Review of Photochemical Smog Pollution in Jakarta Metropolitan, Indonesia

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Abstract: An analysis of photochemical smog pollution in Jakarta was attempted using data from the existing air quality monitoring stations. Ground-level ozone potential is high in Jakarta due to the high traffic emissions of ozone precursors and the favorable surface meteorological conditions. Despite the frequent missing data during the 1996-1999 monitoring, which resulted in lower ozone values, ozone episodes were significantly recorded in 1997-1998. The number of hours on which ozone concentrations exceed the 1-hour standard (100 ppb) at an ambient station located in Kelapa Gading (10 km northeast of city center) was 186 hrs in 1997 and 571 hrs in 1998. El Niño phenomenon in 1997-1998 had affected the local meteorology leading to more favorable conditions for photochemical production of ozone. The annual ozone averages in ambient stations located off the city center have exceeded the 1-year standard limit (15 ppb). Although the annual average and 95-percentile values indicated an increasing trend from 1996 to 1998, the trend remains to be seen in the future as more complete data could be expected from the new monitoring system. The number of hours on which ozone exceed the 1-hour standard and the annual average tend to be increasing since 2001 to 2002 in all 3 newly operated stations. The seasonal variations of ozone indicate that ozone level is highest in the dry season (September-November) and is lowest in the wet season (December-March). Correlation between ozone level and meteorological attributes (solar radiation, relative humidity and temperature) was significant at 0.01 confidence level. The diurnal cycle of ozone and its precursors is clearly shown and is typical for polluted urban areas. Improvement of the database of air quality monitoring is very critical for Jakarta. Through better database management, the development and monitoring of cost-effective air pollution control strategy can be made.

Key words: Ozone, Ozone Precursors, Traffic Emissions, Meteorological Conditions

INTRODUCTION

$$O_3 + NO \rightarrow NO_2 + O_2 \tag{3}$$

Photochemical smog is a condition that develops when primary pollutants (oxides of nitrogen-NO_x and volatile organic compounds-VOC created from burning of fossil fuels and biomass) interact in the presence of sunlight to produce a mixture of hazardous secondary pollutants [1]. The major constituent of photochemical smog is ground-level ozone (O₃), which is not emitted directly into the atmosphere but formed as the product of photochemical reactions of precursors, NO_x and VOC [2]. Basic photochemical cycle of NO, NO₂ and O₃ involves the following reactions: when NO and NO₂ are present in sunlight, ozone formation occurs as a result of the photolysis of NO₂ at wavelength < 424 nm,

$$NO_2 + hv \rightarrow NO + O \tag{1}$$

$$O + O_2 + M \rightarrow O_3 + M \tag{2}$$

where, M represents N_2 or O_2 or another third molecule that absorbs the excess energy and thereby stabilize the O_3 molecule. There are no other significant sources of O_3 in the atmosphere other than reaction (2). Once formed, O_3 reacts with NO to regenerate NO₂. Development of photochemical smog is typically associated with specific climatic conditions. Local meteorological factors that influence the formation of photochemical smog include solar radiation, cloud cover, temperature and precipitation. Other factors such as wind speed, mixing height and topography determine the dispersion of O_3 in the boundary layer. High temperatures are frequently associated with high pressure, stagnant conditions that lead to suppressed vertical mixing and increased O_3 levels.

The database of ground-level ozone observations for urban and suburban areas indicates that at most rural sites, ozone concentrations have been found to vary over a diurnal cycle with a minimum in the early morning hours before dawn and a maximum in the late afternoon [2]. This pattern results from daytime photochemical production or intrusion of ozone from upper layer to the boundary layer, combined with ozone loss by dry deposition and reaction with nitric acid (NO) at night, when photochemical production ceases and vertical transport is inhibited by a temperature inversion. In urban areas the nocturnal minimum in

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thor: Dollaris R. Suhadi, Department of Environmental Management, Faculty of Environmental Studies, Universiti Putra Malaysia, 43400 UPM Serdang, Selangor, Malaysia Tel: +60-12-2641 E-mail: DSuhadi@ssafara.net ozone can be quite evident because of the rapid reaction between O_3 and NO.

This study reviews the status of photochemical smog pollution in Jakarta Metropolitan during a period of 7 years, 1996-2002 with focus on ozone and NO_x pollutants. Average monthly and diurnal concentrations of ozone and its precursor are also evaluated in relation to variations in meteorological conditions.

STUDY AREA AND DATA COLLECTION

Jakarta is the capital city of Indonesia located in the northern part of Java Island, at 106° East and 6° South. The area of Jakarta is approximated at 664 square kilometers with flat topography, close to the shore and has an average elevation of 7 meter above sea level. Jakarta is part of the greater Jabodetabek (Jakarta, Bogor, Depok, Tangerang and Bekasi); bordering with Tangerang in the west, with Bogor and Depok in the south and Bekasi in the east. The total population of Jakarta was estimated at 9 million in 2002, whereas total population of Jabodetabek exceeded 22 million people in 2002. In the meantime, total commuters from Bodetabek to Jakarta today exceed 3 million everyday.

Climate in Indonesia is largely influenced by monsoon systems, both Asia and Australian winter monsoons. During wet months (October-March), wind direction extends from west to southwest transporting moist air from Asia continent; while during dry months (April-September) wind direction is from north up to northeast which transports dry air from southern hemisphere. Besides the monsoon systems, the climate is also affected by global circulation occurring on a time scale of several years known as El Niño/Southern Oscillation (ENSO). In 1997, extreme meteorological values were recorded in Indonesia and the region as a result of ENSO phenomenon. The climate in Jakarta is very hot and humid. Because Jakarta lies so close to the equator, the solar heating during the day and the earth cooling during the night may produce local land-sea breeze. The winds are generally weak. Calm conditions often prevail at night. In the afternoon, winds are quite moderate because of the effects of local winds that modify the monsoon winds.

Jakarta has experienced serious air pollution problem, which is largely contributed from traffic sources accounting for approximately 70% of the total emissions. The vehicular emissions in Jakarta contributed 69% of total NOx emissions, whereas CO and HC emissions from vehicular sources amount to 564,000 ton and 98,000 ton per year respectively [3].

The data used for this study were collected from the Local Environmental Management Agency (BPLHD) of DKI Jakarta and the National Environmental Management Center (EMC). During a period of 1996-1999, data was supplied from an air quality monitoring network which comprises of 6 stations that monitor NO_x , CO, PM_{10} , NMHC, THC, O_3 and SO_2 parameters using designated absorption methods. In 2000, the prevailing stations were overhauled and

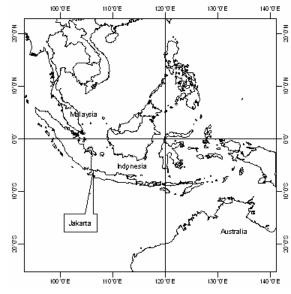


Fig. 1: Map of Indonesia

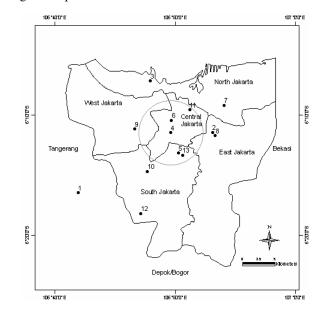


Fig. 2: The Area of Jakarta and Air Quality Monitoring Locations with Station No.4 Being Located in the City Center

replaced with the new monitoring stations, which are operated automatically from the central computers at the BPLHD. Two 2 new stations started operating in 2001 while the remaining 3 stations started operating in the following year. These new stations are equipped with measurement analyzers to monitor NO/ NO₂, SO₂, CO, O₃ and PM₁₀. Specifically, hourly data of O₃, NO, NO₂, CH₄, NMHC and CO from both monitoring network were used as the basis for this analysis. Meteorological data was compiled from 2 major stations, namely Cengkareng International Airport and Curug Airstrip. In addition, *In Situ* meteorological data is also recorded at air quality monitoring

Name	Location	96	97	98	99	00	01	02
EMC (1)	Serpong, southwest of Jakarta. No nearby specific emission sources.	•	•	•	•	•		
Pulogadung (2)	East Jakarta, industrial park.	•	•	•	•			
Pluit (3)	North Jakarta, residential area.	•	•	•	•			
Thamrin (4)	Central Jakarta, at curbside near Thamrin roundabout.	•	•	•	•			
Casablanca (5)	Central Jakarta, commercial and residential area.	•	•	•	•			
Gambir (6)	Central Jakarta, at curbside.	•	•	•				
Kelapa Gading (7)	North Jakarta, residential area.		•					
Pulogadung* (8)	East Jakarta, industrial park.						•	•
Kebon Jeruk* (9)	West Jakarta, commercial and residential area.							•
Senayan* (10)	South Jakarta, commercial area							•
Kemayoran* (11)	Central Jakarta, commercial area							•
Pondok Indah* (12)	South Jakarta, commercial and residential area.							•
BPLHD* (13)	South/central Jakarta, commercial and residential area						•	•

Table 1: Monitoring Locations and Year of Data Recorded

*New monitoring stations (2000-present) Source: BPLHD DKI Jakarta, 2001

Table 2: Emission Standards for O3 and NO2 for Jakarta

	1 hour	24 hour	1 year
NO ₂	$400 \ \mu g \ m^{-3}(200 \ ppb)$	92.5 µg m ⁻³ (50 ppb)	60 µg m ⁻³ (30 ppb)
O_3	$200 \ \mu g \ m^{-3}(100 \ ppb)$	-	$30 \mu\text{g/m}^{-3}(15 \text{ ppb})$
Sourc	e: BPLHD DKI Jakar	ta, 2001	

stations, which are equipped with basic meteorological sensors at 10 m height above the ground.

RESULTS AND DISCUSSION

Status of Ozone Pollution: Table 3 through 6 present a summary of ozone and NO₂ pollution levels for Jakarta during the four-year monitoring period from 1996 to 1999 and the two-year period from 2001 to 2002. In general, the highest ozone was found at ambient station number 7 (Kelapa Gading) located approximately 10 km northeast of the city center. The highest hourly maximum level was recorded at 964 ppb at this station. Although this figure looks highly suspicious as if someone was doing a range check on the ozone instrument, it was noted that the total number of hours on which ozone exceed the standard were quite high reaching 186 hrs and 570 hrs in 1997 and 1998 respectively. Station 7 also held the highest average hours per year of ozone exceeding the standard, followed with station 2 and 3. Where ozone data are available in 1996-1999, 1998 was characterized by the highest ozone values (average and 95 percentile) at station 2. At stations 3 and 7, the highest ozone values were recorded in 1997. The discussion on the relation between ozone episodes and the El Niño phenomenon during 1997-1998 is discussed later in the next section. Lower ozone was detected at curbside station 4 and ambient station 5 of which these stations are located closest to the city center. The major factor that caused low ozone values at these stations was due to destruction of ozone by NO. Since most of the NO_x is emitted in the form of NO, it can be expected that with high emission of NO from traffic in curbside station, ozone level becomes lower (reaction 3). The processes of destruction and formation of ozone in a large urban

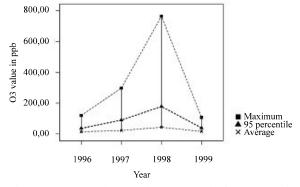


Fig. 3:O₃ Annual Trend Averaged for 3 Stations (Station 2, 3 and 7)

area such as Jakarta should be competing at any location. Over the city, ozone level is found to be relatively high except at the very heavy traffic city center and curbside where the ozone destruction by NO is very apparent. However, it is very difficult to make further assessment as to the transport of ozone and its precursors over the larger area of Jakarta (beyond the 15 km perimeter of the city center) because all stations are located within.

Frequent missing data and limitation of the current two-year monitoring (2001-2002) make the assessment of ozone annual trend very premature. The initial ozone trend averaged for 3 ambient stations (station 2, 3 and 7) indicated an increase of ozone values from 1996 to 1999; but a substantial drop was noted from 1998 to 1999 (Fig. 3). The substantial drop of ozone level was significantly contributed from station 7. Thus, besides the low percentage of data completeness, other factors such as the change in meteorological conditions and in emission of O₃ precursors in the upwind area of station 7 may also play a role in causing the shift of ozone annual trend. In addition, the annual ozone averages in ambient stations located off the city center (station 2, 3 and 7) indicated exceeding the 1-year standard limit (15 ppb). Ozone fluctuation in terms of standard deviation was quite high showing variations in daily and monthly time scales.

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	Number	of hours o	n which O3	exceeds th	e standard		Hourly data count (% data completeness)					
Station	1996	1997	1998	1999	2001	2002	1996	1997	1998	1999	2001	2002
Pulogadung (2) ^a	11	2	268	18			44%	74%	20%	17%		
Pluit (3) ^b	5	42	3	1			83%	75%	22%	27%		
Thamrin (4) ^c	1	0	*	0			94%	58%	-	16%		
Casablanca (5) ^d	**	**	0	0			-	-	30%	95%		
Gambir (6)	**	**	**	**			-	-	-	-		
Kelapa Gading (7) ^e	**	186	567	2			-	23%	58%	68%		
Pulogadung (8)					67	72					91%	85%
Kebon Jeruk (9) ^f					5	174					17%	84%
Senayan (10) ^g					44	190					28%	97%

^aMissing data: June-November 1996; May-November 1998; January-February and April-July and November 1999

^bMissing data: January and November 1997; July-December 1998; January-April and October-December 1999

^cMissing data: April-July and December 1997; January-November 1999

^dMissing data: January-August 1998

^eMissing data: January-June and November-December 1997; September-December 1998; February-March, July, September and December 999

^fMeasurement started in September 2001

^gMeasurement started in August 2001. Missing data: December 2001

*No data available

**No measurement

Table 4: Average Number of Hours per Year on which Ozone Exceeds the Standard (100 ppb)

	Station r	no.(1996-1999)					Station no. (2001-2002)				
Pollution exceeding											
the standard	2	3	4	5	6	7	8	9	10		
Total hours	13	12	1	0	-	573	139	179	234		
Average hours per year*	11	7.58	0.66	0	-	454.62	78.91	176.94	188.89		

* Calculated based on available database

Station 2, station 3 and station 4 based on the two-year data (1996-1997)

Station 5 based on the one-year data (1999)

Station7 based on the two-year data (1998-1999)

The assessment of ozone level in the background area (station 1) was not done due to unavailability of ozone data. It can be expected that O_3 concentration in this background area is fairly high. Rural environments tend to be characterized by relatively high VOC-to-NO_x ratios because of rapid removal of NO_x from distant sources compared to that of VOC [2]. In general, increasing VOC concentrations means more ozone. Transport of ozone from a very photochemically active source region (Jakarta urban center) is also a possible candidate of potential source of ozone enhancement in the background area.

For the current air quality monitoring which uses telemetric equipment, the number of hours on which ozone exceeds the standard indicated a steady increase since 2001 to 2002 in all three new stations. Although the trend remains to be seen in the future, the potential for ozone episodes in Jakarta metropolitan and its surrounding area of Bodetabek is very high. Given the fact that no counter measures are put in place to reduce air pollution at this time, an increasing trend of ground ozone pollution is very likely in the immediate future, as the emission of ozone precursors from traffic source remains high.

Ozone Seasonal Variations and Related Meteorological Conditions: Table 7 presents the total number of hours on which ozone exceeds 100 ppb at stations 2, 3 and 7. High ozone levels were generally recorded in May-September (dry months). Maximum monthly ozone averages at station 2 over 4 years were 47.28±11.49 ppb (in June 1996), 21.11±11.44 ppb (in May 1997), 148.92±201.02 ppb (in December 1998) and 72.61±14.3 ppb (in July 1999). At station 3, maximum monthly average was recorded in July 1996, July 1997, January 1998 and June 1999 with values of 22.78±19.32 ppb, 25.28±25.72 ppb, 19.83±21.60 ppb and 24.91±18.06 ppb respectively. Monitoring at station 7 showed highest monthly average in July 1997 (166.03±217.98 ppb), in September 1998 (77.03±53.23 ppb) and in September 1999 (11.46±7.12 ppb). Overall, standard deviation is high showing high fluctuation of ozone in monthly time scale. Figure 4 shows monthly average ozone value at station 8 in 2001-2002. The selection of this station is primarily due to relatively higher percentage of data completeness for two consecutive years. Although the annual cycle is not clearly shown, maximum ozone value is generally recorded during the September-November period. From December to March (wet season), ozone levels are minimum. Figure 4 also illustrates a constant increase of ozone concentration over two years from 2001 to 2002.

Ozone behavior near the surface may be attributed to local meteorological conditions [4]. During the period of 1997-1999, daily mean temperature was increasing

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Table 5: NO₂ Pollution Level Exceeding the one hour and 24 hour Air Quality Standard (200 ppb and 50 ppb)

	Number	Number of exceedances												
	1996		1997		1998		1999		2001		2002			
Station	Hours	Days	Hours	Days	Hours	Days	Hours	Days	Hours	Days	Hours			
Pulogadung (2) ^a	0	0	1	3	0	6	0	0						
Pluit (3) ^b	1	0	1251	125	295	96	8	0						
Thamrin (4) ^c	24	26	1372	269	427	119	3	1						
Casablanca (5) ^d	1	5	6	21	18	10	2	1						
Gambir (6) ^e	258	48	30	99	5	147	*	*						
Kelapa Gading (7)	**	**	0	3	0	0	*	*						
Pulogadung (8) ^f									0	0	0	0		
Kebon Jeruk (9) ^g									0	0	80	62		
Senayan (10) ^h									*	*	18	39		

Hourly data count (% data completeness)

	-	-				
	1996	1997	1998	1999	2001	2002
Pulogadung (2) ^a	 93%	84%	18%	80%		
Pluit (3) ^b	86%	79%	42%	32%		
Thamrin (4) ^c	97%	72%	39%	70%		
Casablanca $(5)^d$	87%	87%	83%	95%		
Gambir (6) ^e	54%	83%	100%	-		
Kelapa Gading (7)	-	15%	62%	-		
Pulogadung (8) ^f					65%	82%
Kebon Jeruk (9) ^g					29%	78%
Senayan (10) ^h					-	77%

^aMissing data: January-February and May-November 1998

^bMissing data: January and December 1997; January-March and May-June 1998; January-June and October-December 1999

^cMissing data: January-May and October 1998; April-May 1999

^dMissing data: May 1996; April 1997; June-July 1998

^eMissing data: February-March, May and December 1996; April 1997 ^fMissing data: September-December 2002

^gMeasurement started in September 2001. Missing data: August-September 2002

^hNo data available in 2001. Missing data: January-February 2002

*No data available

**No measurement

Table 6: Summary of Annual Ozone Values (1996-1999, 2001-2002)

		Value in ppb					
Station	Year	Average ± 1 SD	Minimum	Maximum	Median	95-percentile	Data count
Pulogadung (2)	1996	17.52±13.63	1	129	14	43	3836
	1997	14.4±13.16	0	161	11	40	6455
	1998	67.36±142.33	0	993	16	363.95	1762
	1999	46.79±20.83	0	122	40	85	1505
Pluit (3)	1996	10.9±11.43	0	115	7	32	7320
	1997	17.77±18.57	0	228	11	53	6564
	1998	12.79±13.45	0	148	9	38	1965
	1999	12.09±14.38	0	101	6	39	2386
Thamrin (4)	1996	9.66±9.91	0	104	7	30	8254
	1997	5.1±5.45	0	78	4	14	5050
	1999	8.08±0.58	0	88	3	31	1352
Casablanca (5)	1998	7.43±11.87	0	88	2	38	2565
	1999	7.51±10.59	0	76	2	31	8326
Kelapa Gading (7)	1997	57.27±122.73	0	964	20.6	366.35	2007
	1998	46.53±82.09	0	923.6	17.7	167.56	5097
	1999	8.52±9.74	0	105.2	5.7	26.6	5944
Pulogadung (8)	2001	16.83±21.47	0	280.25	7	60.05	7967
	2002	25.79±21.12	3.64	279.43	14.57	69.49	7464
Kebon Jeruk (9)	2001	16.45±17.54	0.98	139.60	8.28	61.09	1478
	2002	28.95±25.35	0.26	217.15	17.14	82.97	7384
Senayan (10)	2001	19.14±24.24	0	151.71	7.22	73.60	2475
	2002	26.82±24.19	0.39	203.11	17.98	80.74	8377

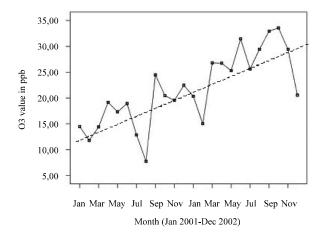


Fig. 4: Monthly Average O₃ Value at Station 8 (2001-2002)

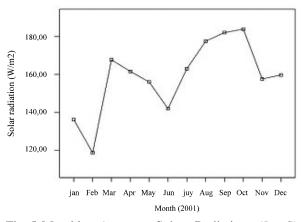


Fig. 5: Monthly Average Solar Radiation (*In Situ* Station 8), 2001

in 1997 and continued to increase towards 1998. Lowest humidity was recorded in October 1997 with 75% of relative humidity. Higher mean temperature and lower relative humidity in 1997 were associated with the El Niño phenomenon in that year. Particularly, strong El Niño extended from July 1997 to April 1998. In 1998 and 1999, the meteorological conditions returned to their normal averages (calculated based on a minimum 10 year meteorological observation).

Wind direction extends mostly from west to southwest during November-March, except in January where wind direction is from west to north. Calm conditions significantly prevail in the months of May through October with frequency ranging from 18 to 26%. In addition, the NE wind balances the calm winds during that period, specifically in the months of July through Sept. The calm conditions are in favor of ozone enhancements in Jakarta during May-October (dry months). Local biomass burning is often active in dry months, especially towards the end of dry season, potentially contributing to maximum ozone.

In addition, *In Situ* meteorological observation at station 8 in 2001 Fig. 5 to 7 may explain the association

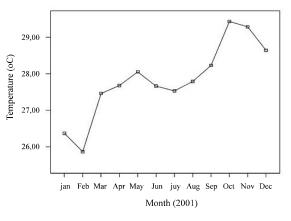


Fig. 6: Monthly Average Temperature (*In Situ* Station 8), 2001

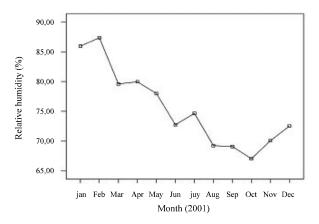


Fig. 7: Monthly Average Relative Humidity (*In Situ* Station 8), 2001

of high ozone levels Fig. 4 with warm temperatures, low humidity and high solar radiation. The local land-sea breeze may potentially influence ozone behavior in the area. Analysis of correlation using the two-year database at station 8 indicated that the correlation between ozone and solar radiation, between ozone and temperature and between ozone and humidity was all significant at the 0.01 level (two-tailed student t test) with the correlation coefficient (r) of 0.64, 0.67 and 0.63, respectively.

Surface meteorological observation indicates that the southerly to NE wind predominates over the period of May-October (dry season) transporting the ozoneenriched air masses towards neighboring Southeast Asian countries. Zhang *et al.* related the high ozone in the dry season in Bangkok to, among others, the regional transport of O_3 and precursors from the Asian continent associated with the NE monsoon.

Ozone Diurnal Cycle: Diurnal variations of ozone and its precursors for the selected stations were analyzed. Stations 8, 9 and 10 were selected for ozone diurnal analysis as they have relatively less missing data. However, the new monitoring system does not measure

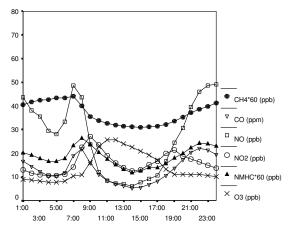


Fig. 8: Daily Average Concentration of Pollutants During the Dec-Mar Period at Station 2 In 1996

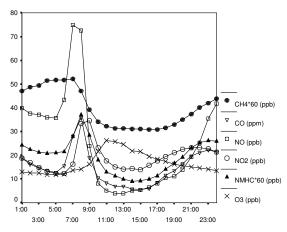


Fig. 9: Daily Average Concentration of Pollutants During the Apr-Aug Period at Station 2 in 1996

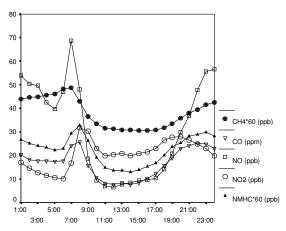


Fig. 10: Daily Average Concentration of Pollutants During the Sep-Nov Period at Station 2 in 1996

 CH_4 and NMHC. Thus, for the analysis of ozone precursors (NO, NO₂, CO, CH_4 , NMHC and O₃), the 1996 data from stations 2, 3 and 4 were used.

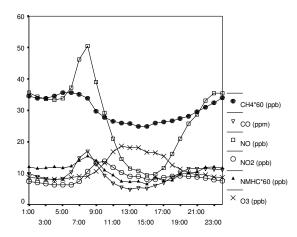


Fig.11: Daily Average Concentration of Pollutants During the Dec-Mar Period at Station 3 in 1996

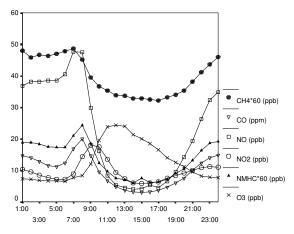


Fig.12: Daily Average Concentration of Pollutants During the Apr-Aug Period at Station 3 in 1996

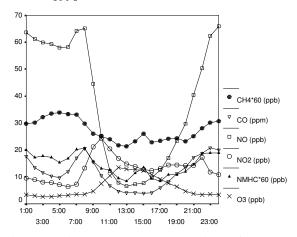


Fig. 13: Daily Average Concentration of Pollutants During the Sep-Nov Period at Station 3 in 1996

The analysis was done for each seasonal period, wet season (Dec-Mar), dry season (Sep-Nov) and transition period (Apr-Aug).

	1996			1997			1998			1999	
	No.2	No.3	No.2	No.3	No.7	No.2	No.3	No.7	No.2	No.3	No.7
Jan	0	0	0	0	-	0	0	0	-	-	0
Feb	0	0	0	0	-	0	0	0	-	-	-
Mar	0	0	0	0	-	2	0	62	0	-	0
Apr	0	0	0	0	-	0	0	42	-	-	0
May	5	0	0	7	-	0	0	178	-	0	0
Jun	6	1	0	8	0	-	2	68	-	1	2
Jul	-	4	0	19	171	4	1	41	1	0	0
Aug	-	0	0	6	0	-	-	92	9	0	0
Sept	-	0	2	2	1	-	-	84	6	0	0
Oct	-	0	0	0	10	-	-	-	1	-	0
Nov	-	0	0	-	4	-	-	-	-	-	0
Dec	0	0	0	-	0	262	-	0	1	-	0

Table 7: Number of Hours Exceeding 100 ppb at Stations 2, 3, and 7 (1996-1999)

Figure 8-13 presents the diurnal variation of pollutants during 1-year monitoring in 1996. The diurnal cycle of pollutants is typically shown for polluted urban areas. Indeed, this pattern is almost similar to the diurnal pattern of photochemical pollutants in Bangkok [5].

When the morning rush hours begin at around 7.00 a.m. to 8.00 a.m.; CO, NO, NMHC and CH₄ also reaches its peak at around 8.00 a.m. due to emissions from traffic. NO peaks at around 7.00 a.m., while NO₂ peaks at around 9.00 a.m. (about two hours after NO). NO concentration drops fast after 8.00 a.m. (after sunrise) as O3 increases reaching its highest at around 11.00-13.00. CO, NMHC, CH_4 and NO_2 also drop when O_3 reaches its maximum during the afternoon hours. The reduction of these pollutant concentrations should be caused by the photochemical reactions that consume the pollutants such as reactions of pollutants with OH and HO₂ radicals and the pollutant dispersion factors such as mixing height and wind speed. As the boundary layer increases gradually due to convective heating during noon hours, the mixing of air in the lower heights with air in the higher heights causes pollutants to be ventilated upwards [5]. The local wind may also contribute to the pollutant dilution.

In the afternoon towards evening, the pollutant concentrations increase steadily as the traffic peaks again but the photochemical production of O_3 ceases due to a minimum-to-zero solar radiation. As the traffic subsides, the concentrations of NO, NO₂, NMHC and CO decrease slowly during night hours until before dawn. Interestingly, CH₄ continues to increase slightly in the early hours, which may be contributed from other sources than road transport, i.e. natural, fossil-fuel related and biospheric carbon sources.

In urban centers, when NO_x predominates over VOC, the OH-NO₂ reaction will remove OH radicals retarding the further O₃ production. Because OH reacts more rapidly with NO₂ than with VOCs, NO_x tends to be removed faster than VOCs. In the absence of fresh NO_x emissions, NO_x is depleted more rapidly leading to increasing ratio of VOC to NO_x. As a result, OH reacts preferentially with VOCs to keep the ozone-forming cycle going [1]. This chemistry plays an important role in the temporal variations of ozone. However, the ratio of VOC to NO_x concentration was not evaluated in this study due to limited VOCs data.

Pollutant daily maximum and average concentrations at station 2 are generally higher than at station 3 for all seasonal periods. Daily average value of NO, NO₂, NMHC, CH₄ and CO at station 2 during the Dec-Mar period are 26.94±15.65 ppb (average±1 SD). 15.94±4.47 ppb, 1114.93±250.5 ppb, 2213±287.28 ppb and 13.11±5.54 ppm respectively; as compared to values at station 3 which are 27.4±12.02 ppb, 8.7±2.02 ppb, 631.15±145.07 ppb, 1810.8±234.87 ppb and 9.33±3.10 ppm respectively. Station 2 is located downwind of station 3 during wet season when westerly wind predominates transporting pollutant-enriched air masses to station 2. In the dry season, wind is mostly from NE bringing air masses from biomass burning in the nearby area (Bekasi) to station 2.

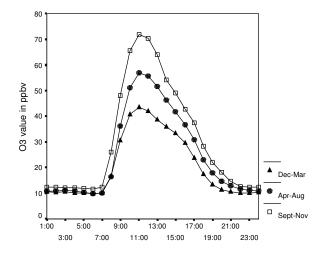


Fig. 14: Diurnal O₃ Concentration Averaged for 3 Stations During Wet Months (Dec-Mar), Transition Period (Apr-Aug) and Dry Months (Sept-Nov) in 2001-2002

This shall explain why higher pollutant concentrations are found in station 2 as compared to station 3, besides the fact that station 3 is located near to the coastal area. Daily average value of pollutants is generally higher in dry months. Figure 14 exhibits a comparison of diurnal O_3 concentration during wet season, transition period and dry season using the 2001-2002 data. It is clearly shown that the Sep-Nov period records the highest diurnal variation of O_3 concentration.

CONCLUSION

Photochemical smog pollution potential in Jakarta metropolitan area is high due to the high traffic emissions of ozone precursors and the favorable surface meteorological conditions, i.e. warm temperature, high solar radiation and calm wind conditions. Over the two-year period from 2001 to 2002, the ground-level ozone indicates an upward trend. The trend remains to be seen in the future as more data is made available. Despite the frequent missing database, which produces uncertainty in analysis, ozone episodes are potentially detected at downwind area off the city center. Further investigation is needed with regard to the transport of ozone and its precursors over the larger area of Jakarta and Bodetabek.

The seasonal variations of ozone illustrate the effect of local meteorological conditions and regional monsoons on ozone formation and accumulation. Ozone level is highest in the dry season (September-November) and is lowest in the wet season (December-March). Correlation between ozone level and meteorological attributes (solar radiation, relative humidity and temperature) is significant. The lower ozone level at curbside stations reflects the destruction of ozone by NO, which is emitted from traffic sources. The diurnal variations of ozone and its precursors are clearly demonstrated and are typical for polluted urban areas.

Improvement of the database of air quality monitoring is very critical for Jakarta. Areas of improvement shall include the development of national guidelines on the standard procedures of air quality monitoring including site location, instrumentation, quality assurance and statistical procedures for data analysis. The local government implementation plans on collection and analysis of data and inventorying emissions from pollution sources is a key to the identification and selection of emissions control measures. This will lead to development of efficient and cost-effective control strategy.

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