

# **Characterization of ambient Ozone (O<sub>3</sub>) and its precursor compounds at a remote site of mid-Brahmaputra valley**

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## **CHAPTER-5**

### **CONCLUSIONS AND FUTURE SCOPE**

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# Chapter 5

## Conclusions and Future Scope

### 5.1: Conclusions

The following conclusions can be drawn from the investigation carried out on ambient O<sub>3</sub> and its precursors over the Brahmaputra Valley:

- The maximum concentrations of O<sub>3</sub>, O<sub>3</sub>*max* and NO<sub>x</sub> were found to be 19 ppb, 43 ppb and 7 ppb respectively during the year 2013-2014; 17 ppb, 41ppb and 7 ppb respectively during the year 2014-2015; 15 ppb, 85 ppb and 10 ppb respectively for the year 2015-2016 and 23.1 ppb, 79.7 ppb and 19.2 ppb during 2016-17 respectively.
- The minimum O<sub>3</sub>, O<sub>3</sub>*max* and NO<sub>x</sub> were found to be 1ppb, 1ppb and 2ppb respectively during the year 2013-2014; 1ppb, 11ppb and 1ppb during the year 2014-2015; 1 ppb, 2 ppb and 1ppb respectively during the year 2015-16 and 1 ppb, 3.8ppb and 2 ppb respectively for the year 2016-2017.
- The calculated OX values were found to be between 174 ppb and 2 ppb in 2013-2014; between 93 ppb and 2ppb in 2014-2015; between 85.4 ppb and 0.9 ppb in 2015-2016 and between 134.7 ppb and 1.3 ppb in 2016-2017.
- For the year 2013-14 a maximum of 26.6 ppb and 33.6 ppb and a minimum of 0.4 ppb and 0.4 ppb were found for NO and NO<sub>2</sub> respectively, while for the year 2014-15 a maximum of 19.4 and 42.3 ppb and a minimum of 0.11 and 0.1 ppb were found for NO and NO<sub>2</sub> respectively. For the year 2015-16 a maximum of 24.2 and 45.3 ppb and a minimum of 0.4 and 0.1 were found for NO and NO<sub>2</sub> respectively while for the year 2016-17 a maximum of 50.8 and 102.8 ppb and a minimum of 1.3 and 0.4 ppb were found for NO and NO<sub>2</sub> respectively.
- The average concentration of ambient O<sub>3</sub> and NO<sub>x</sub> during the whole study period from 2013-2017 were 18 ppb and 11ppb respectively.
- The median concentrations of the species were found to be 17, 19.8, 5, 2, 3 and 21 ppb during the 2013-2014; 15, 18.2, 6, 3, 3 and 18 ppb during the year 2014-2015; 12, 17.7, 10, 5, 3 and 14 ppb during the year 2015-2016 and 19, 37.8, 7.6, 6.7, 10.3 and 31.8 ppb during the year 2016-2017 for O<sub>3</sub>, O<sub>3</sub> max, NO<sub>x</sub>, NO, NO<sub>2</sub> and OX respectively.

## *Conclusions and Future Scope*

- The concentration modes were found to be 12, 15, 3, 3, 3 and 11; 8, 11, 4, 3, 4 and 7; 8, 12, 7, 5, 2 and 21; 15, 25, 6.2, 10.5 and 18 for O<sub>3</sub>, O<sub>3</sub> max, NO<sub>x</sub>, NO, NO<sub>2</sub> and OX respectively for the four consecutive years.
- The days of the week generally showed O<sub>3</sub> trends with single hump (maximum ozone) during the mid-day in all the four years. This site being remote and rural, it has been explicit in NOT showing profound weekday-weekend variation in the trends of O<sub>3</sub> concentrations.
- The O<sub>3</sub> distribution of the days of the week which however shows a mild rise in the concentrations mostly during midweek during the first three years but the rise in concentrations shifts towards weekends in the last year i.e. 2016-2017. The hourly variations of O<sub>3</sub> shows much clear 'bookish' hump around midday in all the four years.
- The monthly distribution of O<sub>3</sub> clearly shows a peak during the month of April (spring) maximum of O<sub>3</sub> concentrations at this remote site like several earlier researchers reported in all the years except in the year 2014-2015 which shows a slight increase in concentrations during the month of February. Least concentrations were observed mainly from June to December in all the years.
- The diel variation of maximum concentration of NO<sub>x</sub> exhibited a slight morning and a prominent evening peak i.e. at around 6 P.M. in all the four years.
- The rise in concentrations of NO<sub>x</sub> was generally seen around midweek i.e. Wednesday and Thursday during the entire study period. The monthly distribution of NO<sub>x</sub> clearly shows a peak during the month of January and February in the year 2013-14, 2014-15 and 2015-16 except in the year 2016-17 the peak was seen during the month of March.
- The daily profile of OX was found to be sharply increasing during the morning hours to the mid-day. Gradual decrease in the concentration of OX was observed after sunset which has a similar trend like that of ozone.
- The maximum concentrations of OX were found during the pre-monsoon season mainly in the month of April. The increased photochemical activities in the presence of intense solar radiation lead to the increase in concentration during the premonsoon season.
- The Highest concentration of ambient O<sub>3</sub> was found during Diwali day in the year 2016. Moreover during the year 2014 and 2016 the effect of Diwali crackers on O<sub>3</sub> concentration during night time was found to be more prominent. The maximum

concentration of O<sub>3</sub> during the year 2013, 2014, 2016 and 2017 were 27 ppb; 35 ppb; 38.5 ppb; 39.8 ppb respectively.

- Maximum concentration of NO<sub>2</sub> during Diwali was generally observed during night hours especially in the 20<sup>th</sup> hour. The maximum concentration of NO<sub>2</sub> during the year 2013, 2014, 2016 and 2017 were 15.8 ppb; 12.1 ppb; 36.7 ppb; 10 ppb respectively.
- During the year 2013 the O<sub>3</sub> and NO<sub>2</sub> shows a strong relationship with r<sup>2</sup> value= 0.123 than the non-Diwali nights showing a poor correlation. Same trend was also observed during the following years. The year 2014 has shown the strongest correlation than the other years during the Diwali night with an r<sup>2</sup> value of 0.5.
- The concentration of O<sub>3</sub> during maximum obscuration for the year 2014 was 36 ppb during eclipse day whereas; during normal days the concentration was 52 ppb for the same hour. However for the year 2015 the concentration was only slight different from the normal day's i.e during eclipse days it was 51 ppb at maximum obscuration level and 52 ppb during normal days at the same hour.
- The changes observed in case of ambient O<sub>3</sub> were pretty much during the first year as compared to the following year. However, slight change in NO<sub>x</sub> concentration was observed during solar eclipse in both the years. Moreover, prominent changes in case of meteorological parameters were observed during solar eclipse hours.
- A positive correlation was seen between O<sub>3</sub> and SR during eclipse period for both the years. However more prominent correlation was seen in the year 2015 (r<sup>2</sup>=0.3) than the previous year (r<sup>2</sup>=0.2).
- An inverse relationship of ambient O<sub>3</sub> with NO<sub>x</sub> was seen during eclipse hours. It was observed that as the concentration of NO<sub>x</sub> increases, the concentration of O<sub>3</sub> decreases thereby leading to an inverse relationship between them. However, the correlation (r<sup>2</sup>=0.3) seen in the first year is more prominent than seen in the next year (r<sup>2</sup>=0.002).
- The polynomial fit curves for NO, NO<sub>2</sub>, and O<sub>3</sub> against NO<sub>x</sub> of the remote rural location (Tezpur) were found NOT to follow the pattern of a highly polluted city (Delhi).
- A very strong relation between O<sub>3</sub> and NO<sub>2</sub>/NO ratio was observed which implied that O<sub>3</sub> concentrations increases with increase in NO<sub>2</sub>/NO ratio. O<sub>3</sub> concentrations tend to increase slowly with lower values of NO<sub>2</sub>/NO ratio and gradually reaches a stable period which explains the photostationary state of O<sub>3</sub>.

## Conclusions and Future Scope

- The concentrations of O<sub>3</sub> increases at low levels of NO<sub>2</sub>/NO ratio which implies that at moderate concentrations of O<sub>3</sub> the reactions involved in its production is dominant.
- The concentration of O<sub>3</sub> decreases with increase of NO. The maximum concentration of O<sub>3</sub> and NO during daytime was found to be 75 ppb and 19.4 ppb respectively and during nighttime the concentrations were 63 ppb and 26.6 ppb respectively. An inverse relation was seen during nighttime which indicates as NO increase during night O<sub>3</sub> concentration decreases.
- The concentration of O<sub>3</sub> increases with increase of NO<sub>2</sub> referring to the fact that NO<sub>2</sub> helps in build of O<sub>3</sub> in the ambient atmosphere. The maximum concentration of O<sub>3</sub> and NO<sub>2</sub> during daytime was found to be 75 ppb and 42.3 ppb respectively and during nighttime the concentrations were 63 ppb and 33.6 ppb respectively.
- A strong relation was seen between NO<sub>x</sub> and NO<sub>2</sub> during Daytime as compared to NO. So it can be said that NO<sub>2</sub> contributes more to O<sub>3</sub> buildup than NO during daytime.
- A very strong relation of NO<sub>x</sub> and NO was observed during nighttime as compared to day. During night the emission of NO is more than that of NO<sub>2</sub> due to more traffic congestion in night hours.
- The average monthly ambient temperature at Tezpur during the period 2013-17 ranged from 18 to 30°C. The minimum temperature recorded was 11°C in January 2014 and 2017 while the maximum temperature recorded was 39°C in August 2016. A positive dependency of O<sub>3</sub> on temperature (max) ( $r^2 = 0.15$ ) is seen.
- The monthly average relative humidity during the whole study period varied between 71 and 86%. The maximum humidity was during June 2016 and 2017 (92%) and the minimum was during April 2014 (55%). An inverse relationship of O<sub>3</sub> with relative humidity (RH) ( $r^2 = 0.2$ ).
- The *net* O<sub>3</sub> (measured O<sub>3</sub> concentration) and  $J_1/k_3$  were plotted against time of day for the different seasons to ascertain how the temporal variation of O<sub>3</sub> behaved relative to  $J_1/k_3$ . The *net* O<sub>3</sub> and  $J_1/k_3$  attained a maximum during midday to the early afternoon hours and then both parameters declined as the day progressed.
- During the pre-monsoon and monsoon seasons, the *net* O<sub>3</sub> and  $J_1/k_3$  attained their maxima and were about equal during midday to afternoon (1130 hr to 1530 hr local time).
- During post-monsoon and winter seasons, the *net* O<sub>3</sub> and  $J_1/k_3$  peaked for a shorter duration at around 14:00 local time. During the post-monsoon *net* O<sub>3</sub> and  $J_1/k_3$  were

nearly equal at ~1400hr local time. However, during the winter, *net* O<sub>3</sub> and  $J_1/k_3$  never became equal. During the peak hour, the levels of *net* O<sub>3</sub> were much higher than the  $J_1/k_3$  during the winter month.

- The ventilation coefficients (VC) calculated for winter month (January) as mixed layer height (MLH) x wind speed (WS), illustrated in showed that the maximum VC was experienced at ~9:00am in the morning and that is then there was steep rise in the winter period concentration of *net*O<sub>3</sub>.
- The polar plots suggest that high O<sub>3</sub> concentrations were found to accompany high windspeed that suggests transport of O<sub>3</sub> to the site. Alternatively, higher concentrations of NO<sub>2</sub> were observed under lower windspeeds. Only moderate and low levels of NO<sub>2</sub> were associated with high windspeeds .It is clear from the plots that higher O<sub>3</sub> and higher NO<sub>2</sub> were seen to be associated with winds from different directions.
- The effects of wind direction on ambient O<sub>3</sub> pollution can be well understood by the use of Pollution rose. Higher concentration of ambient O<sub>3</sub> concentration was dominant towards southeasterly direction in all the seasons. From the plots it is quite evident that the source of O<sub>3</sub> concentration is long range transport or regional transport which is usually seen in case of rural areas.
- The CPF and CBPF results signify extreme ambient O<sub>3</sub> episodes under the influence of wind from south-easterly directions. However, during the year 2016, higher concentrations of ambient O<sub>3</sub> were also seen in the western region along with the eastern regions, which was an exception to the other years and only prominent in the CBPF plots. In addition to the source regions the CBPF plots also shows that higher concentration of ambient O<sub>3</sub> occurs at low wind speed conditions. Moreover, the plots clearly indicate higher concentration of ambient O<sub>3</sub> pollution from eastern direction during the whole study period.
- NO<sub>x</sub>-dependent contribution attributes to the local emissions mainly due to biomass burning and vehicular exhaust, which was found to be ~20ppb. However, the regional contribution attributes to the NO<sub>x</sub>-independent contribution as the concentration of OX during this phase remains unaffected and was found to be ~18ppb. This signifies that the regional to local contribution was 1:1 over mid Brahmaputra Valley.
- OX and NO<sub>2</sub> have a linear relationship which inferred that as NO<sub>2</sub> concentration increases OX concentration also increases.

- Weighted PSCF maps were used to identify the possible source regions contributing to ambient O<sub>3</sub> and NO<sub>2</sub> levels in Tezpur. The PSCF maps were also plotted to study the impact of continental and marine air masses on ambient O<sub>3</sub> and NO<sub>2</sub> concentrations at Tezpur by calculating 3 days back trajectory during winter, premonsoon, monsoon and postmonsoon season for the year 2014-2017 using meteoinfo and trajstat software.
- HYSPLIT air mass trajectories reaching the site were weighted with the concentrations of O<sub>3</sub> and NO<sub>2</sub> and the CWTs were computed. These results suggest that the O<sub>3</sub> concentrations are strongly affected when the trajectories originated or travelled over polluted regions transporting O<sub>3</sub> precursors such as PAN to the area.

## **5.2: Future Scope**

- Continuous monitoring of ambient O<sub>3</sub>, its precursors along with meteorological parameters will enable to determine a detailed study on the trends of ambient O<sub>3</sub> and its precursors and also to identify its sources and sink.
- Further continuous analysis will enable to develop mathematical models for further study of ambient O<sub>3</sub> in this region and thereby determining the air quality of the surrounding areas.
- Extensive study on the emissions of O<sub>3</sub> precursors should be done by studying the population growth, land use change and new technology development.
- Emissions of ambient O<sub>3</sub> and its precursors during festive seasons should be continued to know the emission characteristics during this period.