Evaluation NO2 Detection Using Low-Cost Folded Path Photometer

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Abstract. Nitrogen dioxide (NO2) contributes to several impacts both for human health and the environment. Periodical measurement of NO2 at industrial and residential areas needs comprehensive and reliable instrumentation; long-interference-free measures with minimum maintenance and re-calibration. DOAS can be used as a direct measurement technique based on specific absorption characteristics of NO2 follow Beer-Lambert law. This research applies a low-cost folded path photometer for measurement NO2 in air. Cheap tubular acrylic used as a detection cell with a 3D printed framework makes it compact, modular, and flexible. Evaluation of the DOAS conducted by instrument test responses using NO2 gas. The estimated LOD was ~ 1263 pb using 2 nm resolution of the spectrometer and 6-meter detection cell length. Deviation of the DOAS estimated to be 0.8% at high concentration and 2.85% at a low concentration based on DOAS calibration. An intercomparison between methods used to evaluate instrument performance to measure NO2 using emission from a motorcycle with coefficient correlation (R) 0.649 for paired DOAS-ASTM D1607 Griesz Saltzmann Method 0.846 for paired DOAS-Electrochemical Gas Analyzer. This significant correlation is caused by different respond time between paired methods, while it is still comparable for NO2 measurement.

Keywords: DOAS, intercomparison, nitrogen dioxide, photoabsorption, spectrometer.

1 Introduction

Approximately 4.2 million death per year is caused by poor ambient air quality related to anthropogenic activities [1]. Nitrogen dioxide (NO2), together with Sulfur dioxide (SO2), Ozone (O3), Carbon Monoxide (CO), Particulates, and Lead (Pb) known as one of six criteria air pollutants as it may have an impact on health and ability to degrade ecosystems and agricultures quality [2]. NO2 can directly irritate the human respiratory system in concentrations 50 – 100 pm in 30-minutes exposure. NO2 may contribute to the development of asthma [3]. NO2 and other NOx react well with water, oxygen, and other chemicals in the atmosphere to form acid rain that may harm sensitive ecosystems such as aquatic ecosystems, forests,
and agriculture regions [4]. A comprehensive and reliable data will be an excellent addition to these requirements. There are also specific needs for measurement instruments; long-interference-free measurements with minimum maintenance and re-calibration [5].

Testing laboratories, industry, and local governments use various methods to measure NO₂ in the air. ASTM standards like Griesz Saltzmann are used in the testing laboratory to conduct accurate NO₂ tests, while this test only results in single data in an hour test [6]. Absorption photometry can be used as a direct measurement technique based on specific absorption characteristics of NO₂. Absorption of light would follow Beer-Lambert law below [7].

\[
\frac{I}{I_0} = e^{-\sigma lc} \quad \text{or} \quad c = \frac{1}{\sigma l} \ln \left( \frac{I_0}{I} \right)
\]

(1)

Where \(I_0\) is the intensity of light that passing absorbance cell without NO₂ analyte present, \(I\) is the intensity of light that passing absorbance cell with NO₂ analyte present, \(\sigma\) is the analyte coefficient of extinction (absorption cross-section in cm⁻² molecule⁻¹ for NO₂), \(l\) is the path length of the NO₂ detection cell (cm or m), and \(c\) is the concentration of the NO₂ within detection cell (molec cm⁻³). Arranging Eq.1, absorbance \(\ln \left( \frac{I_0}{I} \right)\), can be measured using a spectrometer-based instrument with a typical precision of \(\sim 3 \times 10^{-6}\). As precision has good agreement with path length, there are several approaches to gain better precision. Several kilometers of the open-path system have been used to measure the pt scale in differential optical absorption spectroscopy (DOAS) [8, 9]. However, this method limits the determination of the spatial distribution of analyte concentration. Another approach, closed-path DOAS equipped with mirrors to fold the light directly through the path length [10, 11]; the White cell is the most common, with typical 100 or more reflections. This White cell has the disadvantages of a long period of flushing time. This research describes a new approach, a low-cost folded path photometer for measurement NO₂ in air. A cheap acrylic tube may replace expensive metallic detection cells like Aluminum since acrylic did not react with NO₂. Two Molybdenum mirrors use in mirror modules at a 45° angle. Construction can be compact enough as several meter path lengths may be folded in one-meter dimension as we fold the path length. A 3D printed material from PLA-based filament used as an entire path length framework made it easy to replicate in another place and further research. The design makes it possible to solve the spatial distribution problem of analyte in an open-path system. Direct absorption at 440 nm technique used as NO₂ measurement using a typical LED available in the market. Measurement of other gas species in the air can be adopted in the future using this modular approach.
2 Method

2.1 Experimental Design

Figure 1 is a schematic diagram of the folded path photometer measuring NO$_2$ concentration in air based on Beer-Lambert law (Eq. 1). An air pump draws analyte with a constant 1 liter per minute volumetric flow through the detection cell. The sample enters the instrument through the particulate filter made of a glass microfiber filter. It prevents particles in the sample from interfering with the absorbance measurement. The sample air enters the detection cell, made by cylindrical acrylic tubes with 30 mm inside diameter and 90 cm long. At the end of each detection cell, two fixed mount Molybdenum mirrors with 45° angle orientation were placed. Two-mirror connected with an air column 10 cm long, so the sample air flowed through them and entered the next detection cell. A total 600 cm path length can be acquired with a 4.5-liter total volume with six detection cell trains.

The light source module used in this research emits light with proper wavelength to selectively absorb NO$_2$ as an analyte. An LED with a wavelength range between 435 nm – 465 nm was utilized to measure NO$_2$. LED light powered by 3.4 DC voltage with 1 Ampere constant current, collimated using 15° convex lens with 10 cm focal length at each end of the first and last detection cell. At the end of the optical train, the light was detected by Thorlabs CCS 200 CCD spectrometer. More than 90% of the light fraction will disappear using absorption spectroscopy since there is a reflection on the cell walls and mirrors [12].
losses have little impact on the measurement since they remain at the constant condition of \( I_0 \) (NO\(_2\) scrubbed) and \( I \) (NO\(_2\) present). The concentration of the NO\(_2\) as the analyte is calculated based on Beer-Lambert law.

### 2.2 Calibration of CCD Spectrometer

Collimated mercury lamp used to analyze the relationship between diffracted light wavelength and pixel number of the CCD camera on Thorlabs CCS 200. Mercury lamp has several lines in the spectral region between 253 nm – 579 nm [13]; Figure S1 describes the calibration of the CCD spectrometer.

### 2.3 Experimental Procedure

In DOAS measurement, background spectrum, \( I_0(\lambda) \) as blank and analyte spectrum, \( I(\lambda) \) needs to be measured. A zero-gas containing \( \text{N}_2 \) is used as blank since it does not have interfering NO\(_2\) absorption in 435 nm – 465 nm. Series of NO\(_2\) gas calibration standard was used as validation measurement, analyzed using the correlative statistic. Co-location measurement deploys by measure three different methods between DOAS, electrochemical gas sensor Alphasense NO2-B4, and ASTM D1607 Griesz Saltzmann as a comparison. Both data were analyzed for the Griesz Saltzmann method, recorded within the same period, and analyzed using correlative analysis.

### 3 Result and Discussion

#### 3.1 Spectral Resolution

Spectral resolution from instrument measured using line spectra using mercury lamp. Mercury lamp has primary line spectra at 546 nm and 578 nm, while secondary spectra at 366 nm, 403 nm, 435.8 nm, 1012 nm, 1125 nm, 1362 nm, 1525 nm, 1688 nm, and 1692 [14]. At the evaluation wavelength range 435 nm – 465 nm, mercury lamps have strong line emissions at 435.8 nm—line spectra at evaluation range in Thorlabs CCS 200 shown in figure S2. Thorlabs CCS 200 have resolution \(< 2\) nm, 1.1 nm deviation of the peak spectra indicate this spectrometer is in proper condition to measure absorbance photometry. According to [15], Bruker 120M FT Fourier transform spectrometer used at 0.01 nm resolution at 240 nm and 0.2 nm at 1000 nm to measure NO\(_2\) absorption cross-section in high resolution used as reference. The cheap blue LED used in this research may have a limited wavelength region compared with previous research. Figure 3(a) showed a high resolution of NO\(_2\) absorption cross-section from Vandaele compared with the evaluation range of this research at 435 nm – 465 nm, overlayed with typical intensity of blue LED absorbance.
Figure 2  (a) Evaluation range of NO$_2$ measurement, (b) Gaussian fitting between different instrument resolution.

The Thorlabs CCS 200 absorption cross-section needs to be adapted to the low resolution of the instrument. Figure 3(b) show that each wavelength range is shown as normal distribution pattern, gaussian fitting may be used as absorption cross-section fitting between two different resolutions. Measured NO$_2$ concentration can be calculated using least square fitting between measured intensity based on Beer-Lambert law and modeled optical density $\sigma'_i(\lambda)$ based on absorption cross-section reference $\sigma'_{Lit}$ [16], as seen on Eq.2. Optical density $\tau$, calculated as logarithmic value of measured spectrum $I(\lambda)$ with background spectrum $I_0(\lambda)$.

$$\tau = \ln \frac{I_0(\lambda)}{I(\lambda)} = \sum_i \sigma'_{Lit}(\lambda) \cdot \bar{c}_i \cdot L + R(\lambda)$$  \hspace{1cm} (2)

Using the least square method, we may minimize residual value $R(\lambda)$, between measurement and model, then NO$_2$ mixing ratio $\bar{c}_i$ can be calculated in molecule per volume. NO$_2$ concentration in pm or pb is calculated as a mixing ratio towards the number of molecules per volume as an ideal gas.

3.2 Limit Detection

The detection limit (LOD) is the lowest concentration that can be measured with statistical significance through a given analytical procedure [17] and can be estimated using Eq.(3).

$$LOD = \mu + 3SD$$  \hspace{1cm} (3)

Blank analyte measured as LOD estimation with data can be seen in Figure 4 data gathered from 1000 blank analyte measurements.
Theoretically, cell detection length would affect LOD and be affected by the spectrometer’s resolution. Ten times lower resolution used in this research than Vandaele’s [15] may imply high LOD compared with a theoretical detection limit of 100 pb for a 6-meter detection length cell. Based on Eq(3) LOD of the DOAS estimated at ~ 1263 pb, this LOD is in step with Devine’s research using Ocean Optics USB 4000 spectrometer with a similar resolution with Thorlabs CCS 200 [18]. With this LOD, calibration of the DOAS may use two points calibrations at a higher concentration than 1.5 pm.

3.3 DOAS Calibration

Calibration is a process to verify and evaluate the accuracy of the DOAS. Several calibration techniques may be used, [19] calibrate DOAS using correlation analysis based on certified reference instruments. [20] use gas calibration standards to check the accuracy of the DOAS. In this research, two different concentrations of NO2 are used, 50 pm and 3.5 pm, each represents high concentration and low concentration [21]. A Zephyr gas dilution system was used to dilute NO2 using N2 to get a NO2 concentration of 3.5 pm. Each concentration was measured at the same condition, calibration data seen in Figure 5. With the uncertainty of the gas calibration standard 5% (marked as the upper limit and lower limit in Figure 5), ten data measurements show that the average concentration of high concentration gas calibration standard is 50.4+0.85 pm 3.4+0.32 pm for low concentration with 95% confidence level. From this measurement, the deviation of the DOAS estimated to be 0.8% at high concentration and 2.85% at low concentration.
Figure 4  DOAS calibration using (a) 50 pm-high concentration and (b) 3.5 pm-low concentration

3.4  Precision Measurement

Intercomparison and co-location analysis between DOAS, electrochemical gas analyzer, and ASTM D1607 reference method was used as a precision test. As a NO$_2$ source, we used NO$_2$ emission from the motorcycle, both DOAS, electrochemical gas analyzer, and ASTM D1607 Griesz Saltzmann data plotted in parallel, as shown in Figure 6.
Statistically, paired sample test was acquired to hypothesize the average measurement between two paired methods; DOAS – Electrochemical Gas Analyzer and DOAS – ASTM D1607 Griesz Saltzmann, as seen in table S3. DOAS – Electrochemical Gas Analyzer seems to be more correlated than DOAS – ASTM D1607 Griesz Saltzmann identified by higher coefficient correlation on it, this significance correlation in between paired method caused by different response time between method as measurement data, DOAS have 150 ms respond time compared with 90 seconds using electrochemical gas analyzer, while ASTM D1607 Griesz Saltzmann has 10 minutes response time for each measured data. As a result, another intercomparison needs to be conducted using an instrument with a short response time compared to DOAS [22]. Based on the paired sample test, we can acquire the hypothesis that both paired methods were comparable by seeing Sig. (2-tailed) a result that higher than < 0.05 indicates no average difference between two paired measurement data and concluded that both paired methods were comparable.

4 Conclusions

There is a good intensity response in DOAS to detect NO2 in the air as seen on the absorbance response in the DOAS instrument. With 2 nm resolution of the spectrometer and 6-meter detection cell length, this setup has a detection limit (LOD) ~ 1263 pb. Deviation of the DOAS estimated to be 0.8% at high concentration and 2.85% at a low concentration based on DOAS calibration. An intercomparison between methods used to evaluate instrument performance to measure NO2 using emission from the motorcycle with coefficient correlation (R) 0.649 for paired DOAS-ASTM D1607 Griesz Saltzmann Method 0.846 for paired DOAS-Electrochemical Gas Analyzer. This different significance correlation is caused by different response times between paired methods. Even
paired sample test indicates both paired methods were comparable for NO$_2$ measurement.

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