

# Fluctuation of Nighttime Ground Level Ozone Due to Artificial Light

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Abstract Ultraviolet (UV) radiation creates ground-level ozone  $(O_3)$  during the day and is subsequently eliminated by nitrogen oxides at night. However, excessive artificial light use, those results in light pollution, may interfere with the chemistry of ground-level O<sub>3</sub> at night by supplying enough energy to start that creation. Therefore, this study aims to identify the effect of artificial light on nighttime ground-level ozone production. Minute average O<sub>3</sub> and NO<sub>2</sub> concentrations with light illumination were measured in two study sites at the USM School of Civil Engineering and its Main Campus. Results of this study suggested that in low-light illumination conditions, no conversions between NO<sub>2</sub> to O<sub>3</sub> were governed due to low-light energy that was unable to break NO<sub>2</sub> bonds. Thus, there are no changes in O<sub>3</sub> concentration trends in the School of Civil Engineering. However, in high-light illumination conditions such as in USM Main Campus, O<sub>3</sub> fluctuated negatively with NO<sub>2</sub> concentrations, and potential conversions were governed.

**Keywords:** nighttime chemistry, ozone production, anthropogenic light

### 1. Introduction

Artificial light pollution and nighttime ground-level ozone are two distinct environmental issues that can have interconnected effects and influences. Nighttime groundlevel ozone (O<sub>3</sub>) has emerged as a significant concern in air pollution research due to its unique characteristics and potential health and environmental impacts. O<sub>3</sub> during daytime was focused compared to nighttime conditions (Awang et al., 2015; Shith et al., 2020; Stolz et al., 2020). Meanwhile, artificial light pollution can indirectly contribute to the formation and accumulation of groundlevel ozone through photochemical reactions. Artificial lighting sources emit light in various wavelengths, including blue and ultraviolet (UV) light. These wavelengths can react with certain air pollutants, such as nitrogen oxides (NO<sub>x</sub>) and volatile organic compounds (VOCs), triggering photochemical reactions that lead to the production of ground-level ozone (Shith et al., 2020).

Other studies have also extensively investigated fluctuations in  $O_3$  concentration at night (Awang & Ramli,

2017; Shith et al., 2020; Wang et al., 2022(a); Wang et al., 2022(b)). Awang and Ramli (2017) reported an interrelation between daytime and nighttime O<sub>3</sub> concentrations in Malaysia. Specifically, the enhanced nighttime depletion chemical removal in O3 concentrations influences the O<sub>3</sub> concentrations for the following day. Shith et al. (2020) detected low nighttime—NO titration during a haze event, indicating that O<sub>3</sub> from anthropogenic sources can stay in ambient air. Meanwhile, Wang et al. 2022(a) focus on understanding the diurnal pattern of  $O_3$ and its interactions with nighttime nitrate radical (NO<sub>3</sub>) and nitrous acid (HONO) in the atmosphere. However, Stark et al. (2011) revealed that this nighttime phenomenon is related to increasing the number of photons the atmosphere and directly enhancing  $O_3$ in photochemical reactions. Therefore, this study aims to identify the effect of artificial light on nighttime groundlevel ozone production.

### 2. Materials and Method

#### 2.1. Study Area

This study took place in two locations of Universiti Sains Malaysia, Penang, the Main Campus and the School of Civil Engineering (Table 1.). The location of the Main Campus of Universiti Sains Malaysia, Penang, is near major transportation routes, including highways and roads, which can result in significant vehicular traffic. Increased transportation activities, including private vehicles, public transportation, and logistics, can lead to emissions of pollutants such as nitrogen oxides (NO<sub>x</sub>), volatile organic compounds (VOCs), which are known as main O<sub>3</sub> precursors, and particulate matter (PM). Meanwhile, the School of Civil Engineering is located inside the Engineering Campus, which exposed Nibong Tebal, Penang, to vehicular emissions.

Table 1. Details of the selected study areas

	No.	Location	Coordinate
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1	Main Campus	5°21'22.5"N
		100°18'28.8"E
2	School of Civil	5°08'48.9"N
	Engineering	100°29'33.5"E

#### 2.2. Measurement and Instruments

The concentration levels of  $O_3$  and  $NO_2$  were monitored at two selected locations using the portable handheld Aeroqual S500 (Aeroqual Limited) monitors and six sensors' heads  $NO_2$  (range 0–1 ppm) and  $O_3$  (range 0–10 ppm). The relatively compact and lightweight Aeroqual S500 monitors were battery-powered at the selected TIs, interchanging the semiconductor and electrochemical sensor, permitting continuous monitoring of the range of selected gases at low mixing ratios (Lin et al., 2017). The instrument was placed nearest the artificial light located. Meanwhile, light Illumination in the selected locations was observed using HI97500 Portable lux meter (range:0.001-199.9 Klux). Figures 1 and 2 show the placement of the instrument with possible gas emissions emitted to the instrument.



Figure 1. The lobby of the School of Civil Engineering



Figure 2. Roadside at Main Campus

## 2.3. Nighttime Data Collection

Continuous data of selected variables are analysed according to the selected locations in this study. The average  $O_3$  concentration and NO concentration visualise the condition of the  $O_3$  reading and symbolise NO titration.

No imputation methods were used in this study, and any missing values discovered during data collection were omitted from the analysis. Time Series fluctuation was conducted to identify the fluctuation of ozone during the nighttime  $O_3$  concentration with existing artificial light. The monitoring duration was recorded per minute averaging data.

## 3. Result and Discussion

## 3.1. Nighttime O<sub>3</sub> and NO Concentration

Figure 3 illustrates the variation of O<sub>3</sub> and NO<sub>2</sub> concentrations and light illumination in the School of Civil Engineering (left) and USM Main Campus (right) during nighttime. For the monitoring in the School of Civil Engineering, freshly new emissions (from a car) were supplied, while no new emissions were supplied during the monitoring in USM Main Campus. The School of Civil Engineering result suggested that O<sub>3</sub> concentration is minimal during nighttime, while NO2 concentration ranges from around 18 ppb to 10 ppb before the new emission; the NO<sub>2</sub> concentration increased significantly as new emissions were released with maximum concentration recorded at 60 ppb. The light illumination at the School of Civil Engineering during the monitoring period is constantly recorded at 28 lux, significantly lower than at USM Main Campus, which is recorded at 64 lux. Conversely, O<sub>3</sub> and NO<sub>2</sub> concentrations in this study area were significantly higher even when no new emissions were supplied.

Studies by Elbayoumi et al. (2014) and Warmiski et al. (2018) elucidate that the chemistry of nighttime ozone differs from that of daytime ozone, with (R1) to (R4) being the most common reactions. However, the primary precursor of O<sub>3</sub>, NO<sub>x</sub>, continues to have the most impact on nighttime O<sub>3</sub> changes. According to Awang et al. (2015), reactions between NO and O<sub>3</sub> concentrations, also known as the NO titration process/reaction, control nighttime O<sub>3</sub> chemistry (R1). The high NO concentrations produced by automotive and industrial emissions stimulate (R1), causing  $O_3$  to be destroyed in the atmosphere. Furthermore, at night (R2), the combination between O<sub>3</sub> and  $NO_2$  can deplete nocturnal  $O_3$  concentrations by converting  $O_3$  to nitrate (NO<sub>3</sub>), which then interacts with  $NO_2$  to generate dinitrogen pentoxide ( $N_2O_5$ ) (R3). The N<sub>2</sub>O<sub>5</sub> concentration combines with water in the ambient air to generate nitric acid, which is then eliminated from the atmosphere as acid rain or precipitation (Seinfeld & Pandis, 2016).

$$NO + O_3 \rightarrow NO_2 + O_2 \tag{R1}$$

$$NO_2 + O_3 \rightarrow NO_3 + O_2 \tag{R2}$$

$$NO_3 + NO_2 + M \iff N_2O_5 + M$$
 (R3)

$$N_2O_5 + H_2O \rightarrow 2HNO_3$$
 (R4)

$$NO_3 + light (\lambda < 590 nm) \rightarrow NO_2 + O$$
 (R7)

Meanwhile, further research has shown that this nightly cleansing action is not as effective as expected because  $NO_3$  is being destroyed by the light reflected in the sky by outdoor lighting on the ground (Stark et al., 2011). In the presence of anthropogenic lights,  $NO_3$  is destroyed, and  $O_3$  concentrations increase through possible chemical reactions occurring at night, as in reactions (R1) and (R2). Reaction (R1) can occur during daytime; however,  $NO_3$  is quickly photolysed by daylight (R3), and  $NO_3$  and  $N_2O_5$  are heavily suppressed during the day.

The results observed in the School of Civil Engineering suggested that in low light illumination conditions, NO<sub>2</sub> will remain as NO<sub>2</sub> even if there are freshly emitted concentrations due to the low light energy available. Thus, it reflected the finding as the O<sub>3</sub> concentration was consistently at minimal concentrations at the monitoring site. Meanwhile, results show interesting trends in USM Main Campus as the light illumination in the monitoring area is relatively higher than in the School of Civil Engineering. The result illustrated that there are opposite trends shown by O<sub>3</sub> concentrations and NO<sub>2</sub> concentrations throughout the study period. This trend might suggest a conversion of NO<sub>2</sub> concentration into O<sub>3</sub> concentrations due to high illumination conditions in the area due to the fluctuation trends shown by O<sub>3</sub> concentrations.

## 4. Conclusion

This study aims to identify the effect of two distinct environmental issues: artificial light's influence on nighttime ground-level ozone production. Previous studies elucidated that there might be chances that artificial light with a specific wavelength and right intensity is potentially initiating nighttime O<sub>3</sub> formations. If these were possible, the issues regarding O<sub>3</sub> pollution would possess more threats from current concerns as their formation is not limited to daytime only but also can form during nighttime. Results of this study suggested that in low-light illumination conditions, no conversions between NO2 to O<sub>3</sub> were governed due to low-light energy that was unable to break NO<sub>2</sub> bonds. Thus, there are no changes in O<sub>3</sub> concentration trends in the School of Civil Engineering. However, in high-light illumination conditions such as USM Main Campus, O<sub>3</sub> fluctuated with NO<sub>2</sub> negatively, concentrations suggesting potential conversions were governed.

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**Figure 3.** Variation of O<sub>3</sub> and NO<sub>2</sub> concentration and light illumination in the School of Civil Engineering (left) and USM Main Campus (right)

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