Impact of land-water sensitivity contrast on MOPITT retrievals and trends over a coastal city

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Abstract.

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We compare MOPITT Version 7 (V7) Level 2 (L2) & Level 3 (L3) carbon monoxide (CO) products for the 1° x 1° L3 gridbox containing the coastal city of Halifax, Canada (longitude = -63.58°, latitude = 44.65°), with a focus on the seasons DJF and JJA, and highlight a limitation in the L3 products that has significant 10 consequences for the temporal trends in near-surface CO identified using those data. Because this gridbox straddles the coastline, the MOPITT L3 products are created from the finer spatial resolution L2 products that are retrieved over both land and water, with a greater contribution from retrievals over water because more of the gridbox lies over water than land. We create alternative L3 products for this gridbox by separately averaging the bounded L2 retrievals over land (L3L) and water (L3W) and demonstrate that profile and total 15 column CO (TCO) concentrations, retrieved at the same time, differ depending on whether the retrieval took place over land or water. These differences (ΔRET) are greatest in the lower troposphere (LT), where mean retrieved VMRs are greater in L3W than L3L, with maximum mean differences of 11.6 % (14.3 ppbv, p = 0.001) at the surface level in JJA. Retrieved CO concentrations are more similar, on average, in the middle and upper troposphere (MT and UT), although large differences (in excess of 40 %) do infrequently occur. 20 TCO is also greater in L3W than L3L in both seasons. By analyzing L3L and L3W retrieval averaging kernels and simulations of these retrievals, we demonstrate that, in JJA, ΔRET is strongly influenced by differences in retrieval sensitivity over land and water, especially close to the surface where L3L has significantly greater information content than L3W. In DJF, land-water differences in retrieval sensitivity are much less pronounced and appear to have less of an impact on ΔRET , which analysis of wind directions suggests is 25 more likely to reflect differences in true profile concentrations (i.e. "real" differences). The original L3 timeseries for the 1° x 1° gridbox containing Halifax (L3O) corresponds much more closely to L3W than L3L, owing to the greater contribution from L2 retrievals over water than land. Thus, in JJA, variability in retrieved CO concentrations close to the surface in L3O is suppressed compared to L3L, and a declining trend detected using Weighted Least Squares (WLS) regression analysis is significantly slower in L3O (strongest 30 surface level trend identifiable = $-1.35 (\pm 0.35)$ ppbv y⁻¹) than L3L (-2.85 (± 0.60) ppbv y⁻¹). This is because contributing L2 retrievals over water are closely tied to a priori CO concentrations used in the retrieval, owing to their lack of near-surface sensitivity in JJA, and these are based on monthly climatological CO profiles from a chemical transport model and therefore have no yearly change (surface level trend in L3W = -0.60

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 (± 0.33) ppbv y⁻¹). Although our analysis focuses on DJF and JJA, we demonstrate that the findings also apply to MAM and SON. The results that we report here suggest that similar analyses be performed for other coastal cities before using MOPITT surface CO.

1 Introduction

The Measurement of Pollution in the Troposphere (MOPITT – Drummond et al., 2010, 2016; Acronyms defined in Appendix A) instrument is one of a large fleet of satellite-borne instruments capable of observing 40 the composition of the Earth's atmosphere from space. The target gas for MOPITT is carbon monoxide (CO), which is emitted from a range of anthropogenic (e.g. fossil fuel use) and natural (e.g. wildfires) sources, produced via the oxidation of methane and other volatile organic compounds, and has an atmospheric lifetime of weeks to months depending on season and location (e.g. Duncan et al., 2007). CO is therefore vital to monitor as a pollutant in its own right, as a tracer of local and transported pollution sources, and also because 45 it plays an important role in atmospheric chemistry, i.e. as a precursor to ozone formation and a primary sink for the hydroxyl radical. While multiple sensors observe CO (see e.g. Worden et al. (2013) for a comparison of CO trends from four satellite instruments), the unique strength of MOPITT lies in its nearly unbroken record of observations since launch in December 1999. This makes MOPITT data very valuable for the analysis of temporal trends in CO concentrations (e.g. He et al., 2013; Worden et al., 2013; Strode et al., 50 2016).

MOPITT retrieves coarse vertical resolution CO profiles in the troposphere by inverting observed upwelling radiances at thermal infrared (TIR) and near infrared (NIR) wavelengths (Deeter et al., 2013). These profiles are integrated to give CO total column amounts (TCO). In addition to several other inputs,
MOPITT's optimal estimation retrieval algorithm requires a priori information – among which is a description of the most probable state of the CO profile and its variability – to obtain physically realistic results (Pan et al., 1998; Rodgers, 2000; the retrieval algorithm is outlined in more detail in Section 2.1.1). The proportion of information about CO concentrations in each individual retrieval that comes directly from the satellite measurement, as opposed to the a priori, is highly variable. It depends on scene-specific factors such as surface temperature, thermal contrast in the lower troposphere, and the actual ("true") CO loading itself, as well as on instrumental noise (e.g. Deeter et al., 2015). This complicates the interpretation of retrievals, thus placing great importance on the analysis of retrieval averaging kernels (AKs), which represent the sensitivity of each retrieved profile point to the true CO profile and quantify the overall information content of the retrieval (as described in detail by, e.g., Deeter et al. (2007, 2015) and Rodgers (2000)). The

65 lower the retrieval information content, the closer the retrieved CO loading will be to the a priori, which is based on a climatological model value. Retrievals with little information content should thus be treated with caution in any analysis.

In general, the greatest information content is associated with daytime retrievals over land, during the summer season (MOPITT Algorithm Development Team, 2017; Deeter et al., 2015). This is where and when thermal contrast conditions are typically greatest, maximizing the instrument's ability to sense CO absorption 70 in the lowermost layers of the troposphere against the hot surface emission background (Deeter et al., 2007; Worden et al., 2010). To ensure that analyses involving MOPITT data are not biased by retrievals that have a heavy reliance on the a priori (in other words, a low information content), it is therefore suggested that users of MOPITT data consider excluding from analysis retrievals obtained during winter months, over water, and also from certain other geographical areas where retrieval information content is known to be low, i.e. 75 over mountainous regions, where the effects of geophysical noise reduce information content relative to flatter terrain (MOPITT Algorithm Development Team, 2017; Deeter et al., 2015). Deeter et al. (2015) specifically emphasizes such filtering in the analysis of long-term CO trends, since inclusion of retrievals with a heavy a priori weighting will weaken any real trends in the data. This occurs because the a priori CO is based on monthly climatologies of modelled CO amounts and is therefore variable by month but not by 80 year (Deeter et al., 2014).

MOPITT data are available as Level 2 (L2) products, where each individual retrieval at 22 x 22 km spatial resolution is available for analysis; and Level 3 (L3) products, which are a 1° x 1° area-averaged version of the individual L2 retrievals that fall within each gridbox (with some filtering criteria applied - see Section 85 2.1.2). At the heart of this study is the fact that some L3 gridboxes straddle the coastline. L3 products for such gridboxes can therefore be based on L2 retrievals over both land and water (see Figure 1), the information content of which can differ greatly (e.g., Deeter et al., 2007). In this study, we demonstrate, for a coastal L3 gridbox, how well known and well characterized differences in retrieval sensitivity over land and water can lead to significant differences in the L2 retrieved profiles that are averaged together to create the L3 products (Section 3.1). We outline the impact that this has on the statistics of the resulting L3 CO 90 profiles, and demonstrate the consequences that it has for temporal trend analysis with the L3 dataset, when compared to the results of the same analysis applied to the underlying L2 data that can be filtered by surface type to maximize information content (Section 3.2). This is an important issue to be aware of for two reasons: firstly, owing to their smaller file size, L3 data are better suited to long timeseries analysis than L2 data (~25 MB vs ~450 MB respectively, for a single daily, global file). Working with L3 data requires fewer computing 95

resources and, arguably, less technical expertise, making the MOPITT data more readily accessible to a greater number of users who are potentially less well-positioned to scrutinize the data. Secondly, 6 of the top

10 and 43 of the top 100 largest agglomerations by population in the world (population data taken from <u>www.citypopulation.de</u>, valid at time of writing) lie within a coastal L3 gridbox, and it is such cities that are

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likely to be targets for analyses of temporal trends in air quality indicators. The results that we report here suggest that similar analyses be performed for other coastal cities before using MOPITT surface CO.

This paper is structured as follows: In section 2 we outline the data and methods used in this study, giving an overview of the MOPITT instrument, retrieval, and surface type classification that is relevant to our work. In Section 3 our results are presented and discussed, and conclusions are drawn in Section 4.

105 **2. Data and Methods**

2.1 MOPITT

2.1.1 Instrument and retrieval overview

MOPITT has been making routine observations almost continuously since March 2000. It is carried on board the polar-orbiting NASA Terra satellite, with a nominal altitude of ~705 km and an equatorial overpass time of $\sim 10:30$ and $\sim 22:30$ local time. The instrument is a nadir-viewing gas correlation radiometer, with a ground 110 resolution of 22 x 22 km. It observes radiances in two CO-sensitive spectral bands: the TIR at 4.7 µm, and the NIR at 2.3 µm. The TIR band is sensitive to both absorption and emission by CO and yields information on its vertical distribution in the troposphere (Pan et al., 1995, 1998). The NIR band measures reflected solar radiation, which constrains the CO total column amount and yields information on CO concentrations in the lower troposphere (LT), to which TIR radiances are typically less sensitive (Pan et al., 1995, 1998). NIR 115 radiances can, however, only be exploited in daytime scenes over land. Our results are based on analysis of the TIR-NIR combined product, owing to its greater sensitivity to LT CO compared to the TIR- and NIRonly products which are also available (e.g. Deeter et al., 2017). Owing to the increased LT sensitivity from NIR radiances being limited to retrievals over land, we expect that the results presented here show an upper bound on the retrieval differences between surface types within our coastal L3 gridbox of focus, and the 120 consequent effects on sample statistics and temporal trends that we outline. Differences are still found in the TIR-only product, however, and we outline these in Supplementary Material 1. We restrict our analysis to daytime-only retrievals (more information on data selection in Section 2.1.3).

Multiple other sources describe MOPITT's CO retrieval algorithm in detail (e.g., Deeter et al., 2003; Francis et al., 2017). Briefly, it employs optimal estimation (Pan et al., 1998; Rogers, 2000) and a fast radiative transfer model (Edwards et al., 1999) to invert radiance measurements performed by the instrument to obtain CO concentrations. Additional inputs required include meteorological data (profiles of temperature and water vapour), surface temperature and emissivity, and satellite viewing geometry for the radiative transfer model; and a priori CO profiles, to constrain the inversion to physically reasonable limits. For latest

- 130 MOPITT product versions, meteorological fields are extracted from the NASA Modern-Era Retrospective Analysis for Research and Applications Version 2 (MERRA-2) reanalysis product; and a priori CO profiles are derived from a monthly CO climatology for the years 2000–2009, simulated with the Community Atmosphere Model with Chemistry (CAM-chem) chemical transport model at a spatial resolution of 1.9° x 2.5° (Lamarque et al., 2012) and then spatially and temporally interpolated to the time and location of the
- MOPITT observation. As it is a multi-year climatology, the a priori features no yearly trend, i.e. values for a given location and day of the year are the same every year. Surface temperature and emissivity values are retrieved from the radiance measurements (the retrieval also requires a priori information for these measurements). Retrievals are only performed for cloud-free scenes, with cloud screening based on collocated Moderate Resolution Imaging Spectroradiometer (MODIS) observations and MOPITT's own radiances. CO profiles are retrieved on 10 vertical levels, with 9 equally spaced pressure levels from 900 to
- 100 hPa (the uppermost level covers the atmospheric layer from 100 to 50 hPa) and a floating surface pressure level. Where the surface pressure is below 900 hPa, less than 10 profile levels are retrieved. Reported values represent the mean CO volume mixing ratio (VMR) in the layer immediately above that level. Retrievals are initially performed on a log₁₀(VMR) scale, owing to large CO variability in the atmosphere.
- Averaging kernels are produced for each retrieval and distributed with the data. The AK matrix (A) quantifies the the sensitivity of the retrieved vertical profile to the "true" vertical profile, and depends on the radiance weighting functions, instrument error covariance matrix, and a priori covariance matrix. Its relationship to the retrieved profile (X_{rtv}), the "true" profile (X_{true}), and the a priori profile (X_{apr}) is expressed as follows (e.g. Deeter et al., 2017):

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$$X_{rtv} = X_{apr} + A(X_{true} - X_{apr})$$
(1)

Thorough analysis of AKs is essential for understanding the physical significance of MOPITT's CO retrievals. We discuss AKs in more detail in section 3.1.2.

The MOPITT retrieval algorithm is subject to continuous development, in line with improvements in understanding of the changing instrumental characteristics and geophysical factors that affect the retrieval sensitivity, and with periodic updates to the radiative transfer model (Worden et al., 2014). This prompts the release of new product versions, with enhanced validation statistics against in situ CO observations. The work presented in this manuscript is based on MOPITT Version 7 (V7) products (Deeter et al., 2017). We analyse both L2 and L3 products (as outlined below). It should be noted that MOPITT Version 8 products have been released very recently, incorporating an improved radiance bias correction method to address a documented drift and geographical variability in retrieval bias compared to in-situ measurements (Deeter et

al., 2019). It remains to be seen whether the impacts of land-water retrieval sensitivity contrasts documented in this study remain in this newest product version.

2.1.2. Surface type classification

Both L2 and L3 data files come with a range of diagnostic fields and values, in addition to the averaging
kernel matrix, that can be used for filtering and interpreting retrievals. Of particular importance is the surface
index flag. Because retrieval information content is variable depending on surface type (Deeter et al., 2007),
each L2 retrieval is tagged according to whether it was performed over land, water, or a combination of the
two ("mixed"). The surface index of each L3 gridbox is then based on the L2 retrievals that fall within the
relevant 1° x 1° grid boundaries (Figure 1). Where more than 75 % of the bounded L2 retrievals have the
same surface index, only those retrievals are used to produce the L3 gridded value (the other L2 retrievals
are discarded) and the L3 surface index is set to that surface type. Otherwise, all L2 retrievals available in
the L3 gridbox are averaged together and the L3 surface index is set to "mixed" (this information is taken
from the MOPITT Version 6 L3 data quality summary¹ – at the time of writing, no V7 L3 data quality

The averaging together of retrievals with significantly different sensitivity profiles – as could be the case when averaging retrievals over land and water – serves to dilute the information coming from the MOPITT observed radiances with information coming from the a priori, thus increasing the dependence of the resulting CO profile values on the a priori profile. In fact, guidelines to maximize the information content of MOPITT data and minimize the influence of the a priori are to restrict analysis to daytime observations over land during the summer season, since this is when thermal contrast conditions are greatest, thus maximizing the

instrument's ability to sense CO in the lowermost layers of the troposphere (MOPITT Algorithm Development Team, 2017; Deeter et al., 2015; Deeter et al., 2007). Unfortunately, such filtering does lead to an overall loss of available retrievals for analysis, reducing the effective temporal and spatial coverage of the data.

185 2.1.3. Study area, time period, and MOPITT data processing in this study

Our analysis is based on MOPITT retrievals over the city of Halifax in Nova Scotia, Canada (Figure 1: longitude = -63.58° , latitude = 44.65°). Situated on the Atlantic coastline, Halifax is the major economic center in Atlantic Canada, with a population in excess of 315,000 in the urban core of Halifax Harbour (from 2016 census statistics). The pollution environment of Halifax, which is an intermediate port city, was

¹ available here: <u>https://eosweb.larc.nasa.gov/sites/default/files/project/mopitt/quality_summaries/mopitt_level3_ver6.pdf</u>

- characterized in detail by Wiacek et al., (2018 trace gases of focus include SO_x, CO₂, CO, NO_x, O₃, HC, and PM); briefly, it showed no exceedances of regulated gaseous contaminants, but nevertheless a substantial contribution of shipping emissions that is comparable to or greater than emissions from the city's vehicle fleet and a nearby 500 MW power plant. All available MOPITT V7 L2 and L3 TIR-NIR files ("MOP02J" and "MOP03J" files, respectively) were downloaded from the NASA Earthdata portal (https://search.earthdata.nasa.gov). There is a small inconsistency in the data record before and after an
- instrumental reconfiguration in 2001 (Drummond et al., 2010); we therefore discard all data prior to this reconfiguration. The remaining data covers the period 2001-08-25 to 2017-03-05. At the time of writing, more recent data are flagged as "beta" files, which await a future retrospective processing after the annual hot calibration becomes available, and their use in scientific analyses is discouraged (Deeter et al., 2017). For
 clarity and brevity, we restrict our main analyses and discussion to the winter (DJF) and summer (JJA) seasons, since these best encapsulate the different thermal contrast conditions over land and water, when compared to the intermediate (MAM and SON) seasons. For completeness, we demonstrate that our findings

also hold for MAM and SON in section 3.3.

- We extract L3 data for the 1° x 1° gridbox that contains the city of Halifax, and retain only the observations that were made during daytime hours. This yields a timeseries with one observation per day, when retrieval data were available within this gridbox. There are no retrievals available on 91 % of all days in DJF and 83 % of all days in JJA for the period covered. This is a result of both 1) MOPITT's polar orbit limiting temporal resolution to ~3 days over most of the globe; and 2) on days when the satellite's swath does encompass Halifax, retrievals either not being made due to cloud coverage, or discarded due to data quality issues. While
- this does not prevent a meaningful comparison of available retrievals, it does mean that caution is needed when using them to draw conclusions about the time period covered as a whole, which is something that we do not attempt to do. For clarity, we refer to the original, "as-downloaded" L3 timeseries as L3O for the remainder of this paper, owing to the way that we process the L2 data (explained below). Because this gridbox straddles the coastline, the L3O surface index varies each day. The surface classification breakdown of the
- L3O timeseries is given in Table 1a. "Water" is the modal classification in both seasons, followed by "mixed". L3O is only classified as "land" on one occasion each season. This is most likely due to the fact that more of the L3 gridbox is situated over water than land (Figure 1). The ratio of water to mixed observations is far greater in DJF than in JJA. This may be due to preferential cloud coverage over land in winter, and/or could be linked to the misidentification of snow/ice coverage on the surface as cloud during cloud screening (identifying the exact cause for this difference is beyond the scope of this paper).
 - We select all L2 retrievals that fall within the 1° x 1° L3 gridbox that contains the city of Halifax (lowerleft corner: -64° E, 44° N; upper right corner: -63° E, 45° N). Because we directly compare the L2 retrievals

to the L3 product that they create, we filter these based on pixel number (each pixel corresponds to one of MOPITT's four along-track detectors) and channel-average signal-to-noise ratio (SNR), as is done at the V7

L3 processing stage to improve L3 information content by excluding observations from specific detector elements on MOPITT's detector array that were found to exhibit greater retrieval noise than the other elements (MOPITT Algorithm Development Team, 2017; Deeter et al. 2017). Specifically, these filters exclude the following: all observations for Pixel 3; and all observations where both (1) the channel 5A SNR < 1000 and (2) the channel 6A SNR < 400. 5A and 6A correspond to the average radiances for MOPITT's length-modulated cell TIR and NIR channels, respectively. Finally, we only retain daytime retrievals, using a solar zenith angle filter of < 80°.

From this subset of L2 retrievals, we take separate area averages for those with a surface index of land and water, creating two timeseries that are effectively new L3 "land only" and "water only" products, for days when MOPITT retrievals over Halifax are available. We herein refer to these as L3L and L3W,
respectively. For clarity of analysis, we discard remaining L2 retrievals with a surface index of mixed (these account for ~5 % of the total L2 retrieval subset). The number of individual L2 retrievals that are averaged together each day to create L3L and L3W is given in Table 1b. From this, it is clear that there are around double the number of L2 retrievals over water than land within the L3 gridbox containing Halifax, which explains the dominance of water in the L3O surface classification (Table 1a), and also means that L2
retrievals over water will have a greater weighting in L3O than L2 retrievals over land on days when the surface index is mixed.

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2.2. Retrieval Simulation

To demonstrate how MOPITT retrieved CO concentrations are affected by retrieval sensitivity (Section 3.1.3), we simulate pairs of L3L and L3W retrieved profiles that are obtained concurrently (i.e. retrieved on the same day – on some days, one of L3L or L3W is missing) as follows:

$$X_{sim} = X_{apr,sim} + A(X_{tr,sim} - X_{apr,sim})$$
⁽²⁾

For each simulated retrieval: $X_{tr,sim}$ is taken from the Copernicus Atmospheric Monitoring Service (CAMS) reanalysis (CAMSRA – see section 2.3), for the model gridbox that contains Halifax, for the corresponding month and year of the observed retrieval (because the CAMSRA data are monthly mean values); $X_{apr,sim}$ is the mean of the a priori fields that correspond to the temporally coincident L3L and L3W pairings; and A is the retrieval averaging kernel from L3L or L3W. Thus, any differences between each pair of simulated retrievals ($X_{sim,L3L}$ and $X_{sim,L3W}$) are solely a result of differences in A, since $X_{tr,sim}$ and $X_{apr,sim}$ are identical

for both. Simulations are initially performed on $log_{10}(VMR)$ for consistency with the MOPITT retrieval algorithm, and then converted back to VMR scale for analysis.

2.3. Additional datasets 255

2003-2016.

The CAMSRA dataset to simulate retrievals is described by Inness et al., (2019). For the CAMSRA gridbox containing Halifax (horizontal resolution = $1^{\circ} \times 1^{\circ}$), we extract CO volume mixing ratios for levels 1000 hPa - 100h Pa at 100h Pa intervals, which correspond to the MOPITT levels of the profile. The CAMSRA dataset has no "surface" level, so we take the 1000 hPa level (the lowest level available in the dataset) to correspond to MOPITT's floating surface level. At the time of writing, CAMSRA data are only available for the years

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Information on mean wind patterns across Nova Scotia and the surrounding area is taken from the European Centre For Medium-Range Weather Forecasts (ECMWF) ERA-Interim dataset (horizontal resolution = $0.75^{\circ} \times 0.75^{\circ}$; see Dee et al., (2011) for a dataset overview). We analyse daily mean u and v vector winds for the following levels: 10-metres (the closest level to the surface for winds in the dataset), 850 265 hPa, and 500 hPa (which correspond roughly to the lower- and mid- troposphere, respectively). In addition, we extract monthly mean temperature profile data (at 100 hPa intervals, plus the skin temperature and 2metre air temperature variables) for the closest model gridboxes to Halifax that exclusively cover land and ocean, in order to illustrate the typical "land-only" and "water-only" temperature profiles that correspond to the MOPITT L2 retrievals over land and water that are analyzed.

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3. Results and Discussion.

3.1. Impact of retrieval sensitivity differences on temporally coincident L3L and L3W retrievals.

In this section we compare the L3L and L3W CO retrievals and demonstrate where and when there are differences in retrieved CO concentrations that are clearly linked to differences in retrieval sensitivity over land and water. We restrict our analysis to days when the L3O surface index is "mixed" and both L3L and 275 L3W retrievals are present, in order to minimize any potential differences in the true profile between land and water (there are a couple of days in the L3O timeseries when one or the other of L3L or L3W is missing, even when the L3O surface index is "mixed", owing to the presence of L2 retrievals with a surface index of "mixed", which we have discarded). Thus, an underlying assumption here is that land-water differences in 280 the true profile for retrievals contributing to L3L and L3W are small, owing to the fact that they are retrieved in close spatial proximity to eachother (i.e. within the same 1° x 1° gridbox) and at the same time. We test this assumption in Sections 3.1.4 and 3.1.5.

3.1.1. Climatology of land-water retrieval and a priori differences.

Figure 2 shows the percentage difference between temporally coincident retrieved VMRs for selected levels of the profile and for CO total column (TCO) amounts (Δ RET) in L3L and L3W. Positive (negative) differences indicate that retrieved VMRs/TCO are greater (less) in L3W than L3L. Differences are expressed as percent values, rather than differences in measurements units, so that we can display profile and TCO retrievals on the same plot (profile units are ppbv, TCO units are molecules cm⁻²).

In both seasons, mean retrieved VMRs are greater in L3W than L3L in the lower troposphere (LT – surface, 900hPa and 800hPa levels), with a maximum mean difference of 11.6 % (14.3 ppbv) at the surface level in JJA, the only profile location where the mean difference is significant (p = 0.001). The spread of ΔRET values is comparable in both seasons at these levels, with a clear skew towards positive values. Thus, although retrieved LT VMRs in L3L may occasionally exceed those in L3W by over 20%, they are usually greater over water than land. Mean ΔRET values are closer to zero and less significant in the MT and UT (represented by the 600 hPa and 300 hPa profile levels respectively), indicating that differences in retrieved VMRs are not as persistent at higher altitudes. However, the spread in ΔRET remains large at these altitudes, with retrieved VMRs in L3L and L3W differing by over ± 40 % on individual days (with outliers exceeding ± 60 %). TCO is greater in L3W than L3L in both seasons, significantly so in JJA. This is consistent with the most persistent VMR differences occurring in the LT, where atmospheric densities are greatest, thus contributing a relatively greater amount to the total column than MT and UT levels.

Our assumption in this section is that L2 retrieved CO concentrations obtained within the same 1° x 1° L3 gridbox should be similar. We may actually expect retrieved CO amounts in L3L to be greater than those in L3W due to CO sources existing on land, particularly within the city of Halifax. One reason for the ΔRET values instead indicating higher concentrations over water could be differences in the a priori profiles (ΔAPR)
used in the corresponding retrievals. The L2 retrievals over land and water have different a priori profiles owing to spatial interpolation of the 1° x 1° model climatology to the 22 x 22 km footprint of the MOPITT L2 retrieval. However, as Figure 2 demonstrates, ΔAPR values are small in comparison to ΔRET, with mean difference values very close to zero and a maximum range of 10.4 % (occurring at the surface level in JJA). Moreover, the sign of mean ΔAPR does not match that of mean ΔRET at several levels in both seasons (i.e.
retrieved VMRs in L3W are greater than in L3L, but a priori VMRs are less). It therefore appears unlikely

that a priori profile differences are responsible for the observed differences in retrieved CO concentrations in L3L and L3W.

3.1.2. Climatology of land-water retrieval sensitivity differences.

An alternative explanation for the observed ΔRET could be differences in retrieval sensitivity over land and water, quantified by the retrieval AK matrix. Figure 3 compares the mean AKs corresponding to the retrieved 315 profiles in L3L and L3W analysed in the previous section. Each curve corresponds to a row of the AK matrix and represents the sensitivity of the corresponding level of the retrieved profile to each level of the true CO profile, with the widest part of each AK in the x-direction (when a peak is evident) indicating the portion of the true profile that the corresponding level of the retrieved profile is most sensitive to. The sum of the elements in each AK row represents the overall sensitivity of the retrieved profile at the corresponding 320 pressure level to the whole true profile; values close to zero indicate that the retrieval is relatively insensitive to the true profile and therefore closely tied to the a priori profile, while the converse is true as the rowsum approaches one. The mathematical trace of the AK matrix (i.e. sum of the diagonals) gives the degrees of freedom for signal (DFS) of the retrieval, which is a measure of the number of independent pieces of 325 information (in other words, "information content") in the retrieval from the measurement, with respect to the true profile. When DFS values approach two, this is interpreted as the retrieval being able to resolve CO in two independent atmospheric layers.

There are some clear differences between the mean AKs over land (from L3L) and water (from L3W) shown in Figure 3. In DJF, AKs for LT (especially surface level) retrievals reach greater values in the LT in L3W than L3L, indicating that sensitivity to the true profile at these levels is actually greater in L3W (surface 330 level AK peak = 0.14 for water, 0.09 for land, both at the surface level). This is reflected in greater rowsum values for LT AKs in L3W than L3L. Differences in MT (600 hPa) and UT (300 hPa) AKs are much less pronounced, with sensitivity to the true profile actually becoming slightly greater in L3L than L3W higher up in the troposphere. In JJA, the mean LT and MT AKs are qualitatively much more different between L3L and L3W than in DJF. LT rowsums are significantly greater in L3L (p < 0.001 at the surface, 900 hPa and 335 800 hPa profile levels, see Supplementary Material 2) and closer to one, signifying that these retrievals contain much more true profile information than those in L3W. The AK shapes also indicate that the respective retrievals are sensitive to different parts of the true profile. In L3L, surface and LT AKs peak at either the surface (surface level AK) or at 900 hPa (both the 900 hPa and 800 hPa level AKs) and decline towards the UT, while MT AKs indicate relatively equal sensitivity throughout the LT, MT and the lower 340 levels of the UT. In L3W on the other hand, LT and MT AKs (excluding the largely insensitive surface level) indicate relatively little sensitivity in the lowest profile levels and peak in the MT. The surface level AK in L3W actually indicates close to zero sensitivity throughout the profile, aside from a very weak peak at the surface level, which is around 3 times lower than that over land. As in DJF, UT AKs are quite similar, except

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for relatively small differences at the surface, where the 300 hPa AK over land actually indicates negative sensitivity. Differences in mean DFS values for retrievals in L3L and L3W are greater in JJA (-0.12) than DJF (0.07), highlighting the greater land-water sensitivity contrast in JJA and also reflecting the switch in surface type exhibiting the greatest LT retrieval sensitivity between the seasons.

- The differences in surface and LT AKs for MOPITT retrievals in L3L and L3W discussed above can be
 accounted for primarily by the differing LT thermal contrast conditions over land and water, as explored in
 detail by Deeter et al. (2007). The sensitivity of MOPITT retrievals to CO in the LT is predominantly
 controlled by the thermal contrast between the surface skin temperature (T_{skin}) and the surface air temperature
 (T_{sfc}), and by the tropospheric temperature profile. Seasonal mean temperature profile data from ERA-Interim
 for the nearest land-only and water-only model gridboxes to Halifax show clear differences in DJF and JJA
 (Figure 4). In DJF, T_{skin} is around 6° K warmer than the 2-meter air temperature (T_{2m}, which we use as a
 proxy for T_{sfc} as this is the lowest model level) over water, and a further degree warmer than the air at 1000
 hPa, whereas the temperature gradient is weak/slightly inverted over land, with T_{skin} less than a degree
 warmer than T_{2m}, which is actually slightly cooler than the air at 1000 hPa. Correspondingly, the lowest
- L3L) in DJF (Figure 3). Temperature profiles converge towards the MT, as do AKs. In JJA on the other hand, there is a clear gradient between T_{skin} and the overlying air on land, while the ocean surface is actually cooler than the air above, up to a height of 900 hPa. As a result, surface and LT sensitivity is greater in L3L than in L3W in JJA, with the sensitivity of retrievals in L3W approaching zero close to the surface owing to this inverted temperature profile. The T_{skin} increase approaching 20° K between DJF and JJA likely also accounts
 for the relatively greater overall true profile sensitivity (indicated by DFS values) in JJA for L3L than in DJF for L3W. Since our analysis is conducted using the joint TIR-NIR product, it is important to bear in mind that the benefit of enhanced LT sensitivity due to the incorporation of NIR is limited to retrievals over land, so this will also have an impact on the AK differences presented above. However, a land-water retrieval sensitivity contrast of comparable magnitude to that presented here is also evident in the TIR-only product, reinforcing the primary role of thermal contrast differences (see Supplementary Material 1).

That the retrieval sensitivity contrast between L3L and L3W is most pronounced in the LT is consistent with the finding that retrieved CO profiles in L3L and L3W show the greatest differences in the LT. Although mean retrieval sensitivity in L3L and L3W converges with altitude, differences do exist from day-to-day but they are neither as large, nor as skewed in favour of retrievals over land or water (depending on season) as in the LT (see Supplementary Material 2), where there is a well understood thermal contrast mechanism

creating the systematic land-water sensitivity contrast. Likely causes for this could be day-to-day changes in atmospheric conditions (i.e. temperature or water vapour profiles), or random instrumental or retrieval noise.

3.1.3. Control of $\triangle RET$ by land-water sensitivity differences

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To demonstrate that retrieval sensitivity differences over land and water can lead to the observed differences between CO profiles in L3L and L3W that are retrieved at the same time, we simulate and compare the pairs of retrieved CO profiles in L3L and L3W that are analysed in the preceding sections, using the transformation outlined in Section 2.2. Recall that any differences between each pair of simulated retrievals are solely a result of differences in AKs. Because available CAMSRA data only covers the years 2003-2016, only a subset of the retrieval pairings considered in earlier sections (which span the period 2001-2017) are simulated.

We first demonstrate the sensitivity effect with a case study on 2013-08-18. Profiles (a priori (X_{apr,sim}) and "truth" (X_{tr,sim})) and surface level AKs used in the simulation, and the resulting X_{sim,L3L} and X_{sim,L3W}, are given in Table 2. For brevity, we only focus on the surface level. In this example, X_{tr,sim} is considerably lower than X_{apr,sim} at the surface level, and several features of the AKs indicate greater sensitivity to X_{tr,sim} in L3L
and L3W: the AK value is significantly greater at the surface in L3L than L3W, and the rowsum is over 5 times as high. Correspondingly, X_{sim,L3L} is much lower than X_{sim,L3W} at the surface and much closer to X_{tr,sim} (32.21 ppbv higher vs 57.13 ppbv higher). Both X_{sim,L3L} and X_{sim,L3W} indicate that X_{tr,sim} is lower than X_{apr,sim} at the surface, but X_{sim,L3L} gives the closer estimate, as would be expected. In both cases, a portion of the overall departure from X_{apr,sim} at the surface level (in other words, "value added" over the a priori) originates at other levels of the profile, and not the surface level itself. This is a result of the surface level AK being nonzero at other levels, and is a function of the inter-level correlation of the original retrieval, which is linked to the a priori covariance matrix used in the retrieval (Deeter et al., 2010).

We now consider differences between all X_{sim,L3W} and X_{sim,L3L} pairings (ΔSIM) throughout the profile and compare these to the observed differences between temporally coincident retrievals in L3L and L3W
(ΔRET) discussed previously (Figure 5a). For ease of comparison, ΔRET values are overlaid (faint lines) for the shorter time period matching CAMSRA data availability (the 2003-2016 ΔRET patterns are very similar to those seen in Figure 2). It should be noted that we cannot expect ΔSIM to match ΔRET exactly, owing to (possibly large) differences between the X_{tr,sim} profiles used in the simulations and the (unknown) true profiles at the time of the actual MOPITT retrievals (X_{true}). For instance, X_{tr,sim} is a monthly mean value from a reanalysis model, whereas X_{true} varies by day, which should result in less variance in ΔSIM than ΔRET. In DJF, mean ΔSIM is negligible at all profile levels shown, and the range of values is far smaller than seen for ΔRET. In JJA on the other hand, ΔSIM reaches considerably larger values than in DJF and mean values are

significantly different (p < 0.05) over land and water at all but one level shown (300 hPa). LT and MT Δ SIM distributions are a remarkably good match for ΔRET , given the likely substantial differences between X_{tr,sim}

and X_{true} . In the UT, Δ SIM values are somewhat smaller, although this isn't unexpected given the smaller

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land-water AK differences evident in Figure 3.

That L3L and L3W simulated and retrieved profile differences over land and water are comparable in JJA is clear evidence that ΔRET is strongly influenced by the land-water sensitivity contrast in summer months, at least in the LT. However, while Δ SIM values are of a much smaller magnitude in DJF than in JJA, Δ RET 415 is actually of a similar magnitude in both seasons. The question therefore arises as to why Δ SIM is so different from ΔRET in DJF, and why it is of much smaller magnitude in DJF than JJA. Considering the terms of Equation 2, there are two possible explanations. Firstly, differences between L3L and L3W LT and MT AKs (A in Equation 2) are much smaller in DJF than in JJA (Figure 3). This means that the deviation of X_{sim,L3W} and $X_{sim,L3L}$ from $X_{apr,sim}$ will be more similar in DJF (resulting in small Δ SIM), as opposed to in JJA where, as was seen in the case study discussed above, X_{sim,L3L} can deviate more from X_{apr,sim} than X_{sim,L3W} owing to 420 increased sensitivity in L3L in the lower profile levels (resulting in greater Δ SIM than in DJF). In the UT, AK differences are comparable in both seasons, and Δ SIM is correspondingly similar. Secondly, the magnitude of X_{tr,sim} – X_{apr,sim} from Equation 2 is, on average, around 4 (3) times greater in JJA than in DJF at the surface (900 hPa) level (Figure 5b). X_{sim,L3L} therefore deviates more from X_{apr,sim} than X_{sim,L3W} 425 throughout the LT and MT in JJA owing to strong contrasts in near-surface sensitivity at these profile levels, thus yielding large Δ SIM. Conversely, closer X_{tr,sim} and X_{apr,sim} profiles combine with small sensitivity differences to limit Δ SIM in DJF. A final, alternative explanation for Δ SIM being a poor match for Δ RET in DJF, unlike in JJA, is that sensitivity differences in L3L and L3W could have less of an impact on ΔRET in DJF than in JJA, and that something else is responsible – for example, "real" differences between the true CO profile over land and water. This is something we explore in detail in the following sections.

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3.1.4. Regional land-water contrast in L2 data

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To further evaluate whether $\triangle RET$ within the L3 gridbox containing Halifax is a function of land-water sensitivity contrasts, or actually due to real gradients in the true CO profile (for example, due to offshore transport of emissions from the city of Halifax or marine-land chemistry differences), we analyse the characteristics of retrieved profiles over the broader geographical region surrounding Halifax. If ΔRET is linked to sensitivity, then we would expect there to be a clear land-sea contrast across the whole region. This is exactly what we see. Figure 6 shows seasonal median L2 retrieved and a priori CO concentrations for the surface level of the profile, where $\triangle RET$ and L3L-L3W AK differences are greatest and most significant, for

the Canadian maritime provinces and a small portion of northeast USA (note that we show seasonal median fields here as the spatial patterns are clearer than for plots consisting only of the subset of days analysed in 440 the rest of this section. The corresponding plot for the subset of days is shown in Supplementary Material 3, and the main findings are unchanged). Difference fields (RET-APR) are also shown. In JJA, a land-sea contrast is remarkably clear in both the retrieved and RET-APR fields. The a priori field shows elevated CO amounts emanating from the west-southwest (indicative of CO sources in northeast USA and around the Great Lakes) and decreasing quite smoothly towards the north and east. The west-southwest maxima is 445 replicated in the retrieved field, but the smoothly decreasing gradient is clearly broken, with the land in the image characterized by lower CO values than the adjacent ocean. This contrast is enhanced further in the RET-APR field. RET-APR values close to zero indicate either very low retrieval sensitivity or closely matching retrieved and a priori values. The analysis of averaging kernels in Section 3.1.2 demonstrated the lack of retrieval sensitivity at the surface over water (in L3W) in JJA. On average, RET-APR is 11.5 ppbv 450 lower over land than over water (determined by binning the L2 data for the region shown according to surface classification). This reinforces our earlier interpretation that, in JJA, LT retrievals in L3W are weighted more heavily towards an a priori profile in which CO concentrations are too high than LT retrievals in L3L.

The land-sea contrast is less clear in DJF, although the RET-APR field does indicate generally positive values over water and negative values over land. The contrast being less apparent in DJF compared to JJA is 455 consistent with the smaller L3L-L3W AK differences in DJF compared to JJA. However, it is surprising that RET-APR changes sign from land (generally negative) to water (generally positive). This may be linked to some factor other than retrieval sensitivity to the true CO profile, such as errors in retrieved/a priori surface temperatures or emissivities, which are important components of the radiative transfer model used in MOPITT's CO retrieval algorithm. If this were the case over the sea, it could explain the maxima in RET-460 APR values to the northwest of Halifax in the Bay of Fundy, where the water is relatively shallow and the tidal range is the highest in the world at 16.3 m, transporting large amounts of suspended sediments (which will affect emissivity). Alternatively, the difference could reflect a physical process causing elevated CO over the ocean, although this seems unlikely, as we expect atmospheric transport to minimize such a contrast. 465 Corresponding maps for selected other levels of the profile and TCO are shown in Supplementary Material 4. In JJA, a land-sea contrast is qualitatively evident at all levels and for TCO, with the exception

of 800 hPa; and in DJF a contrast is evident at 800 hPa and for TCO.

3.1.5. Can \triangle RET be explained by circulation-driven horizontal gradients in the true CO profile?

The preceding sections presented evidence that differences in temporally coincident L3L and L3W retrieved profiles are linked to sensitivity contrasts over land and water, especially in JJA and in the LT. However, these differences could also be a result of horizontal gradients in the true CO profile. It is plausible that retrieved LT VMRs are greater in L3W than in L3L due to, e.g., offshore transportation of CO by regional winds either from Halifax or from the large polluting areas on the northeast coast of the US and around the Great Lakes (as seen in the general decline in CO amounts from the west-southwest towards the northeast in Figure 6). Winds generally tend from the west/northwest/southwest in this area (Figure 7), which will lead to offshore transport of continental pollution.

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We compare composite mean wind patterns across Nova Scotia using ERA-Interim data for days when retrieved surface level VMRs in L3W are greater than in L3L (L3W > L3L) and days when they are less (L3W < L3L), since a clear shift in wind direction on these days would support the case that atmospheric transport plays a role in generating differences in retrieved CO amounts over land and water. These are shown 480 in Figure 7 (10-metre winds are used). There is a clear circulation difference evident in DJF. When L3W >L3L (12 days of the 18 in Figure 2) the wind is in an offshore direction, whereas when L3W < L3L (6 days of the 18 in Figure 2) the wind is alongshore from the west-southwest (and noticeably weaker). This is evidence to suggest that the LT ARET patterns in DJF are linked to the horizontal gradients in the true CO profile, although the small sample sizes involved here dictate caution. In contrast to DJF, however, there is 485 no clear circulation difference in JJA; the winds are generally from the west-southwest irrespective of whether L3W > L3L or L3W < L3L. This lends further support to the conclusion that LT ΔRET in JJA is strongly linked to the demonstrated land-water sensitivity contrast. The seasonal difference in these results could explain why LT (and especially near-surface) ΔRET is of comparable magnitudes in DJF and JJA (Section 3.1.1), despite smaller L3L-L3W retrieval sensitivity contrasts and Δ SIM values in DJF than JJA 490 (Sections 3.1.2 and 3.1.3 respectively). In other words, in DJF, ΔRET is more indicative of differences in true CO concentrations, while in JJA it is more strongly tied to differences in retrieval sensitivity. This is not to say contrasts in retrieval sensitivity do not influence LT ΔRET in DJF; just that the effect is not as strong as in JJA, when the LT sensitivity contrast is greater. We only consider the surface profile level here; the 495 findings are consistent higher up in the LT, but the DJF circulation difference is absent in the MT (see Supplementary Material 5), consistent with ΔRET being much smaller in the MT and above, on average.

While there is no obvious circulation difference at the surface in JJA between days when L3W > L3L and L3W < L3L, there is a difference in the distribution of these days throughout the analysed MOPITT timeseries, which spans 16 JJA seasons from 2001 to 2016. 16 of the 19 days when L3W < L3L occur in the first half of the timeseries (i.e. before 2009), whereas days when L3W > L3L are spread more evenly throughout (33 of the 65 days (51%) occur before 2009). In DJF there is no such difference, with roughly 50% of days occurring before and after 2009 in each case. This is something we explore further in Section 3.2.2.

3.2. Consequences for L3O timeseries

In this section we demonstrate how the statistics of the L3O timeseries, and the results of a typical trend 505 analysis using those data, are affected by the loss of loss of LT retrieval information from L2 products over land when the L3 products for the coastal gridbox containing Halifax are created. We do this through comparison with the L3L and L3W timeseries. Because users of L3 data are advised to filter according to surface index in order to limit their analysis to retrievals with maximal information content, we consider L3O subsets that remain after filtering the timeseries for days with a surface index of "water" and "mixed" 510 ("L3O_{(water})" and "L3O_{(mixed})"), as well as the unfiltered L3O timeseries to evaluate the full range of options available to users of the products. L3O only has a surface index of "land" once each season, so this subset is omitted. We focus on the LT profile levels since this is where retrieval sensitivity differences are greatest and can be linked to differences in retrieved CO values, as shown in the previous analyses.

3.2.1. Impact on seasonal data distribution 515

Seasonal surface level VMR distributions for L3L, L3W and all L3O subsets are shown in Figure 8. Most strikingly, L3L is the clear outlier in JJA. Mean VMRs are significantly lower than in all other timeseries (p < 0.1 in all cases) and the spread of values is around twice as large, both in terms of the interquartile range and overall range (excluding outliers), this difference mostly coming from the lower end of the distribution. This is unsurprising when comparing L3L with the timeseries that are based purely on retrievals over water 520 (L3W and L3O_(water)), given the demonstration in previous sections that retrievals over water have significantly lower information content than over land in the summer months and are therefore more closely tied to the a priori CO concentration (i.e. retrieved VMRs will vary less). However, it clearly shows how the valuable additional information on true CO content that is available in L2 retrievals over land is diluted by their averaging with retrievals over water for $L3O_{(mixed)}$ days (mean surface level AK rowsum = 0.38 for 525 L3O_(mixed) vs 0.69 for L3L and 0.14 for L3W, see Figure 3), effectively creating a high bias in the resulting gridded mean VMRs. The loss of retrieval information from L2 to L3 is actually exacerbated for this 1° x 1° L3 coastal gridbox containing Halifax, given that a greater number of L2 retrievals over water contribute to the gridded averages than L2 retrievals over land (as previously outlined in Table 1), primarily because more of the surface within the L3 gridbox is water than land (see Figure 1). Consequently, the L3O(mixed) 530 distribution more closely resembles L3W than L3L (although the L3W-L3O_(mixed) mean difference is still statistically significant (p < 0.1)). Owing to the lack of days when the L3O timeseries is created only from retrievals over land, L3O_(mixed) represents the best option for quantifying surface level CO in JJA that is available to users of the original L3 product in this case. The optimal retrievals for this task are only available in the L2 data products, a direct result of the way that the L3 products are created.

- For the same reasons discussed above, L3L also represents the outlier distribution in DJF. Unlike in JJA however, the spread of VMR values is similar in L3L and L3W, likely reflecting the fact that there is some surface level sensitivity in retrievals over both land and water in DJF, allowing for a similar degree of departure from the a priori. The main difference in the distributions is that VMRs in L3L are offset towards
 lower values. However, the mean difference is not significant between L3L and any of the other timeseries. Also unlike in JJA, L3L does not necessarily represent the optimal timeseries for analyzing surface level CO
- in DJF, since retrieval sensitivity is actually higher over water in this season. Information loss resulting from the way L3 products are created is therefore less of an issue than in JJA, owing to the dominance of retrievals over water on the L3O timeseries (mean surface level AK rowsum = 0.52 for L3O_(mixed) vs 0.43 for L3L and 0.53 for L3W, see Figure 3). It is worth noting, however, that L3W offers ~25% more days with data than L3O_(water) in DJF, due to the fact that it "gains" the retrievals over water that go into L3O_(mixed). This is potentially valuable additional temporal information for users of MOPITT products.

Although the sample sizes considered here are different, because L2 retrievals over land and water are not necessarily always present on the same days (e.g. due to variable cloud coverage), the differences in seasonal data distribution discussed in this section hold if the analysis is restricted to only days when L3L and L3W are both present (see Supplementary Material 6). We have only presented analysis for the surface level of the profile here as this is where L3L-L3W differences in retrieved VMRs and retrieval sensitivity are greatest. Plots of other levels are given in Supplementary Material 7.

3.2.2. Consequences for temporal trend analysis

- To identify and compare temporal trends in the timeseries considered above, we perform weighted least squares (WLS) regression analyses on respective seasonal mean profile and TCO values, weighted by the standard deviation of the measurements used in the seasonal mean. For seasons that contain just a single measurement, we use the data record standard deviation scaled by a factor of 100 so as to de-weight these seasons in the fit. All trends identified are detailed in Table 3, and WLS best-fit lines, along with boxplots of seasonal data distributions, are presented for the surface level in Figure 9. Here, the decreasing trend identified in L3L is over four times stronger than the trend in L3W, a highly significant difference (p < 0.01). This is a direct consequence of surface level retrievals over water being tied closely to the a priori owing to their negligible sensitivity, which has no yearly change. This effectively masks the full magnitude of the decrease in CO that appears to be occurring, and which is better detected by retrievals over land owing to their greater sensitivity. Consequently, trends in all L3O subsets are also significantly weaker than the trend
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in L3L. Since it has some contribution from retrievals over land, L3O_(mixed) provides the closest approximation of the trend in L3L, but it is still over 50% weaker – representing a decrease of 10% (19 ppbv) over the 15-year period covered by the analysis vs 21% (40 ppbv) in L3L. Compared to JJA, trends in surface level VMRs in DJF are far more similar across all the timeseries, with no significant differences between any

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of the trends identified. This is attributable to the much smaller differences in retrieval sensitivity over land and water in this season. Thus, while the greater number of days corresponding to the L3W timeseries makes it of potentially greater value than L3O_(water), at least for temporal trend analysis it has no statistical benefit (in fact, users of the original L3 product would seem to get comparable results by performing the analysis on the unfiltered version of L3O).

- Although the land-water retrieval sensitivity contrast remains large at the 900hPa level in JJA, the trend in L3O_(mixed) is a closer match to L3L than at the surface, and the difference loses statistical significance. The loss of information content available in retrievals at the 900hPa level over land during the creation of L3O (mean 900hPa AK rowsum = 0.91 for L3L vs 0.64 for L3O_(mixed)) therefore does not have a statistically significant impact on temporal trends identified using L3O_(mixed), when compared to L3L. Although the L3L-
- L3W trend difference is smaller than at the surface level, L3L is still twice as strong as L3W and the difference remains statistically significant. This could be a result of either the retrieval over water still lacking sufficient information to deviate as far from the a priori as the retrieval over land, despite the increase in information content relative to the surface level (mean AK rowsums = 0.15 and 0.44 for the surface and 900hPa levels respectively); and/or the retrieval at the 900hPa level over water having a sensitivity peak higher up in the troposphere where CO concentrations may be decreasing at a slower rate than they are closer to the surface, where sensitivity peaks for the 900hPa level over land (see Figure 3). By 800hPa, trends in all timeseries in JJA have converged, consistent with the further weakening of the land-water sensitivity contrast.
- Moving away from the LT, the trends outlined in Table 3 indicate that all timeseries generally agree on the broad picture: in both seasons, CO concentrations are decreasing in the LT and MT, increasing in the UT,
 while TCO shows a decrease in DJF and no significant trend in JJA. Although not as pronounced or significant as at the surface in JJA, in all cases there are differences in the magnitude of the identified trend. This is not unexpected given that the seasonal means being regressed differ between timeseries. As outlined in Sections 3.1.1 and 3.1.2, temporally coincident retrieved VMRs over land and water do differ in levels of the profile above the LT despite similar retrieval sensitivity at these levels; the differences are just not as systematic as in the LT. However, in addition to the surface and 900hPa levels in JJA, there are 2 other instances where trends in L3L and L3W are significantly different: 600hPa in JJA and 300hPa in DJF (in both cases there is no statistically significant trend identified in L3L, whereas the trend in L3W is significant). The cause of these discrepancies is not readily apparent given that retrieval sensitivity over land and water is

highly comparable in both cases, so further investigation would be needed before we can say with confidence

600 whether or not they have consequences for analyses using the L3O timeseries, as we have been able to do for the LT.

The main differences in trend discussed above remain if the WLS regression analysis is restricted to only days when L3L and L3W are both present. Results from this restricted analysis are shown in Supplementary Material 8. It is important to note that MOPITT profile measurements are known to have a drift (Deeter et al., 2017), and this should be corrected for in the data if the focus of analysis is to use them to quantify temporal changes in CO over time. Since the intention of the WLS trend analysis presented here is more illustrative, namely to demonstrate trend differences in the data, we have not corrected for this drift. The results should therefore not be taken out of this context (as well as bias correction, verification against a range of other datasets would be required, especially given the large proportion of missing data). We do however provide the reported drift values in Table 3 for context, which shows that the majority of the trends that we have identified appear to be stronger than the measurement drift (at least for the dataset that has greatest retrieval sensitivity at the respective level of the profile). As noted in Section 2.1.1, the measurement drift has been significantly reduced in the latest version of the MOPITT products to be released (Version 8; Deeter et al., 2019).

615 **3.3. Consideration of MAM and SON**

In this paper we have focused only on DJF and JJA for brevity and clarity. The main findings discussed also hold for MAM and SON, however. In MAM, there is a strong sensitivity contrast in the LT between L3L and L3W, similar to that seen in JJA with retrieval sensitivity much greater over land than water (Figure 10a). As in JJA, the decreasing trend detected in WLS regression analysis of L3L surface level VMRs in MAM is strongly underestimated in L3W and, consequently, in all L3O subsets (Figure 10b). In SON on the other hand, AKs indicate a much more comparable degree of LT retrieval sensitivity over both land and water; correspondingly, the detected temporal trends are similar in L3L, L3W and L3O subsets.

4. Conclusions

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Users of MOPITT products are advised to filter the data before analysis of profile values in order to maximize the influence of satellite measurements and minimize the impact of a priori CO concentrations on results (MOPITT Algorithm Development Team, 2017; Deeter et al., 2015). In particular, it is advised that retrievals over water, which are known to have lower information content than retrievals over land, are discarded. This is especially so for the analysis of temporal trends in CO concentrations, owing to the year-to-year stationarity

of the a priori. However, for L3 gridboxes that straddle the coastline, the ability to apply such filtering is limited since the products will generally have some contribution from L2 retrievals that take place over both 630 land and water. This is a direct consequence of the way that L3 products are created.

As we have explicitly demonstrated for the 1° x 1° L3 gridbox containing the coastal city of Halifax, Canada, the L2 retrieved CO concentrations, from which the L3 products are created, differ depending on whether the retrieval took place over land or water. In JJA, and especially near to the surface, this is directly linked to differences in the sensitivity of the retrievals to the true CO profile. The merging of these retrievals 635 to create the L3 product can significantly affect the statistics of the dataset and the results of temporal trend analysis with the data, with the largest and most statistically significant effects at the surface, where landwater sensitivity contrasts are greatest. As we show, results that are more representative of changes of true CO concentrations close to the surface within the L3 gridbox containing Halifax can only currently be obtained by use of the L2 products, which can be filtered by surface type to maximize information content.

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Our results suggest that L2 retrievals over land and water should not both contribute to L3 products in coastal gridboxes. This is consistent with previous data filtering recommendations (MOPITT Algorithm Development Team, 2017; Deeter et al., 2015). The horizontally averaged L3L and L3W timeseries that we have analysed in this paper are effectively L3 "land-only" and L3 "water-only" datasets, and these offer an alternative in this respect that preserves the benefits of available L3 products - namely, less computing 645 resources and expertise required for their analysis compared to L2 products, which broadens access to the data – but offers users the flexibility to select over which surface the contributing retrievals were performed in order to maximize the information content of L3 data in coastal gridboxes. Although our study has only focused on the city of Halifax, the results suggest that similar studies be performed for other coastal L3 gridboxes before using MOPITT surface CO, since these contain 6 of the top 10 and 43 of the top 100 650 agglomerations by population and are therefore likely targets for analysis of temporal changes in air pollution indicators such as CO, especially near to the surface. The degree of information content loss in the L3 data will depend on the relative contributions of L2 retrievals over land and water to each specific L3 gridbox, as well as on the strength of the land-water retrieval sensitivity difference, which in turn depends on scenespecific geophysical variables such as surface temperature and emissivity. Work is currently ongoing to 655 compare the results of analyses conducted with MOPITT L2 and L3 CO data over these cities.

Data Availability

MOPITT data were downloaded from the NASA Earthdata portal (<u>https://search.earthdata.nasa.gov/</u>). CAMSRA and ERA-Interim data were downloaded from the ECMWF public datasets portal (<u>https://apps.ecmwf.int/datasets/</u>).

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Appendix A: List of acronyms

	ΔRET	Percentage differences in retrieved CO concentrations in $L3_L$ and $L3_W$
	Δ SIM	Percentage differences in simulated retrieved CO concentrations in $L3_L$ and $L3_W$
	AK	Averaging kernel
665	CAM-chem	Community Atmosphere Model with Chemistry
	CAMS	Copernicus Atmospheric Monitoring Service
	CAMSRA	CAMS Reanalysis
	CO	Carbon Monoxide
	DFS	Degrees of freedom for signal
670	DJF	December, January, February
	JJA	June, July, August
	L2	Level 2 data products
	L3	Level 3 data products
	L3L	Area-averaged L2 data for the L3 gridbox containing Halifax, for L2 retrievals over
675		land only
	L3O	Original, "as-downloaded" L3 timeseries for the 1° x 1° L3 gridbox containing
		Halifax
	L3W	Area-averaged L2 data for the L3 gridbox containing Halifax, for L2 retrievals over
		water only
680	LT	Lower troposphere (surface – 800hPa levels of the profile in MOPITT data)
	MERRA-2	NASA Modern-Era Retrospective Analysis for Research and Applications V2
	MOPITT	Measurement of Pollution in the Troposphere (instrument)
	MT	Mid-troposphere (700hPa – 500hPa levels of the profile in MOPITT data)
	NIR	Near infrared
685	SNR	Signal-to-noise ratio
	TCO	Total Column CO
	TIR	Thermal infrared

	T_{sfc}	Surface air temperature
	T_{skin}	Surface skin temperature
690	UT	Lower troposphere (400hPa – 100hPa profile levels in MOPITT data)
	V7	MOPITT Version 7 data products
	VMR	Volume Mixing Ratio
	X _{apr}	A priori CO VMR profile
	X _{apr,sim}	A priori CO VMR profile used in retrieval simulation
695	X _{rtv}	Retrieved CO VMR profile
	X_{sim}	Simulated retrieved CO VMR profile
	X _{sim,L3L}	Simulated retrieved CO VMR profile from L3L
	X _{sim,L3W}	Simulated retrieved CO VMR profile from L3W
	X _{true}	"True" CO VMR profile
700	X _{tr,sim}	"True" CO VMR profile used in retrieval simulation

Author contributions

IA and AW jointly conceived of and designed the study. IA performed data analysis; both authors examined and interpreted the results, and prepared the manuscript.

Competing interests

The authors declare that they have no conflict of interest.

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Worden, H. M., Deeter, M. N., Edwards, D. P., Gille, J., Drummond, J., Emmons, L. K., Francis, G., and Martínez-Alonso, S. (2014). 13 years of MOPITT operations: Lessons from MOPITT retrieval algorithm development. Annals of Geophysics, 56(FAST TRACK 1), 1–5. https://doi.org/10.4401/ag-6330 810 Table 1 (a) Surface classification of L3 data for the 1° x 1° gridbox containing Halifax (L3O) for the period 2001-08-25 to 2017-03-05. Note that no retrievals are available for this L3 gridbox on 91 % of days in DJF and 83 % of days in JJA. (b) Mean number of individual L2 retrievals with "land" and "water" surface index that contribute to the L3L and L3W timeseries, respectively (the standard deviation is in brackets).

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(a)					(b)			
Season	Water	Land	Mixed	Total	L3L	L3W		
DJF	101	1	23	125	1.7 (±1.1)	4.0 (±1.7)		
JJA	136	1	98	235	2.1 (±1.2)	4.1 (±1.8)		

Table 2 Case study information for simulated L3L and L3W profile retrievals ($X_{sim,L3L}$ and $X_{sim,L3W}$ respectively) for 2013-08-18. $\Delta = X_{apr,sim} - X_{tr,sim}$. Values that are underlined indicate the maxima (by magnitude) for that column. Simulated profile values for levels 900-100 hPa are shown in brackets because AKs and calculations for these levels are not discussed in the text and AKs for the 900 – 100 hPa levels are not shown.

		Input profil	es	Surface (== A	level AK A[1,i])	Simulated profiles (X_{sim})		
Profile level	X _{apr,sim}	X _{tr,sim}	Δ	L3W	L3L	$X_{sim,L3W}$	$X_{sim,L3L}$	
Surface	188.81	128.45	-60.36	0.01	0.23	185.58	160.67	
900 hPa	148.88	119.53	-29.34	0.03	0.20	(141.49)	(124.02)	
800 hPa	118.07	109.80	-8.27	0.02	0.15	(110.76)	(100.25)	
700 hPa	102.72	97.64	-5.08	0.03	0.14	(95.37)	(88.72)	
600 hPa	95.49	90.34	-5.15	0.03	0.12	(87.92)	(83.64)	
500 hPa	92.00	87.10	-4.91	0.03	0.08	(84.57)	(82.57)	
400 hPa	89.64	83.85	-5.78	0.02	0.02	(83.46)	(84.12)	
300 hPa	84.76	81.42	-3.34	-0.01	-0.07	(81.73)	(83.91)	
200 hPa	64.45	65.20	0.75	-0.01	-0.04	(63.77)	(64.75)	
100 hPa	26.51	28.71	2.20	0.00	0.00	(26.41)	(26.52)	
			Rowsum \rightarrow	0.15	0.84			

Table 3 Results from WLS regression analysis of seasonal mean L3W, L3L, L3O, L3O_(water) and L3O_(mixed)
timeseries for selected profile levels in DJF and JJA. Trend corresponds to the gradient of the WLS best-fit line; SE = standard error of trend; P-value = probability that the trend is zero; % change y⁻¹= mean percentage change in retrieved CO per year, calculated from WLS regression model predicted values as follows:

% change
$$y^{-1} = \left\{ \left[\left(\frac{\text{Predicted}_{\text{last}}}{\text{Predicted}_{\text{first}}} \right) * 100 \right] - 100 \right\} / ny$$

where ny = number of years. The penultimate two columns correspond to the result of a significance test
performed on the difference between that row's trend and the trend in L3L and L3W, respectively, as follows:

$$Z = \frac{Trend_1 - Trend_2}{\sqrt{SE_1^2 + SE_2^2}}$$

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where SE₁ and SE₂ correspond to the standard errors of Trend₁ and Trend₂ respectively, and Z is the test statistic. Where Z is greater (less) than 1.645 (-1.645) the trend difference is statistically significant to at least 90 % (i.e. p < 0.1). Drift = the measurement drift values given in Deeter et al., (2017). No values are given for the 900 and 300 hPa levels of the profile: we therefore cite values for the 400 and 200 hPa levels to give context to the 300 hPa trends we show; and we expect that the 900 hPa level drift is somewhere between that of the surface and 800 hPa levels, which are both shown.

[†] Units for TCO are mol cm⁻²

LX Discrete Discrete <thdiscrete< th=""> Discrete Di</thdiscrete<>	DIF	Level	Timeseries	Trend (ppby)	Standard Error	P-value	% change v ⁻¹	Sig. di	iff. to:	Drift $(\% v^{-1})$
n13: = 31 n33: = 31 n34: = 31 n35: = 31				(()	(ppbv)		70 change y	L3L	L3W	Dirit (70 y)
Math 2 31 Lit -2.11 0.00 0.000 -1.03 0.00	nL3W = 122	Surface	L3W	-1.63	0.33	0.000	-0.84	No	n/a	
ILBD ILD ILD <thild< th=""> <thild< th=""></thild<></thild<>	nL3L = 31		L3L	-2.11	0.6	0.006	-1.13	n/a	NO	0.60 + 0.40
Indequent 1.2.2 0.2.3 0.0.2 1.0.1 1.00 B13D _{mend} 1.2.3 0.2.8 0.001 -0.68 No No B13D _{mend} 1.2.3 0.2.8 0.001 -0.68 No No B13D _{mend} 1.2.4 0.44 0.48 0.000 -1.2.6 No No B13D _{mend} 2.44 0.48 0.000 -1.2.6 No No No B13D _{mend} -2.49 0.27 0.000 -1.48 No No No No B13D _{mend} -2.49 0.27 0.000 -1.48 No No -1.04 1.01 -0.33 0.007 -1.58 No No -1.04 1.01 -0.33 0.07 -1.58 No No -1.04 0.01 -0.03 0.027 -0.75 No No -1.04 0.01 -0.03 No No -1.04 0.01 -0.03 No No -1.04 0.01 -0.03<	nl 30 101		130.	-1.74	0.31	0.000	-0.30	No	No	-0.69 ± 0.10
Intropage 12 Expansion Last Count Last Count Last No No 900h9 13 -3.14 0.48 0.000 -1.26 No No No 130 -2.51 0.29 0.000 -1.26 No No No 130 -2.51 0.29 0.000 -1.41 No No No 130 -2.47 0.29 0.000 -1.41 No No -1.04 to 1.1 130 -2.47 0.29 0.000 -1.48 No No -1.04 to 1.1 130 -2.47 0.29 0.000 -1.48 No No -1.04 to 1.1 130 -2.12 0.41 0.011 0.39 No No -3.3 to .9 130 -2.12 0.41 0.021 0.38 0.35 0.466 -0.567 No No No -3.3 to .9 130 -2.30 0.60 0.69 No No </td <td>$nL3O_{(water)} = 101$</td> <td></td> <td>L3O_(water)</td> <td>-1.92</td> <td>0.33</td> <td>0.000</td> <td>-0.52</td> <td>No</td> <td>No</td> <td></td>	$nL3O_{(water)} = 101$		L3O _(water)	-1.92	0.33	0.000	-0.52	No	No	
900hPa 13W -2.39 0.29 0.000 -1.26 No n/a 130 -2.51 0.29 0.000 -1.32 Na No No 130 -2.51 0.29 0.000 -1.32 Na No No 130 -2.9 0.35 0.000 -1.61 No No 130 -2.9 0.35 0.000 -1.74 No No -1.64 0.17 130 -2.49 0.27 0.000 -1.48 No No -1.64 0.11 130 -1.21 0.63 0.007 -1.58 n/a No No 0.164 0.11 -1.03 No No 0.164 0.11 1.03 0.104 0.11 0.03 0.037 0.15 No No No No No 1.04 0.11 1.01 1.03 1.01 1.03 1.01 1.01 1.03 1.01 1.01 1.03 1.00 <t< td=""><td>mixed) = 25</td><td></td><td>LSO (mixed)</td><td>1.25</td><td>0.20</td><td>0.001</td><td>0.00</td><td>NO</td><td>NO</td><td></td></t<>	mixed) = 25		LSO (mixed)	1.25	0.20	0.001	0.00	NO	NO	
131 -3.14 0.28 0.000 -1.72 n/a Moo 130 -2.51 0.29 0.000 -1.32 No No 130 -2.4 0.25 0.000 -1.45 No No 130 -2.47 0.29 0.000 -1.47 No n/a 130 -2.47 0.29 0.000 -1.48 No No No 130 -2.47 0.29 0.000 -1.48 No No No No 130 -2.47 0.29 0.000 -1.48 No No No No 130 -1.21 0.41 0.011 -0.33 No No No No 130 134 -0.71 0.40 0.3 0.34 0.87 Yes n/a 130 0.51 0.45 0.45 0.59 No No No No 130 0.51 0.45 0.001 -0.8		900hPa	L3W	-2.39	0.29	0.000	-1.26	No	n/a	
130 -2.51 0.23 0.000 -1.24 NO NO NO 130 _{ment} -3.05 0.35 0.000 -1.61 NO NO 130 _{ment} -2.36 0.35 0.000 -1.44 NO NO 130 -2.29 0.35 0.000 -1.44 NO NO 130 -2.29 0.37 0.000 -1.44 NO NO 130 -2.17 0.03 0.027 0.75 NO NO NO 130 -1.21 0.41 0.021 -0.80 NO NO NO 0.33 ± 0.09 130 -1.21 0.41 0.021 -0.80 NO NO NO NO NO 0.33 ± 0.09 130 -1.24 0.41 0.021 -0.80 NO NO NO NO NO NO NO 1.15 ± 0.12 130 -2.55 ± 4 7.07 0.55 NO NO NO NO			L3L	-3.14	0.48	0.000	-1.72	n/a	NO	
L3Durent -2.4 0.23 0.000 -1.61 NO NO 800hPa L3U -2.9 0.35 0.000 -1.41 NO NO L3D -2.47 0.23 0.000 -1.48 NO NO -1.04 NO L3D -2.47 0.23 0.000 -1.48 NO NO -1.04 1.01 L3Durent -3.56 0.42 0.000 -1.78 NO NO Yes 600hPa L3W -0.962 0.39 0.027 -0.75 No r/a L3D -1.21 0.41 0.011 -0.38 NO NO NO 0.33 ± 0.09 L3D 0.35 0.122 0.41 0.011 NO NO NO NO 1.03 ± 0.35 1.15 NO NO NO 1.03 ± 0.35 1.15 ± 0.12 1.15 ± 0.12 1.15 ± 0.12 1.15 ± 0.12 1.15 ± 0.12 1.15 ± 0.12 1.15 ± 0.12 1.15 ± 0.12 1.15 ± 0.12 1.15 ± 0.12			130	-2.51	0.29	0.000	-1.32	NO	NO	Not given
Index Index <th< td=""><td></td><td></td><td>L3O_(water)</td><td>-2.4</td><td>0.25</td><td>0.000</td><td>-1.20</td><td>No</td><td>No</td><td></td></th<>			L3O _(water)	-2.4	0.25	0.000	-1.20	No	No	
BODIPa L3W -2.36 0.31 0.000 -1.44 No n/a n/a L3D -2.47 0.29 0.000 -1.48 No No 1.04 ± 0.11 L3D -2.47 0.27 0.000 -1.48 No No 1.04 ± 0.11 L3D -3.56 0.42 0.000 -2.08 No YEs 600h7a L3W -0.62 0.33 0.027 -0.75 No n/a L3D -1.21 0.41 0.011 -0.93 No No -0.33 ± 0.09 L3D -1.48 0.43 0.007 -1.18 No No No L3D -1.48 0.43 0.057 -0.59 No No No 1.15 = 0.12 200hPa L3L -0.271+6 0.001 0.031 0.071 No No 1.14 = 0.13 L3D -2.561+16 J.0001 -0.95 No No 0.001 + 0.01 L3D			L3O (mixed)	-3.05	0.55	0.000	-1.01	NO	NO	
I = I = I = I = I = I = I = I = I = I =		800hPa	L3W	-2.36	0.3	0.000	-1.41	No	n/a	
Lio -2.49 0.27 0.000 -1.48 No No -1.0 ±0.11 L30 _{mment} -3.56 0.42 0.000 -2.08 No Yes 600hPa L3W -0.962 0.39 0.027 -0.75 No n ² n ³ L30 -1.21 0.41 0.011 -0.93 No No -0.33 ±0.09 L30 _{mment} -1.48 0.43 0.007 -1.18 No No -0.33 ±0.09 L30 _{mment} -1.48 0.43 0.007 -1.18 No No -0.33 ±0.09 300hPa L3W 0.704 0.3 0.034 0.87 Yes n ³ L30 -0.507 0.55 0.456 -0.56 0.74 Yes L30 -0.507 0.55 0.456 -0.56 0.74 Yes L30 -0.507 0.035 0.456 0.55 0.74 Yes L30 -0.567 0.475 0.59 No No L30 _{mment} 0.483 0.55 0.475 0.59 No No L30 _{mment} -2.08E+15 5.000 -0.89 No No L30 _{mment} -2.08E+15 5.000 -0.49 No No L30 _{mment} -2.08E+15 5.000 -0.11 No No L30 _{mment} -2.08E+15 5.000 -1.19 No Yes No No L30 _{mment} -2.08E+15 5.000 -0.68 Yes No No 0.001 ±0.01 L30 _{mment} -2.05 0.031 0.058 -0.35 Yes n/9 nL30 -2.25 L30 -0.69 0.33 0.058 -0.32 Yes No No 0.69 ±0.10 nL30 _{mment} -1.35 0.35 0.002 -0.68 Yes No No 0.69 ±0.10 nL30 _{mment} -1.35 0.411 0.002 -0.68 Yes No No 0.69 ±0.10 nL30 _{mment} -1.14 0.31 0.001 -0.81 Yes n/9 L30 _{mment} -1.14 0.31 0.001 -0.81 Yes No No tgiven L30 _{mment} -1.14 0.31 0.001 -0.81 Yes No No tgiven L30 _{mment} -1.14 0.31 0.001 -0.77 No No L30 _{mment} -1.14 0.31 0.002 -1.10 No L30 _{mment} -1.14 0.31 0.002 -1.10 No No L30 _{mment} -1.142 0.44 0.010 -1.22 No No No tgiven L30 _{mment} -1.142 0.45 0.002 -1.10 No L30 _{mment} -1.142 0.45 0.002 -1.10 No L30 _{mment} -1.142 0.45 0.002 -1.12 No No L30 _{mment} -1.142 0.45 0.002 -1.12 No No L30 _{mment} -1.142 0.45 0.002 -1.10 No No L30 _{mment} -1.142 0.45 0.002 -1.02 No No L30 _{mm}			L3L	-2.9	0.35	0.000	-1.79	n/a	No	
ISO (mean) -2.49 -3.56 0.22 -0.000 -2.03 -2.03 No Yes 600 hPa 13W -0.962 0.39 0.027 -0.75 No n/a 120 1.21 0.63 0.007 -1.18 No No -0.33 0.037 120 1.21 0.61 0.007 -1.18 No No -0.33 0.001 130 meal 1.48 0.41 0.011 -0.93 No No No 300hPa 13W 0.704 0.3 0.034 0.87 Yes n/a 130 meal 1.48 0.55 0.475 0.59 No No 1.15 ± 0.12 130 meal 0.493 0.65 0.475 0.59 No No 1.45 ± 0.13 130 meal 0.493 0.65 0.475 0.000 -0.81 No n/a 130 meal 3.256+16 4.707+15 0.000 -0.49 No No 0.001 ± 0.01 130 meal			120	-2.47	0.29	0.000	-1.48	NO	NO	-1.04 ± 0.11
ISO(march) 5.3.0 0.5.2 0.000 7.4.0 No no 600hPa L3W -0.60 0.039 0.027 -0.75 No no L3O -1.21 0.43 0.011 0.033 No No No L3O -1.21 0.41 0.012 -0.93 No No No L3O -1.48 0.43 0.077 -1.18 No No A00 Pb; L3O 0.59 0.55 0.456 -0.56 n/4 No No 100 Pb; L3O 0.58 0.35 0.122 0.71 No No 100 Pb; 100P			L3O _(water)	-2.49	0.27	0.000	-1.47	No	NO	
600hPa I.3W -0.962 0.39 0.027 -0.75 No n/a Na I30 -1.21 0.41 0.011 -0.93 No No 0.33 ± 0.09 I30 -1.21 0.41 0.0125 -0.80 No No 0.03 ± 0.09 J300hPa I.3W 0.704 0.3 0.034 0.87 Yes n/a 400 hPa: I31 -0.507 0.65 0.456 -0.55 n/a Yes 0.00 hPa: 115 ± 0.12 0.00 hPa: 13 ± 0.20 0.71 No No 1.49 ± 0.13 I30 0.58 0.355 0.475 0.59 No No 1.49 ± 0.13 I30 -2.56±16 4.70±15 0.000 -0.81 No n/a 1.49 ± 0.13 I30 -2.56±16 5.10±15 0.000 -1.19 No 0.01 ± 0.01 I30 -2.35±16 5.00±15 0.000 -1.42 Na No No 0.02 0.01 ± 0.01 <td></td> <td></td> <td>L3O(mixed)</td> <td>-3.50</td> <td>0.42</td> <td>0.000</td> <td>-2.06</td> <td>NU</td> <td>res</td> <td></td>			L3O(mixed)	-3.50	0.42	0.000	-2.06	NU	res	
IBA -2.1 0.63 0.007 -1.58 n/a No No 0.033 ± 0.05 I3Opment -1.04 0.41 0.025 -0.80 No No 0.033 ± 0.05 300hPa I3W 0.704 0.3 0.034 0.37 Yes n/a 400 hPs: I3D -0.507 0.65 0.456 -0.56 n/a 400 hPs: 1.15 ± 0.12 200 hPs: 1.15 ± 0.12 200 hPs: 1.15 ± 0.12 200 hPs: 1.42 ± 0.13 200 hPs: 1.43 ± 0.23 200 hPs: 1.43 ± 0.23 200 hPs: 1.43 ± 0.23 200 hPs: 1.44 ± 0.13 200 hPs: 1.43		600hPa	L3W	-0.962	0.39	0.027	-0.75	No	n/a	
ISO 1.14 0.41 0.011 -0.93 No No -0.33 ± 0.09 I3O _{persen} -1.04 0.41 0.025 -0.80 No No No 300hPa I3W 0.704 0.3 0.034 0.37 Yes No No 300hPa I3W 0.704 0.65 0.656 0.56 n/6 n/6 n/8 0.00 hPa: 1.15 ± 0.12 0.00 hPa: 1.49 ± 0.13 1.49 ± 0.14 1.49 ± 0.13 1.49 ± 0.13 1.49 ± 0.13 1.49 ±			L3L	-2.1	0.63	0.007	-1.58	n/a	No	
I.330 _{metern} -1.48 0.43 0.025 -0.80 No No 300hP9 I3W 0.704 0.3 0.034 0.87 Yes n/a 400 hPa: I3D 0.577 0.65 0.456 -0.56 n/a Yes 1.15 ± 0.12 I3O 0.58 0.35 0.422 0.71 No No No 1.45 ± 0.12 I3O _{metern} 0.49 0.20 0.030 0.59 No No No 1.49 ± 0.13 I3O _{metern} 0.493 0.65 0.475 0.59 No No 1.49 ± 0.13 I3O _{metern} 0.483 0.65 0.475 0.59 No No 0.001 ± 0.01 I3O -2.56±16 4.70±15 0.000 -1.19 No No No 0.01 ± 0.01 I3O -3.24±16 5.06±15 0.000 -1.42 n/a No No 0.69 ± 0.10 nI3W -1.51 0.32 0.68 0.022			L30	-1.21	0.41	0.011	-0.93	NO	NO	-0.33 ±0.09
JAP 1.48 0.43 0.007 1.18 NO NO 300hPa L3W 0.704 0.3 0.034 0.87 Yes n/a No			L3O _(water)	-1.04	0.41	0.025	-0.80	NO	NO	
300hPa L3W 0.704 0.3 0.034 0.87 Yes n/a 400hPs: 1.50.12 L30 0.58 0.35 0.122 0.71 No No No L30 0.483 0.65 0.759 No No No L30 0.483 0.65 0.755 No No No L30 -2.166+16 4.706+15 0.000 -0.81 No No No 1.49±0.13 L30 -2.266+16 5.106+15 0.000 -0.89 No No No 0.001±0.001 L30 -2.268+16 5.106+15 0.000 -1.19 No Ves No JJA -3.246+16 3.506+15 0.000 -1.19 No Ves No -0.69±0.13 nL30_matel 1.34 0.31 0.001 -0.87 Yes No -0.69±0.10 nL30_matel 1.35 0.43 0.001 -0.21 Yes No -0.69±0.10 <td></td> <td></td> <td>L3O(mixed)</td> <td>-1.48</td> <td>0.43</td> <td>0.007</td> <td>-1.18</td> <td>NO</td> <td>NO</td> <td></td>			L3O(mixed)	-1.48	0.43	0.007	-1.18	NO	NO	
L31 -0.507 0.65 0.456 -0.56 -0.71 N0 N0 N0 L3200hPa: L300hPa: L300hPa: L300mmed L30 0.49 0.20 0.030 0.59 N0 N0 N0 L30 0.49 0.20 0.030 0.59 N0 N0 L30 -2.76+16 4.70E+15 0.000 -0.81 N0 n/a L30 -2.56E+16 4.70E+15 0.000 -0.95 N0 N0 0.011±0.001 L30(med) -2.08E+16 5.10E+15 0.000 -1.19 N0 Ves JJA 0.31 -2.56 0.60 0.000 -1.42 N/a N0 0.69±0.10 L30(med) -2.08E+16 5.10E+15 0.000 -1.19 N0 0.0.9±0.10 L30 -2.85 0.60 0.000 -1.42 N/a N0 0.0.9±0.10 L30(med) -1.35 0.35 0.001 -1.42 N0 N0 0.0.9±0.10 L30(mee		300hPa	L3W	0.704	0.3	0.034	0.87	Yes	n/a	400 h Day
L30 0.58 0.32 0.71 N No No 200 hrs. L30(person) 0.49 0.20 0.030 0.59 No No No No No 1.49±0.13 TCO L3W -2.16E+16 4.70E+15 0.000 -0.81 No No No 0.001±0.001 L30 -2.25E+16 6.00E+15 0.000 -0.81 No No No 0.001±0.001 L30 -2.25E+16 5.10E+15 0.000 -1.19 No No No 0.001±0.001 L30(person) -3.24E+16 3.50E+15 0.000 -1.19 No Yes No -0.69±0.13 0.000 -1.19 No Yes No -0.69±0.10 nL30 E30 -0.59 0.63 0.003 -0.50 Yes No -0.69±0.10 nL31 E30 0.51 0.60 0.001 -1.19 No Yes No -0.69±0.10 nL30 1.32 <td></td> <td></td> <td>L3L</td> <td>-0.507</td> <td>0.65</td> <td>0.456</td> <td>-0.56</td> <td>n/a</td> <td>Yes</td> <td>400 IIPa. 1 15 +0 12</td>			L3L	-0.507	0.65	0.456	-0.56	n/a	Yes	400 IIPa. 1 15 +0 12
I30 _(mixed) 0.49 0.20 0.030 0.59 No No 1.49±0.13 I30 _(mixed) 0.483 0.65 0.475 0.59 No No No I31 -2.27E+16 6.00E+15 0.000 -0.88 No No 0.001±0.001 I30 -2.56E+16 4.70E+15 0.000 -0.89 No No 0.001±0.001 I30 -2.56E+16 5.10E+15 0.000 -1.19 No No No I30 -2.56E+16 3.50E+15 0.000 -1.19 No Yes JA 131 -2.285 0.60 0.000 -1.42 n/a No No nL30 = 101 131 -0.62 0.31 0.069 -0.32 Yes No No nL30 (mixel) 1.34 0.35 0.001 -0.61 Yes No nL30 (mixel) 1.30 0.31 0.001 -0.51 Yes No No No No			L30	0.58	0.35	0.122	0.71	No	No	200 hPa:
ISOmmedie 0.483 0.65 0.475 0.59 No No TCO I3W -2.16E+16 4.70E+15 0.000 -0.89 n/a No n/a I3O -2.27E+16 4.70E+15 0.000 -0.95 No No 0.001±0.001 I3O -2.56E+16 4.70E+15 0.000 -1.19 No Yes nI3W -2.24E+16 3.50E+15 0.001 -1.19 No Yes nI3D -3.24E+16 3.50E+15 0.000 -1.42 n/a Yes nI3D 131 -2.85 0.60 0.003 -0.52 Yes No nI3O 130 -0.99 0.28 0.003 -0.68 Yes No nI3O 130 -1.34 0.31 0.001 -1.68 Yes No I3O -1.34 0.31 0.001 -1.55 n/a Ne No I3O -1.43 0.35 0.002 -1.06			L3O _(water)	0.49	0.20	0.030	0.59	No	No	1.49 ± 0.13
TCO I3W -2.16E+16 4.70E+15 0.000 -0.81 No n/a No I3U -2.27E+16 6.00E+15 0.000 -0.85 No No 0.001±0.001 I3O -2.56E+16 5.10E+15 0.001 -0.79 No No I3O -2.36E+16 3.50E+15 0.000 -1.19 No Yes JJA -2.08E+16 3.50E+15 0.000 -1.19 No Yes nL3W = 231 Surface I3W -0.69 0.33 0.058 -0.35 Yes No nL3O_exter) 136 I3O -0.99 0.28 0.003 -0.50 Yes No nL3O_exter) 130 -1.34 0.31 0.001 -0.81 Yes No nL3D_exter) -0.62 0.51 Yes No Not given I3O -1.34 0.33 0.002 -1.10 No No I3O -1.43 0.35			L3O _(mixed)	0.483	0.65	0.475	0.59	No	No	
13L -2.27E+15 6.000 -0.95 No No 0.001±0.001 13O -2.56E+16 5.10E+15 0.000 -0.79 No No 0.001±0.001 13O -2.68E+16 5.10E+15 0.000 -1.19 No No No JJA -2.08E+16 5.10E+15 0.000 -1.19 No Yes JJA N13W = 231 Surface L3W -0.69 0.33 0.058 -0.35 Yes n/a n130 = 235 L3O -0.99 0.28 0.003 -0.69 -0.69±0.10 n130-gwater) = 136 L3O(water) -0.62 0.31 0.069 -0.32 Yes No nL30-gwater) = 136 L3O(water) -0.62 0.31 0.061 -1.55 n/a nL30-gwater) = 136 L3O(water) -0.91 0.38 0.032 -0.68 Yes No 130 (mated) -2.05 0.45 0.001 -1.23 No No 130 (mated)		тсо	L3W	-2.16E+16	4.70E+15	0.000	-0.81	No	n/a	
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$			L3L	-2.27E+16	6.00E+15	0.000	-0.89	n/a	No	
L3O _(mater) -2.08E+16 5.10E+15 0.001 -0.79 No No JJA nL3W = 231 Surface L3W -0.69 0.33 0.058 -0.43 res nL3W = 231 Surface L3W -0.69 0.33 0.058 -0.50 Yes n/a nL3t = 101 L3L -2.85 0.60 0.000 -1.12 n/a Yes No -0.69 ± 0.10 nL3G(mater) -1.62 0.31 0.069 -0.32 Yes No -0.69 ± 0.10 nL3D(mater) -1.62 0.31 0.069 -0.32 Yes No 900hPa L3W -1.34 0.31 0.001 -1.55 n/a Yes No Not given L3O (mater) -0.91 0.38 0.022 -1.10 No n/a L3O (mater) -1.01 0.46 0.001 -1.22 n/a No L3O (mater) -1.01 0.46 0.001 -1.22 <th< td=""><td></td><td></td><td>L30</td><td>-2.56E+16</td><td>4.70E+15</td><td>0.000</td><td>-0.95</td><td>No</td><td>No</td><td>0.001 ± 0.001</td></th<>			L30	-2.56E+16	4.70E+15	0.000	-0.95	No	No	0.001 ± 0.001
I30 -3.24E+16 3.50E+15 0.000 -1.19 No Yes JJA nL3W = 231 Surface L3W -0.69 0.33 0.058 -0.35 Yes n/a nL3L = 101 L3L -2.85 0.60 0.000 -1.42 n/a r/a Yes nL30_sets] 136 L30 -0.99 0.28 0.000 -0.69 Yes No -0.69 ± 0.10 nL30_weter] 136 L30 -0.62 0.31 0.069 -0.32 Yes No -0.69 ± 0.10 nL30_mixed] 1.35 0.35 0.002 -0.68 Yes No Net given L30 -1.34 0.31 0.001 -1.81 Yes No Not given L30 -1.43 0.35 0.001 -1.23 No No 1.04 ± 0.11 L30 -1.47 0.39 0.002 -1.10 No n/a No L30 -1.47 0.39			L3O _(water)	-2.08E+16	5.10E+15	0.001	-0.79	No	No	
JJA			L3O _(mixed)	-3.24E+16	3.50E+15	0.000	-1.19	No	Yes	
$\begin{array}{c c c c c c c c c c c c c c c c c c c $	IIΔ									
n130 = 101 Surface L3W - 0.69 0.33 0.058 0.035 0.45 0.45 0.45 0.45 0.45 0.45 0.45 0.4	33	.			0.00		0.05		,	
$\begin{array}{cccccccccccccccccccccccccccccccccccc$	nL3W = 231	Surface	L3W	-0.69	0.33	0.058	-0.35	Yes	n/a	
n130 user) 136 133 (uster) -0.62 0.31 0.069 -0.32 Yes No n130 (uster) -1.35 0.35 0.002 -0.68 Yes No 900hPa L3W -1.34 0.31 0.001 -0.81 Yes No 130 (uster) -0.62 0.33 0.001 -0.81 Yes No 900hPa L3W -1.34 0.31 0.001 -0.81 Yes No L30 -1.43 0.35 0.001 -1.55 n/a Yes No Not given L30 -1.43 0.35 0.001 -1.23 No No Not given L30 (uster) -0.91 0.38 0.032 -1.01 No n/a L30 (uster) -1.01 0.46 0.002 -1.10 No n/a L30 (uster) -1.01 0.46 0.049 -0.75 No No L30 (uster) -1.01 0.46 0.071 -0.83 No n/a No L30 (uster) -0.13 <t< td=""><td>nL3L = 101 nL3O = 235</td><td></td><td>130</td><td>-2.85</td><td>0.60</td><td>0.000</td><td>-1.42</td><td>li/d Voc</td><td>No</td><td>0.60 ± 0.10</td></t<>	nL3L = 101 nL3O = 235		130	-2.85	0.60	0.000	-1.42	li/d Voc	No	0.60 ± 0.10
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	nl 30(130(-0.62	0.20	0.005	-0.32	Yes	No	-0.09±0.10
900hPa L3W -1.34 0.31 0.001 -0.81 Yes n/a I30 -1.43 0.35 0.001 -0.87 Yes No Not given I30 _(water) -0.91 0.38 0.032 -0.56 Yes No I30 _(mixed) -2.05 0.45 0.001 -1.23 No No 800hPa L3W -1.55 0.41 0.002 -1.10 No n/a I31 -1.62 0.54 0.010 -1.22 n/a No I30 -1.47 0.39 0.002 -1.07 No No -1.04±0.11 I30 _(water) -1.01 0.46 0.049 -0.75 No No I30 _(water) -1.01 0.46 0.071 -0.83 No n/a I30 -0.75 0.43 0.105 -0.70 No No -0.33±0.09 I30 _(water) -0.41 0.49 0.424 -0.39 No No 300hPa L3W 1.30 0.43 0.010 1.40 No n/a I30 -0.75 0.43 0.105 -0.70 No No -0.33±0.09 I30 _(water) -1.12 0.45 0.026 -1.02 No No 130 _(water) -1.12 0.45 0.026 -1.02 No No 130 _(water) -1.12 0.45 0.026 -1.02 No No 130 _(water) 0.86 0.42 0.060 0.92 No No 130 _(water) 0.86 0.42 0.060 0.92 No No 1430 -3.38 0.47 0.011 1.52 No No 1430 -3.38 0.47 0.011 1.52 No No 1430 -1.38 0.47 0.011 1.52 No No 149 -0.13 149 +0.13 149 +0.13 149 +0.13 149 +0.13 149 +0.13 149 +0.13 149 +0.13 150 -3.75 0.43 0.001 2.52 No No 149 +0.13 149 +0.13 149 +0.13 150 -3.75 0.43 0.001 2.52 No No 149 +0.13 149 +0.13 150 -3.75 0.43 0.001 2.52 No No 149 +0.13 150 -3.75 0.441 0.49 -0.40 No n/a 140 +0.74 No 130 -8.75 +15 0.109 -0.40 No n/a 140 +0.01 +0.0	$nL3O_{(mixed)} = 98$		L30 (mixed)	-1.35	0.35	0.002	-0.68	Yes	No	
900hPa L3W -1.34 0.31 0.001 -0.81 Yes n/a L3L -2.58 0.60 0.001 -1.55 n/a Yes L3O -1.43 0.35 0.001 -0.87 Yes No Not given L3O (1.43 0.35 0.001 -1.23 No No L3O (mixed) -2.05 0.45 0.001 -1.23 No No 800hPa L3W -1.55 0.41 0.002 -1.10 No n/a L3L -1.62 0.54 0.010 -1.22 n/a No L3O -1.47 0.39 0.002 -1.07 No No -1.04 \pm 0.11 L3O (mixed) -1.89 0.48 0.002 -1.07 No No -1.04 \pm 0.11 L3O (mixed) -1.89 0.48 0.002 -1.38 No No 600hPa L3W -0.91 0.46 0.071 -0.83 No n/a L3O (mixed) -1.89 0.48 0.002 -1.38 No No 600hPa L3W -0.91 0.46 0.071 -0.83 No n/a L3O (mixed) -1.12 0.45 0.026 -1.02 No No -0.33 \pm 0.09 L3O (mixed) -1.12 0.45 0.026 -1.02 No No L3O (mixed) -1.12 0.45 0.026 -1.02 No No 130 1.38 0.47 0.011 1.52 No No 1.30 1.38 0.47 0.011 1.52 No No 1.49 \pm 0.13 1.49 \pm 0.13 1.30 0.02 1.02 No No 1.30 0.12 1.00 No No 1.30 0.12 1.00 No No 1.30 0.875E+15 0.109 -0.40 No n/a 1.49 \pm 0.13 1.30 0.01 \pm 0.00 No No 0.001 \pm 0.00 No 1.30 -8.75E+15 6.40E+15 0.192 -0.36 No No 0.001 \pm 0.0001 \pm 0.001 \pm 0.001 \pm 0.0001 \pm 0.001 \pm 0.0001 \pm 0.001 \pm 0.001 \pm 0.0001 \pm 0.001 \pm 0.0001 \pm 0.001 \pm 0.0001 \pm 0.00001 \pm 0.0	(fince)		(mixed)						,	
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$		900hPa	L3W	-1.34	0.31	0.001	-0.81	Yes	n/a	
LOC 1-1-3 0.33 0.001 -0.57 NO NO NO green L3O _(water) -0.91 0.38 0.032 -0.56 Yes No L3O _(mixed) -2.05 0.45 0.001 -1.23 No No 800hPa L3W -1.55 0.41 0.002 -1.10 No n/a L31 -1.62 0.54 0.010 -1.22 n/a No L30 -1.47 0.39 0.002 -1.07 No No -1.04±0.11 L30 _(water) -1.01 0.46 0.049 -0.75 No No L30 _(mixed) -1.89 0.48 0.002 -1.38 No n/a L30 -0.75 0.43 0.105 -0.70 No No -0.33±0.09 L30 _(water) -0.41 0.49 0.424 -0.39 No No L30 _(water) -1.12 0.45 0.026 -1.02 No No L30 _(water) -1.12 0.45 0.026 -1.02 No No 130 _(mixed) -1.12 0.45 0.026 -1.02 No No 130 _(water) 0.86 0.42 0.060 0.92 No No L30 _(water) 0.86 0.42 0.060 0.92 No No TCO L3W -9.84E+15 5.70E+15 0.109 -0.40 No n/a L30 -8.75E+15 6.40E+15 0.192 -0.36 No No 0.001±0.001 L30 _(water) -1.12E 0.45 0.109 -0.40 No n/a L30 -0.75 0.86 0.42 0.060 0.92 No No 1.49±0.13 L30 _(water) -0.84E+15 5.70E+15 0.109 -0.40 No n/a L30 -8.75E+15 6.40E+15 0.192 -0.36 No No 0.001±0.001 L30 _(water) -1.12E 0.45 0.282 -0.32 No No			130	-2.56	0.60	0.001	-1.55	li/d Voc	No	Not given
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$			130(-0.91	0.35	0.032	-0.56	Yes	No	NOT BIVEI
800hPa L3W -1.55 0.41 0.002 -1.10 No n/a L3L -1.62 0.54 0.010 -1.22 n/a No L3O -1.47 0.39 0.002 -1.07 No No L3O(water) -1.01 0.46 0.049 -0.75 No No L3O(mbed) -1.89 0.48 0.002 -1.38 No No 600hPa L3W -0.91 0.46 0.071 -0.83 No n/a L3L -0.38 0.47 0.440 -0.36 n/a No L3O -0.75 0.43 0.105 -0.70 No No -0.33 ± 0.09 L3O(mbed) -1.12 0.45 0.026 -1.02 No No -1.15 ± 0.12 L3O(mbed) -1.12 0.45 0.026 -1.02 No No 1.15 ± 0.12 L3O(mbed) -1.12 0.45 0.026 -1.02 No No 1.15 ± 0.12 L3O(mbed) -1.12 0.45 0.026 -1.02			L30 (mixed)	-2.05	0.45	0.001	-1.23	No	No	
800hPa L3W -1.55 0.41 0.002 -1.10 No n/a L3L -1.62 0.54 0.010 -1.22 n/a No L3O -1.47 0.39 0.002 -1.07 No No -1.04±0.11 L3O -1.47 0.39 0.002 -1.07 No No -1.04±0.11 L3O -1.49 0.46 0.049 -0.75 No No L3O -1.89 0.48 0.002 -1.38 No No 600hPa L3W -0.91 0.46 0.071 -0.83 No n/a L3O -0.75 0.43 0.105 -0.70 No No -0.33±0.09 L3O -0.75 0.43 0.105 -0.70 No No -0.33±0.09 L3O 1.30 0.43 0.010 1.40 No n/a No L3O 1.38 0.47 0.011 1.52 No No 1.15±0.12 200 hPa: L3O 1.38 0.47 0.011			- (mixed)				-	-		
$\begin{array}{cccccccccccccccccccccccccccccccccccc$		800hPa	L3W	-1.55	0.41	0.002	-1.10	No	n/a	
$\begin{array}{cccccccccccccccccccccccccccccccccccc$			130	-1.62	0.54	0.010	-1.22	n/a No	NO	1 04 ±0 11
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$			130	-1.47	0.39	0.002	-0.75	No	No	-1.04 ±0.11
600hPa L3W -0.91 0.46 0.071 -0.83 No n/a L3L -0.38 0.47 0.440 -0.36 n/a No L3O -0.75 0.43 0.105 -0.70 No No -0.33 ± 0.09 L3O(water) -0.41 0.49 0.424 -0.39 No No L3O(mixed) -1.12 0.45 0.026 -1.02 No No 300hPa L3W 1.30 0.43 0.010 1.40 No n/a L3L 2.51 1.30 0.072 2.90 n/a No 1.15±0.12 L3O 1.38 0.47 0.011 1.52 No No 1.49±0.13 L3O(water) 0.86 0.42 0.060 0.92 No 1.49±0.13 L3O(mixed) 2.24 0.53 0.001 2.52 No 1.49±0.13 L3O(water) 0.86 0.422 0.60 0.92 No 1.49±0.13 L3O(water) 0.84E+15 5.70E+15 0.109 -0.40 No			L3O (water)	-1.89	0.48	0.002	-1.38	No	No	
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$			(mixed)							
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$		600hPa	L3W	-0.91	0.46	0.071	-0.83	No	n/a	
L30 (water) -0.41 0.49 0.424 -0.39 No No L30 (mixed) -1.12 0.45 0.026 -1.02 No No 300hPa L3W 1.30 0.43 0.010 1.40 No n/a L3L 2.51 1.30 0.072 2.90 n/a No 400 hPa: L3O 1.38 0.47 0.011 1.52 No No 200 hPa: L3O (water) 0.86 0.42 0.060 0.92 No No L3O (mixed) 2.24 0.53 0.001 2.52 No No TCO L3W -9.84E+15 5.70E+15 0.109 -0.40 No n/a L3L -1.01E+16 1.00E+16 0.331 -0.42 n/a No L3O (water) -7.62E+15 6.80E+15 0.282 -0.32 No No			130	-0.38	0.47	0.440	-0.36	n/a No	NO	0.22 + 0.00
L3O _(mixed) -1.12 0.45 0.026 -1.02 No No 300hPa L3W 1.30 0.43 0.010 1.40 No n/a L3L 2.51 1.30 0.072 2.90 n/a No 400 hPa: L3O 1.38 0.47 0.011 1.52 No No 1.15 ± 0.12 200 hPa: L3O _(witer) 0.86 0.42 0.060 0.92 No No 1.49 ± 0.13 L3O _(mixed) 2.24 0.53 0.001 2.52 No No TCO L3W -9.84E+15 5.70E+15 0.109 -0.40 No n/a L3L -1.01E+16 1.00E+16 0.331 -0.42 n/a No L3O -8.75E+15 6.40E+15 0.192 -0.36 No No 0.001 ± 0.001 L3O _(witer) -7.62E+15 6.80E+15 0.282 -0.32 No No			L30()	-0.41	0.49	0.424	-0.39	No	No	-0.33 ± 0.05
300hPa L3W 1.30 0.43 0.010 1.40 No n/a L3L 2.51 1.30 0.072 2.90 n/a No 400 hPa: L3O 1.38 0.47 0.011 1.52 No No 1.15 ± 0.12 L3O 1.38 0.47 0.011 1.52 No No 1.15 ± 0.12 L3O 1.38 0.47 0.011 1.52 No No 1.15 ± 0.12 L3O(water) 0.86 0.42 0.060 0.92 No No 1.49 ± 0.13 L3O(mixed) 2.24 0.53 0.001 2.52 No No L3U -1.01E+16 1.00E+16 0.331 -0.42 n/a No L3O -8.75E+15 6.40E+15 0.192 -0.36 No No 0.001±0.001 L3O -8.75E+15 6.80E+15 0.282 -0.32 No No L3O(water) -7.62E+15 6.80E+15 0.109 -0.48 No No			L3O (water)	-1.12	0.45	0.026	-1.02	No	No	
300hPa L3W 1.30 0.43 0.010 1.40 No n/a L3L 2.51 1.30 0.072 2.90 n/a No 400 hPa: L3O 1.38 0.47 0.011 1.52 No No 1.15 ± 0.12 L3O 0.86 0.42 0.060 0.92 No No 1.49 ± 0.13 L3O _(mixed) 2.24 0.53 0.001 2.52 No No L30 -9.84E+15 5.70E+15 0.109 -0.40 No n/a L31 -1.01E+16 1.00E+16 0.331 -0.42 n/a No L30 -8.75E+15 6.40E+15 0.192 -0.36 No 0.001 ± 0.001 L30 -8.75E+15 6.80E+15 0.282 -0.32 No No L30 -1.18E+16 6.90E+15 0.109 -0.48 No No			(mixeu)						,	
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$		300hPa	L3W	1.30	0.43	0.010	1.40	No n/r	n/a	400 hPa:
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$			130	2.51	1.30 0.47	0.072	2.90	n/a No	NO No	1.15 ± 0.12
L30 (mixed) 2.24 0.53 0.001 2.52 No No TCO L3W -9.84E+15 5.70E+15 0.109 -0.40 No n/a L3L -1.01E+16 1.00E+16 0.331 -0.42 n/a No L3O -8.75E+15 6.40E+15 0.192 -0.36 No No 0.001±0.001 L3O (water) -7.62E+15 6.80E+15 0.282 -0.32 No No L3O (mixed) -1.18E+16 6.90E+15 0.109 -0.48 No No			L30	0.86	0.47	0.060	0.92	No	No	200 hPa:
TCO L3W -9.84E+15 5.70E+15 0.109 -0.40 No n/a L3L -1.01E+16 1.00E+16 0.331 -0.42 n/a No L3O -8.75E+15 6.40E+15 0.192 -0.36 No No 0.001±0.001 L3O _(water) -7.62E+15 6.80E+15 0.282 -0.32 No No L3O _(mixed) -1.18E+16 6.90E+15 0.109 -0.48 No No			L30(mixed)	2.24	0.53	0.001	2,52	No	No	1.49 ± 0.13
$\begin{array}{cccccccccccccccccccccccccccccccccccc$			(mixeu)							
L3L -1.01E+10 1.00E+10 0.331 -0.42 n/a NO L3O -8.75E+15 6.40E+15 0.192 -0.36 No No 0.001±0.001 L3O _(water) -7.62E+15 6.80E+15 0.282 -0.32 No No L3O _(mixed) -1.18E+16 6.90E+15 0.109 -0.48 No No		TCO	L3W	-9.84E+15	5.70E+15	0.109	-0.40	No n/-	n/a	
L30 (water) -7.62E+15 6.80E+15 0.192 -0.32 No No 0.001±0.001 L30(water) -7.62E+15 6.80E+15 0.282 -0.32 No No L30(mixed) -1.18E+16 6.90E+15 0.109 -0.48 No No			L3L 130	-1.U1E+16 -8 75F±15	1.00E+16 6 40F+15	0.331	-0.42	n/a No	NO No	0.001 ± 0.001
L30 _(mixed) -1.18E+16 6.90E+15 0.109 -0.48 No No			L30	-7.62F+15	6.80F+15	0.282	-0.32	No	No	0.001 ± 0.001
			L30(mixed)	-1.18E+16	6.90E+15	0.109	-0.48	No	No	



Figure 1 (a) Map of Nova scotia and the surrounding region in Atlantic Canada. Lighter colours on land indicate higher elevation. The pink icon shows the location of Halifax, the coastal city that we focus on in this study (see Section 2.1.3), and the red box shows the MOPITT L3 1° x 1° gridbox containing Halifax. (b) Map zoomed to the MOPITT L3 gridbox containing Halifax (red dashed box), with the approximate location of individual L2 retrieval footprints shown (blue boxes = L2 surface index of water, green boxes = L2 surface index of land). L2 retrieval footprints with a midpoint that falls within the boundaries of the L3 gridbox will be averaged together to create the L3 data, according to certain rules – see Section 2.1.2 for full explanation.



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Figure 2 Distribution of percentage difference[†] between temporally coincident retrieved VMRs (selected profile levels) and CO total column (TCO) from L3W and L3L. Squares = mean differences, with the p value associated with each mean difference (from a 2-tailed Student's t test) given on the right-hand side y axis. Plus symbols = outliers[‡]. Red triangles = mean percentage difference between a priori values. Red lines = range of a priori difference values (where barely visible this means range is very small).

[†]Method for calculating percentage differences: $\left\{ \left(\frac{L3W}{L3L} \right) * 100 \right\} - 100$

[‡]Outliers defined as: above (below) percentile 75 (25) + (-) 1.5 * interquartile range



Figure 3 Mean retrieval averaging kernel (AK) rows and DFS values for L3L ("Land", left column) and L3W ("Water", right column) in DJF (top row) and JJA (bottom row), selected profile levels only. Filled diamonds indicate diagonal value location for that AK. Numbers in legend indicate corresponding retrieval level of AK and show the mean rowsum for that AK level.



Figure 4 Mean temperature profile from ERA-Interim for the closest all-land and all-water model gridboxes to Halifax. Errorbars = standard deviation. Data are the mean of the 12:00 UTC and 18:00 UTC timesteps (this corresponds to 08:00 and 14:00 local time, the closest ERA Interim timesteps to local MOPITT overpass time of ~10:30), for 2001-2017.

Figure 5 (a) Distribution of percentage difference[†] between simulated temporally coincident VMR retrievals in L3W (SIM_{L3W}) and L3L (SIM_{L3L}). Squares = mean differences. The p value associated with each mean difference (from a 2-tailed Student's t test) is given on the right-hand side y axis (top row = Δ SIM; bottom 895 row = ΔRET). Plus symbols = outliers[‡]. Faint shading = corresponding ΔRET boxplots for comparison. Note that the sample size is different to Figure 2 owing to the CAMSRA data used as the X_{tr,sim} profile only covering a subset of the MOPITT years (2003-2016 vs 2001-2017 in Figure 2). The ∆RET boxplots overlaid cover this shortened period. (b) Distribution of the $log_{10}X_{tr,sim} - log_{10}X_{apr,sim}$ values calculated during the simulation of retrieved profiles (see Equation 2) in DJF (top row) and JJA (bottom row). 900

[†]Method for calculating percentage differences: $\left\{ \left(\frac{L3W}{L3L} \right) * 100 \right\} - 100$

[‡]Outliers defined as: above (below) percentile 75 (25) + (-) 1.5 * interquartile range

Figure 6 Seasonal median L2[†] retrieved VMR (left column), a priori VMR (centre column), and RET-APR (right column) at the surface profile level in DJF (top row) and JJA (bottom row). Values to the right above RET-APR plots = $\overline{(L2 \text{ retrievals over water})} - \overline{(L2 \text{ retrievals over land})}$ for plotted area (data were first binned according to L2 surface index); numbers in brackets correspond to significance of mean difference using a 2-tailed Student's t test. Blue or green dashed square = outline of L3 grid box that contains Halifax.

[†]These maps were created from L2 data that were interpolated to a regular 0.25° x 0.25° grid for ease of plotting.

Figure 7 Mean ERA-Interim 10-metre winds (vectors) and MOPITT L2[†] VMR at the surface profile level (shading) for days when retrieved surface level VMRs in L3W are greater than in L3L (L3W > L3L) and days when they are less (L3W < L3L). Top row = DJF; bottom row = JJA. Blue dashed square = outline of L3 grid box that contains Halifax.

[†]These maps were created from L2 data that were interpolated to a regular 0.25° x 0.25° grid for ease of plotting.

Figure 8 Boxplots showing the seasonal distribution of retrieved surface level VMR values from L3W, L3L, L3O, $L3O_{(mixed)}$ and $L3O_{(water)}$. Squares = mean values; Red triangles = corresponding mean a priori values. Sample sizes are given below the top x axis.

Figure 9 WLS regression best-fit lines calculated from seasonal mean retrieved surface VMR timeseries in DJF (top row) and JJA (bottom row). Left column = L3L (green) and L3W (blue); Right column = L3O (black), L3O_(mixed) (brown) and L3O_(water) (blue). The daily observations corresponding to each seasonal mean value are represented by colour-coded boxplots each year, and the seasonal mean value is represented by filled squares. The dashed red line is the mean of the corresponding seasonal mean a priori data from each of the timeseries in the respective panel. Colour-coded values below the top x axis correspond to the number of observations each season. Values in the legend are the value, standard error, and probability of zero value of the trend, respectively.

Figure 10 (a) As Figure 3 but for MAM and SON; (b) As Figure 9 but for MAM and SON