Traffic-related air pollution near roadways: discerning local impacts

from background

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26 Abstract. Adverse health outcomes related to exposure to air pollution have gained much attention in recent years, with a 27 particular emphasis on traffic-related pollutants near roadways, where concentrations tend to be most severe. As such, many 28 projects around the world are being initiated to routinely monitor pollution near major roads. Understanding the extent to 29 which local on-road traffic directly affects these measurements, however, is a challenging problem, and a more thorough 30 comprehension of it is necessary to properly assess its impact on near-road air quality. In this study, a set of commonly 31 measured air pollutants (black carbon; carbon dioxide; carbon monoxide; fine particulate matter, PM_{2.5}; nitrogen oxides; ozone; 32 and ultrafine particle concentrations) were monitored continuously between June 01st, 2015 and March 31st, 2017 at six stations 33 in Canada: two near-road and two urban background stations in Toronto, Ontario, and one near-road and one urban background 34 station in Vancouver, British Columbia. Three methods of differentiating between local and background concentrations at 35 near-road locations were tested: 1) differences in average pollutant concentrations between near-road and urban background 36 station pairs, 2) differences in downwind and upwind pollutant averages, and 3) interpolation of rolling minima to infer 37 background concentrations. The latter two methods use near-road data only, and were compared with method 1, where an 38 explicit difference was measured, to assess accuracy and robustness. It was found that method 2 produced average local 39 concentrations that were biased high by a factor of between 1.4 and 1.7 when compared with method 1 and was not universally 40 feasible, whereas method 3 produced concentrations that were in good agreement with method 1 for all pollutants except ozone 41 and $PM_{2.5}$, which are generally secondary and regional in nature. The results of this comparison are intended to aid researchers 42 in the analysis of data procured in future near-road monitoring studies. Lastly, upon determining these local pollutant 43 concentrations as a function of time, their variability with respect to wind speed (WS) and wind direction (WD) was assessed 44 relative to the mean values measured at the specific sites. This normalization allowed generalisation across the pollutants and 45 made the values from different sites more comparable. With the exception of ozone and PM_{2.5}, local pollutant concentrations 46 at these near-road locations were enhanced by a factor of 2 relative to their mean in the case of stagnant winds and were shown to be proportional to WS^{-0.6}. Downwind conditions enhanced local concentrations by a factor of ~2 relative to their mean, while 47 upwind conditions suppressed them by a factor of ~4. Site specific factors such as distance from roadway and local meteorology 48 49 should be taken into consideration when generalizing these factors. The methods used to determine these local concentrations, 50 however, have been shown to be applicable across pollutants and different near-road monitoring environments.

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60 1 Introduction

61 Traffic-related air pollutants (TRAPs) are of concern because on-road traffic is often a major source of air pollution in urban 62 environments (Belis et al., 2013; Molina and Molina, 2004; Pant and Harrison, 2013) where population densities are greatest— 63 in Canada, it is estimated that one third of the population live within 250 m of a major roadway (Evans et al., 2011)—and it is 64 within these near-road regions TRAP concentrations are generally highest (Baldwin et al., 2015; Jeong et al., 2015; Kimbrough 65 et al., 2018; Saha et al., 2018).

As such, there is a growing interest in measuring air pollutant concentrations near roadways in order to better understand TRAP exposure levels in these environments. However, in order to isolate the underlying sources and reasons for elevated concentrations, further processing of raw measurement data is necessary. In general, near-road TRAP concentrations are influenced by both regional and local emissions, and being able to distinguish the contributions of these sources allows their relative impacts to be more properly assessed. Of particular importance to near-road measurements is understanding the role of on-road traffic. For TRAPs whose source(s) cannot be readily identified from their measurement at a singular location, concurrent samples at various locations and/or algorithmic methods can be used to enable apportionment.

73 Often, determining TRAP background concentrations is accomplished through monitoring at remote, representative locations 74 that are minimally impacted by nearby sources; properly siting background stations in urban environments is in itself a 75 challenge, and not always feasible. This practice, while useful in providing confidence in information regarding background 76 air quality, is expensive because it requires additional monitoring stations and personnel to maintain them. The value of these 77 background stations is lessened if similar knowledge is extractable from near-road locations alone. Various time-series analysis 78 algorithms have been proposed for this purpose, many of which make use of the inverse relation between source proximity 79 and signal frequency. For example, the technique of interpolating minima across time windows of varying length has been 80 applied successfully to data from both mobile laboratories (Brantley et al., 2014; Shairsingh et al., 2018) and stationary 81 measurements (Wang et al., 2018) for the purposes of estimating urban background pollutant concentrations. Additionally, 82 work by Klems et al., (2010) and Sabaliauskas et al., (2014) made use of the discrete wavelet transform, an algorithm used 83 widely in signal compression and denoising, to ultrafine particle time-series data to determine the time-dependent contribution 84 of local sources to roadside concentrations. Another technique, statistical clustering of air quality data in urban environments, 85 was utilized by Gomez-Losada et al. (2018) to characterize background air quality. Indeed, there are many promising avenues 86 of background-subtracting near-road air quality data.

Given the diversity of techniques available for differentiating local and background pollutant concentrations, as well as the large variety of instrumentation available, it is not clear which approaches are most generalizable or applicable, or whether it is necessary to invest in concurrent measurements at many versus few locations. In addition, the exact definition of what is background air quality is somewhat unclear, and in the context of this study, given the spatial separation between sites (on the order of 10 km or less), it is assumed to be a measure of background air quality in the urban airshed. Ma and Birmili (2015), in a study of ultrafine particle nucleation, defined measurement locations in their study which were 4.5 km and 40 km from an 93 urban roadside station as urban background and regional background, respectively. The former was presumed to be a measure 94 of regional air quality superimposed with diffuse urban emissions, and it is this definition that best characterizes the background 95 air quality measured in this study. To evaluate whether information regarding this urban background was attainable from near-96 road measurements alone, two strategies for quantifying the contribution of local on-road traffic to near-road air quality were 97 compared, and their reliability and accuracy were assessed through comparison with tandem measurements in both 98 environments.

In this study, data were collected continuously at three near-road and three urban background monitoring locations for close to two years (namely, between June 01st, 2015, and March 31st, 2017). Various gas and particle-phase pollutants along with meteorological parameters were measured using an array of instrumentation. Concentrations in excess of the urban background were calculated from the near-road data using three techniques, one of which calculated an explicit difference between sites, whereas the other two made use of only near-road data. Comparison of these methodologies addresses whether information regarding background air quality is readily inferable from measurements made in the near-road environment.

105 2 Methods

106 2.1 Measurement locations

107 Data were collected from six separate monitoring locations: four of which were in Toronto, Ontario (two situated near 108 roadways and two in urban background environments), with the remaining two located in Vancouver, British Columbia (one 109 situated near a roadway and another in the urban background). The location of each station, along with information regarding 110 the major roadway next to which they were located (for the near-road sites), is summarized in Table 1. The two near-road 111 stations in Toronto, NR-TOR-1 (43.7111, -79.5433) and NR-TOR-2 (43.6590, -79.3954), and their respective instrumentation 112 setups have been utilized and reported by others and are described therein (Sabaliauskas et al., 2012; Sofowote et al., 2018; 113 Wang et al., 2015). The NR-TOR-1 site was positioned 10 m from Highway 401, the busiest highway in North America in 114 terms of Annual Average Daily Traffic (AADT) with over 400,000 vehicles per day distributed across eight eastbound and 115 eight westbound lanes. The Southern Ontario Centre for Atmospheric Aerosol Research (SOCAAR) served as the second near-116 road site (NR-TOR-2), and was located 15 m from College Street in downtown Toronto which experienced traffic volumes of 117 17,200 vehicles per day on average. The northernmost station in Toronto, BG-TOR-1, was located at Environment and Climate 118 Change Canada (43.7806, -79.4675), 180 m from the nearest roadway, and the measurements from this station served as an 119 urban background/baseline for NR-TOR-1, which was located 9.8 km to the southwest of it. The second background station, 120 BG-TOR-2, was positioned on the southernmost point of the Toronto Islands on Lake Ontario (43.6122, -79.3887), and was 121 5.2 km south of NR-TOR-2. Since vehicular traffic on the Toronto Islands was limited to a small number of service vehicles, 122 the BG-TOR-2 station was well removed from tailpipe emissions.

The near-road station in Vancouver, NR-VAN, was situated 6 m from Clark Drive (49.2603, -123.0778), a major roadway that experienced on average 33,100 vehicles per day across four southbound and three northbound lanes. Additionally, located 65 m south of the station was a major intersection, Clark Drive and 12th avenue, at which there were two gas stations located on the northwest and northeast sides. The effect this intersection had on traffic patterns (stop-and-go, especially) directly next to the station, and its effect on measured TRAP concentrations are explored in this study. Lastly, the urban background station in Vancouver, BG-VAN, was located 2.2 km east of NR-VAN at Sunny Hill Children's Hospital (49.2529, -123.0492). This area was relatively removed from traffic emissions because it was located within a neighbourhood zoned predominately for

130 single unit family dwellings.

131 2.2 Instrumentation

132 A common suite of instrumentation was employed at all stations. Gas-phase pollutants measured include: carbon dioxide (CO₂; 133 840A, LI-COR Biosciences; attenuation of infrared radiation at wavelengths of 4.26 µm and 2.95 µm for H₂O differentiation), 134 carbon monoxide (CO; 48i, Thermo Scientific; attenuation of infrared radiation at a wavelength of 4.6 μ m), ozone (O₃; 49i, 135 Thermo Scientific; attenuation of ultraviolet radiation at a wavelength of 254 nm), and nitrogen oxides (NO_x; 42i, Thermo 136 Scientific; infrared chemiluminescence). Particle-phase pollutant properties measured include: mass concentration of particles less than 2.5 microns in diameter (PM_{2.5}; SHARP 5030, Thermo Scientific; beta attenuation and light scattering); particle 137 138 number concentration (UFP; 651, Teledyne API; water-based condensation particle counting); and black carbon (BC; AE33, 139 Magee Scientific; filter-based attenuation of 880 nm wavelength light) mass concentration. Additionally, a meteorological 140 sensor (WXT520, Vaisala; ultrasonic anemometer) recorded wind direction, wind speed, ambient temperature, pressure, and 141 relative humidity at each station. Traffic intensities, velocities, and approximate vehicle lengths were measured continuously 142 (SmartSensor HD, Wavetronix; dual beam radar) at the three near-road stations.

143 Gas-phase instruments were calibrated on-site every two months using cylinders of compressed gasses at certified 144 concentrations (Linde). One cylinder contained SO_2 , CO_2 , and CO_2 , while the other contained NO; both contained N_2 as an 145 inert makeup gas. Dilution and mixing of the gasses was accomplished using a dynamic gas calibrator (146i, Thermo Scientific; 146 6100, Environics) to produce zero checks and span concentrations that were similar to ambient ranges. Additionally, these 147 dynamic gas-phase calibrators contained ultraviolet (UV) based O₃ generators which were used to calibrate the 49i monitors 148 as well as test the efficiencies of the molybdenum NO₂ converters in the 42i monitors. SHARP 5030 instruments were zero 149 checked using a HEPA filter, had their temperature and relative humidity sensors calibrated, and were span checked using 150 mass standards supplied by Thermo Fisher Scientific twice annually. In addition to recommended monthly maintenance 151 procedures for the API 651, each instrument underwent routine annual calibration by the manufacturer. Flow rates at each 152 station were verified on a monthly basis, and a variable flow rate pump was attached to a stainless steel particle manifold, from 153 which all particle-phase instruments sampled, to ensure a constant flow rate of 16.7 LPM to satisfy the 2.5 µm cut-off 154 conditions of the inlet cyclone.

155 3 Data analysis

156 Data acquisition was accomplished using Envidas Ultimate software (DR DAS Ltd.). Quality assurance of the data was 157 performed by the primary operators of each station. This included, among other things: discounting data in which instrument 158 diagnostic parameters were outside of acceptable ranges, omitting calibration times, and flagging suspect periods. Data from 159 this study was acquired at a one-minute resolution, and further averaged to hourly resolution. Only hours containing at least 160 45 minutes (\geq 75%) of valid data are reported. Data processing and analysis was done through a combination of SOL 161 (Microsoft), SAS 9.4 (SAS Institute Inc.), and IGOR Pro 6.37 (Wavemetrics Inc.) software. Using the hourly concentrations 162 in the finalized dataset, three methods of separating local and background concentrations from the near-road measurements 163 were tested. One of these methods made use of the urban background measurements to explicitly infer background concentrations, whereas the other two, downwind/upwind comparison and interpolation of minimum concentrations, estimated 164 165 background concentrations from the near-road measurements alone.

166 3.1 Average site differences

167 The first method for determining local pollutant concentrations explored in this paper, henceforth referred to as method 1, is 168 through the difference between concentrations measured at a near-road location, C_{NR} , and at the nearest urban background 169 location, C_{BG} , for some concurrent observation, i. Concentrations associated with local influences determined using method 1, 170 C_{L-1} , rely on the assumption:

$$171 \quad C_{NR}[i] = C_{L,1}[i] + C_{BG}[i]. \tag{1}$$

172 Average C_{L,1} values for each near-road location were then determined using Eq. (2):

173
$$\bar{C}_{L,1} = \frac{1}{N} \sum_{i=1}^{N} (C_{NR}[i] - C_{BG}[i]),$$
 (2)

again, $C_{NR}[i]$ and $C_{BG}[i]$ are near-road and urban background measurements, respectively, made over a concurrent time interval, i, As N, the number of observations used in calculating the temporal average increases, the calculated average difference will encompass more of the variability from meteorological and traffic conditions, and therefore be more representative of an average site difference.

178 **3.2 Downwind-upwind analysis**

Through association with meteorology at a near-road measurement location, it is possible to assess traffic's influence on TRAP concentrations from the differences between downwind and upwind conditions. For example, Galvis et al. (2013) utilized average downwind and upwind concentrations of CO_2 , BC, and $PM_{2.5}$ from a railyard to calculate local pollutant concentrations for use in fuel-based emission factor calculations. A similar approach is used here to isolate concentrations emitted from a roadway, henceforth referred to as method 2. Defining ranges of wind directions as corresponding to downwind and upwind of the major street next to which a station is located, average local concentrations from method 2, $C_{L,2}$, can be estimated using Eq. (3):

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$$\bar{C}_{L,2} = \frac{1}{N} \sum_{i=1}^{N} C_{DW}[i] - \frac{1}{M} \sum_{i=1}^{M} C_{UW}[i],$$
 (3)

where C_{DW} and C_{UW} are near-road TRAP concentrations measured when winds originate from downwind and upwind of the 187 188 major roadway, respectively. Note that the number of points used to compute the averages of these conditions, N and M, are 189 not necessarily equivalent, and the times that comprise these two averages are mutually exclusive by definition. For example, 190 if the prevailing wind at a site is downwind of the roadway, then downwind data will naturally occur more frequently than 191 upwind. Fig. S1 in the supplementary information shows wind frequency data as measured at each near-road site throughout the monitoring campaign. Similar to method 1, as the averaging time for both conditions is increased, confidence in $C_{1,2}$ will 192 193 improve. It is also important to note that because these two meteorological scenarios encompass different time frames, it is 194 possible for certain times of day, etc. to be overrepresented in either average.

In all analyses in which meteorological data are utilized, stagnant periods (wind speed (WS) $< 1.0 \text{ m s}^{-1}$) were omitted. Local concentrations cannot be estimated as a function of time using this method, as downwind and upwind concentrations cannot be measured simultaneously with a single near-road station. Also, stagnant time periods, as well as time periods that are not within the downwind/upwind ranges are omitted, thereby increasing the amount of time needed to attain a representative average. Lastly, an inherent assumption to this method is that upwind concentrations on either side of the roadway are similar. Depending on the site, however, this assumption may not be accurate.

201 3.2.1 Wind sector definitions at NR-TOR-1

Defining downwind and upwind sectors at NR-TOR-1 was straightforward, owing to the flat terrain of the area and the lack of nearby TRAP sources excluding those from Highway 401. Hence, 90° quadrants perpendicular to the highway axis were chosen. These definitions were further supported by average ambient CO₂ concentrations—an indicator of combustion associated with traffic emissions—measured as a function of wind direction, shown in Fig. 1. Thus, downwind conditions at NR-TOR-1 were defined as WD \geq 295° or WD \leq 25° and upwind as 115° \leq WD \leq 205°, where WD denotes wind direction as measured locally at the station atop a 10 m mast.

208 3.2.2 Wind sector definitions at NR-TOR-2

209 Unlike the NR-TOR-1 site, wind dynamics at NR-TOR-2 were complicated by urban topography; namely, the roadside inlet 210 was within an urban canyon (aspect ratio of ~0.5: building heights of ~20 m on either side and a street width of ~40 m) resulting 211 in more stagnant conditions roadside and introducing micrometeorological effects such as in-canyon vortices (Oke, 1988). The 212 effect of urban canyon geometry on micrometeorology is an effect that has been known for some time, and in general, for city-213 scale wind patterns perpendicular to the street axis, ground-level winds tend to be opposite to those above the urban canopy 214 (Vardoulakis et al., 2003). Given the urban canyon's effect on ground-level wind direction, downwind/upwind quadrants at NR-TOR-2 were determined based on wind direction measurements made above the urban canopy, and are defined as: $WD \ge 300^{\circ}$ or $WD \le 30^{\circ}$ and $120^{\circ} \le WD \le 210^{\circ}$ for downwind and upwind conditions, respectively. Figure 2 shows a satellite image of the site with these respective quadrant definitions, along with average CO₂ concentrations as a function of wind direction, similar to Fig. 1. From the range of CO₂ concentrations seen here, it is clear that obtaining a precise definition of what exactly is downwind or upwind of College Street is non-trivial. Impact from the intersection southwest (winds from ~230°) of the receptor is somewhat apparent in Fig. 2, also.

222 3.2.3 Wind sector definitions at NR-VAN

223 While the presence of 2-3 story buildings within the immediate vicinity of the NR-VAN station may have complicated 224 meteorological measurements to some extent, the role of wind direction on the impact of local traffic emissions was much 225 more evident at this site than it was at NR-TOR-2. Other streets in the vicinity of Clark Drive affected the driving patterns near the station—a major intersection (Clark Drive and 12th Avenue) approximately 65 m south of the station had an impact on 226 227 average measured CO₂ concentrations (Fig. 3) originating from the SSE direction. Because of this, the downwind and upwind sector definitions for this site were not taken to be orthogonal: instead, downwind was defined as $135^{\circ} \leq WD \leq 195^{\circ}$ and 228 upwind as $235^{\circ} \leq WD \leq 315^{\circ}$; these definitions were chosen in accordance with surrounding land usage. While the upwind 229 definition does include 12th avenue, a major roadway within 120 m of the station, it is suspected that lower TRAP 230 concentrations from this sector are due to: lower traffic volumes on 12th compared with Clark Drive, truck restrictions on 12th, 231 and mechanical mixing from surface roughness (i.e. winds carrying TRAPs emitted on 12th being pushed up over the densely 232 233 spaced buildings between the roadway and monitor, resulting in diluted or no TRAPs measured at ground-level). Contrasting 234 this upwind definition with measurements from the sector 315°-345° in Fig. 3, which includes the major roadway Broadway 235 250 m from the receptor (farther than 12^{th}), there is a difference in average CO₂ concentrations of about 15 ppm, and this 236 difference is likely due to reduced surface roughness NNW of the receptor. Both NR-TOR-2 and NR-VAN provide examples 237 of the complexity of siting near-road stations, and how site-specific considerations must be made when associating data with 238 meteorology.

239 **3.3 Background subtraction using time series data**

Extracting information from one-dimensional ambient pollution time-series data (i.e. concentration as a function of time) for the purpose of source apportionment is appealing as it allows the possibility of obtaining local and background estimates without the need for more rigorous chemical analysis, computationally expensive multivariate analyses, or measurements made at multiple locations. Such algorithms make use of the underlying principle that signal frequency is inversely related to source distance. Regional or background sources (farther away from a receptor) produce slower varying, lower frequency signals, whereas local (nearby) sources, such as traffic, produce faster varying, higher frequency signals (Tchepel and Borrego, 2010). The frequency at which data is acquired limits the highest frequencies separable by such a method. Daily averages, for example, are too lengthy to capture processes whose time scales are much shorter—a plume from a nearby on-road vehicle, for example, would have a characteristic time on the order of seconds to minutes. Therefore, in order to isolate these local temporal fluctuations, relatively high time resolution data are necessary. A technique recently developed by Wang et al. (2018) applied to hourly near-road measurements in order to determine above-background pollutant concentrations for use in calculating fleetaveraged emission factors is explored further in this paper.

252 **3.3.1 Interpolation of windowed minima**

The algorithm explored in this paper is an interpolation of minimum values across a variable time window, the duration of which effectively defines, in a sense, a cut-off frequency for local and urban background signal differentiation. This algorithm was developed, validated, and utilized by Wang et al. (2018), and is described in full detail therein along with code compatible with IGOR Pro 6.37.

The background-determining function, ψ , takes as arguments: near-road pollutant concentrations as a function of time, C_{NR}, a window length in hours, W, and a smoothing factor α . Its output is an inferred baseline for the near-road environment, **b**:

259
$$\boldsymbol{b} = \psi(C_{NR}, W, \alpha), \qquad \alpha \ge 1, W \ge 3$$
, (4)

In the case for which the smoothing factor, α , is equal to 1, the baseline function, **b**, simplifies to an interpolation of minimum values determined across M windows of width W, where M is the total number of measurements divided by W. In order to account for the detection of minima being biased by the range of each window, this process is repeated three times, in which the window is offset in time by *floor*(W/3) each time. This yields three separate functions, **b**₁, **b**₂, and **b**₃, with the final baseline, **b**, determined from the average:

265
$$\boldsymbol{b} = \psi(C_{NR}, W, \alpha = 1) = \frac{1}{3} \cdot \sum_{i=1}^{3} \boldsymbol{b}_i$$
, (5)

For the case in which $\alpha > 1$, the process in Eq. (4) and Eq. (5) is repeated α times, and the window for determining minimum values increases by a factor of W each time, giving window lengths of: W, 2W, ..., α W. Then, the final baseline function becomes the mean of α *W baseline functions, **b**_{i,j}:

269
$$\boldsymbol{b} = \psi(C_{NR}, W, \alpha) = \frac{1}{3\alpha} \cdot \sum_{j=1}^{\alpha} \sum_{i=1}^{3} \boldsymbol{b}_{i,j} , \qquad (6)$$

270 Thus, in addition to creating a smoother baseline output, the magnitude of the parameter α , in conjunction with that of W, 271 determines how slowly-varying the resultant baseline, **b**, becomes. The effect of these input parameters can be observed in 272 Fig. 4, in which ψ is applied to CO₂ data at NR-TOR-2 for various values of α and W. If the resulting baseline function, **b**, is 273 greater than C_{NR} for any point in time, it is instead set equal to C_{NR}. Henceforth, this algorithm shall be referred to as method 3. This method yields a baseline function, **b**, based on input nearroad concentrations, C_{NR} , constrained to yield non-negative solutions for each observation, i. Average local concentrations from method 3, $C_{L,3}$, were then calculated using Eq. (7) and Eq. (8):

277
$$C_{L,3}[i] = C_{NR}[i] - \boldsymbol{b}[i], \qquad \boldsymbol{b}[i] \le C_{NR}[i] \forall i$$
, (7)

278
$$\bar{C}_{L,3} = \frac{1}{N} \sum_{i=1}^{N} C_{L,3}[i]$$
, (8)

279 Again, $\mathbf{b}[i]$ are background concentrations determined algorithmically, and are a function of C_{NR} , whereas C_{BG} , as in Sect. 3.1, 280 are physically measured concentrations. It is worth noting that while the constraint $\mathbf{b} \leq C_{NR}$ was applied in this algorithm, it is 281 not always the case that a background station will measure less than a near-road station during a given hour for a number of 282 different reasons. For example, Sofowote et al. (2018) showed that a receptor 167 m from the edge of Highway 401 measured $PM_{2.5}$ concentrations that exceeded concurrent measurements at NR-TOR-1 (10 m from the edge of the highway) ~5% of the 283 284 time based on half-hourly measurements. Regardless, the impact of this assumption on estimated average local concentration 285 is likely minimal. In using this algorithm, the width of the averaging window will affect the resulting baseline—windows that 286 are shorter in duration will result in more temporally varying baselines, while longer windows will result in flatter baselines. For information regarding function input parameters please refer to Wang et al. (2018). This study used the parameters $\alpha = 4$ 287 288 and W = 8 hr.

289 **3.3.2** Application to near-road ozone concentrations

Near roadways O_3 concentrations, unlike most other pollutants considered in this study, are generally less than background concentrations. This is because O_3 is formed through secondary chemistry in the troposphere, and one of its sinks is through reaction with NO, which is a primary pollutant emitted by vehicles and is therefore often abundant near roadways. Hence, transient emissions of NO from passing vehicle plumes will result in decreases in O_3 concentrations during a similar time scale. Background O_3 concentrations in the near-road environment were instead estimated by interpolating maximum values rather than minima. A baseline for $-O_3(t)$ was established, and the resulting output's sign flipped, effectively yielding an interpolation of maxima.

297 4 Results

298 4.1 Average differences between near-road and background sites

Over the duration of the study period average $C_{L,1}$ values were calculated using method 1, as described in Sect. 3.1, with resulting differences summarized in Tables 2-4. Note that no CO₂ difference was calculated between Vancouver stations because CO₂ was not measured at BG-VAN. 302 The background-subtracted differences were smallest at NR-TOR-2; for every TRAP measured, both NR-TOR-1 and NR-303 VAN saw greater $C_{1,1}$ concentrations in comparison. This pattern is consistent with the lower traffic volumes at NR-TOR-2. 304 Surprisingly, despite the drastic difference in traffic intensities between NR-VAN and NR-TOR-1, C_{L1} values at both sites 305 were remarkably similar for most TRAPs. This similarity was in part due to NR-VAN's closer proximity to the roadway (6 m) 306 compared with NR-TOR-1 (10 m), in conjunction with the significant fraction of diesel vehicles passing along Clark Drive 307 (Wang et al., 2018). While most $C_{1,1}$ concentrations were similar between these two locations, UFPs at NR-TOR-1 were significantly greater (3.0E+4 vs. 1.2E+4 cm⁻³). However, this may in part be due to seasonal bias in UFP data availability 308 309 (Table S1) between NR-TOR-1 and BG-TOR-1 (note especially the lack of concurrent data during summer months when 310 ambient UFP concentrations are often lowest).

The NO₂/NO_x ratios for $C_{L,1}$ at NR-TOR-2 were also markedly lower than the other near-road sites; these ratios at NR-VAN, NR-TOR-1, and NR-TOR-2 were, on average, 0.18, 0.29, and 0.61, respectively. A potential explanation for this is the relative residence times of vehicle plumes prior to detection at each site: because NR-VAN was positioned closest to the roadway, it is likely that vehicle plumes were fresher upon detection, whereas NR-TOR-2 sampled within an urban canyon where air tends to stagnate and recirculate. These results emphasize an important implication for near-road monitoring policies: while NO₂ alone is often regulated because of associated health effects, measurements of only NO₂ may not be a reliable metric for assessing near-road health impacts, as characteristics of the site may result in NO₂ being a negligible fraction of total NO_x.

The average differences for O_3 were negative, indicating that ozone concentrations tend to be lower near major roads. Ozone is presumably being titrated due to the higher near-road concentrations of NO. Furthermore, O_3 production in downtown Toronto and metropolitan Vancouver generally occurs in a VOC-limited regime, meaning that the additional NO_x near roads does not enhance local ozone formation (Ainslie et al., 2013; Geddes et al., 2009).

322 While $PM_{2.5}$ is generally considered to be a more regional and homogenous pollutant in urban environments, the observed values of C_{1.1} (1.48, 0.27, and 2.26 µg m⁻³ at NR-TOR-1, NR-TOR-2, and NR-VAN, respectively) were found to be 323 324 significantly greater than zero, and may be indicative of both primary tailpipe and non-tailpipe (e.g. brake wear, road dust 325 resuspension, etc.) emissions. A recent study by Jeong et al. (2019) characterized the sources and composition of PM_{2.5} at both 326 NR-TOR-1 and NR-TOR-2 using an X-ray fluorescence continuous metals monitor. They found that while concentrations of 327 aged organic aerosol, sulfate, and nitrate were similar between the two sites, contributions from sources such as traffic exhaust, 328 brake wear, and road dust differed significantly, and were the primary factors responsible for differences in average PM_{2.5} 329 concentrations. Another study by Sofowote et al. (2018), examined in more detail the reasons for elevated PM_{2.5} constituents 330 at NR-TOR-1, with particular emphasis on BC, relative to another receptor 167 m from Highway 401.

331 4.2 Downwind-upwind pollutant differences

As stated previously, NR-TOR-1 was the most ideal near-road monitoring location in this study for associating TRAP measurements with local meteorology, as it was positioned on flat terrain, and the major roadway which it was stationed next to was the only significant source of TRAPs in the immediate area. Thus, the direction of wind at this site had a significant 335 impact on measured pollutant concentrations (Fig. 1). Using the methods described in Sect. 3.2, hourly TRAP concentrations 336 were aggregated based on wind direction, and were classified as either being downwind, upwind, or neither. Downwind and 337 upwind averages were calculated across the entirety of the study period and their differences, $C_{1,2}$ are summarized also in 338 Tables 2-4. Additional information regarding the number of downwind/upwind hours and confidence intervals are provided in 339 the supplementary information (Sect. S2). Note that downwind and upwind conditions were generally not uniform with respect 340 to time of day (Fig. S2); however, it was found that even if downwind and upwind data occurred uniformly with respect to 341 time of day the impact it would have on average $C_{L,2}$ values is minimal for most pollutants (Tables S5 and S6). 342 The C_{L2} values reported in Table 2 for NR-TOR-1 correspond relatively well with, but are higher than, respective C_{L1} values.

This is true for most pollutants, with the exception of O_3 and $PM_{2.5}$. The reason local concentrations generated via method 2 ($C_{L,2}$) are generally greater than those generated via method 1 ($C_{L,1}$) is believed to be due to the following: when a site is directly downwind from a road it will generally experience greatest TRAP concentrations, as it is this case in which there is the smallest distance for dilution between the road and the site. In contrast, $C_{L,1}$ values were averaged across all meteorological scenarios. The fundamental differences between methods 1 and 2 is explored further in Sect. S3 in the supplementary information.

349 Unlike NR-TOR-1, NR-TOR-2 was not an ideal site for applying method 2 in a straightforward manner, as it measured air 350 samples within an urban canyon where micrometeorology was complicated by vortices, stagnation, and recirculation effects. 351 Using the downwind and upwind sector definitions in Sect. 3.2.2, $C_{1,2}$ values were calculated at NR-TOR-2 and are 352 summarized in Table 3. This methodology of contrasting downwind and upwind pollutant averages at NR-TOR-2 was unable 353 to produce meaningful differences and the resulting disagreement with the near-road-urban-background differences ($C_{L,1}$) is 354 evident. Associating ground-level TRAP concentrations with city-scale meteorology at this site was complicated by 355 surrounding urban architecture and the presence of an intersection approximately 50 m SW of the receptor. In actuality, the 356 difference calculated for this site was between that of leeward and windward in-canyon concentrations, and this difference was 357 not as substantial as the NR-TOR-2 and BG-TOR-2 average site difference. For these reasons, associating near-road pollutant 358 concentrations with meteorological data was not an effective way of differentiating between local and regional influences on 359 pollutant concentrations at this particular near-road site. In general, in order to attain this differentiation for measurements 360 made in urban canyons, more complicated meteorological models are necessary; hence, simple downwind/upwind differences 361 are not universally applicable to near-road monitoring data, especially for locations in heavily urbanized landscapes.

Lastly, the siting of NR-VAN was somewhere between NR-TOR-1 and NR-TOR-2 in terms of complexity in associating TRAP concentrations with meteorology. The presence of densely spaced residential buildings within the immediate vicinity of the measurement station resulted in surface roughness having an effect on winds carrying TRAPs from major roadways farther away. Despite this, the differences between average downwind and upwind TRAP concentrations at NR-VAN were similar to, albeit larger, than the NR-VAN/BG-VAN differences in Table 4, a result similar to that for NR-TOR-1. The fact that consistent results were seen for NR-VAN and NR-TOR-1 but not NR-TOR-2 underlines the importance of a station's location, surrounding obstructions to winds, and location of traffic sources, and that associating near-road TRAP

- 369 concentrations with meteorological variability should be done with caution, taking into account the subtleties of each site's
- 370 environ. The apparent stronger influence of the intersection rather than traffic directly next to NR-VAN (i.e. winds originating
- 371 from 90°; see Fig. 3), despite Clark Drive being 6 m vs the intersection being 65 m away, may seem paradoxical. We speculate
- 372 that the acceleration of southbound traffic along Clark Drive at this intersection was the main source of emissions, while
- 373 coasting past the site, particularly when slowing down for the stop light, would have contributed much less.

374 **4.3 Local concentrations inferred from baseline subtraction**

Method 3, as described in Sect. 3.3.1, was applied to hourly pollutant concentrations, and the algorithm input parameters used were $\alpha = 4$ and W = 8 hr. From the output, C_{L,3} was determined as a function of time, and then averaged across the entirety of the measurement campaign; the resultant averages are summarized in Table 2-4 for each near-road site.

A benefit to this method was that it was able to estimate local and background CO_2 concentrations at NR-VAN, where CO_2 measurements were made only in the near-road environment and not at the background site. This emphasizes a key advantage to approaches such as these: traffic-related signal can be isolated from near-road measurements alone, without the need for background or even meteorological measurements. Furthermore, this differentiation was performed on an hourly basis, thereby retaining information in the time domain, which was not possible with method 2.

383 Across all near-road locations, average C_{L3} concentrations were quite similar to respective average C_{L1} values, implying that 384 method 3, which uses only near-road data, is a robust means of estimating urban background and local traffic-related pollutant 385 concentrations. This was true even for NR-TOR-2, where micrometeorology complicated analysis using method 2. Fine 386 particulate matter was an exception to this, however. Regarding $PM_{2.5}$, because its signal was largely dominated by regional-387 scale sources and dynamics, temporal fluctuations in roadside PM2.5 concentrations generally varied more slowly than those 388 of primary pollutants such as NO or BC, for example. Furthermore, this variability is generally meteorologically-driven and 389 occurs homogeneously over large areas (10s of kilometres); we posit that these variabilities associated with meteorology were 390 falsely attributed to local signal, causing local PM_{2.5} concentrations ascertained through this method to be much higher than respective $C_{1,1}$ concentrations. Lastly, for ambient concentrations < 80 µg m⁻³, the hourly precision of the SHARP 5030 is ±2 391 392 μg m⁻³. So, the average site differences between near-road and background sites, which are all around 2 μg m⁻³ or less, are 393 likely too small for method 3 to isolate as the signal-to-noise ratio on an hourly basis is quite small.

The choice of time window parameter, when comparing results obtained from method 1, is both site-specific and pollutantdependent. For example, shorter time windows will produce results that are in better agreement with stations that are closer in proximity. Further, the role of secondary chemistry will affect agreement between method 1 and method 3. Variability in $C_{L,3}$ is shown in Table S9, where average $C_{L,3}$ values are reported for W = 6 and W = 14. When comparing average $C_{L,3}$ values to average $C_{L,1}$ values as a function of W, it appears as though some pollutants produce better agreement for smaller W values (e.g. CO₂ and PM_{2.5}), whereas others agree better for larger values of W (e.g. UFPs). This is likely due to the relative homogeneity of PM_{2.5} and CO₂ and heterogeneity of UFP concentrations in urban environments. Generally, however, it 401 appears that the values $\alpha = 4$ and W = 8 hr are an appropriate middle-ground for the pollutants considered in this study, and

402 likely represent an urban background spatial scale of between 5 and 10 km.

Although application of method 3 was less suitable for some pollutants (i.e. PM_{2.5}), it appears to behave in an accurate and 403 404 robust manner for most others. Comparing $C_{L,1}$ and $C_{L,3}$ values in Tables 2-4, it appears that method 3 produced similar results 405 when compared with method 1, with the added benefit of retaining information in the time domain and not requiring a second 406 site. It is worth emphasizing that method 3 was an independently developed method for background-subtracting near-road data without the need for concurrent background measurements. The parameters $\alpha = 4$ and W = 8 hr were originally chosen to be 407 408 generalizable for near-road measurements, and to differentiate similar local/regional scales. While a direct comparison with 409 method 1 to assess the accuracy of method 3 is tempting, method 1 is not without its own limitations (i.e. differences in distance 410 between near-road and background stations, difficulty in removing background stations from local sources, etc.). Thus, while 411 this comparison is useful for understanding the spatial scales of different pollutants, background-subtraction parameters should 412 not necessarily be chosen based on this alone.

413 4.4 Comparison of background subtraction methods

414 Three techniques were applied to the near-road monitoring locations in this study to extract information regarding local TRAP 415 concentrations: 1. Average differences between near-road and urban background locations, 2. Downwind-upwind differences 416 in near-road measurements, and 3. Average concentrations inferred through time-series analysis of near-road data. Generally, 417 methods 1 and 3 agreed well with one another, whereas method 2 produced values that were high in comparison with the other 418 two methods at NR-TOR-1 and NR-VAN, and generated results that were close to zero at NR-TOR-2. A comparison of the 419 three methodologies is summarized graphically in the supplementary information (Fig. S4-S6). The close agreement of 420 methods 1 and 3, which describe the average concentrations attributed to local traffic, is encouraging, suggesting a background 421 is inferable from near-road data alone using method 3. Method 2 was able to isolate traffic-related pollutant signal for NR-422 VAN and NR-TOR-1, but was not feasible for NR-TOR-2, thus highlighting a drawback of relying exclusively on wind 423 direction data for source apportionment efforts. It is believed that method 2, while useful for isolating traffic-related pollution, 424 is less relevant for epidemiological purposes as it only considers certain meteorological scenarios.

425 **4.5 Application of local concentrations**

Subtraction of background concentrations allows the influences of local traffic on near road TRAP concentrations to be assessed. The benefits in terms of improved understanding were examined and illustrated by applying the local concentrations thereby derived in two ways. The degree to which traffic influences TRAP concentrations beside a road can vary day-to-day depending on the prevailing meteorology. Using the local signal allowed the magnitude of this source of variability to be assessed in a manner that is consistent across most TRAPs and across all near-road sites. In contrast, the contribution of traffic to the total concentration will differ across pollutants. For example, some pollutants such as NO may be predominantly from traffic while others such as CO₂ will be dominated by the background. Separating the local and background concentrations allowed assessment of how the portion from local traffic varied between sites and across the pollutants. Effectively, the
background subtraction methodology provided estimates that illustrate how much concentrations beside a road would drop if

435 all the traffic on that road were to be removed, as concentrations would converge to that of the urban background in that case.

436 4.5.1 Effect of meteorology on local TRAP variability

437 Using the hourly values of $C_{L,3}$ at each near-road station determined using method 3 in Sect. 3.3.1, the roles of individual 438 meteorological parameters on the variability of these local concentrations were explored. While roadside concentrations are 439 affected by meteorology in a number of ways, local pollutant quantities—of interest are those from vehicular exhaust—are 440 expected to behave in a more predictable manner in comparison, and indeed there are many means in which to predict the 441 evolution of these exhaust plumes, from simple dispersion models to computational fluid dynamics. Here, however, a more 442 simplified means of underlining the effect of wind on above-background TRAP concentrations was utilized: local TRAP 443 concentrations normalized to their mean values were associated with both the direction and speed of local winds, the former 444 showing the effect of downwind/upwind variability and the latter showing that of dilution. Normalization allowed results to 445 be more comparable between sites and pollutants where mean emission rates of TRAPs may differ. While different receptor 446 distances from a roadway will lead to different absolute concentrations measured, it is assumed here that when these 447 concentrations are normalized to their mean that the trends with respect to meteorology will be similar. Because NR-TOR-2 448 was situated within an urban canyon, the effect of meteorology on its measured concentrations was not relatable to the other 449 two stations in this study; for this reason it is omitted from this section.

450 4.5.2 Wind direction

451 Wind direction can have a large influence on roadside TRAP concentrations. Shown in Fig. 5 is the dependence of normalized 452 local pollutant concentrations on wind direction at both NR-VAN and NR-TOR-1. Generally, downwind measurements have 453 the effect of enhancing local concentrations by a factor of ~ 1.5 -2.0, whereas upwind conditions suppress local concentrations 454 by a factor of ~4.0, with respect to the mean. Note that these upwind concentrations did not necessarily converge to zero as 455 hourly averages were used to create these trends. It is also conceivable that during upwind periods, local turbulence from traffic 456 and/or brief shifts in wind direction resulted in some degree of plume capture. It would appear that, on an hourly-averaged 457 basis, traffic's contribution to local TRAP variability (i.e. irrespective of background pollution) at a near-road receptor may 458 change by a factor of six to eight depending on the average direction of wind.

As shown in Fig. 5, a clear sinusoidal wind direction dependency is apparent at NR-VAN and NR-TOR-1, with similar ranges in enhancement and suppression at both sites. However, at NR-VAN, there appears to be two modes in concentration enhancement. The Clark Drive and 12th Avenue intersection, located approximately 65 m from the receptor, had an influence on local TRAPs originating from the south. However, given its distance, west/eastbound traffic along 12th avenue should not have had an influence similar to that of Clark Drive which was only 6 m away. We postulate that the traffic lights at the intersection caused stop-and-go patterns in which southbound traffic on Clark Drive was often backed up to the monitoring 465 location, and it is these driving patterns that are believed to be associated with the enhancement seen between the wind 466 directions of $100^{\circ}-200^{\circ}$ at NR-VAN.

When comparing methods of background subtraction, it was shown that method 2 yielded higher estimates of the local concentrations in comparison with the other two methodologies, as further explored in Sect. S3 of the supplementary data. Across pollutants, it was found that on average this downwind/upwind difference resulted in local TRAP concentrations that were factors of 1.3 and 1.4 times greater than those inferred from method 1 at NR-VAN and NR-TOR-1, respectively (Table S8). In short, this corresponds well with above-average normalized local pollutant concentrations during downwind conditions at both sites (Fig. 5), during which conditions values of $C_{L,3}$ were found to be similar factors greater than the mean at both sites (Table S8).

474 Lastly, it is of interest to note that hourly upwind $C_{L,3}$ concentrations at either site yielded non-zero local concentrations. It is 475 indeed likely that at an hourly time-resolution some plume capture will occur during predominately upwind conditions; 476 however, this seems to carry with it the implication that upwind analysis at a near-road location may overestimate background 477 concentrations. To test this, average upwind concentrations were compared with average concentrations measured at each 478 nearest background location, the results of which are summarized in Table S7. Generally, the two appear to agree well with 479 one another, and so any plume capture during upwind conditions apparently produced a negligible impact on total 480 concentrations.

481 **4.5.3 Wind speed**

Similar to the analysis in the previous section, the effect of wind speed on roadside TRAP concentrations was explored at NR-TOR-1 and NR-VAN, and consistent results were found between them. Under stagnant conditions (wind speeds of ~1.0 m s⁻¹), local pollutant quantities were found to be enhanced by factors of ~2.0 and ~1.7 at NR-VAN and NR-TOR-1, respectively, and high wind speeds (> 10 m s⁻¹) suppressed these quantities by a factor of ~2.0 at both sites (Fig. 6), giving an overall influence factor of 3.4 to 4. The maximum levels of enhancement and suppression were slightly smaller than the results found for wind direction, implying a slightly smaller or equivalent importance on local TRAP concentrations at a given roadside receptor. The relation used to model the effect of wind speed on normalized local concentrations was the following:

$$489 \quad \frac{c_{L,3}}{\bar{c}_{L,3}} = \frac{c_1}{WS^{c_2}},\tag{9}$$

where $C_{L,3}$ are local pollutant concentrations determined through method 3, c_1 and c_2 are regression parameters, and WS is wind speed as measured at the station. Indeed, more involved models have been shown to better represent the wind speed dependency of specific pollutants (Jones et al., 2010); however, simplicity is preferred here so as to generalize results across sites and pollutants.

On average, the regression parameters c_1 and c_2 were found to be ~2.0 and ~0.6 for NR-VAN, and ~1.6 and ~0.5 for NR-TOR-1, respectively (Table S10). Section S5.1 in the supplementary information compares these results between weekdays and weekends. While different c_1 parameters were determined for both sites, presumably due to their difference in roadway 497 proximity, similar c₂ parameters between 0.5-0.6 were found. The c₂ parameter, which embodies the wind speed-pollutant 498 decay relationship, is expected to be independent of a station's proximity to the roadway. As with the wind direction analysis in the previous section, these associations with respect to wind speed were averaged from two years of hourly data across the entire study domain, meaning they were acquired from a range of pollutants, traffic conditions, wind directions, and times of day. While less descriptive from a mechanistic perspective, these results are intended to be more representative of the ranges of variability in average above-background exposure levels in the immediate area.

503 **4.6 Fraction of near-road pollution attributable to local sources**

504 The time-series based estimates of the background concentrations were also applied to estimate the portion of the pollutant 505 concentrations that were due to local traffic. For example approximately half of total BC concentrations were estimated to be 506 due to local sources at NR-TOR-1 with lower and higher percent contributions at NR-TOR-2 and NR-VAN, respectively (Fig. 507 7). The contribution of local sources varied across the pollutants; NO had the highest local contribution at the near road sites while CO₂ had the lowest (Fig. 8). Further, this methodology was able to replicate trends in weekday/weekend background 508 509 pollution variability—shown in Fig. 7 is BC, for example, with others in the supplementary (Fig. S7-S12). Local components 510 of air pollution showed far greater differences between weekdays and weekends at each near-road monitoring location, 511 emphasizing the effect of different on-road traffic conditions between these two sets of days. Generally, TRAP concentrations 512 measured at urban background sites were slightly higher on weekdays compared to weekends, and this change in regional 513 pollution was captured in the background contributions extracted from the near-road data. It should be expected that average 514 concentrations measured at BG-TOR-1 should match the background elements of NR-TOR-1 reasonably well, with a similar 515 argument to be made for BG-TOR-2 and NR-TOR-2; however, these urban background concentrations are likely not perfectly 516 homogeneous throughout the city. The spatial difference between BG-TOR-1 in north Toronto and BG-TOR-2 in south 517 Toronto was 20 km, and the difference in average pollutant levels between the two reflects this.

518 5 Conclusions

519 In this study TRAP concentrations were measured continuously at time resolutions of one hour or finer for over two years at 520 three near-road and three urban background locations. Three methods were explored for estimating the contribution of local 521 and regional/background sources on near-road measurements: differences between average measurements taken near-road and 522 at a nearby urban background location, downwind-upwind analysis at the near-road location, and time-series analysis of near-523 road pollutant data. Generally, the near-road vs urban background and time-series analysis methods produced results that were 524 in good agreement; these values represent contributions to TRAP due to local traffic averaged over all wind directions. The 525 downwind-upwind method yielded local concentrations that were higher than the average station differences by approximately 526 40%; this was attributable to the downwind/upwind analysis isolating the conditions where traffic has the greatest impact on 527 a site while the average differences included data across all wind conditions.

The time-series analysis method was an accurate and robust means of differentiating local and regional signal, with the added benefits of being applicable across all near-road sites, not being constrained to certain meteorological scenarios or requiring a separate background site, and retaining information in the time domain. This methodology is recommended for future use in applications such as: determining the impact of local on-road traffic to a roadside receptor, isolating background concentrations from ambient data for use in dispersion modelling, and obtaining above-background concentrations for fleet emission factor calculations, for example.

534 Lastly, to demonstrate the value in isolating the influence of local sources at an hourly time resolution, local TRAP concentrations determined using time-series analysis were compared with meteorological variables at two of the near-road 535 536 sites, NR-VAN and NR-TOR-1. This analysis yielded trends that were similar between sites and generalizable across all 537 measured pollutants, with the exception of $PM_{2.5}$ and O_3 . Wind direction had a factor of influence of approximately seven at 538 both near-road sites, while the effect of wind speed was found to be slightly smaller, varying local hourly concentrations by a 539 factor of four, with highest concentrations seen during stagnant conditions and lowest concentrations as wind speed became 540 large. Both sites exhibited similar decays in local concentration with respect to wind speed; proportionality to wind speed was 541 found to be between WS^{-0.5} and WS^{-0.6}.

542 Author contribution

543 AM, LW, CA, DH, JRB, and GJE designed and initiated the near-road monitoring study. Data collection and quality assurance 544 from Torontonian stations was performed by: NH, JMW, CHJ, RMH, US, JD, YS, and MN, while GD was responsible for the 545 two stations in Vancouver. NH prepared the manuscript, with contributions from all co-authors, and performed all data 546 analysis.

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552 Competing interests

- 553 The authors declare they have no conflict of interest.
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557 References

Ainslie, B., Steyn, D. G., Reuten, C., and Jackson, P. L.: A Retrospective Analysis of Ozone Formation in the Lower Fraser
Valley, British Columbia, Canada. Part II: Influence of Emissions Reductions on Ozone Formation, Atmos. Ocean., 51:2, 170186, doi:10.1080/07055900.2013.782264, 2013.

561

Andersen, Z. J., Hvidberg, M., Jensen, S. S., Ketzel, M., Loft, S., Sorensen, M., Tjonneland, A., Overvad, K., and RaaschouNielsen, O.: Chronic Obstructive Pulmonary Disease and Long-Term Exposure to Traffic-related Air Pollution, Am. J. Resp.
Crit. Care., 183, 455-461, doi:10.1164/rccm.201006-0937OC, 2011.

565

Baldwin, N., Gilani, O., Raja, S., Batterman, S., Ganguly, R., Hopke, P., Berrocal, V., Robins, T., and Hoogterp, S.: Factors 566 567 affecting pollutant concentrations in the near-road environment, Atmos. Env., 115. 223-235. 568 doi:10.1016/j.atmosenv.2015.05.024, 2015.

569

Belis, C. A., Karaguilian, F., Larsen, B. R., and Hopke, P. K.: Critical review and meta-analysis of ambient particulate matter
source apportionment using receptor models in Europe, Atmos. Env., 69, 94-108, doi:10.1016/j.atmosenv.2012.11.009, 2012.

Brantley, H. L., Hagler, G. S. W., Kimbrough, E. S., Williams, R. W., Mukerjee, S., and Neas, L. M.: Mobile air monitoring
data-processing strategies and effects on spatial air pollution trends, Atmos. Meas. Tech., 7, 2169-2183, doi:10.5194/amt-72169-2014, 2014.

576

Evans, G. J., Jeong, C-H, Sabaliauskas, K., Jadidian, P., Aldersley, S., Larocque, H., and Herod, D.: Design of a Near-Road
Monitoring Strategy for Canada, A Final Report to Environment Canada, SOCAAR, Toronto, 1-60, 2011.

579

Galvis, B., Bergin, M., and Russell, A.: Fuel-based fine particulate and black carbon emission factors from a railyard in Atlanta,
J. Air. Waste. Manage., 63, 648-658, doi:10.1080/10962247.2013.776507, 2013.

582

Geddes, J. A., Murphy, J. G., and Wang, D. K.: Long term changes in nitrogen oxides and volatile organic compounds in
Toronto and the challenges facing local ozone control, Atmos. Env., 43, 3407-3415, doi:10.1016/j.atmosenv.2009.03.053,
2009.

586

Gomez-Losada, A., Pires, J. C. M., and Pino-Mejias, R.: Modelling background air pollution exposure in urban environments:
Implications for epidemiological research, Environ. Modell. Softw., 106, 13-21, doi:10.1016/j.envsoft.2018.02.011, 2018.

- Jeong, C-H., Evans, G. J., Healy, R. M., Jadidian, P., Wentzell, J., Liggio, J., and Brook, J. R.: Rapid physical and chemical
 transformation of traffic-related atmospheric particles near a highway, Atmos. Pollut. Res., 6, 662-672,
 doi:10.5094/APR.2015.075, 2015.
- 593

Jeong, C-H., Wang, J. M., Hilker, N., Debosz, J., Sofowote, U., Su, Y., Noble, M., Healy, R. M., Munoz, T., Dabek-Zlotorzynska, E., Celo, V., White, L., Audette, C., Herod, D., and Evans, G. J.: Temporal and spatial variability of trafficrelated PM_{2.5} sources: Comparison of exhaust and non-exhaust emissions, Atmos. Env., 198, 55-69, doi:10.1016/j.atmosenv.2018.10.038, 2019.

- 598
- Jones, A. M., Harrison, R. M., and Baker, J.: The wind speed dependency of the concentrations of airborne particulate matter and NO_x, Atmos. Env., 44, 1682-1690, doi:10.1016/j.atmosenv.2010.01.007, 2010.
- 601
- Kimbrough, S., Hanley, T., Hagler, G., Baldauf, R., Snyder, M., and Brantley, H.: Influential factors affecting black carbon
 trends at four sites of differing distance from a major highway in Las Vegas, Air. Qual. Atmos. Hlth., 11, 181-196,
 doi:10.1007/s11869-017-0519-3, 2018.
- 605
- Klems, J. P., Pennington, M. R., Zordan, C. A., and Johnston, M. V.: Ultrafine Particles Near a Roadway Intersection: Origin
 and Apportionment of Fast Changes in Concentration, Environ. Sci. Technol., 44, 7903-7907, doi:10.1021/es102009e, 2010.
- Ma, N. and Birmili, W.: Estimating the contribution of photochemical particle formation to ultrafine particle number averages
 in an urban atmosphere, Sci. Total. Environ., 512-513, 154-166, doi:10.1016/j.scitotenv.2015.01.009, 2015.
- 611
- Molina, M. J. and Molina, L. T.: Megacities and Atmospheric Pollution, J. Air. Waste. Manage., 54, 644-680,
 doi:10.1080/10473289.2004.10470936, 2004.
- 614
- 615 Oke, T. R.: Street Design and Urban Canopy Layer Climate, Energ. Buildings., 11, 103-113, doi:10.1016/0378-616 7788(88)90026-6, 1988.
- 617
- Pant, P. and Harrison, R. M.: Estimation of the contribution of road traffic emissions to particulate matter concentrations from
 field measurements: A review, Atmos. Env., 77, 78-97, doi:10.1016/j.atmosenv.2013.04.028, 2013.
- 620
- Sabaliauskas, K., Jeong, C-H., Yao, X., Jun, Y-S., Jadidian, P., and Evans, G. J.: Five-year roadside measurements of ultrafine
 particle in a major Canadian city, Atmos. Env., 49, 245-256, doi:10.1016/j.atmosenv.2011.11.052, 2012.
- 623

- Sabaliauskas, K., Jeong, C-H., Yao, X., and Evans, G. J.: The application of wavelet decomposition to quantify the local and
 regional sources of ultrafine particles in cities, Atmos. Env., 95, 249-257, doi:10.1016/j.atmosenv.2014.05.035, 2014.
- Saha, P. K., Khlystov, A., Snyder, M. G., and Grieshop, A. P.: Characterization of air pollutant concentrations, fleet emission
 factors, and dispersion near a North Carolina interstate freeway across two seasons, Atmos. Env., 177, 143-153,
 doi:10.1016/j.atmosenv.2018.01.019, 2018.
- 630
- 631 Shairsingh, K. K., Jeong, C-H., Wang, J. M., and Evans, G. J.: Characterizing the spatial variability of local and background 632 concentration signals for air pollution the neighbourhood 183. 57-68. at scale. Atmos. Env.. 633 doi:10.1016/j.atmosenv.2018.04.010, 2018.
- 634
- Sofowote, U. M., Healy, R. M., Su, Y., Debosz, J., Noble, M., Munoz, A., Jeong, C-H., Wang, J. M., Hilker, N., Evans, G. J.,
 and Hopke, P. K.: Understanding the PM_{2.5} imbalance between a far and near-road location: Results of high temporal frequency
 source apportionment and parameterization of black carbon, Atmos. Env., 173, 277-288, doi:10.1016/j.atmosenv.2017.10.063,
 2018.
- 639
- Tchepel, O. and Borrego, C.: Frequency analysis of air quality time series for traffic related pollutants, J. Environ. Monitor.,
 12, 544-550, doi:10.1039/b913797a, 2010.
- 642
- Vardoulakis, S., Fisher, B. E. A., Pericleous, K., and Gonzalez-Flesca, N.: Modelling air quality in street canyons: a review,
 Atmos. Env., 37, 155-182, doi:10.1016/S1352-2310(02)00857-9, 2003.
- 645
- Wang, J. M., Jeong, C-H, Zimmerman, N., Healy, R. M., Wang, D. K., Ke, F., Evans, G. J.: Plume-based analysis of vehicle
 fleet air pollutant emissions and the contribution from high emitters, Atmos. Meas. Tech., 8, 3263-3275, doi:10.5194/amt-83263-2015, 2015.
- 649

Wang, J. M., Jeong, C-H., Hilker, N., Shairsingh, K. K., Healy, R. M., Sofowote, U., Debosz, J., Su, Y., McGaughey, M.,
Doerksen, G., Munoz, T., White, L., Herod, D., and Evans, G. J.: Near-Road Air Pollutant Measurements: Accounting for
Inter-Site Variability Using Emission Factors, Environ. Sci. Technol., 52, 9495-9504, doi:10.1021/acs.est.8b01914, 2018.

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Station ID	Latitude	Longitude	Major Roadway	Annual Average Daily Traffic (AADT)	Distance from Roadway [m]
NR-TOR-1	43.7111	-79.5433	Highway 401	405,500	10
BG-TOR-1	43.7806	-79.4675	-	-	-
NR-TOR-2	43.6590	-79.3954	College Street	17,200	15
BG-TOR-2	43.6122	-79.3887	-	-	-
NR-VAN	49.2603	-123.0778	Clark Drive	33,100	6

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658	Table 1: IDs.	locations.	name of mai	or roadway.	and averag	e dailv tr	affic intensi	tv for eacl	1 monitoring	location.
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BG-VAN

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684 Table 2: Mean local pollutant concentrations at NR-TOR-1 determined using each background-subtraction method.

Pollutant	М	ethod 1	Method 2			Method 3	
	N (hours)	$C_{L,1} \pm 95\% CI$	C _{DW}	C _{UW}	C _{L,2}	N (hours)	$C_{L,3} \pm 95\% CI$
NO [ppb]	14169	21.5 ± 0.4	37.8	2.9	34.9	15524	18.3 ± 0.4
NO ₂ [ppb]	13765	8.7 ± 0.1	21.2	10.7	10.5	15087	9.2 ± 0.1
CO [ppb]	6479	103.2 ± 2.7	364.4	226.6	137.9	13008	114.6 ± 2.2
CO ₂ [ppm]	7900	14.4 ± 0.6	437.3	416.4	20.9	14812	$19.6\ \pm 0.4$
O ₃ [ppb]	13753	$\textbf{-5.9}\pm0.1$	15.3	33.2	-17.9	15181	-12.3 ± 0.2
PM _{2.5} [µg m ⁻³]	14170	1.48 ± 0.06	7.68	9.01	-1.33	15484	4.30 ± 0.08
UFP [cm ⁻³]	5212	29600 ± 800	57000	15300	41700	12683	22754 ±449
BC [µg m ⁻³]	8036	1.03 ± 0.03	2.13	0.73	1.4	15443	$1.01 \hspace{0.1 cm} \pm \hspace{0.1 cm} 0.02$

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702	Table 3: Mean local	ollutant concentrations at NR-TOR-2 determined using each background-	subtraction method.

Pollutant	Method 1			Method 2		Method 3	
	N (hours)	$C_{L,1}\pm95\%CI$	C_{DW}	$C_{\rm UW}$	$C_{L,2}$	N (hours)	$C_{L,3} \pm 95\% CI$
NO [ppb]	13768	3.5 ± 0.1	6	3.2	2.8	14937	3.8 ± 0.1
NO ₂ [ppb]	11211	5.4 ± 0.1	8.5	10.4	-1.9	12359	5.3 ± 0.1
CO [ppb]	13603	72.3 ± 1.5	247.9	246.8	1.1	15152	68.7 ± 1.3
CO ₂ [ppm]	10686	10.6 ± 0.4	423.1	421.4	1.7	14626	13.3 ± 0.2
O ₃ [ppb]	15109	-2.9 ± 0.1	24.2	28.7	-4.5	15827	$-9.0\ \pm 0.1$
PM _{2.5} [µg m ⁻³]	15193	0.27 ± 0.05	3.8	9.01	-5.21	15730	2.92 ± 0.06
UFP [cm ⁻³]	7400	7400 ± 200	12900	16700	-3800	14931	7088 ± 108
BC [μg m ⁻³]	14740	0.34 ± 0.01	0.63	0.81	-0.18	15451	0.41 ± 0.01

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720	Table 4: Mean local pollutant concentrations at NR-VAN determined using each background-subtraction method.	
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Pollutant	Method 1			Method 2		Method 3	
	N (hours)	$C_{L,1} \pm 95\% CI$	$C_{\rm DW}$	C _{UW}	$C_{L,2}$	N (hours)	$C_{L,3} \pm 95\%$ CI
NO [ppb]	10647	23.0 ± 0.5	56.6	9.7	46.8	15134	27.6 ± 0.6
NO ₂ [ppb]	10666	5.1 ± 0.1	21.9	11.5	10.4	15148	9.7 ± 0.1
CO [ppb]	9435	95.7 ± 2.3	414.3	210.1	204.2	13935	153.3 ± 3.4
CO ₂ [ppm]	-	-	461.6	414.5	47.1	13503	39.0 ± 0.7
O ₃ [ppb]	10535	-3.9 ± 0.1	9.4	19.7	-10.3	15016	-10.6 ± 0.1
PM _{2.5} [µg m ⁻³]	10491	2.26 ± 0.07	8.81	5.57	3.23	14879	3.99 ± 0.10
UFP [cm ⁻³]	9452	11600 ± 300	30000	14000	16000	14463	15252 ± 251
BC [µg m ⁻³]	10728	1.18 ± 0.02	2.48	0.84	1.64	15312	1.26 ± 0.02



Figure 1: Satellite image of the NR-TOR-1 site, along with upwind (blue) and downwind (red) quadrant definitions. Meteorological measurements were taken on top of a 10 m mast at the location of the station (labelled: NR-TOR-1) (a). Average ambient CO₂ concentrations by wind direction, with upwind and downwind definitions again highlighted in blue and red, respectively. Error bars

- 732 are 95% confidence intervals on the mean (b).

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Figure 2: Satellite image of the NR-TOR-2 site, along with upwind (blue) and downwind (red) quadrant definitions. Meteorological measurements were recorded on the roof of the facility (labelled: NR-TOR-2) (a). Average ambient CO₂ concentrations by wind direction, with upwind and downwind definitions again highlighted in blue and red, respectively. Error bars are 95% confidence intervals on the mean (b).







Figure 3: Satellite image of the NR-VAN site, along with upwind (blue) and downwind (red) sector definitions. Meteorological measurements were recorded on a 10 m mast above the station's location (labelled: NR-VAN) (a). Average ambient CO₂ concentrations by wind direction, with upwind and downwind definitions again highlighted in blue and red, respectively. Error bars are 95% confidence intervals on the mean (b)

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Figure 4: Method 3 applied to hourly CO₂ concentrations (black) measured at NR-TOR-2. The effect of varying the input parameters
 α and W are shown in blue, orange, and green.

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- TRAPs.







Figure 6: Normalized local pollutant concentrations determined using method 3 as a function of wind speed at NR-VAN (a) and NR TOR-1 (b). Solid lines indicate the average trend amongst all TRAPs, and shaded areas indicate the range of variability between
 TRAPs.

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Figure 7: Black carbon concentrations at each monitoring location in this study. Each site is separated by weekday and weekend, and bars are stacked according to concentrations attributed to local and regional sources. Background stations are presumed fully regional and therefore contain no local component.

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Figure 8: Average fraction of near-road measurements attributed to local sources, as determined by method 3, for each near-road monitoring location.