission prevention and reduction and their performance and costs as documented in the technical background report prepared by the Task Force on the Assessment of Abatement Options/Techniques for NQ which includes information provided to the lead country by plant operators, producers of NQ control equipment and members of the Task Force. The technical background report is also based on information from official documentation of the Executive Body and its subsidiary bodies, e.g. the documentation of the Sixth Seminar on Control Technologies for Emissions from Stationary Sources held in 1996 in Budapest, and other published information. It gives the present status (April 1998) of development and application of the relevant options and techniques and related costs for the reduction of NQ emissions from stationary sources in the main emitting source categories.

4. 'Best available techniques' means the most effective and advanced stage in the development of activities and their methods of operation which indicate the practical suitability of particular techniques for providing in principle the basis for limit values designed to prevent and, where that is not practicable, generally to reduce emissions and the impact on the environment as a whole:

(i) 'Techniques' includes both the technology used and the way in which the installation is designed, built, maintained, operated and decommissioned;

(ii) 'Available' techniques means those techniques developed on a scale which allows implementation in the relevant industrial sector, under economically and technically viable conditions, taking into consideration the costs and advantages, whether or not the techniques are used or produced inside the Party in question, as long as they are reasonably accessible to the operator; (iii) 'Best' means most effective in achieving a high general level of protection of the environment as a whole.

In determining the best available techniques, special consideration should be given, generally or in specific cases, to the factors below bearing in mind the likely costs and benefits of a measure and the principles of precaution and prevention:

(a) The use of low-waste technology;

(b) The use of less hazardous substances;

(c) The furthering of recovery and recycling of substances generated and used in the process and of waste where appropriate;

(d) Comparable processes, facilities or methods which have been tried with success on an industrial scale;

(e) Technological advances and changes in scientific knowledge and understanding;

(f) The nature, effects and volume of the emissions concerned;

(g) The commissioning dates for new or existing installations;

(h) The length of time needed to introduce the best available technique;

(i) The consumption and nature of raw materials (including water) used in the process and their energy efficiency;

(j) The need to prevent or to reduce to a minimum the overall impact of the emissions on the environment and the risks to it;

(k) The need to prevent accidents and to minimize the consequences for the environment.

5. This document addresses the control of NQ emissions considered as the sum of nitrogen oxide (NO) and nitrogen dioxide (NQ) expressed as NO₂. However, when measures or techniques are planned for NQ sources emitting other components and, in particular, sulphur oxides (SQ), volatile organic compounds (VOCs), ammonia (NH₃), greenhouse gases (such as CQ and N₂O), particulates (including heavy metals), and persistent organic pollutants (POPs), it is worthwhile to consider them together with such other pollutant-specific control options in order to maximize the abatement effect and minimize the impact on the environment. Respective trade-offs between different pollutants have to be accounted for. This is particularly important for multi-pollutant/multi-effect approaches.

I. GENERAL ISSUES

6. Several possibilities exist to control or to prevent NQ emissions from stationary sources. Primary, secondary (add-on or end-of-pipe) and structural measures are generally distinguished. If not stated otherwise, measures are applicable to existing and new installations. The following list gives a general outline of available measures, which may also be combined:

- (a) Enhanced effectiveness of existing NQ control technologies;
- (b) Energy management (efficient and rational use of energy);
- (c) Appropriate boiler design;
- (d) Improved combustion technologies;
- (e) Combustion modification (primary measures);
- (f) New concepts for combustion technologies;
- (g) Flue gas cleaning (secondary measures);
- (h) Good housekeeping (e.g. good maintenance, good control).

A. Structural measures

7. The use of clean fuels and the rational use of energy result in a reduction in NO_x emissions. Usually, the use of certain fuels is governed by a country's energy supply structure. Thus, the use of low NQ producing fuels is usually limited. Although there is a substantial technical potential for reducing NO_x by fuel switching, achieving this potential will depend on country-specific conditions such as its infrastructure and policy. On the other hand, substantial reductions in energy consumption may be achieved in many production processes through energy management, e.g. energy conservation, technology switch, and demand-side management. The costs of energy management options may be lower than the costs of additional energy supply. Energy management can contribute substantially to the mitigation of air pollution. Fuel cleaning for fuel nitrogen removal is not a commercial option. Increasing the application of hydroprocessing in refineries, however, also brings about a reduction in the nitrogen content of the end product.

B. Primary and secondary measures

8. To achieve the most efficient NQ reduction, beyond energy management measures, a combination of technical options (fuel switching, other combustion technologies, process and combustion modifications, flue gas treatment) should be considered. Furthermore, to identify the best combinations of combustion modifications and flue gas treatment, site-specific evaluation is needed, in order to meet given emission targets.

9. Process and combustion modifications are applied to reduce the formation of NO_x during combustion. They include the control of combustion air ratio, flame temperature, fuel-to-air ratio, etc. Some of the options are typical for retrofit; others are typical for new installations, but may also be applied to retrofit. Efficiency and applicability could have some limitations. These measures are widely implemented, either singly or in combination:

- (a) Low excess air combustion (LEA);
- (b) Reduced air preheat (RAP); $\frac{2}{}$
- (c) Burner-out-of-service (BOOS) $i^{2/2}$
- (d) Biased-burner-firing (BBF); $i^{2/2}$
- (e) Low NO_x burners (LNB);
- (f) Flue gas recirculation (FGR);
- (g) Over fire air combustion (OFA);
- (h) In-furnace NO_x reduction or reburning (IFNR);
- (i) Water/steam injection and lean/premixed combustion. $\frac{2}{r}$

10. NO_x control technologies such as reburning are well developed and available for all combustible fuels. The achievable removal efficiency is reported to reach 70 - 80%, alone or in combination with other primary measures. Reburning offers some advantages such as compatibility with other primary NO_x emission reduction measures, simple installation, use of a standard fuel (oil, gas) as reducing agent, no additives and little additional energy required. Especially the successful implementation of reburning in large oil-fired units has to be mentioned, as well as the experience with coal as a reduction agent. The trend nowadays is to use the same fuel as fuel and reducing agent.

11. NO_x emissions can be also reduced by using inherent low NQ control technologies such as fluidized bed combustion. This technology is applicable to a large range of fuels (coal, biomass, residues, etc.). Due to the rather low combustion temperature (about $85 \times 10^{\circ}$ and an inherent air staging, this technology achieves low NQ emissions and can generally be used without secondary measures. Oxycombustion is another way of abating NQ emissions; until now, industrial applications of this technique have been limited to glass manufacturing.

12. Basic combustion modifications are incorporated mainly into boiler and burner design. For example, modern furnace designs incorporate OFA devices. The latest generation of LNB combines both air-staging and fuel-staging (reburning at burner level).

13. Unlike most combustion processes, the application of combustion and/or process modifications in industrial processes with combustion has many process-specific limitations. In cement kilns or glass melting furnaces, for example, certain high temperatures and homogeneous temperature distributions are necessary to ensure product quality. Typical combustion modifications being used are staged combustion/low NQ burners, flue gas recirculation and process optimization (e.g. precalcination in cement kilns).

The selective catalytic reduction process (SCR) is the most mature and 14. widely implemented flue gas treatment process with a high removal efficiency (in some sectors, this efficiency can, depending on the case, reach up to 95% and high availability. The SCR process usually uses ammonia or urea as a reducing agent, but the use of high-pressure stored anhydrous ammonia is the most common. Many SCR installations using these additives are being operated successfully in Europe, mostly for boilers. Application of SCR at gas-fired plants can reduce NQ emissions to very low levels. It has been implemented at gas, oil, and coal-fired installations. Catalyst lifetimes are much higher than initially projected and have reached as much as 6 - 10 years for coalfired units, and 8 - 12 years for gas- and oil-fired units, the lower values being achieved for high-dust configuration and the higher for tail-gas configuration. SCR is applicable also to smaller combustion installations and it is a well established de-NQ technology for combustion in boilers and for certain industrial processes, such as:

- (a) Nitric acid plants;
- (b) Glass smelters;

(c) Cement production (the applicability of SCR is currently being tested on a pilot scale);

- (d) Refinery furnaces;
- (e) Combustion of hazardous wastes (usually in rotary kilns);

(f) Combustion of municipal wastes (usually in grate furnaces);

(g) Combustion of hospital and other special wastes in public or industrial units (rotary kilns, pyrolysis plants, fluidized bed incineration of industrial wastes like sludges, rejects, production residues) also in rather small units (< 10 MW_h).

15. The selective non-catalytic reduction process (SNCR) is applicable to small and medium-sized installations with medium raw gas NQ contents. The SNCR process is implemented in a variety of combustion installations and process furnaces (glass and cement production) as well as in waste incinerators. The SNCR process is in general capable of 30 - 70% reductions. Combined with flue gas recirculation, it is an attractive and reliable technology for moderate NQ removal (50 - 80%), especially for smaller combustion plants and industrial processes.

16. Other flue gas treatment technologies are the combined NQ/SO_2 removal processes. The activated carbon process (AC) is an option used in few cases only, since it is expensive and leads to limited NQ reduction (around 60%). The SNOX process, which removes both sulphur oxides and nitrogen oxides, offers some advantages in the case of high-sulphur feedstocks.

C. <u>Costs</u>

17. The estimation of investments and operating costs for NQ emission reduction options/techniques is important when choosing from the wide range of measures and, on a more macroeconomic level, when developing an emission control strategy on national or regional level. A detailed assessment of cost data in terms of consistent investment and operating costs is presented in the background document on best available techniques of the Task Force on the

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Assessment of Abatement Options/Techniques for NQ. The respective data are based on practical experience. Data have been collected for existing and new plants; for the latter, the investment can usually be considerably lower.

18. Examples of investments and operating costs are given in table 2 for selected relevant combinations of primary and secondary measures, which are applicable in the considered sectors. The examples are specific for detailed relevant parameters, such as yearly operating hours, waste gas volume flow and pollutant concentration therein, capacity of the base installation, fuel type, etc. Thus, the given data serve an illustrative purpose and cannot be taken as generally applicable values. In table 3, the most important parameters are given for the determination of operating costs for the example of SCR. These examples illustrate the methodology followed for the cost assessment in the technical background report of the Task Force on the Assessment of Abatement Options/Techniques for NQ.

Table 2: Investments and operating costs of control options for the abatement of NO_x emissions (retrofit of existing installations)

Characteristics of reference installation	Control options	Invest- ments ^{a)} [ECU]	Operating Costs ^{b)} [ECU/a]	Abated mass flow [Mg
				$NO_x/a]$
Boilers		13,300,000	1,400,000	
Public power and district heating;	Low NO _x burner	13,300,000	1,400,000	3,560
Boiler;	Low NO_x burner & over-fire	16,100,000	1,500,000	4,100
Fuel: hard coal;	air			
Capacity:	Low NO_x burner & SCR	61,300,000	6,400,000	8,110
1,500 MW _{th} ;	Low NO_x burner& over-fire	17,500,000	2,200,000	6,770
Operating time:	air & SNCR	64 100 000	6 500 000	0.000
5,500 h/a	Low NO _x burner & over-fire air & SCR	64,100,000	6,500,000	8,200
Public power and	Low NO _x burner	7,000,000	430,000	390
district heating;				
Boiler;	Low NO_x burner & over-fire air	8,500,000	505,000	450
Capacity: 600 MW _{th} ; Fuel: hard coal;	Low NO_x burner & SCR	32,000,000	1,930,000	885
Operating time:	Low NO _x burner& over-fire	9,200,000	610,000	740
1,500 h/a	air & SNCR Low NO _x burner & over-fire	33,500,000	2,000,000	895
	air & SCR	33,500,000	2,000,000	695
Industry; Boiler; Fuel:	Low NO_x burner	2,800,000	170,000	140
natural gas;				
Capacity: 160 MW _{th} ;				
Operating time:				
6,000 h/a				

Characteristics of	Control options	Invest-	Operating	Abated
reference	Concret of crows	$ments^{\underline{a}}$	$Costs^{\underline{b}}$	mass
installation		[ECU]	[ECU/a]	flow
				[Mq
				NO _x /a]
Industry;	Low NO _x burner & SNCR	1,500,000	240,000	830
Boiler;	Low NO _x burner & SCR	3,400,000	560,000	1,040
Fuel: plant gas;	Flue gas recirculation &	1,300,000	210,000	780
Capacity: 160 MW _{th} ;	SNCR			
Operating time:	Flue gas recirculation &	3,200,000	530,000	1,030
6,000 h/a	SCR			
	Low NO _x burner & flue gas recirculation & SNCR	2,000,000	310,000	920
	Low NO_x burner & flue gas	3,850,000	630,000	1,070
	recirculation & SCR			
Public power,	No measure	10,700,000	830,000 <u>≏</u>	760 <u>°</u> ∕
district heating,		<u>c</u> /		
industry;				
Circulating	SNCR	290,000	74,000	152
fluidized bed combustion:				
Fuel: coal;				
Capacity: 160 MW _{th} ;				
Operating time:				
6,000 h/a				
Commercial and	Low NO _x burner	245,000	13,000	3.1
institutional				
heating;	Over-fire air	51,000	2,500	0.75
Boiler;				
Fuel: heavy fuel	Low NO_x burner & overfire	295,000	15,700	3.6
oil; Capacity: 5	air		,	
MW _{th} ;	Low NO _x burner & flue gas	295,000	16,500	4.5
Operating time:2,000 h/a	recirculation	,	.,	
Gas turbines				
Combined cycle;	PM	2,050,000	370,000	670
Output capacity: 150		_,,	2.0,000	5,5
MW _{el} ;				
Fuel: natural gas;	SCR	27,000,000	3,110,000	1,145
Working time: 6,000				
h/a				
Simple cycle;	PM	360,000	36,000	70
Output capacity: 30	SCR	5,400,000	380,000	115
MW _{el} ; Fuel: natural		5,100,000	500,000	110
gas; Working time:	PM & SCR	4,000,000	300,000	125
2,000 h/a	24		F	
Cogeneration plant	PM	550,000	50,000	685
in industry; Output capacity of	SCR	4,500,000	600,000	1,165
gas turbine:		Ŧ,500,000	000,000	т,тор
25 MW _{el} ; Fuel: diesel	PM & SCR	5,050,000	610,000	1,265
oil; Working time:		2,200,000		_,200
8,000 h/a				
,				-

Characteristics of	Control options	Invest-	Operating	Abated
	Concror operons		Costs ^{\underline{b}}	
reference		ments ^{a)}		mass
installation		[ECU]	[ECU/a]	flow
				[Mg
- a a a d/		5 400 000	E 00.000	NO _x /a]
IGCC ^d /;	PM	5,400,000	700,000	4,000
Output capacity: 450	SCR	81,000,000	10,000,000	6,900
MW _{el} ; Fuel: heavy fuel oil;				
Working time: 8,000				
h/a				
Stationary engines	Terr burn	10 000	2 000	4.0
Old rich burn gas	Lean burn	10,000	- 3,000	40
engine;				
Fuel: natural gas;		[4 0 0 0 0]		[40]
Output capacity: 600	[NSCR]	[40,000]	[2,500]	[40]
kW _{el} ; Working time:				
5,000 h/a				
Improved lean burn	SCR	50,000 ≌⁄	15,000	5
gas engine; Fuel:				
natural gas; Output				
capacity: 600 kW _{el} ;				
Working time: 5,000				
h/a				
Diesel engine; Fuel:	SCR	230,000	40,000	650
heavy fuel oil;				
Output capacity: 20				
MW _{el} ; Working time:				
5,000 h/a				
Diesel engine;	Exhaust gas recirculation	15,000	5,000	26
Fuel: diesel oil;	SCR	230,000	40,000	78
Output capacity:	Exhaust gas recirculation &	245,000	40,000	80
3 MW _{el} ; Working time:	SCR			
5,000 h/a				
Diesel engine; Fuel:	SCR	60,000	9,000	100
heavy fuel oil;				
Output capacity: 20				
MW _{el} ; Working time:				
5,000 h/a				
Iron and steel produc	tion: sinter plants			1
Travelling grate	Flue gas recirculation	5,000,000	- 200,000 <u>f</u> /	2,000
sinter machine;	-			
Fuel: coke breeze;	SCR	50,000,000	10,000,000	6,600
Production output:	Flue gas recirculation &	48,000,000	5,000,000	6,700
12,000 Mg sinter/d;	SCR			
Working time: 8,400				
h/a				
,				

Characteristics of	Control ontions	Transat	Onomating	Nhatad
reference	Control options	Invest- ments ^{a)}	Operating Costs ^{b)}	Abated
installation		[ECU]	[ECU/a]	mass flow
Installation		[EC0]	[ECO/a]	[Mq
				NO _x /a]
Cement production				no _x , aj
Dry process with	Low NO_x burner and staged	3,600,000	240,000	340
preheater/precalcine	combustion for precalciner			
r; Fuel: coal;	kiln			
Production capacity:	Low NO_x burner and SNCR	2,175,000	340,000	460
2,000 Mg/D; Working				
time: 5,000 h/a				
Dry process with	Low NO_x burner and SCR	6,400,000	820,000	960
preheater/precalcine				
r; Fuel: coal;				
Production capacity:				
2,000 Mg/d; Working				
time: 8,000 h/a				
Glass production: con				
Production of flint	CM g/	600,000	30,000	300
and coloured glass;	Reburning	250,000	200,000	470
Regeneratively	$LoNO_x$ melter	2,000,000	100,000	550
heated furnace; Fuel: natural gas;	Oxycombustion	- 1,200,000	1,200,000	630
Production capacity:	Oxycombustion	- 1,200,000 <u>h</u> /	1,200,000	630
400 Mg/d;	SNCR	1,000,000	130,000	390
Working time:	SCR	4,000,000	500,000	550
8,760 h/a	CM & SNCR	1,580,000	161,000	550
	CM & SCR	4,390,000	536,000	650
Production of flint	СМ д/	360,000	18,000	76
and coloured glass;				
Recuperatively				
heated furnace;				
<pre>Fuel: natural gas;</pre>				
Production capacity:				
400 Mg/d; Working				
time:				
8,760 h/a				
Nitric acid productio				
Production of medium	SCR	2,000,000	300,000	950
concen-tration				
nitric acid				
High dual pressure				
plant; Production				
capacity: 1,000				
Mg/d; Working time:				
8,400 h/a				
Production of high	SCR	1,200,000	200,000	800
concentration nitric				
acid; Production				
capacity: 500 Mg/d;				
Working time: 8,400				
h/a				

Characteristics of reference installation	Control options	Invest- ments ^{a)} [ECU]	Operating Costs ^{b)} [ECU/a]	Abated mass flow
				[Mq
				NO _x /a]
Waste incineration: i	ncineration of domestic or mu	micipal waste	S	
Moving grate	PM ⁱ /	300,000	30,000	100
incinerator;				
rotary kiln	SNCR	1,000,000		350
Capacity: 30 Mg/d;	SCR	26,000,000		430
Operating time:	PM & SNCR	1,300,000		380
8,400 h/a	PM & SCR	26,300,000		440
<u>a</u> / Depending on e.g. waste gas flow rate, production capacity, peripheral conditions <u>b</u> / Depending on e.g. waste gas flow rate, NO _x inlet concentration in the waste gas, annual operating time, NQ reduction rate, etc. <u>c</u> / Compared with conventional pulverized coal combustion. <u>d</u> / Limited operating experience. <u>e</u> / Including a small oxidation catalyser <u>f</u> / Due to reduced coke breeze consumption. <u>g</u> / E.g. low excess air, reduced air preheating, staged combustion. <u>h</u> / Compared to conventional air combustion.				
PM = Primary Measure				
CM = Combustion Modification				
SNCR = Selective Non Catalytic Reduction SCR = Selective Catalytic Reduction				
	Catalytic Reduction (three-wa	ay catalyser)		

Table 3: Major cost components and related parameters considered for the assessment of operating costs (ECU/a) for the SCR technology (for aqueous ammonia)

Parameters 6 Main cost components 9	Waste gas flow rate	C (NO _x , in)	Annual working time	NO _x reduction	Share of NO in total NO _x	Rela- tive impor-
			(h/a)			tance
Direct consumption rela	ted operat:	ing costs				
Ammonia consumption ^{a/}	i	ļ	ļ	ļ	!	++
Electricity	!		!			++
consumption ^{a/}						
Fuel consumption ^{b/}	ļ		ļ			+
Catalyst replacement ^{a/}	!		!			+++
Personnel			!			+
Investment-related oper-	ating cost:	5				
Maintenance and repair	!	ļ		!		++
Taxes	ļ	i		ļ		+
Insurance	_!	ļ		ļ		+
C (NO _x , in) = inlet NO _x -	concentrat	lon in flue	gas	-	-	
<u>a</u> / Dominating items to	gether wit	h capital c	osts (not t	aken into a	ccount in	

 $\underline{\mbox{a}'}$ Dominating items together with capital costs (not taken into account in table 3).

 $\frac{b}{2}$ To be considered in the case of tail gas or low dust configuration

Concerning the investment-related costs, it should be taken into account that depreciation and interest (capital costs) represent the most important cost items (not taken into account in table 3) for the SCR technology.

19. To derive the abatement cost per sector for a given country, it is necessary to consider parameters such as:

- Capacity distribution of production processes;
- Age distribution and final lifetime of production processes;
- Transition periods/implementation schedules with regard to technology changes;
- Production and abatement technologies already in place according to current legislation;
- Future activity rates of sectors;
- Nature of raw materials, process design and operation.

To derive this information, substantial further analysis than table 3 provides is needed. The background document provides the reference installation methodology and the necessary data (investments, annual operating costs) taking into account the set of relevant parameters mentioned above to assess these costs.

20. To improve national cost functions, they should adequately reflect the individual country's situation. Therefore, in a bottom-up approach, the country-specific ranking of options for the various sectors should be combined with country-specific activity levels. Finally, the national cost functions should be the result of an optimization consideration how to adapt the production system over time to different NQ reduction requirements. This includes not only end-of-pipe measures, but also structural changes within the sectors.

D. <u>Side effects</u>

21. Side effects of emission abatement options/techniques should be accounted for. Particular attention should be paid to the influence of NQ control measures on related air pollution issues such as ammonia emissions released from certain emission reduction facilities. Moreover, cross-media aspects should be considered in terms of water pollution and waste generation which may be induced or increased by certain NQ emission reduction options/techniques. However, these side effects can generally be limited by proper design and operation of the facilities. More specifically, the side effects to be considered with different emission reduction techniques are:

(a) Combustion modifications: possible side effects are decrease in overall energy efficiency, increased CO formation and hydrocarbon emissions, corrosion due to reducing atmosphere, increase of unburnt carbon in fly ash;

(b) FBC: this technique also brings about a considerable reduction in SQ emissions. A possible drawback in FBC systems may be the increased formation of N_2O under certain process conditions. The handling of the ashes needs consideration in relation to their possible use and/or disposal;

(c) SCR: some possible side effects are ammonia slip in the exhaust gas, ammonia content in the fly ash, formation of ammonium salts on downstream facilities, deactivation of the catalyst and increased conversion of SQ to

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 SO_3 (corrosion). By the controlled operation of the plant, the fly ash quality can, however, be guaranteed and the formation of ammonia salt reduced. In terms of by-products, deactivated catalysts from the SCR process may be the only relevant products, although this has become a minor problem since the catalyst lifetime has been improved and reprocessing options exist;

(d) SNCR: side effects to be considered are ammonia in the exhaust gas, formation of ammonium salts on downstream facilities, the formation of NO, when urea, for instance, is used as a component of the reducing mixture, and CO releases;

(e) The reagent production of ammonia and urea for flue gas treatment processes involves a number of separate steps which require energy and reactants. The storage systems for ammonia are subject to the relevant safety legislation and such systems are designed to operate as totally closed systems, with a resultant minimum of ammonia emissions. The use of NH is still considered appropriate, even when taking into account the indirect emissions related to the production and transport of NH.

III. <u>SECTOR-SPECIFIC ISSUES</u>

A. Combustion installations

22. This section covers boilers, (small: 1 - 10 MW_h, medium: 10 - 50 MW_{th}, and large: > 50 MW_{th}), gas turbines (> 1 MW_{el}), and stationary engines (> 100 kW_{el}). Given capacity classes in terms of rated thermal input refer to the lower heating value (LHV) of the respective fuel.

23. New concepts for combustion technologies with improved thermal efficiency and reduced NQ emissions include combustion turbines, atmospheric and pressurized fluidized bed combustion, integrated gasification combined cycle, combined cycle gas turbines, cogeneration and supercritical boilers:

(a) Stationary combustion turbines can also be integrated in existing conventional power plants (known as topping). The overall efficiency can be increased by 5 - 6%, but the achievable NQ reduction will depend on sitespecific conditions. Major alterations to the existing boiler system may become necessary;

(b) Fluidized bed combustion (FBC) is a combustion technology for burning hard coal and lignite, but also low-grade fuels such as waste, peat and wood. Emissions can be further reduced through integrated combustion control in the system. Within the sector of energy conversion, atmospheric fluidized bed combustion is a well established commercial technology. There are more circulating fluidized bed combustors than bubbling fluidized bed combustors. Currently, more than 200 circulating fluidized bed systems with a total capacity of about 26,000 MW_h and 15 pressurized fluidized bed systems (representing about 5,000 MW_h) are in operation worldwide;

(c) The integrated gasification combined cycle (IGCC) process incorporates coal gasification and combined cycle power generation in a gas and a steam turbine. The gasified coal is burned in the combustion chamber of the gas turbine. The technology also exists for heavy oil residue. The installed capacity at present is about 1,000 MW_h (5 plants). However, this process is not yet fully commercialized;

(d) Combined cycle gas power stations using advanced gas turbines with an overall energy efficiency significantly higher than 55 % and with reduced NQ emissions are being built;

(e) The combined generation of electricity and heat in so-called cogeneration plants represents a possibility to save fuel (up to 50% reduction in energy consumption compared to separate generation of electricity and heat). Although the electrical efficiency is decreased by the extraction of steam, the overall efficiency of the cogeneration system ranges between 70 and 90%;

(f) A further measure to increase conventional power plant efficiency is the use of supercritical steam cycle conditions. Total system net energy efficiencies can achieve 43% for hard coal fired plants, 41% for lignitefired plants, and 56% for gas-fired combined cycle plants.

24. Table 4 gives a selection of applicable abatement options and the respective clean gas concentrations achievable by typical emission sources.

Emission source	Combination of control	Clean gas
	measures	concentration
		(mg/Nm^3)
Small boilers 1 - 10 M	$M_{ m h}$, medium boilers 10 - 50) MW _{th}
Dry bottom boiler; Fuel: hard	Primary measures (PM)	400 - 600
coal (> 10 MW_{th})		
Boiler; Fuel: light fuel oil	PM	150 - 300
Boiler; Fuel: heavy fuel oil	PM	300 - 600
Boiler; Fuel: natural gas	PM	50 - 150
Circulating FBC; Fuel: coal,	no further measure	150 - 300
peat, biomass, etc.		
Bubbling FBC; Fuel: coal, bark,	no further measure	200 - 400
oil, sediment, etc.		
Industrial boiler; Fuel: process	PM	100 - 300
gas		
Large bo	$pilers > 50 MW_{th}$	
Dry bottom boiler	РМ	300 - 600
Fuel: hard coal	PM and SCR (many	80 - 150
	applications)	
	PM and SCR (many	
	applications	
Boiler	PM (without reburning)	250 -500
Fuel: heavy fuel oil	PM (incl. reburning)	# 200

Table 4:Emission sources and selected NQ control measures with theirrespective clean gas concentrations for combustion installations

Emission source	Combination of control	Clean gas
	measures	concentration
	meabureb	(mg/Nm ³)
	PM and SCR	60 - 150
		00 100
Boiler; Fuel: natural gas	PM	50 - 200
Wet bottom boiler	PM and SCR (tail gas)	# 150
		"
Fuel: hard coal	PM and SNCR	# 200
Pressurized FBC	no further measure	150 - 200
Fuel: hard coal	SCR and/or SNCR	# 100
	ben and, or bron	<i>"</i> 100
Circulating FBC	no further measure	150 - 300
Fuel: coal, peat, biomass, etc.	SNCR	100 - 200
Bubbling FBC; Fuel:	no further measure	200 - 400
and have all addiment at a	andb	130 - 200
coal, bark, oil, sediment, etc.	SNCR	130 - 200
Industrial boiler	PM	100 - 300
		100 300
Fuel: process gas	PM and SCR	100 - 200
Ga	s turbines	
Simple cycle, combined cycle, cog	generation (prior to supple	ementary
firing), mechanical drive		
Fuel: natural gas	PM	50 - 150
		[400]*
Twoli diaral ail an marana war	SCR	10 - 50 100 - 200
	Wet controls	100 - 200
Fuel: diesel oil or process gas	CCD	
	SCR	20 - 100
IGCC		20 - 100
	Nitrogen and steam	
IGCC Fuel: coal or heavy fuel oil		20 - 100
IGCC Fuel: coal or heavy fuel oil Statio	Nitrogen and steam injection	20 - 100
IGCC Fuel: coal or heavy fuel oil	Nitrogen and steam injection onary engines	20 - 100
IGCC Fuel: coal or heavy fuel oil Static Spark ignition (= Otto) engines,	Nitrogen and steam injection onary engines 4-stroke	20 - 100
IGCC Fuel: coal or heavy fuel oil Station Spark ignition (= Otto) engines,	Nitrogen and steam injection onary engines 4-stroke NSCR (three-way	20 - 100
IGCC Fuel: coal or heavy fuel oil Static Spark ignition (= Otto) engines, Old rich burn	Nitrogen and steam injection onary engines 4-stroke NSCR (three-way catalyser)	20 - 100 50 - 100 350
IGCC Fuel: coal or heavy fuel oil Static Spark ignition (= Otto) engines, Old rich burn Improved lean burn Compression ignition (= Diesel)	Nitrogen and steam injection onary engines 4-stroke NSCR (three-way catalyser) No measure	20 - 100 50 - 100 350 300 - 550
IGCC Fuel: coal or heavy fuel oil Static Spark ignition (= Otto) engines, Old rich burn Improved lean burn	Nitrogen and steam injection onary engines 4-stroke NSCR (three-way catalyser) No measure SCR	20 - 100 50 - 100 350 300 - 550 100 [200; 400 -
IGCC Fuel: coal or heavy fuel oil Static Spark ignition (= Otto) engines, Old rich burn Improved lean burn Compression ignition (= Diesel) Fuel: heavy fuel oil	Nitrogen and steam injection onary engines 4-stroke NSCR (three-way catalyser) No measure SCR engines SCR	20 - 100 50 - 100 350 300 - 550 100 [200; 400 - 1,000]
IGCC Fuel: coal or heavy fuel oil Static Spark ignition (= Otto) engines, Old rich burn Improved lean burn Compression ignition (= Diesel)	Nitrogen and steam injection onary engines 4-stroke NSCR (three-way catalyser) No measure SCR engines SCR SCR	20 - 100 50 - 100 350 300 - 550 100 [200; 400 - 1,000] 360 - 500
IGCC Fuel: coal or heavy fuel oil Static Spark ignition (= Otto) engines, Old rich burn Improved lean burn Compression ignition (= Diesel) Fuel: heavy fuel oil	Nitrogen and steam injection onary engines 4-stroke NSCR (three-way catalyser) No measure SCR engines SCR	20 - 100 50 - 100 350 300 - 550 100 [200; 400 - 1,000] 360 - 500
IGCC Fuel: coal or heavy fuel oil Station Spark ignition (= Otto) engines, Old rich burn Improved lean burn Compression ignition (= Diesel) Fuel: heavy fuel oil Fuel: diesel oil Boilers: Solid fuels: 6% excess Q; Liquid	Nitrogen and steam injection onary engines 4-stroke NSCR (three-way catalyser) No measure SCR engines SCR SCR SCR Exhaust gas recirculation and SCR I fuels: 3% excess Q; Gaseous fue	20 - 100 50 - 100 350 300 - 550 100 [200; 400 - 1,000] 360 - 500 180 - 240
IGCC Fuel: coal or heavy fuel oil Static Spark ignition (= Otto) engines, Old rich burn Improved lean burn Compression ignition (= Diesel) Fuel: heavy fuel oil Fuel: diesel oil Boilers: Solid fuels: 6% excess Q; Liquid Gas turbines: 15% excess Q; Stationary en	Nitrogen and steam injection onary engines 4-stroke NSCR (three-way catalyser) No measure SCR engines SCR SCR Exhaust gas recirculation and SCR 4 fuels: 3% excess Q; Gaseous fue ogines: 5% excess Q	20 - 100 50 - 100 350 300 - 550 100 [200; 400 - 1,000] 360 - 500 180 - 240 ls: 3% excess Q
IGCC Fuel: coal or heavy fuel oil Static Spark ignition (= Otto) engines, Old rich burn Improved lean burn Compression ignition (= Diesel) Fuel: heavy fuel oil Fuel: diesel oil Boilers: Solid fuels: 6% excess Q; Liquid Gas turbines: 15% excess Q; Stationary en Values apply to different raw gas concent considered utilities.	Nitrogen and steam injection onary engines 4-stroke NSCR (three-way catalyser) No measure SCR engines SCR SCR Exhaust gas recirculation and SCR 4 fuels: 3% excess Q; Gaseous fue ogines: 5% excess Q	20 - 100 50 - 100 350 300 - 550 100 [200; 400 - 1,000] 360 - 500 180 - 240 ls: 3% excess Q ad operation of the

* For simple cycle

B. Mineral oil refineries

25. This section addresses combustion processes in mineral oil refineries without contact between flame or flue gas and products. The most relevant sources of NO_x emissions in mineral oil refineries are process heaters for the heating of crude oil and petroleum products.

26. A significant part of the fuel used for process heaters is provided by refinery gas. Various processes contribute a large variety of compounds to the refinery gas, resulting in varying NQ emissions. Other fuels in use in mineral oil refineries are natural gas, petroleum coke, heavy fuel oil, or other residues originating from atmospheric and vacuum distillation, fluid catalytic cracking (FCC) and thermofor catalytic cracking (TCC). Table 5 gives a selection of applicable abatement options and the respective clean gas concentrations achievable by typical emission sources.

Table 5:Emission sources and selected NQ control measures withrespective clean gas concentrations in mineral oil refineries

Emission source	Combination of control	Clean gas
	measures	concentration
		(mg/Nm^3)
Process heater; Fuel:	Primary measures (PM)	# 200
petroleum coke		
Process heater; Fuel:	PM	250 - 600
heavy fuel oil		
Process heater; Fuel:	PM	50 - 200
natural gas		
Process heater; Fuel:	РМ	100 - 300
process gas		
FCC	SCR	100 - 200
Solid fuels: 6% excess Q;	liquid fuels: 3% excess Q; ga	seous fuels:
3% excess O ₂		
Values apply to different	raw gas concentrations.	

C. <u>Coke oven furnaces</u>

27. This section deals with emissions originating from coke oven furnaces in iron and steel production. Table 6 gives a selection of applicable abatement options and the respective clean gas concentrations achievable by typical emission sources. Table 6:Emission sources and selected control measures with respectiveclean gas concentrations for coke oven furnaces in iron and steel production

Emission source	Combination of control	Clean gas		
	measures	concentration		
		(mg/Nm^3)		
Fuel: coke oven gas	Combustion modification* and	# 60		
	SCR			
Fuel: blast furnace gas	SCR	# 10		
Gaseous fuels: 3% excess Q. Values apply to different raw gas concentrations. * Waste gas recirculation, air-staged combustion, lowering the coking				
temperature.	,	5 5		

D. <u>Production and processing of metals</u>

28. This section deals with emissions originating from sinter plants, blast furnace cowpers, and reheating furnaces in iron and steel production. Until now, no NO_x -removal option has been applied to blast furnace cowpers. Direct reduction and direct smelting are currently under development and may reduce the need for sinter plants and blast furnaces in the future. The application of these technologies depends on the ore characteristics and requires the resulting product to be processed in an electric arc furnace. Table 7 gives a selection of applicable abatement options and the respective clean gas concentrations achievable by typical emission sources.

Table 7:Emission sources and selected control measures with respectiveclean gas concentrations for sinter plants and reheating furnaces in iron andsteel production

Emission source	Combination of control	Clean gas		
	measures	concentration		
		(mg/Nm^3)		
	Sinter plants			
Travelling grate; Fuel:	Flue gas recirculation and	# 230		
coke breeze	SCR			
	Reheating furnaces			
Fuel: blast furnace gas	Low NO _x burner	# 390		
Fuel: coke oven gas,	Low NO _x burner	# 1,100		
heavy fuel oil				
Fuel: natural gas, gas	Low NO_x burner	# 250		
oil				
Solid fuels: 6% excess Q; liquid fuels: 3% excess Q; gaseous fuels: 3%				
excess O _{2.}				
Values apply to different raw gas concentrations.				

E. <u>Cement production</u>

29. Cement kilns make use of fossil as well as secondary fuels such a waste oil or waste tyres. For the production of clinker, several kiln types are available, showing different NQ emission levels: long wet rotary kiln, long dry rotary kiln, dry rotary kiln with cyclone/grate preheater, dry rotary kiln with cyclone/grate preheater and precalciner, shaft furnace. In terms of energy demand and emission control opportunities, dry rotary kilns with cyclone/grate preheater and precalciner are preferable. In Europe, the dry process is mostly used, especially in new plants. In modern dry process kiln systems, the precalciner technology is used to further improve, among other things, thermal efficiency and production capacity.

30. Table 8 gives a selection of applicable abatement options and the respective clean gas concentrations achievable by typical emission sources. In the last years, the selective catalytic reduction process has been implemented at several demonstration plants on a pilot scale. SCR is not yet current practice in the cement manufacturing sector. However, given the recent positive experience, it may become an applicable control option in the future.

Table 8:Emission sources and selected NQ control measures with theirrespective reduction efficiencies and clean gas concentrations in cementproduction

Emission source	Combination of control	Clean gas		
	measures	concentration		
		(mg/Nm^3)		
Dry process with preheater/precalciner				
Fuel: coal	Low NO_x burner and staged	# 1,000		
	combustion for precalciner			
	kiln			
	Low NO_x burner and SNCR	200 - 800		
	Low NO_x burner and SCR	100 - 200		
Solid fuels: 10% excess Q ₂ .				
Values apply to different	raw gas concentrations.			

F. <u>Glass production</u>

31. This section deals with the production of flat and container glass, glasswool, commodity glass (TV screen, lighting) and domestic glassware. In glass production, several parameters have a significant influence on the NQ emission levels: the type of fuel used (natural gas, heavy fuel oil), the furnace type (cross-fired, end-fired furnaces; regenerative, recuperative air preheating) and the type of glass produced (flint glass, clear glass, tint glass).

32. Table 9 gives a selection of applicable abatement options and the respective clean gas concentrations achievable by typical emission sources. In the last years, some promising abatement technologies have emerged: reburning and oxycombustion. When using oxycombustion, special care has to be

taken with regard to energy efficiency so as not to reduce the NQ emission abatement potential.

Table 9:Emission sources and selected NQ control measures with theirrespective reduction efficiencies and clean gas concentrations in glassproduction

Emission source	Combination of control measures	Clean gas concentration (mg/Nm ³)	
	Flat glass		
Clear glass			
Cross-fired furnace with	Combustion modification ^{a/} and	# 500	
regenerative preheating;	low NO_x burner and SCR		
Fuel: natural gas			
Cross-fired furnace with	Reburning	# 500	
regenerative preheating;			
Fuel: natural gas or			
heavy fuel oil			
Cross-fired furnace with	Combustion modification	# 600	
regenerative preheating;	(primary measures)		
Fuel: heavy fuel oil			
Tinted glass	·		
Cross-fired furnace with	Combustion modification $\mathbb{A}^{1/2}$ and	# 700	
regenerative preheating;	low NO_x burner and SCR		
Fuel: natural gas	X		
Cross-fired furnace with	Reburning	# 500	
regenerative preheating;	5		
Fuel: natural gas or			
heavy fuel oil			
	Container glass		
Regeneratively heated	Low NO_x burner and SCR	# 350	
furnace; Fuel: natural	Combustion modification \underline{P} and	# 600	
gas	low NO_x burner and SNCR		
	Oxycombustion ^{b/}	# 400	
	Reburning	# 500	
Regeneratively heated	Combustion modification $\underline{\mathbb{P}}^{\underline{A}}$ and	# 450	
furnace;	low NO_x burner and SNCR		
Fuel: heavy fuel oil	Oxycombustion ^{b/}	# 300	
	Reburning	# 500	
Recuperatively heated	Combustion modification	# 350	
furnace; Fuel: natural gas	(primary measures)		
	Glass wool		
Recuperatively heated	Oxycombustion ^{b/}	# 300	
furnace; Fuel: natural	-		
gas			
Regeneratively heated	Oxycombustion ^{b/}	# 250	
furnace; Fuel: heavy fuel			
oil			

Emission source	Combination of control measures	Clean gas concentration (mg/Nm ³)
	Special glass	
Commodity glass		
Recuperatively heated furnace;	Combustion modification ${\tt M}$ and low ${\tt NO}_{x}$ burner and SNCR	# 600
Fuel: natural gas	Oxycombustion ^{b/}	# 400
Recuperatively heated furnace;	Combustion modification ^{μ} and low NO _x burner and SNCR	# 450
Fuel: heavy fuel oil	Oxycombustion ^{b/}	# 300
Solid, liquid and gaseous fuels: Values apply to different raw gas ^a ∠ E.g. low excess air, reduced ^b ∠ Related to total NO ₂ emissio	s concentrations. 1 air preheating, staged combustion.	

G. <u>Nitric acid production</u>

33. For the production of nitric acid, two production methods exist: the first produces medium-concentration nitric acid (50 - 75 wt.-%), the second leads to high-strength nitric acid (98 wt.-%), which can be achieved either by direct process or extractive distillation (via medium-concentration nitric acid). The most relevant process in terms of NQ emissions is the production of medium-concentration nitric acid. Traditionally, medium-concentration nitric acid production plants are designed either as low-pressure, medium-pressure or high-pressure. The state of the art for new plants is medium-pressure equipped with selective catalytic reduction, and high-pressure plants.

34. Table 10 gives achievable clean gas concentrations for typical emission sources when applying selective catalytic reduction.

Table 10:Emission sources and selected NQ control measures with theirrespective clean gas concentrations in nitric acid production

Emission source	Combination of control measures	Clean gas concentration (mg/Nm ³)		
Production of	f medium concentration nitric	acid		
Medium dual pressure	SCR	# 400		
High dual pressure	SCR	# 100		
Production of medium- and high-concentration nitric acid (extractive distillation process)				
Medium dual pressure	SCR	# 240		
Production of high-concentration nitric acid				
Direct process or extractive distillation	SCR	# 180		
The given concentrations refer to the production step only (not to the concentration step) and to 3% Q.				

H. <u>Waste incineration</u>

35. This section addresses the incineration of municipal (or domestic), hazardous and medical wastes as well as the incineration of sludges from waste-water treatment. Different incineration technologies generate different levels of NO_x emissions (e.g. moving grate, rotary kiln, fluidized bed combustion, electric infrared).

36. Emissions of NQ_x can generally be reduced by reducing the amount of incinerated waste. This can be accomplished through various waste management stra-tegies, including recycling programmes and composting of organic materials.

37. The most relevant technologies for reducing NQ emissions are flue gas recirculation, air-staged combustion, SCR and SNCR. Table 11 gives a selection of applicable abatement options and the respective clean gas concentrations achievable by typical emission sources.

Table 11:	Emission	sources an	nd selected	NQ cont	rol measure	s with	their
respective	clean gas c	oncentratio	ons in waste	e incine	ration		

~ 1 ' · · '

Emission source	Combination of control	Clean gas	
	measures	concentration	
		(mg/Nm^3)	
Domestic	or municipal waste incineration	n	
Moving grate, rotary kiln	Primary measures (air	# 250	
	staging)		
	Flue gas recirculation or	# 140	
	air-staged combustion and		
	SNCR		
	Flue gas recirculation or	# 70	
	air-staged combustion and SCR		
Fluidized bed combustion	Flue gas recirculation or	# 80	
	air-staged combustion and		
	SNCR		
	Flue gas recirculation or	# 40	
	air-staged combustion and SCR		
Ind	ustrial waste incineration		
Grate furnace, rotary	Flue gas recirculation or	# 140	
kiln, fluidized bed	air-staged combustion and		
combustion	SNCR		
	Flue gas recirculation or	# 70	
	air-staged combustion and SCR		
Incineration of	of sludges from waste-water tre	atment	
Rotary kiln, fluidized	Flue gas recirculation or	# 140	
bed combustion,	air-staged combustion and		
multiple hearth furnace	SNCR		
	Flue gas recirculation or	# 70	
	air-staged combustion and SCR		
All fuels: 11% excess Q. Values apply to different raw gas concentrations.			

<u>Endnote</u>

 $\underline{1}/$ Recovery and recycling are to be understood in the broad sense, including reutilization off-site.

- <u>2</u>/ For retrofit only.
- $\underline{3}$ / For combustion turbines.

ADDITIONAL PROPOSALS BY THE NETHERLANDS

<u>Table 6</u>:

Primary measures should be taken into account as well.

<u>Table 7</u>:

The value given for sinter plants does not require SCR.

Table 8:

Table 8: Emission sources and selected NQ control measures with their respective reduction efficiencies and clean gas concentrations in cement production

Emission source	Combination of control	Clean gas		
	measures	concentration		
		(mg/Nm^3)		
Dry process with preheater	/precalciner			
Fuel: coal	Low NO _x burner	600 - 1,000		
		400 - 600		
	Low NO_x burner and and SNCR	300 - 500		
	Low NO_x burner and SCR	100 - 200		
Solid fuels: 10% excess Q.				
Values apply to different raw gas concentrations.				