

## Oil Refineries Emissions Impact on Urban Localities Using AERMOD

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**Abstract: Problem statement:** The absolute necessity of compulsory fuel utilities, no matter small or big has resulted into substantial high hazards pollutants. Petroleum refineries are major industrial installations that are necessary for providing the best suited fuel for various necessary utilities, but are responsible of the emission of several hazardous pollutants into the atmosphere. Hydrocarbons are among the most perilous air pollutants that are emitted from almost all refining processes in petroleum refineries. **Approach:** Every day leaks and gaseous discharge from relief valves and liquid discharge, which are often directed to knock-out drums, are flared to minimize the impact of hydrocarbons emissions. But these flares are not that efficient and result into partial discharge of pollutants that have severe impact on the industrial area and urban localities in the vicinity of industrial refining complex. **Results:** In the present study, a thorough investigation has been completed to estimate the total emissions of sulfur dioxide SO<sub>2</sub> and non methane hydrocarbons NMHC (VOCs) and to assess their impact on the air quality in industrial and suburban areas. The latest version AMS/EPA Regulatory Model (AERMOD) specially designed to support the US Environmental Protection Agency (EPA)'s was used to predict the ground level concentrations of SO<sub>2</sub>, VOCs from AL-Ahmadi and Al-Shuiba Refineries of total refining capacity of 646 thousand barrels/day. **Conclusion/Recommendations:** These concentrations are compared with EPA standards to indicate the ambient air quality. The dispersion model was corroborated with extensive one year hourly record of the surface and upper air meteorological data for year 2006 and emission rates of the specified pollutants, with detailed refinery stacks parameters, such as stack height, diameter, exit flue gas velocity and temperature to determine the fraction of total study area in the vicinity of refineries that had substantially high concentration of these pollutants. It's found that 10 % of selected area under study has exceedance for SO<sub>2</sub>, 13 % exceedance for non-methane hydrocarbons NMHC, and there is no exceedance for NO<sub>2</sub> which is considered about 50 % of total NO<sub>x</sub> emissions.

**Key words:** Hydrogen sulfide, Volatile Organic Compound (VOC), meteorological parameters, sulfur dioxide, Environmental Protection Agency (EPA), Industrial Source Complex (ISC), urban localities, oil refineries, AMS/EPA Regulatory Model (AERMOD)

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### INTRODUCTION

Air pollution is a major problem that has been recognized throughout the world for almost a century. The lifestyle that we take for granted is made possible through the usage of the fuels (i.e., LPG, motor gasoline, diesel fuels, jet fuels, aviation gasoline, heating oil's, heavy fuel oils,) and specialty products (i.e., lubricants, waxes, asphalts and solvents), as well as the petrochemical industry feed stocks produced by the petroleum refining industry from associated gas and crude oils. The petroleum refineries and petrochemical

plants are the largest sources of air pollution in the region. Their operation is associated with the emission of various organic compounds into the atmosphere. (Cetin *et al.*, 2003; Al-Hamad *et al.*, 2008).

The flare system is designed to provide safe receipt and disposal of unwanted, toxic gases/vapors released from process equipment during normal operations to minimize the impact of emissions, but these flares are not that efficient and result into partial discharge of hydrocarbons and other emissions of carbon dioxide, carbon monoxide, NO<sub>x</sub> and SO<sub>x</sub> at

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elevated temperature that have severe influence on the industrial area and urban localities in the vicinity of industrial refining complex.

The most two controversial pollutants are SO<sub>2</sub> and NMHC. These atmospheric pollutants once emitted into the atmosphere may cause a pollution problem on local scale and a major health risk to the population in the urban areas in the vicinity of oil installation. The regional problem like acid rain or photochemical ozone production in the troposphere initiated by the reaction of hydrocarbons with OH radicals in the presence of nitrogen oxides and sunlight originate smog and visibility problems. (Kalabokas *et al.*, 2001; Ramadan *et al.*, 2008).

Most of the Hydrocarbons emitted by Petroleum refineries play major role in the physicochemical processes of the troposphere as they largely contribute to the formation of ozone and other photochemical oxidants (Kalabokas *et al.*, 2001). Moreover, some hydrocarbons are highly toxic or carcinogenic. Hydrocarbon are also generated as a result of human activities, arising mainly from motor vehicle exhausts and other combustion processes utilizing fossil fuels, petrol storage and distribution, solvent usage and other industrial processes (Al-Hamad *et al.*, 2008).

Sulfur dioxide causes a wide variety of health risk and its environmental impact has been known as acid rain over a century. Particularly sensitive groups include people with asthma who are active outdoors and children, the elderly with heart or lung diseases. SO<sub>2</sub> and the other associated pollutants such as sulfate particles can be transported over long distances and deposited far from the point of origin. This means that problems with SO<sub>2</sub> are not confined to areas where it is emitted (Al-Jahdali and Bisher, 2008).

## MATERIALS AND METHODS

The aim of the present study is the determination of atmospheric pollution levels, a thorough investigation has been completed to estimate the total emissions of SO<sub>2</sub> and VOCs and assess their impact in the air quality in industrial and suburban areas. The latest version AMS/EPA Regulatory Model (AERMOD) (Venkatram *et al.*, 2004) specially designed to support the US Environmental Protection Agency (EPA)'s was used to predict the ground level concentrations of SO<sub>2</sub>, NMHC from AL-Ahmadi and Al-Shuiba Refineries of total refining capacity of 646 thousand barrels/day. The dispersion model was corroborated with extensive one year hourly record of the surface and upper air meteorological data for year 2006 and emission rates of the specified pollutants, with detailed refinery stack parameters, such as stack height, diameter, exit flue gas velocity and temperature to determine the ground level concentrations of SO<sub>2</sub> and NMHC and assess the fraction of the total study area under exceedence of EPA standards.



Fig. 1: Satellite image showing urban areas in the vicinity of oil refineries

**Area description under investigation:** The area under study in this study covers the oil refinery areas in the state of Kuwait. Kuwait is surrounded by hot deserts in the north, west and south and in the east it is bordered by sea. Kuwait has the third largest oil reserves in the world after Saudi Arabia and Iraq. The oil reserve of Kuwait is estimated to around 10% of the total world reserve. The production capacity of Kuwait's refineries altogether amounts to approximately 936 thousand barrels/day, distribute as follows: Mina Ahmadi Refinery (446 thousand barrels/day) consists of four flaring stacks (MAFP, FUP, RMP and OR), Shuaiba Refinery (200 thousand barrels/day) has two identical flares in all respect (ST29-01 and ST29-02) And Mina Abdulla Refinery (470 thousand barrels/day).

The urban areas surrounding oil refineries are Fahaheel, Ahmadi and Omm Hayman Areas as shown in Fig. 1. These urban areas include all the human facilities like schools, hospitals, hotels, shopping malls and recreational activities in clubs and stadiums. The receptor grid specified covers an area of 500 km<sup>2</sup> keeping the predominant emission sources in the center.

An average composition of Kuwaiti crude oil is 2.44% by weight Sulfur, 0.14% Nitrogen, 7.7 ppm Nickel, 28 ppm Vanadium, 22.7% Naphtha fraction (boiling pt. from 20-205°C), 77.3% High boiling fraction (boiling pt. above 205°C), 23.3% Aromatics, 20.9% Paraffin and 3.5% Insoluble. That's refined to produce various types of fuels for domestic and industrial use. The total emissions for oil refinery facilities are mainly from boilers, fired heaters, hydrogen reforming, hydro-cracking, hydro-treating, hydro-desulphurization units and other distillation processes.

**Model application:** For this purpose, The AMS/EPA Regulatory Model (AERMOD) specially designed to support EPA's regulatory modeling programs. AERMOD is regulatory steady state plume modeling system with three separate components: AERMOD (AERMIC Dispersion Model), AERMAP (AERMOD Terrain Preprocessor) and AERMET (AERMOD Meteorological Preprocessor). The AERMOD model includes a wide range options for modeling air quality impacts of pollution sources, making it popular choice among the modeling community for a variety of applications. (Holmes and Morawska, 2006).

AERMOD has an improved approach for characterizing the fundamental boundary layer parameters and vertical profile of the atmosphere along with better representation of plume buoyancy, penetration and urban nighttime boundary layer, it provides variable urban treatment of vertical dispersion as a function of city populations as compared to Industrial Source Complex (ISC) which is mainly limited to regulatory purposed. (Kesarkar *et al.*, 2007; Abdul-Wahba *et al.*, 2002).

The input data that describe both emission source such as pollutant emission rate ( $\text{gs}^{-1}$ ); Base elevation from sea level (m); Stack height (m); Stack coordinate location; Exit stack inner diameter (m); Exit stack gas speed ( $\text{ms}^{-1}$ ); and Exit stack gas temperature (K); and meteorology provide a comprehensive set of information which can be used to run AERMOD model and thus simulate the ground level concentration of the pollutants.

Meteorological conditions play a major role in the dispersion of pollutants emitted from the refineries in the state of Kuwait (Isakov *et al.*, 2007); the Area under study affected by the effect of sea and land breeze and hence affects the ground level concentration of  $\text{SO}_2$  and VOC in the residential areas. A one year hourly record of the surface and upper air meteorological data for year 2006 obtained for Ahmadi and Shuiba refineries is used in this study for simulation of the dispersion of these pollutants from the flares.

The study area consists of  $21 \times 24$  km rectangular having all emission sources in the center covering the industrial and urban areas.

The AERMET program is a meteorological preprocessor which prepares hourly surface data and upper air data for use in the U.S. EPA AERMOD short term air quality dispersion model. AERMET processes meteorological data in three stages and from this process two files are generated for use with the AERMOD model:

- A surface file of hourly boundary layer parameter estimates

- A profile files of multiple-level observations of wind speed, wind direction, temperature and standard deviation of the fluctuating wind components

Other assumptions which were used in the AERMOD model include the following:

- Steady state conditions
- emission inventories are time independent
- no interactions among various emission sources
- emission losses due to chemical reaction, absorption or deposition are negligible
- Effects of various structures located in the vicinity of emitting sources have negligible effect

## RESULTS AND DISCUSSION

The emission rates of the point sources, locations and other physical parameters were fed from Al-Ahmadi and Al-suiba refineries. A  $500 \text{ km}^2$  grid consisting of  $21 \times 21$  nodes equally spaced 2.5km apart has been selected in combination with a finer grid of 441 receptor locations and 70 discrete most sensitive receptors distributed in the urban areas. The AERMOD model was executed for total  $\text{SO}_2$  emissions and the impact of predicted ground level concentration in the study area. The model was recomputed with VOCs emissions to assess the impact of ground level concentration of total VOCs in the study area. The AERMOD model results were composed of the maximum ground level concentration of both  $\text{SO}_2$  and VOCS in the selected grid area. Their emissions impact was assessed for a period of 8760 h equivalent to one year using hourly synoptic meteorological data for a year 2006. The average hourly, daily and annual maximum ground level concentrations of both  $\text{SO}_2$  and VOC's were compared with the Kuwait Environmental Public Authority, EPA standards.

The isopleths plot was generated; (Fig. 3) which showed the highest hourly average ground-level concentration of  $\text{SO}_2$  around the refineries. From the total area under investigation 18% was above the KEPA limits at immediate neighborhood of the industries, which is  $90.25 \text{ km}^2$ , 6.75 km E, 2.5 km W, 7 km S and 2.5 km N with respect to MAFP flare. Highest maximum concentration predicted hourly of  $\text{SO}_2$  was  $1224.46 \mu\text{g m}^3$  on 15th February at 1.00 AM in south east SE direction with respect to MAFP with distance 5.8 km.

In Kuwait the prevailing wind is almost all the time from north west NW as shown in Fig. 2 confirming the highest ground level  $\text{SO}_2$  concentration to be the south east SE from the predominate source.

The model was run considering only AL-Shuiba refinery emission sources and its contribution to the highest ground level concentration has been assessed.

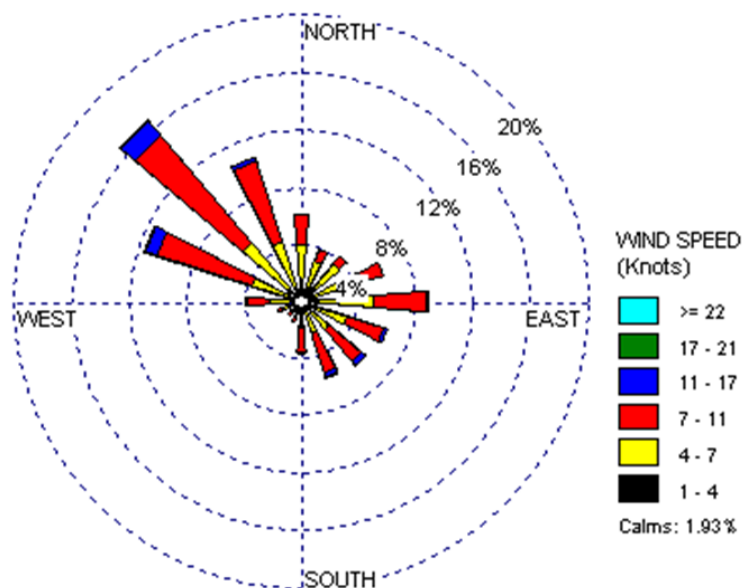


Fig. 2: Wind rose for Coastal Meteorological Data of year 2006 in Kuwait

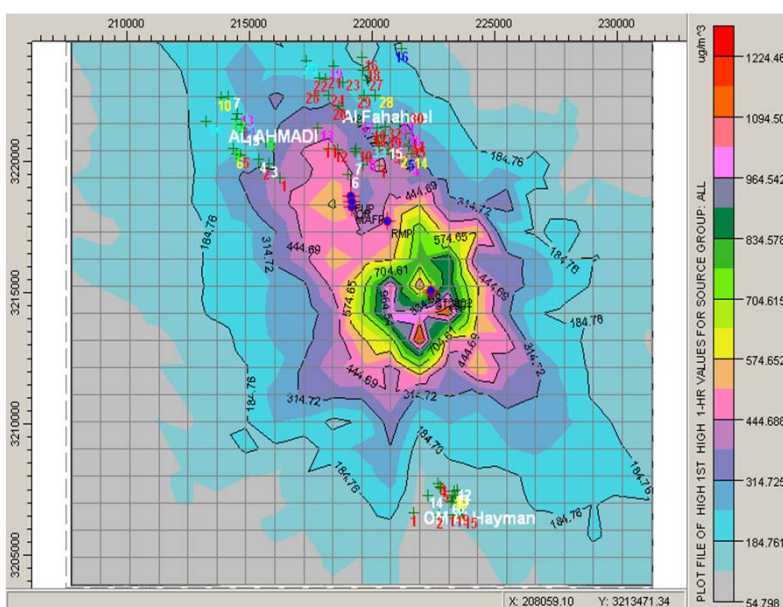


Fig. 3: Isopleth showing the hourly average ground Level SO<sub>2</sub> concentration

The first maximum concentration predicted hourly from Al-Shuiba refinery alone was 1219.96  $\mu\text{g m}^{-3}$ . And this contributes 99.63% of the total ground level concentration of SO<sub>2</sub> indicating very little impact due to Al-Ahmadi refinery. The average sulfur content in the gas flared from Al-Ahmadi 0.88% by volume hydrogen sulfide H<sub>2</sub>S while the average sulfur content of flared gas from Al-Shuiba refinery is 6.1% by V H<sub>2</sub>S showing the predominate influence on ground level concentration.

All the maximum values were in cold months of winter having low temperature and low conversion layer controlling adversely the dispersion phenomena.

Al-Shuiba refinery consists of two refinery stacks, ST29-01 and ST29-02 and AL- Ahmadi refinery consists of four flaring stacks, MAFP, RMP, OR and FUP. We compared between the first maximum concentration predicted hourly of SO<sub>2</sub> from individual flares it's found that ST29-01 and ST29-02 of Al-Shuiba refinery have

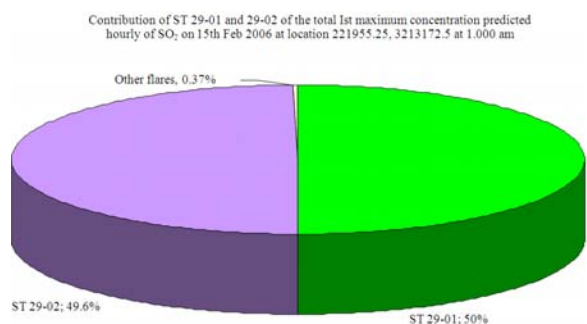


Fig. 4: Contribution of flaring sources in the hourly predicted of the highest maximum ground level concentration of SO<sub>2</sub>

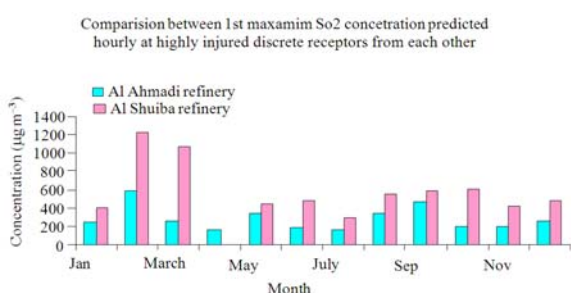


Fig. 5: Comparison of the highest maximum concentration of SO<sub>2</sub> predicted hourly from Al-Ahmadi and Al-Shuiba refineries

the highest impact then MAFP, OR, RMP and FUP due to low sulfur emissions respectively.

Also Fig. 4 illustrated the Contribution of flaring sources in the hourly predicted of the highest maximum ground level concentration of SO<sub>2</sub>

As shown Fig. 5 the concentrations predicted hourly from Al-Shuiba refinery of SO<sub>2</sub> is higher than Al-Ahmadi refinery in all the months except April where Shuiba refinery was shut down for one month. All the maximum values were in cold months of winter due to low temperature that adversely affected the dispersion phenomena.

The highest maximum concentration predicted daily of SO<sub>2</sub> was 177.58 µg m<sup>-3</sup> which exceed EPA value on 14th February at midnight 24.00 PM in direction south east SE with respect to MAFP with distance 3.4 km from MAFP flare. It was found that 0.04% of the total area under investigation was above the KEPA limit which is 0.18 km<sup>2</sup> at immediate neighborhood of the industries, that's confirming again the prevailing North West NW wind as indicating in Fig. 6.

Highest maximum concentration predicted annually of SO<sub>2</sub> was 9.97 µg/m<sup>3</sup> in the direction of south east SE with respect to MAFP with distance 1.8 km as shown in Fig. 7.

Table 1 shows the first maximum concentrations of SO<sub>2</sub> predicted hourly and daily monthly in each month for year 2006. Winter months had high pollutants concentrations than summer months due to the least dispersion influenced by low temperature and low inversion layer. Sometimes the emission rates in summer are high due to the higher capacity of production and market demand that has resulted into higher ground level concentrations as shown in Table 2.

Table 3 showing that the 50 values of maximum concentration of SO<sub>2</sub> predicted hourly from Al-Ahmadi and Al-Shuiba Refineries are mostly in south east SE direction as the prevailing wind is almost all the time from north west NW as shown in Fig. 2. Most of the 50 values in winter months at early morning hours.

Highly influenced discrete receptors of the 70 points from SO<sub>2</sub> Concentration predicted hourly were number NO.16 (primary school), No.26 (primary school), NO.21 (Ministry of Electricity), NO.19 (Public Park) with concentrations levels 494.15, 480.92, 466.42 and 450.87 (µg m<sup>-3</sup>) respectively. These values exceed EPA and are located in direction NE, NW, NW and NE respectively with respect to reference source MAFP at distance 1.9, 2.4, 1.25 and 2.8 km. These receptors are near to Al-Ahmadi refinery and has strong influence from Al-Ahmadi emissions as shown in Fig. 8. The receptor NO.19 (Public Park) has showed high concentration that was influenced from Al-Shuiba refinery on 14th August at 21.00 as the wind direction was from south west SW direction blow up towards Al-Fahaheel area.

The isopleths plot was generated, (Fig. 9) which showed the highest 3 h. maximum ground-level concentrations of VOCs around the refineries. From the total study area under investigation 14.85% was above the KEPA limits at immediate neighborhood of the industries, which is 74.3 km<sup>2</sup>, 5.5 km E, 5.3 km W, 3.6 km S and 3 km N with respect to MAFP flare. First maximum concentration predicted per 3HR was 992.4 µg m<sup>-3</sup>, (0.47 ppm) on 14th February at 6.00 am in North West NW direction with respect to MAFP with distance 2 km. because in early morning there is low temperature and low inversion layer. The wind condition is not strongly influencing the plume touchdown which is in the NW 2 km from MAFP.

The model was run considering only Al-Ahmadi refinery emission sources and its contribution to the highest ground level concentration has been assessed. The first maximum concentration predicted for 3HR from Al-Ahmadi refinery alone was 935.05 µg m<sup>-3</sup>, (0.46 ppm). And this contributes 99.84% of the total ground level concentration indicating almost negligible impact due to Al-Shuiba refinery. The average content of NMHC in the gas flared from Al-Ahmadi refinery is 30% by V while the average NMHC content of flared gas from Al-Shuiba refinery is 10.3% by V showing the predominate influence on ground level concentration.

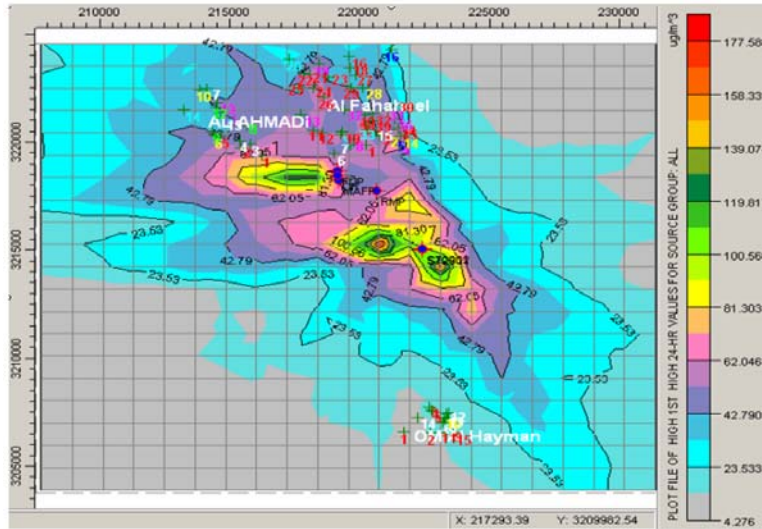


Fig. 6: Isopleth showing the daily maximum average ground Level SO<sub>2</sub> concentration

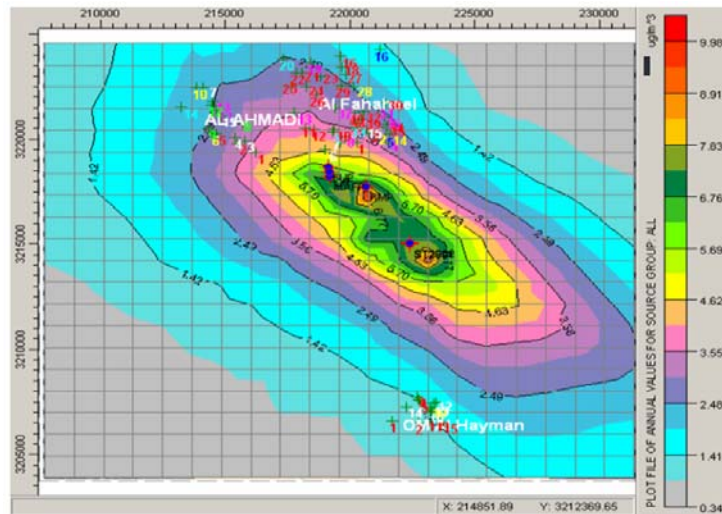


Fig. 7: Isopleth showing the annual maximum average ground Level SO<sub>2</sub> concentration

Table 1: First maximum concentrations of SO<sub>2</sub> predicted hourly and daily in each month

	1st hourly ( $\mu\text{g m}^{-3}$ )	Direction	Distance (km)	Date	1st Daily ( $\mu\text{g m}^{-3}$ )	Direction	Distance (km)	Date
Jan	467.39	SE	5.63	12/01/2006 01:00	38.24	SE	5.63	12/01/2006 24:00
Feb	1224.5	SE	5.76	15/02/2006 01:00	177.59	SE	3.38	14/02/2006 24:00
March	1080.9	SE	3.38	04/03/2006 07:00	64.20	SE	3.38	04/03/2006 24:00
April	168.75	NW	2.32	20/04/2006 22:00	11.79	NW	2.32	20/04/2006 24:00
May	441.02	SE	4.88	12/05/2006 21:00	29.69	SE	4.32	06/05/2006 24:00
June	484.41	SE	4.94	08/06/2006 22:00	27.27	SE	1.99	18/06/2006 24:00
July	296.56	SE	6.51	21/07/2006 21:00	33.68	SE	5.63	07/07/2006 24:00
Aug	548.21	SE	4.88	17/08/2006 01:00	56.09	SE	4.94	13/08/2006 24:00
Sep	646.80	SE	5.63	17/09/2006 21:00	40.16	SE	5.63	17/09/2006 24:00
Oct	605.09	SE	4.94	23/10/2006 01:00	70.26	SE	4.94	22/10/2006 24:00
Nov	439.11	SE	5.63	13/11/2006 07:00	38.38	SE	5.63	06/11/2006 24:00
Dec	482.35	SE	4.94	15/12/2006 20:00	45.85	SE	1.83	31/12/2006 24:00

All maximum values of VOC concentrations were in cold months of winter facilitated by low temperature and low inversion layer controlling adversely the dispersion phenomena.

1st maximum concentration predicted daily of VOC was  $260.16 \mu\text{g m}^{-3}$  on 14th February at 24.00 PM in North West NW direction with respect to MAFP at a distance 8 km (Fig. 10).

Highest maximum concentration predicted annually of VOC was  $15.33 \mu\text{g m}^{-3}$  in the North West NW direction with respect to MAFP at distance 1 km. (Fig. 11).

Table 4 show the maximum concentrations of VOC predicted per 3HR and daily for each month of year 2006.

Table 6 show the 50 maximum concentration of VOC predicted hourly from Al-Ahmadi and Al-Shuiba Refineries. From the 50 maximum concentrations of VOC predicted per 3HR we found only 14 values exceed EPA standards for 6.00-9.00 h while all other high concentration values were almost from 21.00-24.00 h.

Discrete receptors located at schools, clinics, shopping malls showed the highest VOCs concentrations at 21.00-24.00 HR in September month as indicated at Table 7, this due the high emissions rate from OR flare as indicating in Table 5.

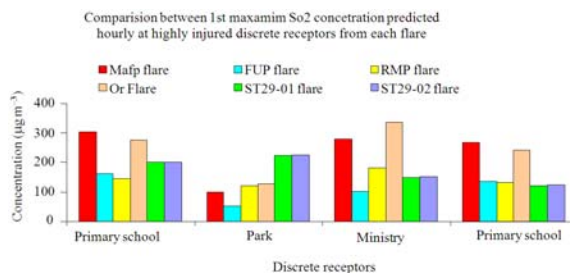


Fig. 8: Individual flares impact on highly influenced discrete receptors

Table 2: Influence of different flares on predicted first maximum ground level concentration of SO<sub>2</sub>

	Month of highest max Conc. of SO <sub>2</sub> hourly	Month of highest max Conc. of SO <sub>2</sub> daily
Real Situation	Feb	Feb
Ahmadi Refinery	Feb	Feb
Shuiba Refinery	Feb	Feb
MAFP	Feb	Feb
FUP	Feb	Feb
RMP	Dec	Dec
OR	Sep	Sep
ST29_01	Feb	Feb
ST29_02	Feb	Fe

Table 3: The 50 maximum concentrations of SO<sub>2</sub> predicted hourly from both Al-Ahmadi and Al-Shuiba Refineries

The maximum 50 1-HR average concentration values of SO <sub>2</sub> for source group: both Ahmadi and Shuiba refineries ( $\mu\text{g m}^{-3}$ )					The maximum 50 1-HR average concentration values of SO <sub>2</sub> for source group: both Ahmadi and Shuiba refineries ( $\mu\text{g m}^{-3}$ )				
Date (YYMMDDHH)	Distance (km)	Direction	Rank		Date (YYMMDDHH)	Distance (km)	Direction	Rank	
06021501	5.7	SE	26	1224.5	06030420	4.4	SE		604.50
06021222	5.6	SE	27	1169.4	06021219	5.6	SE		594.46
06021218	5.6	SE	28	1121.1	06021402	7.9	NW		591.21
06030407	3.4	SE	29	1080.9	06090124	3.4	SE		588.85
06021402	3.4	SE	30	1065.2	06030421	5.9	SE		583.04
06031406	4.9	SE	31	1023.1	06102223	4.9	SE		581.55
06031902	4.3	SE	32	999.48	06102224	4.9	SE		581.33
06022201	4.4	SE	33	909.95	06021222	7.9	SE		578.39
06033023	5.8	SE	34	909.65	06022805	2.8	NE		573.11
06030421	4.9	SE	35	889.22	06030724	6.5	SE		570.33
06022804	3.4	SE	36	795.73	06022320	5.2	SE		565.42
06022704	3.4	SE	37	775.42	06090923	5.7	SE		562.45
06022805	2.9	SE	38	760.41	06022222	5.1	SE		559.01
06021406	3.4	SE	39	719.88	06022124	3.4	SE		548.88
06022124	2.9	SE	40	707.55	06081701	4.8	SE		548.21
06031807	4.1	SE	41	694.57	06030719	2.5	SE		544.65
06021501	6.7	SE	42	690.80	06021406	2.1	SW		543.63
06020419	5.3	SE	43	668.34	06022201	5.2	SE		528.41
06030407	3.1	SE	44	666.16	06021406	7.9	NW		527.38
06091721	5.7	SE	45	646.79	06021423	3.4	SE		525.81
06091302	5.7	SE	46	639.38	06091223	4.9	SE		523.69
06022222	4.3	SE	47	633.64	06093023	4.9	SE		523.13
06030406	5.3	SE	48	632.86	06030720	4.1	SE		521.80
06031402	4.4	SE	49	610.65	06022320	4.4	SE		519.34
06102301	4.9	SE	50	605.09	06101105	3.4	SE		508.10

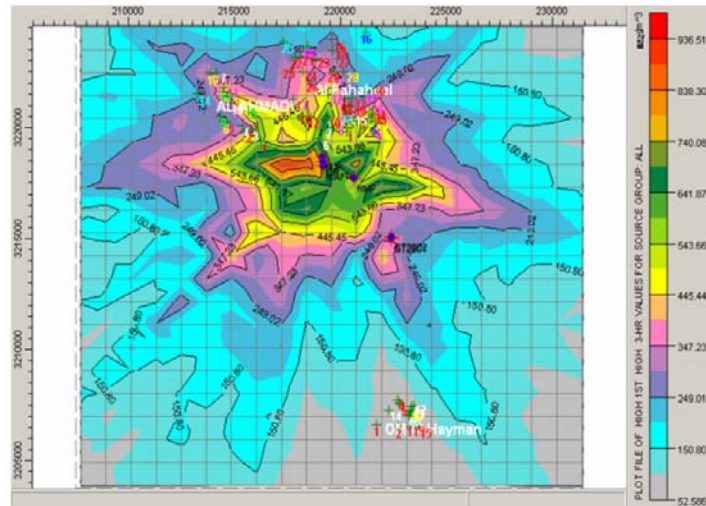


Fig. 9: Isopleth showing the maximum 3HR average ground Level VOC concentrations

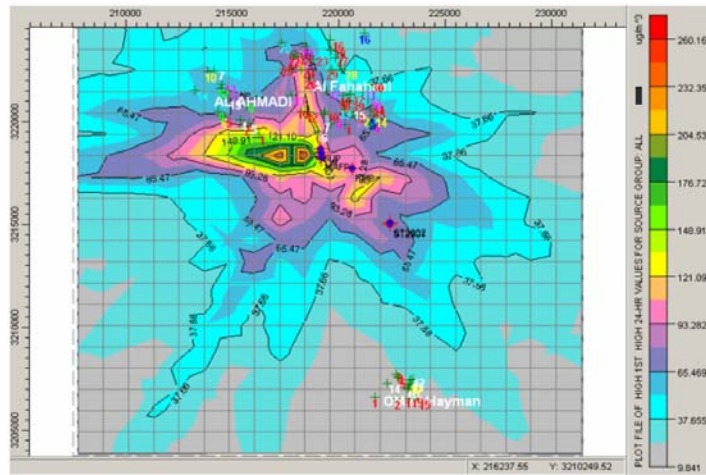


Fig. 10: Isopleth showing the daily maximum average ground Level VOC concentration

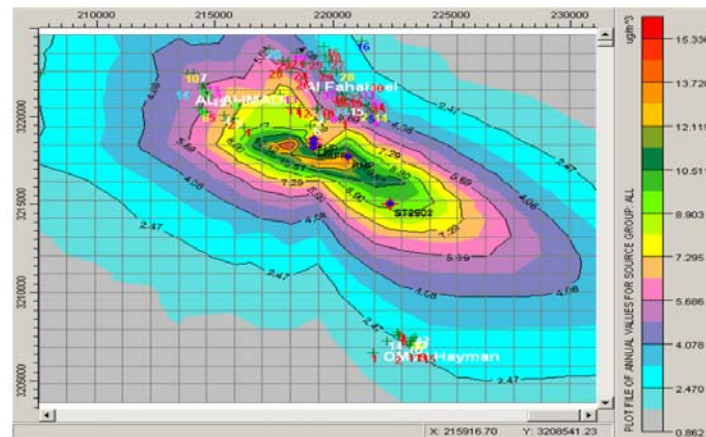


Fig. 11: Isopleth showing the annual maximum average ground Level VOC concentration



Table 4: Maximum concentrations of VOC predicted per 3HR and daily for each month of year 2006

	1st 3HR		Direction	Distance		1st Daily		Direction	Distance	
	( $\mu\text{g m}^{-3}$ )	ppm		(km)	Date	( $\mu\text{g m}^{-3}$ )	ppm		(km)	Date
Jan	370.27	0.19	NE	1.6	05/01/2006 24:00	81.85	0.04	NW	1.3	11/01/2006 24:00
Feb	936.52	0.47	NW	1.9	14/02/2006 06:00	260.16	0.13	NW	0.8	14/02/2006 24:00
March	707.11	0.36	SW	2.1	30/03/2006 24:00	98.26	0.05	SW	2.1	30/03/2006 24:00
April	624.37	0.31	NW	1.9	11/04/2006 06:00	127.75	0.06	NW	2.3	20/04/2006 24:00
May	533.76	0.27	NW	1.2	26/05/2006 24:00	109.07	0.05	SE	1.1	26/05/2006 24:00
June	322.28	0.16	NW	1.4	06/06/2006 24:00	54.76	0.03	NE	1.9	05/06/2006 24:00
July	323.42	0.16	SE	2.9	21/07/2006 21:00	66.91	0.03	SE	1.1	05/07/2006 24:00
Aug	497.46	0.25	NW	1.3	27/08/2006 24:00	124.36	0.06	NW	1.2	27/08/2006 24:00
Sep	774.62	0.39	NE	1.6	08/09/2006 03:00	121.79	0.06	SE	2.9	13/09/2006 24:00
Oct	362.55	0.18	NW	2.4	08/10/2006 24:00	71.93	0.04	NW	4.4	21/10/2006 24:00
Nov	328.74	0.17	SE	1.8	11/11/2006 24:00	56.44	0.03	SE	1.9	11/11/2006 24:00
Dec	290.81	0.15	NW	1.4	06/12/2006 03:00	62.18	0.03	NW	2.9	05/12/2006 24:00

Table 5: Influence of different flares on predicted first maximum ground level concentration of VOC

	Month of 1st max Conc of VOC 3 HRS
Real Situation	Feb
Al-Ahmadi refinery	Feb
Al-Shuiba refinery	Feb
MAFP	Feb
FUP	Nov
RMP	Dec
OR	Sep
ST29_01	Feb
ST29_02	Feb

Table 6: 50 maximum concentration of VOC predicted hourly from Al-Ahmadi and Al-Shuiba Refineries

The maximum 50 1-HR average concentration N values of VOC for source group: both Ahmadi and Shuiba refineries						The maximum 50 1-HR average concentration N values of VOC for source group: both Ahmadi and Shuiba refineries				
( $\mu\text{g m}^{-3}$ )	ppm	Date (YYMMDDHH)	Distance (km)	Direction	Rank	( $\mu\text{g m}^{-3}$ )	ppm	Date (YYMMDDHH)	Distance (km)	Direction
936.51	0.47	06021406	1.9	NW	26	567.95	0.29	06021406	4.3	NW
897.75	0.45	06021406	0.8	NW	27	567.64	0.29	06042024	2.5	NW
774.61	0.39	06090803	1.6	NE	28	567.33	0.29	06091706	0.4	NE
736.14	0.37	06090803	2.9	SE	29	565.48	0.28	06091721	3.2	NW
724.41	0.36	06021406	3.1	NW	30	562.59	0.28	06040103	1.8	SE
707.11	0.36	06033024	2.1	SW	31	560.34	0.28	06091721	3.3	NW
703.28	0.35	06091706	1.5	NE	32	558.68	0.28	06041921	3.9	NW
687.58	0.35	06090803	4.1	SE	33	547.49	0.28	06090921	3.3	NE
682.63	0.34	06021403	0.8	NW	34	540.92	0.27	06041921	4.3	NW
682.06	0.34	06090124	2.1	NE	35	539.58	0.27	06093024	2	NE
667.11	0.34	06090806	2.2	SW	36	534.94	0.27	06091921	2.2	NE
664.06	0.33	06090124	2.2	NE	37	533.76	0.27	06052624	1.3	NW
656.12	0.33	06030406	2.8	SW	38	529.63	0.27	06090321	1.5	NE
644.48	0.32	06090921	2.7	NE	39	529.36	0.27	06030406	4.4	SW
634.24	0.32	06091221	1.2	NW	40	528.93	0.27	06021224	2.5	SE
632.29	0.32	06090803	0.4	NE	41	528.37	0.27	06091924	1.6	NE
631.92	0.32	06090306	1.1	SE	42	528.24	0.27	06091224	3	NE
624.37	0.31	06041106	2	NW	43	522.01	0.26	06090124	1.3	NW
613.16	0.31	06090306	2.5	SE	44	520.96	0.26	06090921	3.3	NE
598.17	0.3	06021403	2	NW	45	520.01	0.26	06090921	3.2	NE
595.56	0.3	06042024	2.3	NW	46	519.55	0.26	06091303	2.9	SE
590.11	0.3	06090124	2.2	NE	47	517.69	0.26	06091221	2.1	NE
576.38	0.29	06031406	2	SE	48	514.99	0.26	06091224	1.9	NE
575.56	0.29	06041724	1	SE	49	508.80	0.26	06090803	5.2	SE
573.82	0.29	06030406	1.2	SW	50	499.52	0.25	06090724	1.8	SE

Table 7: Highly influenced discrete receptors from VOC ground level concentrations for 3 h

Area	Type	Distance (km)	Direction	1st-3HR Conc.	ppm	Date (YYMMDDHH)
Al Fahaheel	Clinic	2.1	NE	682.06	0.343	06090124
Al Fahaheel	Primary school	2.2	NE	664.06	0.334	06090124
Al Fahaheel	Mall	2.7	NE	644.48	0.324	06090921
Al Fahaheel	Ministry	1.2	NW	634.24	0.319	06091221
Al Fahaheel	Primary school	2.5	NW	567.64	0.285	06042024
Al Ahmadi	Secondary school	3.2	NW	565.48	0.284	06091721
Al Fahaheel	Nursery	3.3	NE	547.47	0.275	06090921
Al Ahmadi	Mosque	4.3	NW	540.92	0.272	06041921
Al Fahaheel	Secondary school	1.9	NE	539.58	0.271	06093024
Al Fahaheel	Primary school	3.4	NE	520.96	0.262	06090921
Al Fahaheel	Mall	3.2	NE	520.01	0.261	06090921
Al Fahaheel	Homes	1.6	NE	489.86	0.246	06091921
Al Fahaheel	Hotel	2.8	NE	488.66	0.246	06090921

### CONCLUSION

- Al shuiba refinery contributes 99.63% of the total ground level concentration of SO<sub>2</sub> indicating very little impact due to Al-Ahmadi refinery. The average sulfur content in the gas flared from Al-Ahmadi 0.88% by V hydrogen sulfide H<sub>2</sub>S while the average sulfur content of flared gas from Al-Shuiba refinery is 6.1% V H<sub>2</sub>S showing the predominate influence on ground level concentration
- AL-Ahmadi refinery contributes 99.84% of the total ground level concentration of NMHC indicating very little impact due to Al-Shuiba refinery. The average content of NMHC in the gas flared from Al-Ahmadi refinery is 30% V, while the average NMHC content of flared gas from Al-Shuiba refinery is 10.3% V reflecting the predominate influence on ground level concentration
- In Kuwait the prevailing wind is almost all the time from north west NW confirming the highest ground level SO<sub>2</sub> concentration to be the south east SE from the predominate source
- All the maximum values were in cold months of winter having low temperature and low inversion layer controlling adversely the dispersion phenomena. In the month of September there were high emissions of VOCs from Al-Ahmadi refinery producing high level concentrations at various discrete receptors
- About 50 maximum highest values of SO<sub>2</sub> exceeded EPA standards at locations within 4-5 km around sources.
- From the 50 maximum concentrations of VOC predicted per 3HR we found only 14 values exceed EPA standards for 6.00-9.00 h while all other high concentration values were almost from 21.00-24.00 h

- Model validation with actual ground level measurements can authenticate the results and application of model including all emission sources can provide exact gas concentrations in and around the petroleum refineries

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