# STUDY ON FLARING SYSTEM FOR SOUR GASES IN OIL FIELDS IN INDONESIA

by

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#### ABSTRACT

Many of the oil fields in East Java, Indonesia, are known to contain sulfur in amounts sufficiently high that the crude oil or the associated gases are considered sour. The sour gases have to be handled accordingly in order to prevent the workers and the surrounding communities suffering from the toxic gas.

The flaring system for disposing of the sour gases normally applied in oil fields in Indonesia has been evaluated in this study. Flare Stacks typically attempt to convert hydrogen sulfide  $(H_2S)$  in sour gas streams into sulfur dioxide  $(SO_2)$  and water. The amount of  $SO_2$  emitted does not solely depend on the  $H_2S$  content of the gas, but also on the internal physical factors, such as flare gas flow rate, flare gas heat content, flare gas exit velocity, and external factors such as cross wind velocity.

This study focuses on the evaluation of the influence of the physical factors, especially flare gas flow rate and wind velocity, on  $SO_2$  emission. In order to comply with the regulation the study also attempts to search the Government of Indonesia Regulations which are suitable for this purpose.

Key words : sour gas, flaring, hydrogen sulfide

#### I. INTRODUCTION

Flaring has been used primarily to dispose of gases in oilfields where the gas volumes are too small or the sites too remote, to make pipelining and gas processing economic. Due to government and industry commitments to reduce the amount of flaring, as well as the rising economic value of natural gas, more of the gas is now being pipe to processing plants.

Flare Stacks typically attempt to convert hydrogen sulfide ( $H_2S$ ) in sour gas streams into sulfur dioxide ( $SO_2$ ) and water. Since most of the gas streams in question contain carbon dioxide ( $CO_2$ ), a fire retardant, the result is an incomplete process and leads to no overall reduction in the toxicity of the gas stream.

At some facilities, flares have been replaced with more efficient incinerators. Previously incineration had been used primarily at smaller gas processing plants where the gas volumes or  $H_2S$  concentrations were too low to make sulfur recovery economic.

Due to a world oversupply of sulfur relative to demand, and resulting low prices, sulfur extraction has become less attractive economically. Sulfur stockpiles at gas processing plants represent an expense and a potential disposal problem. This provides an additional incentive for sour gas producers to seek alternatives such as acid gas injection.

Many of the oil field in East Java, Indonesia, are known to contain sulfur in amounts sufficiently high so that the crude oil or the associated gases are considered sour. The sour gases have to be handled accordingly in order to prevent the workers and the surrounding communities suffering from the toxic gas. Flare Stacks typically attempt to convert hydrogen sulfide ( $H_2S$ ) in sour gas streams into sulfur dioxide ( $SO_2$ ) and water. The amount of  $SO_2$  emitted does not solely depend on the  $H_2S$  content of the gas, but also on the physical factors, such as flare gas flow rate, flare gas heat content, flare gas exit velocity, cross wind velocity, etc.

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### **II. METHODOLOGY OF THE STUDY**

#### A. Regulation search and interviewing

Regulation search is conducted by searching available Government of Indonesia (GOI) Regulations (*Peraturan Pemerintah*, PP), including Ministerial Decrees (*Keputusan Menteri, Kep. Men.*). Interviewing is conducted with Directorate General of Oil and Gas, cq. Directorate of Techniques and Environment.

#### **B.** Dispersion calculation

Using gas composition, stack parameters and atmospheric factors, calculation of SO2 content are conducted using parameters as follows:

- Pollutant Emission Concentration
- Downwind maximum concentration of the pollutant. Equation used for the calculation is based on the formula suggested by Pasquill and modified by Gifford<sup>1</sup>.
- Emission rate (g/s)
- Stack height (m), and
- Meteorological Parameters (Wind speed, m/s, Wind direction, and Atmospheric stability)

#### III. RESULTS AND DISCUSSIONS

# A. Emission Standards

There is no specific regulation in Indonesia concerning the flaring of sour gases. Upon searching the available regulations, it is found that the Decree of The State Minister of Environment, (Kep. MenLH No. 129/2003<sup>2</sup>), describing Emission Standards for Oil and Gas Activities, is suitable for evaluating the flare system. It is stated in the Attachment-I of the regulation that opacity is the only parameter that has to be complied to. Unfortunately, no other compliance parameters such as  $H_2S$ ,  $SO_2$ , or  $NO_3$ , are described in the regulation.

Other regulations that can be referred to for the emission standard of those gases is Kep. MENLH No. 13/1995<sup>3</sup> describing Emission Standards for Point Sources (Attachment VB, Emission Standard for Point Sources of Other Activities). The Attachment VB of the regulation does not explicitly describe provisions for oil and gas activity, instead other industries that are not covered by the Attachment IB (Iron and Steel Industry), Attachment IIB (Pulp and Paper Industry), Attachment IIIB (Coal Generated Power Plant), and Attachment IVB (Cement Industry). Table 1 shows the emission standard as described in the Attachment VB of the regulation.

# **B.** Ambient Air Standards

Upon flaring the sour gas, hydrogen sulfide  $(H_2S)$  will be burned to sulfur dioxide  $(SO_2)$ . In the worst case when the flare extinguishes,  $H_2S$  will be dispersed to the ambient air. This gas is heavier than air and it will be distributed downward. Since  $H_2S$  is toxic, it is worthy to refer to the existing regulation that describes the maximum concentration level of the gases  $(H_2S)$  and  $SO_2$  in the ambient air.

Emission standard as described in Rep. MENETING. 15/1995			
No.	Parameters	Units	Threshold Limit
1	Ammonia (NH <sub>3</sub> )	mg/m <sup>3</sup>	0.5
2	Chlorine (Cl <sub>2</sub> )	mg/m <sup>3</sup>	10.0
3	Hydrochloric Acid (HCl)	mg/m <sup>3</sup>	5.0
4	Hydrofluoric Acid (HF)	mg/m <sup>3</sup>	10.0
5	Nitrogen Dioxide (NO <sub>2</sub> )	mg/m <sup>3</sup>	1000
6	Opacity	%	35
7	Particulate	mg/m <sup>3</sup>	350
8	Sulfur Dioxide (SO <sub>2</sub> )	mg/m <sup>3</sup>	800
9	Total Reduced Sulfur as $H_2S$	mg/m <sup>3</sup>	35
10	Mercury (Hg)	mg/m <sup>3</sup>	5.0
11	Arsenic (As)	mg/m <sup>3</sup>	8.0
12	Antimon (Sb)	mg/m <sup>3</sup>	8.0
13	Cadmium (Cd)	mg/m <sup>3</sup>	8.0
14	Zinc (Zn)	mg/m <sup>3</sup>	50.0
15	Lead (Pb)	mg/m <sup>3</sup>	12.0
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Emission standard as described in Kep. MENLH No.	13/1995

No.	Parameters	Units	Exposure (hour)	Threshold Limit		
1	Sulfur Dioxide (SO <sub>2</sub> )	<i>u</i> g/Nm <sup>3</sup>	24	365		
2	Carbon Monoxide (CO)	<i>u</i> g/Nm <sup>3</sup>	24	10.000		
3	Nitrogen Dioxide (NO <sub>2</sub> )	<i>u</i> g/Nm <sup>3</sup>	24	150		
4	O3 (Ozone)	<i>u</i> g/Nm <sup>3</sup>	1	235		
5	Hydrocarbon (HC)	<i>u</i> g/Nm <sup>3</sup>	3	160		
6	Particulate < 10 um (PM10)	<i>u</i> g/Nm <sup>3</sup>	24	150		
7	Particulate < 2.5 um (PM2.5)	<i>u</i> g/Nm <sup>3</sup>	24	65		
8	Total Suspended Particulate (TSP)	<i>u</i> g/Nm <sup>3</sup>	24	230		
9	Lead (Pb)	<i>u</i> g/Nm <sup>3</sup>	24	2		
10	Ductfell	<i>u</i> g/Nm <sup>3</sup> 30 day	20 day	10 tons/Km2 (Settlement)		
10	Dustian		ug/initi 50 day	ug/Mill So day	ug/Mill 50 day	
11	Total Fluorides	<i>u</i> g/Nm <sup>3</sup>	24	3		
12	Fluor Index	<i>u</i> g/Nm <sup>3</sup>	30 day	40 ug/100 cm2		
13	Chlorine & Chlorine oxide	<i>u</i> g/Nm <sup>3</sup>	24	150		
14	Sulphate Index	<i>u</i> g/Nm <sup>3</sup>	30 day	1 mg SO3/100 cm3		
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Table 2 National Ambient Air Standard

Government of Indonesia Regulation PP No. 41/ 1999<sup>4</sup> describing Air Pollution Control can be referred to for this purposes. Table 2 shows the threshold limits of the contaminant parameters. Unfortunately, the regulation does not include H<sub>2</sub>S as the contaminant parameter. Article 58 of the regulation states that "all regulations describing air pollution control that still exist and does not controverts to this regulation and it has not been replaced by this regulation is still valid". It is therefore, in referring to H<sub>2</sub>S level in the ambient air, Kep. MenKLH No. 02/19885 is still suitable. This Ministerial Decree states that H<sub>2</sub>S concentration in the ambient air is allowed not to exceed 0.03 ppm  $(42 \text{ ug/m}^3)$ . It should be noted, however, that this Kep. MenKLH No. 02/1988 is only valid for H<sub>2</sub>S in ambient air, and not valid for evaluating emission gases since it has been replaced by Kep. MenLH No. 13/ 1995.

Since this study is focused on  $H_2S$  and  $SO_2$ , we attempt to summarize the regulation for these parameters. Table 3 shows the threshold concentrations of these parameters either for emission or ambient standard. It is clearly stated that for flaring activity in oil fields only opacity is the only factor to comply to.

Meanwhile, if we have to follow the provision of  $H_2S$  and  $SO_2$  parameters for emission standards, we should refer to the Emission Standard for Point Source (Kep. MenLH No. 13/1995) that describes the threshold limit for SO<sub>2</sub> and  $H_2S$ , which are 800 mg/m<sup>3</sup> and 35 mg/m<sup>3</sup>, respectively.

It has to be noted, however, that Emission Standard for Point Source described in the Kep. MenLH No. 13/1995 is only valid for waste gases, resulted from the combustion of the materials, which are ready to be emitted to the ambient air. This provision clearly can not be applied to the flare system in which the waste gas is being burned at the end of the gas stream. The gas which has been burned in the flare system is then dispersed to the atmosphere. Emission gas measurement is not possible to be conducted either at the gas stream or at the location nearest to the combustion process. For this purpose, emission gas measurement can only be carried out at certain distance from the combustion source. Once the gas has been emitted to the atmosphere, the only possibility to measure the pollutant concentration is at the ambient air as the receiving media. Therefore, provision for gas emission in flare system is worthy to refer to the National Ambient Air Standard. Ambient standard for SO<sub>2</sub> refers to PP 41/1999 that requires SO<sub>2</sub> concentration in ambient air not to exceed 365 ug/ m<sup>3</sup>, while those of H<sub>2</sub>S concentration refers to Kep. MenKLH No. 02/1988 that requires H<sub>2</sub>S concentration in ambient air not to exceed 42 ug/m<sup>3</sup>.

Both of the pollutant concentrations in the ambient air can be measured directly using  $SO_2$  and  $H_2S$ analyzers with regard to the distance of the receptors to the emission source.  $SO_2$  and  $H_2S$  concentrations in the ambient air can also be calculated using a dispersion equation model. This calculation is written in this report in Section 3.3: dispersion calculation.

#### C. Dispersion calculation

Dispersion calculations of the pollutants are carried out using a Gaussian plume model that incorporates source related and meteorological factors (or parameters) to estimate pollutant concentrations from continuous sources. It is assumed that the pollutant does not undergo any chemical reactions, and that no other removal processes, such as wet or dry deposition, act on the plume during its transport from the source. As mentioned in the methodology, equation used for the calculation is based on the formula suggested by Pasquill and modified by Gifford.

In such case when the flare is extinguished, the pollutant will be assumed solely as hydrogen sulfide. When the flare is activated, meaning that the waste gas can be burned, the pollutants will be a mixture of  $SO_2$  resulting from the oxidation of  $H_2S$  and unburned  $H_2S$ . The amount of  $SO_2$  and unburned  $H_2S$  is a func-

tion of the  $CO_2$  content in the gas that determines the combustion efficiency.

According to the available data, the dispersion of the pollutants will be calculated using this data (Table 4).

At a standard condition (0 °C, 1 atm), mass of  $H_2S$  can be calculated using the formula below:

$$m_{H_2S} = \frac{Q_{H_2S}}{V_c} x M_{H_2S}$$

Where:

 $V_{s}$  molar volume of the gas= 22.414 m<sup>3</sup>/kmol

 $M_{H2S}$  molecular weight of H2S = 34.082 kg/kmol Hence  $m_{H2S}$  is 1,271 kg/d, and mass of H<sub>2</sub>S being burned is also 1,271 kg/d.

Physical factors used in the dispersion calculation are assumed to be: Effective stack height is 20 m, Wind speed is 3 m/s, and Atmospheric stability is Moderate/Stable (C class).

#### **Determination of SO**<sub>2</sub> emmission:

Combustion rate of  $H_2S$  (combustion efficiency is considered to be 100%)

Reaction:  $H_2S + 3/2O_2 = SO_2 + H_2O (1 \text{ mol of } H_2S \text{ produces } 1 \text{ mol of } SO_2)$ 

Mol of $H_2S$ is	37	kmol/d
Mol of SO <sub>2</sub> produced is	37	kmol/d
Rate of SO <sub>2</sub> produced is	2,390	kg/d SO <sub>2</sub>
Hence Q <sub>SO2</sub> is	28	g/s

Anowable maximum concentration of $\Pi_2$ 5 and SO <sub>2</sub> for emission gases and ambient air					
	Concentration				
	Emiss	sion Gas	Ambient Air		
Parameters	Kep. MenLH No. 13/1995	Kep. MenLH No. 129/2003	PP. No. 41/1999	Kep. MenKLH No. 02/1988	
	(Attachment VB)	(Attachment I)	(Attachment I)	(Attachment III)	
	Emission Standard for Point Source	Emission Standard for Oil and Gas Exploration and Production	National Ambient Standard	Ambient Standard	
SO <sub>2</sub>	800 mg/m <sup>3</sup>	-	365 ug/m <sup>3</sup>	-	
H <sub>2</sub> S	35 mg/m <sup>3</sup>	-	-	42 ug/m <sup>3</sup>	
Opasitas (Flare Stack)	-	40%	-	-	

Iablel 3			
Allowable Maximum Concentration of H	S and SO	, for emission ga	ases and ambient air

Maximum concentration of SO2 at certa	ain distance	
Nearest settlement to the flare	1,000 m	
Lateral diffusion coefficient s <sub>v</sub>	120	т
Vertical diffusion coefficient s <sub>z</sub>	70	т
$C_{max}$ of SO <sub>2</sub> at the distance of 1 km	0.00033542	2 g/m3
	335	ug/m3
	0.12	ppm-vol
Ambient threshold limit of SO <sub>2</sub> (PP.4	41/1999)	365 ug/m3
	0.13	ppm-vol
When flare extinguished		
Q H <sub>2</sub> S	37 <i>k</i>	mol/d
	1,271	kg/d
	14.7	g/s
$C_{max}$ of $H_2S$ at the distance of 1 km	0.0001	7844 <i>g/m3</i>
	178	ug/m3
	0.12	ppm-vol
Ambient threshold limit of $H_2S$		
(Kep. MNLH 02/88)	42	ug/m3
	0.03	ppm-vol
OSHA Standard for H <sub>2</sub> S		
(Odor Threshold)	0.02	ppm-vol

Results of the dispersion calculation indicate that only certain value of production rate of crude oil (hence the production of associated gases to be burned) can use flare system. In this study it is found that for a production rate of crude oil at 5000 bpd (barrels per day), the SO<sub>2</sub> concentration in the ambient air with a distance of 1 km from the emission source is 335 ug/m<sup>3</sup>. This value can be regarded as a limit for the crude oil production rate, otherwise the SO<sub>2</sub> concentration in the ambient air will exceed the ambient air standard. 1 Km distance has been chosen as a distance limit, since in the populated area such as Blora and Bojonegoro Regencies the public settlements are very close to the oil fields.

#### **D.** Alternatives technologies

Alternative technologies are needed when a flare system for sour gas is not suitable, or crude oil production rate would be increased. There are two approaches that can be used to alternate direct flaring technique. This would be sulfur removal/recovery system and sub-surface injection6.

One of the sulfur removal or recovery unit uses Clauss process, in which  $H_2S$  is oxidized to free sulfur. The elemental sulfur formed is sold to customers and is used mainly in the manufacturing of phosphate fertilizers. Some sulfur is also used in making products such as pharmaceuticals, plastics and matches.

Environmental concern over global warming due to greenhouse gas emission has given ever rising importance to the re-injection of waste gases in oil fields; either for reutilization to Enhance Oil Recovery (EOR) or just simple disposal to a depleted reservoir to avoid atmospheric venting. Moreover, with the growing acceptance of  $H_2S$  re-injection as a feasible alternative to costly sulfur, a number of very sour gas wells can be reconsidered as exploitable to produce much needed natural gas.

Acid gas re-injection to a disposal reservoir will provide the simplification and cost reduction to make exploitation of the sour gas reserves attractive even in a low gas price scenario. Neither  $CO_2$  nor sulfur emissions are emitted to the atmosphere. Re-injection of sour gases is very promising; nevertheless the discussion of this topic is beyond the scope of our study.

# IV. CONCLUSIONS AND RECOMMENDATIONS

Upon completing the study on flaring system for sour gases some conclusions and recommendations can be made as follows:

# A. Conclusions

- There are no recent Indonesian Regulations specifically regulating the flaring of sour gases. The only parameter that should be complied to is opacity as described in

Table 4Crude oil and gas data				
Parameters Concentration Unit				
Gas Oil Ratio (GOR)	381	scf/STB		
Q (flow rate) oil production	5	bpd		
Q gas	1,905,000	scf/d		
	53,944	m³/d		
H <sub>2</sub> S content	1.55	%-mol		
$Q_{H2S} = Q_{gas} \times C_{H2S}$	836	m <sup>3</sup> /d		

Kep. MenLH No. 129/2003), Emission Standard for Oil and Gas Exploration and Production.

- No other parameters, such as H<sub>2</sub>S and SO<sub>2</sub> are described in the regulation. For a flaring system of H<sub>2</sub>S and SO<sub>2</sub> emissions it is worthy to refer to ambient standard as described in Kep. MenKLH No. 02/1988 and PP. No. 41/1999, respectively, rather than emission standard as described in Kep. MenLH No. 13/1995.
- When a flaring system has to be used for emitted sour gases, especially in a populated area, only limited portion of the gas can be burned depending on the gas oil ratio (GOR) and the  $H_2S$  content, otherwise the concentration of the emission gases in the ambient air will exceed the allowable concentration.
- H<sub>2</sub>S and SO<sub>2</sub> concentration in the ambient air can be measured directly using analyzers or can be calculated using a dispersion equation model.

# **B.** Recommendation

- It is recommended to review the existing regulations for the emission of H<sub>2</sub>S and SO<sub>2</sub> gases in oil field flaring systems in Indonesia.
- Re-injection is an alternative technology to a flare

system of sour gases and it is recommended to be investigated in more detail.

# REFERENCES

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