

Assessing the influence of long-range transport of aerosols on the PM_{2.5} chemical composition and concentration in the Aburrá Valley

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Abstract. Assessing long-range transport (LRT) of pollutants recognizes that multiple sources of varying scale and location can impact air quality. In the Aburrá Valley (AV), Colombia, and other cities in Northern South America, biomass burning (BB), dust, and volcanic degassing have been identified as sources of LRT of aerosols. However, the impact of these sources on air quality and their characterization have yet to be thoroughly studied. This work investigates the influence of these sources on the chemical composition of PM_{2.5} during annual and intra-annual high-load aerosol events in the AV. We identified, tracked, and meteorologically characterized LRT events and evaluated their influence on PM_{2.5} concentration and chemical composition. Annually, we found that LRT of aerosols from BB, dust and volcanic degassing influence approximately 13%, 8% and 13% of days, respectively. We ran a Positive Matrix Factorization (PMF) for each kind of event, identifying high contribution in organic carbon (OC1, OC2), F⁻ and secondary aerosols trace SO₄²⁻ and NO₃⁻ for the BB events, crustal mineral along with Ti and Ca contribution for dust events and SO₄²⁻, Na, Al and Ca for volcanic events. The increasing concentration of some ions and toxic heavy metals (Cr, Mn, Cd, and Ni) were also related to BB and volcanic degassing influence. ~~During these LRT events, the BB fraction BB exhibited the highest contribution of PM_{2.5} dominates by frequency and amount, averaging 11.14 µg/m³ (38%). On average, within the events (~ 11 µg/m³), while aerosols from dust and volcanic degassing contribute 6.77 µg/m³ (34%) and 6.46 µg/m³ (30%) of the concentrations. Of the three, dust events showed fewer affected days. events were also significant (< 7 µg/m³).~~ The study highlighted hotspot zones such as the Orinoco and Middle Magdalena Valley for BB aerosols, the Caribbean for dust, and the Nevado del Ruíz volcano for volcanic aerosols. This study gives insights for future chemical transport modeling studies in the region and supports strategies to manage internal and external pollution sources and effects for the ~~Aburrá Valley AV~~ and the region.

1 Introduction

Long-range transport (LRT) of aerosols influences the chemical composition of air over thousands of kilometers (Kaneyasu et al., 2014; Wang et al., 2014; Wang et al., 2015; Rincón-Riveros et al., 2020) and plays a crucial role in the biogeochemical cycle of some components, such as dust and biomass aerosols, which distribute iron and phosphorus across the oceans and continents

(Okin et al., 2004; Boyd and Ellwood, 2010). Furthermore, aerosols interact with solar radiation, influencing cloud formation and light scattering (absorbance) with a cooling (warming) effect on the planet (Choobari et al., 2014). In particular, dust and black carbon can contribute to albedo reduction and accelerate melting in snow-covered zones, significantly affecting climate worldwide (Kaspari et al., 2014).

LRT of aerosols also increases human health risks, particularly in urban areas with high local emissions. Above all, fine aerosols, represented by PM_{2.5} (particulate matter with a diameter less than 2.5 μm), penetrate deeply into the human body and trigger cerebrovascular and heart diseases, lung cancer and obstruction, and respiratory infections (Xie et al., 2021; Lippmann et al., 2013). Indeed, LRT of PM_{2.5} enlarges overall concentration to more dangerous levels and can increase specific components with particular effects on human health. For instance, carbonaceous dominant particles are assumed to be more toxic than crustal components (Tuomisto et al., 2008). Increased OC (organic carbon) and EC (elemental carbon) associated with biomass burning episodes drive cardiovascular diseases (Hwang et al., 2017).

The accountability of PM_{2.5} from LRT has urged cooperation between cities and countries to identify and control pollution increases and population vulnerability. For decades, in Europe and Asia, LRT of pollutants has been recognized as a significant factor for air pollution (Kulshrestha et al., 2014), and accounted into policy plans. For instance, the Task Force on Hemispheric Transport of Air Pollution (TF HTAP) is a collaboration led by the European Union and the United States that targets inter-continental and north-hemisphere transport of PM and O₃, acknowledging their influence in meeting air quality goals (UN, 2010). The collaboration has contributed to identifying regional sources, tracking, and basing more effective emission reduction strategies (Liang et al., 2018; Zhao et al., 2021; Dong et al., 2018). On a smaller scale, Hong Kong/Guangdong Cooperation addresses transboundary air pollution in the Pearl River Delta Region (Zhong et al., 2013). The collaboration task consists of monitoring pollution level changes, evaluating the effectiveness of control measures, and providing training to stakeholders.

In Northern South America (NSA), LRT of pollutants is a more recently recognized public problem, and regional cooperation is gaining importance and momentum. Three primary kinds of LRT of aerosols have been identified in the region: biomass burning, desert dust, and volcanic emissions. Although for South America, biomass burning around the Amazon basin is one of the primary sources of aerosols (Ballesteros-González et al., 2020), atmospheric circulation patterns and substantial precipitation (i.e. aerosol wet deposition) means there is limited LTR of aerosols towards Colombia (Hamburger et al., 2013). ~~So~~ Therefore, transboundary emissions from open fires in the Orinoco basin and the Caribbean are significant drivers of intra-annual periods of hazardous air quality for Colombian cities such as Bogotá, Medellín, Arauca, Yopal, Bucaramanga and Villavicencio (~~Mendez-Espinosa et al., 2019; Henao et al., 2021; Hernandez et al., 2019; Rodríguez-Gómez et al., 2022~~) (Mendez-Espinosa et al., 2019; Rincón-Riveros et al., 2020; Henao et al., 2021; Rodríguez-Gómez et al., 2022) and on Pico Espejo in Venezuela (Hamburger et al., 2013).

Likewise, LRT of dust has also been reported in NSA, mainly produced from Sahara and Sahel deserts (Prospero et al., 2020). These primarily affect the Caribbean region, as reported in islands as Barbados, Guadalupe, Virginia, and Tobago Island, and the continental land with the Cayenne (French Guiana) (Prospero et al., 2014; Kumar et al., 2014). African dust effects have also been reported in Colombia, with the analysis of high particle loads on June 2014 and 2020 (Bolaño-Ortiz et al., 2023a; Mendez Espinosa et al., 2018; Bedoya et al., 2016). Notably, the LRT of African dust in June 2014 covered 95% of the country

(dust concentrations $> 90\text{kgkm}^{-2}$) (Mendez Espinosa et al., 2018). Few long-term assessments have also been done in the Caribbean and Andean regions (Bolaño-Ortiz et al., 2023b; Arregocés et al., 2023).

60 Volcanic activities (eruption and degassing) also play a crucial role in the LRT of pollutants in NSA and are part of the natural ecosystem in the region since it belongs to the Andean volcanic belt. Few studies have been conducted on air quality, which has mainly focussed on the Sangay volcano in Ecuador and Nevado del Ruíz in Colombia (Casallas et al., 2024; Moran-Zuloaga et al., 2023). These studies track $\text{PM}_{2.5}$ and Sulfur dioxide (SO_2) in the air and road particles from the volcanoes (Casallas et al., 2024; Cuesta-Mosquera et al., 2020; Trejos et al., 2021). The degassing activities for the Nevado del Ruíz have been
65 especially highlighted for their magnitude and frequency (Carn et al., 2016).

In Colombia, the Aburrá Valley (AV) has made substantial progress the last few years, different studies in Colombia have made substantial contributions in monitoring and identifying agents of the state of air quality in the territory, managing to report significant affectation driven by external sources (SIATA, 2021). In the territory, as on the national scale, impacts caused by open fire emissions have been intensely investigated in the last few years (Ballesteros-González et al., 2020; Mendez-Espinosa et al., 2019;
70 ~~In contrast~~ on air quality (see e.g., Hernandez et al., 2019; Mendez-Espinosa et al., 2019; Ballesteros-González et al., 2020; Rincón-Rivera et al., 2020).
Nonetheless, dust and volcanic aerosols studies have been limited to episodic high-pollution events (Mendez Espinosa et al., 2018) and do not provide comprehensive information on the sources' influence and the aerosols transport (Liu et al., 2022). Even though, for the three kinds of emissions sources, there are still significant gaps in understanding the representative influence on regional air quality throughout the year and their impact on aerosol composition, particularly for dust and volcanic
75 aerosols, which are frequently overlooked due to their uncontrolled nature (Woo et al., 2020; Pouliot et al., 2012). ~~To the AV,~~ obtaining the Obtaining the annual representation of the LRT of aerosols would contribute to the characterization of concentrations at local and national scales. It enables decision-makers to develop cooperative and effective projects to manage the risk of air pollution involving natural sources (Gómez Peláez et al., 2020; Jiao et al., 2021). Furthermore, advances in the characterization of the chemical composition of air give rise to strategies to reduce the exposure of the population to high concentrations
80 of certain species such as toxic heavy metals and carbonaceous matter (Gómez Peláez et al., 2020; Briffa et al., 2020; Allajbeu et al., 2017).

This study aims to analyze the impact of LRT of biomass burning (BB-LRT), dust (Dust-LRT), and volcanic aerosols (Volcanic-LRT) on $\text{PM}_{2.5}$ concentrations and chemical composition through year-frequent high aerosol load events in the ~~AV~~ Aburrá Valley (AV), which is one of the most populated metropolitan areas in Colombia, situated over the Andean Mountains.
85 We characterize meteorologically favorable conditions for transport to the valley and involve information from one of the largest registered $\text{PM}_{2.5}$ chemical characterization campaigns conducted in the territory (April 2019 to October 2022), as well as in-situ $\text{PM}_{2.5}$ measurements and satellite-based products. We identify representative events, analyze atmospheric transport, and highlight potential sources. Finally, we evaluate the impact of these sources on ground-level $\text{PM}_{2.5}$ concentrations and chemical composition in the AV.

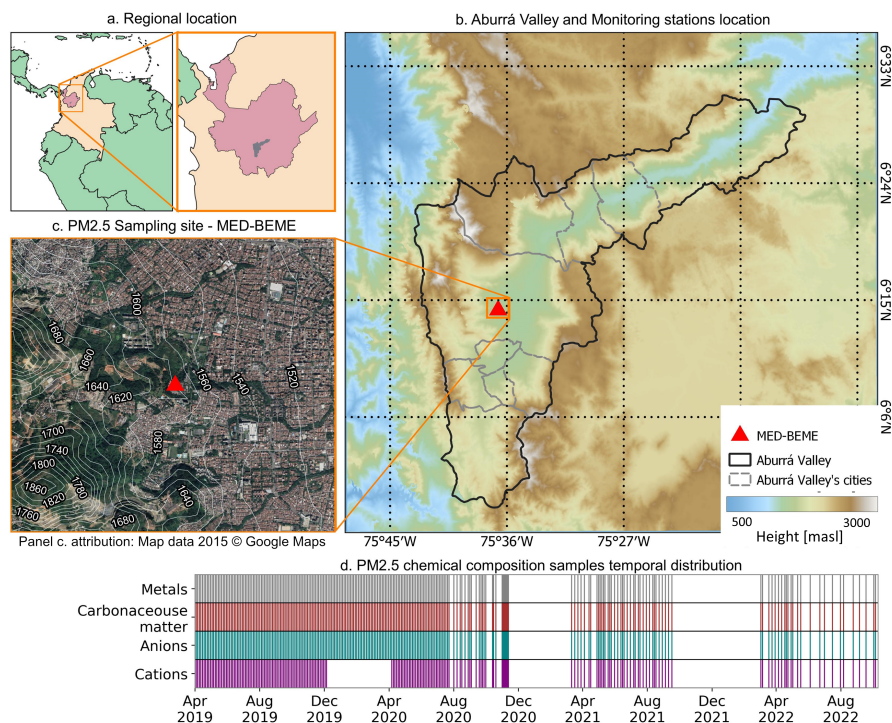


Figure 1. a. Regional location of the AV, b. Geomorphology of the AV and the distribution of the ten municipalities in black lines; c. Location of the PM_{2.5} chemical sampling campaign station (MED-BEME), represented by the red triangle on b and c., and d. Temporal coverage of the PM_{2.5} sampling campaign with white patches denoting no-sampling days.

90 2 Data and Methods

2.1 Study region

The AV, illustrated in Fig. 1b, is a 1.152 km² natural river basin located in the northeast of Colombia (see Fig. 1a). The territory contains ten cities, with Medellín as the largest city. The AV is in the central mountain range of the Colombian Andes, and its height ranges from 1300 masl (meters above sea level) in the valley to 2800 masl at the western mountaintop. By 2018, the population in the AV reached 3.73 million inhabitants, with a dense conurbation. Due to the accelerated urban expansion (Echeverri and Orsini, 2011; Salazar Hernandez et al., 2022), the increasing vehicular fleet (Corrales Espinosa et al., 2016), and the limited air pollutant dispersion, the ~~valley air quality status represented by average daily concentrations of~~ PM_{2.5}, frequently exceed ~~strict local and international air quality 24-h national and international~~ standards (WHO, 2021, 15 $\mu\text{g}/\text{m}^3$) in the valley, with more than 60% of days in the year exceeding this limit at most stations located in the urban areas of the AV (see Supplementary Table S1). Hence, the addition of external pollution sources has caused severe air quality episodes in the valley (SIATA, 2021).

The AV has two rainy periods responding to the latitudinal migration of the Intertropical Convergence Zone (ITCZ), with maximum precipitation during April and November and minimum during January and July. In the transition period, February and March are characterized by persistent atmospheric stability and a thinner atmospheric boundary layer enhancing the accumulation of air pollutants (Herrera-Mejía and Hoyos, 2019). During this period of dry conditions, fires are more likely to spread in NSA, which affects air quality in the AV and neighboring regions (Mendez-Espinosa et al., 2019; Henao et al., 2021). These conditions modulate the intra-annual variability of $PM_{2.5}$ and PM_{10} . In this period, the transport and environmental local authority (Área Metropolitana del Valle de Aburrá) implemented special control on mobile sources, limiting daily vehicle use.

2.2 Data sources

2.2.1 $PM_{2.5}$ chemical campaign

A sampling campaign was conducted at an urban background station from April 3, 2019, to October 5, 2022, to characterize chemicals on $PM_{2.5}$ filters. A total of 247 daily noon-to-noon $PM_{2.5}$ samples were analyzed and described. While April 2019 to July 2020 represented an intense sampling campaign with samples every three days, the frequency of the surface site observations became ~~more sporadic less intense~~ after July 23, 2020, ~~due to several factors. This resulted in several large data gaps in the second half of the time series (i.e. up to a maximum of 2 weeks during periods of routine sampling).~~ However, ~~the sampling in this later period was typically between 3 to 14 days. Therefore, while the temporal sampling resolution did decrease with time, we still have periods of intense sampling and measurements across the majority of the period. Thus, we still have retrieved sufficient~~ there were two extended gaps in the campaign from November 2020 to mid-March 2021 and suitable quality data for our analysis mid-September 2021 to March 2022. Despite the decrease in sampling frequency, the measurements still provide sufficient temporal coverage to get robust seasonal and annual information on aerosol concentration level and composition in this study.

The samples were taken in Belén, Medellín, on the western slope of the valley by the laboratory GYGHAM, close to MED-BEME, a station of the official air quality monitoring network (see Fig. 1c). Although some brick factories exist, the zone's economy is based on service and small business. ~~Both the~~ The residential and closer brick factory areas are around 250 m and 620 m from the station, respectively. According to Gómez-Marín et al. (2021), the primary sources of $PM_{2.5}$ in MED-BEME are the ceramic industry (22.2%), emissions from BB (21.9%), diesel (13.6%), gasoline combustion (12.8%), incineration (1.7%), and coal-fired boilers (16.3%).

The campaign considered minerals (Be, Na, Mg, Al, Si, K, Ca, Ti, V, Cr, Mn, Fe, Co, Ni, Cu, Zn, As, Se, Mo, Ag, Cd, Sb, Ba, Hg, Pb), anions (F^- , SO_4^{2-} , NO_3^- , Cl^-), cations (K^+ , Mg^{2+} , Ca^{2+} , Na^+), and carbonaceous matter species (OC1, OC2, OC3, OC4, OC5, Pyrogenic Carbon- PyC, EC1, EC2, EC3, EC4, EC5, EC6, OC, EC, C). In particular, cations were measured only in 207 of the 247 sampling days. Fig. 1d shows the temporal distribution of the samples throughout the study period. An exception for minerals is Na, Mg, K, Se, Ca, and Hg, whose measurements were finished in March 2021. The elements were analyzed from an 8"x10" quartz filter in a High Volume High-volume $PM_{2.5}$ ambient air sampler (Reference: $PM_{2.5}HV$ -Tisch Environmental) TE-6070, TISH), following Australian/New Zealand Standard 3580.9.14:2013 Method 9.14 (AS/NZS, 2013)

135 . An inductively coupled plasma mass spectrometry and thermal/optical transmission (TOT by NIOSH 5040) were used to measure minerals and carbonaceous matter. ~~and~~ Ionic chromatography (IC) was used for both anions and cations. ~~Official campaign concentrations of PM_{2.5} were measured by a Low Volume~~ The PM_{2.5} was additionally sampled by a Low-volume sampler (Reference: Wilbur TE-WILBUR - Tish). Since these measurements followed the reference method described by the CFR 40 Appendix L Part 50 reference method suggested by the US-EPA (2011) and adopted by Colombian regulations (MinAmbiente-Colombia, 2010), the measured 24-hour PM_{2.5} ambient air sampler concentrations from the Low-volume are used for the positive matrix factorization model. Further details on the techniques and the support of laboratory protocols are described in Gómez-Marín et al. (2021).

~~In addition to the~~ To complement the characterization of carbonaceous matter, ~~species measured included the~~ secondary organic carbon (SOC) ~~, which is used in this study. The was calculated using the~~ elemental carbon trace methodology (Hunter et al., 1986) ~~was used for calculating the SOC.~~ This method assumes that the organic carbon measured comes from the background, primary, and secondary sources and hence follows the relation described by Equations 1 and 2.

$$POC = OC_{back} + EC \times (OC/EC)_{pri} \quad (1)$$

$$SOC = OC_{Total} - POC \quad (2)$$

150 Where POC represents the primary organic carbon, OC_{back} is the background, OC_{total} is the consolidated OC measured in the campaign, and $(OC/EC)_{pri}$ is the primary ratio between OC and EC.

Equation 1 describes a linear relation of primary carbon matter compositions. The linear model was calculated using the 20th percentile OC and EC concentrations, aligned with Yao et al. (2020); Lin et al. (2009). The process was repeated for each hydrologic trimester (DJF, MAM, JJA, and SON) due to the influence of weather conditions in the region. The resulting slope is interpreted as $(OC/EC)_{pri}$, and the intercept is the OC_{back} . After calculating the $(OC/EC)_{pri}$ and OC_{back} , Equation 1 was used for calculating every sampling day POC using the sampled EC. Then, the SOC was obtained with Equation 2.

This study involved creating four different linear regression models (Equation 1) for the two yearly dry (June-August and December-February) and rainy (March-May and September-November) periods. The square of correlation coefficient (R^2) showed high performance of the models with values ≥ 0.95 .

160 2.2.2 PM_{2.5} concentrations

In addition to the campaign data, ~~we used we used daily for the station MED-BEME, we downloaded hourly~~ PM_{2.5} data come concentrations from the official air quality monitoring network operated by the local early-warning system – SIATA (Sistema-Sistema de Alerta Temprana de Medellín y el Valle de Aburrá, <https://siata.gov.co/>). ~~These hourly~~ From this, the PM_{2.5} concentrations ~~, from April 2019 to October 2022 in our study, from at~~ the MED-BEME station coincided with the period and site of the chemical sampling campaign (illustrated in Fig. 1b). Daily average concentrations were calculated matching the

sampling schedule (noon-to-noon). Only days with at least 75% of the hourly data were processed. Within our study period, we derived valid daily average concentrations for 96.7% of the days investigated. ~~The Pearson correlation coefficient between the~~

170 ~~We found a good agreement between the campaign (low-vol sampler) instrument and the official MED-BEME automatic station. For the study period, $PM_{2.5}$ measured by the campaign and the network was 0.84, concentrations from the automatic instrument had a minor overestimation against the reference method with a mean absolute percentage error of bias error of $-0.76 \mu g/m^3$. The corresponding mean absolute error (MAE) was 21.5%. For $PM_{2.5}$ measurements, the low-volume as a reference method provides better precision and accuracy than the MED-BEME sensor (Tasić et al., 2012), which follows equivalent methods. Despite this, the official sensor provides continuous measurements that are used in this study for more robust comparisons. Regarding temporal variability, the Pearson correlation coefficient was 0.84, highlighting good consistency~~
175 ~~between them.~~

2.2.3 CAMS Reanalysis dataset

The European Centre for Medium-Range Weather Forecasts (ECMWF) provides atmospheric composition datasets through the Copernicus Atmospheric Monitoring Service (CAMS; Inness et al., 2019). For this research, we utilized the ECMWF Atmospheric Composition Reanalysis-4 (EAC4) dataset. It is a global atmospheric chemistry model, simulating a range of
180 key tracers of composition, which uses a 4-dimensional variational data assimilation to assimilate satellite retrievals of, e.g., aerosol optical depth (AOD), carbon monoxide (CO), nitrogen dioxide (NO_2) and ozone (O_3) (Inness et al., 2019).

CAMS simulates different aerosols, which complete the total AOD at 550 nm and different gases. To identify considerable loads from biomass burning (BB), dust, and volcanic aerosols, respectively, this study considers two AOD fractions: dust AOD (Du-AOD) and organic matter (OM-AOD), in addition to the total column sulfur dioxide ($TCSO_2$). The products were
185 downloaded daily at the original 3-hourly resolution and then resampled for local noon-to-noon periods aligned with the sampling campaign schedule. Only pixel information over the sampling site was used.

2.2.4 Meteorological data

Meteorological data come from the ECMWF Reanalysis v5 (ERA5; Hersbach et al., 2020). Fields of zonal and meridional winds, temperature, and moisture were obtained at temporal and spatial resolutions of 3 hours and 0.25° (approximately 27
190 km), respectively. ERA5 data were up-sampled to a daily frequency to guarantee consistency with the chemical sampling. Due to poor performance in representing precipitation in the region by reanalysis (Posada-Marín et al., 2019), satellite-based fields were retrieved from the Global Precipitation Measurement (GPM) daily final precipitation product (Huffman et al., 2019) with a spatial resolution of 0.1° .

Further, a back trajectories dataset was built from the chemical sampling point. 8-day trajectories starting at heights of 800
195 hPa, 750 hPa, and 700 hPa were run every 3 hours with a time step of 3 hours. The model only uses wind data from the products 4h-daily pressure levels U-wind, V-wind, and omega (with a spatial resolution of approximately 2.5°) from NCEP-NCAR Reanalysis 1 (<https://psl.noaa.gov>). The model follows Equation 3. for estimating every back position.

$$X(t - \Delta t) = X(t) - V(X, t)\Delta t \quad (3)$$

Where X is the location with vertical and horizontal coordinators, V is the wind vector, and t is the time.

200 2.3 Identification and tracking of aerosol events

2.3.1 Identification of LRT events

Despite the selected CAMS products covering the whole atmospheric columns and not only the surface, CAMS has a high predictive capability for $PM_{2.5}$ in the AV (Pérez-Carrasquilla et al., 2023). Furthermore, its products relatively reasonably reproduce $PM_{2.5}$ tendencies and extreme events in the territory (Casallas et al., 2022). Therefore, the magnitude of CAMS
205 products OM-AOD, Du-AOD, and TCSO₂ in the AV were utilized to identify possible BB-LRT, Dust-LRT, and Volcanic-LRT events, respectively.

To help identify $PM_{2.5}$ days that were subject to long-range transport, the time series of the CAMS data sets were standardized. Here, for each day, the time-series average (mean) was subtracted from the daily value and then normalized by the time-series standard deviation (i.e., 1σ). This then allowed for the identification of LRT events using a range of subjective thresholds and the quantification of the corresponding $PM_{2.5}$ concentrations and composition. The thresholds investigated ranged from 0.8 to 2.0 σ , but in this study, we present the results for thresholds of 0.8, 1.5, and 2.0 σ . ~~To help accurately identify A 7-day rolling window was used to accurately identify prolonged and intense periods of LRT events and remove short-term sporadic events, a 7-day running window was used. Here, if less than four days with values greater than the specified threshold were detected, then they were classed as outliers and removed (i.e., we are focussing on LRT events, which we define as~~
215 ~~lasting more than half a week).~~ Within this window, at least 4 days had to have values above the respective thresholds to be classified as a LRT event. We subjectively chose 4 days of elevated values due to the sampling frequency of the campaign. Here, campaign temporal sampling was ≥ 3 days, so these criteria were required to get representative samples of the aerosol characterization for the PMF analysis.

Observe that continuous days marked as peaks are possible for longer-lasting events. Finally, for each peak event (DtE), the
220 surrounding days, relative to the peak event centered in the 7-day window (i.e., DtE₀), were labeled. For instance, the LRT events ranged from three days before (DtE₋₃) to three days after (DtE₃) the peak event. The days outside of three-day windows around a DtE₀ are generally identified as no-event days, and no significant/lasting event affectation is supposed. Although the process described above was run for every event and threshold, further assessments are based on 0.8 σ for BB-LRT and 1.5 σ for Dust-LRT and Volcanic-LRT, as is described in Section 3.1.

225 2.3.2 Regional meteorological analysis

A regional analysis of meteorological fields was performed during pollution LRT events to identify conditions that favor aerosol transport from the sources of interest. We computed anomalies for each pixel of the meteorological fields by subtracting the

average of the corresponding month from the original values in the study period. Subsequently, composites during days with events (DtE₋₃ to DtE₃) were performed with meteorological fields and anomalies for each source. This approach helps reveal particular conditions during days with aerosol LRT events. The results are also supported by analyzing air masses arriving at the AV using back-trajectories during the events. The number of times trajectories pass through a pixel was counted to estimate a percentage indicating the influence area.

2.3.3 Influence on local PM_{2.5} concentrations

The local measurements of PM_{2.5} were used to assess the influence of CAMS-based identified events on air quality in the AV. The daily PM_{2.5} datasets were labeled with the days to the closest event peak (DtE) to compare concentrations before and after the events. The days before the event constitute from 15 to 4 days before the event peak (DtE₋₁₅ to DtE₋₄), while the days after the event peak consisted of 4 to 15 days after the event peak (DtE₄ to DtE₁₅). These ranges were determined to have enough data from the chemical campaign on similar weather conditions for comparison between the days affected by the LRT of aerosols and the days before and after. We used the Mann-Whitney U test to compare these periods. The null hypothesis is denied with the confidence of 90% (p-value ≤ 0.1) and 95% (p-value ≤ 0.05).

The COVID-19 lockdown period (April 1st to June 1st, 2020) was filtered from our record to avoid perturbation out of standard emission patterns in the region.

2.3.4 PMF and PM_{2.5} chemical composition

The Positive Matrix Factorization (PMF) is a receptor model developed by the EPA (Environmental Protection Agency) to analyze water and air samples. The model expresses observations of PM_{2.5} as a sum of the contributions from a number of source profiles (non-time dependent). In this case, we are using the PMF model and our campaign measurement data of PM_{2.5} chemical composition and PM_{2.5} concentrations to identify the dominant sources (e.g., coal combustion, vehicular emissions, secondary pollution) during different LRT classed events (i.e., BB, dust or volcanic LRT events). For the AV, 5 to 7 factors (i.e., sources) have been suggested (see Gómez-Marín et al. (2021)).

The PMF is mathematically expressed as (Paatero and Tapper, 1994):

$$X_{ij} = \sum_{k=1}^N g_{ik} f_{kj} + e_{ij} \quad (4)$$

The objective of PMF is to determine the values of g_{ik} , f_{kj} , and N (i.e. the number of factors) that best reproduce X_{ij} . Here, X represents the measurement data for sample i (daily temporal resolution) and chemical component j . An inversion approach is used for this matrix problem, exploiting an iterative scheme to converge on the solutions for g_{ik} and f_{kj} . In the factorization problem, f_{kj} contains the concentration of each chemical component (j) in the unit profile for the factor k (i.e. PM_{2.5} source). The matrix $g_{i,k}$, the contribution factor, defines how much of the profile is counted in the total concentration in day i . $e_{i,j}$ contains the residual for each compound/sampling day. Outputs from the PMF include statistical metrics that help

to evaluate the model performance, such as the correlation coefficient (R^2), the recuperated mass (% RM), and the objective function value (Q). The PMF adjusts g_{ik} and f_{kj} to minimize the function Q in Equation 5 (Paatero, 1997).

$$260 \quad Q = \sum_{i=1}^n \sum_{j=1}^m \left[\frac{X_{ij} - \sum_{k=1}^N g_{ik} f_{kj}}{\mu_{ij}} \right]^2 \quad (5)$$

Where μ_{ij} contains the uncertainty compound/sampling day.

The model uses two input uncertainties. The first is the species uncertainty (μ_{ij}), which does not change in the process and is according to the method unique for every species sample. In this study, μ_{ij} considers the measurement and analytical errors, comprising uncertainties of the volume of air processed, filter area, and species mass, in addition to the uncertainty of the
 265 detection limit of the measurer method. The last one represents the analytical uncertainty and was calculated following Noris and Duvall (2014) by using the methods detection limit suggested by Eugene Kim and Qin (2005). However, for PM_{2.5}, we followed the approach of Eugene Kim and Edgerton (2003) who set μ_{ij} (where j represents PM_{2.5} for day i) greater than the daily PM_{2.5} concentration (i.e., three times the concentration in this study). The second uncertainty in μ_{ij} is recognized as the "Extra modeling uncertainty". It is added to the model as a percentage to cover other inherent errors, such as variations of
 270 source profiles and chemical transformations in the atmosphere. A range from 10% to 16% was tested for the models of every kind of LRT event; this aligns with other studies (Callén et al., 2009; Shin et al., 2022; Salim et al., 2019).

During the modeling process, species can be classified as "strong" "weak," or "poor" based on individual statistics. The "weak" category triples the uncertainty μ_{ij} , and the "bad" excludes the species from the model. Two statistics are key for categorizing the species. First, the signal-to-noise ratio (S/N) represents overall uncertainty per concentration. $S/N < 0.5$ is
 275 generally classified as "bad" because the uncertainty surpasses two times the concentrations (Noris and Duvall, 2014). Second, the regression diagnostic where R^2 informs about the linear relation of every species' observed and modeled concentration.

The model convergence evaluation statistics considers this classification with the $Q/Q_{expected}$ ratio. $Q_{expected}$ is the theoretical value of Q (Equation 5), expressed as Equation 6.

$$Q_{expected} = (n \times m_s) - ((N \times n) + (m_s \times N)) \quad (6)$$

280 where n is the number of sampling days, and m_s is the number of "strong" species ~~use-used~~ for modeling (Noris and Duvall, 2014).

For the PMF we anticipate small datasets since the target events may not significantly impact the entire PM_{2.5} chemical campaign. Nonetheless, multiple studies with PMF samples ranging from 14 to 30 have reported useful and meaningful results (Yu et al., 2015; Haghazari et al., 2022; Via et al., 2022). As the sample dataset decreases, rotational ambiguity caused
 285 by infinite valid solutions strongly affects the results and increases overall uncertainty (Manousakas et al., 2017). To mitigate the error, the software EPA PMF v 5.0 allows for estimating the effect of random errors and rotational ambiguity in the dataset using bootstrapping (BS) and Displacement (DISP) tools. While BS evaluates random errors by performing 100 runs with randomly relocated blocks of observation of the original dataset, DISP focuses on indicating rotational ambiguity by adjusting

Table 1. Statistics of OM-AOD, Du-AOD, and TCSO₂ representing BB-LRT, Dust-LRT, and Volcanic-LRT

Threshold	Events stats.	OM-AOD	Du-AOD	TCSO ₂
0.8 σ	Samples	31	31	59
1.5 σ	Samples	18	19	32
2.0 σ	Samples	10	4	21
0.8 σ	Limit	0.22	0.01	0.70mg/m ²
1.5 σ	Limit	0.28	0.02	0.87mg/m ²
2.0 σ	Limit	0.32	0.02	0.99mg/m ²
0.8 σ	Average	0.25	0.02	0.78mg/m ²
1.5 σ	Average	0.29	0.02	0.88mg/m ²
2.0 σ	Average	0.32	0.03	0.97mg/m ²

"Samples" represent the number of sampling days available for the PMF characterization.

Each threshold limit defines the calculated absolute magnitude. Note, AOD is a dimensionless quantity.

290 up and down all values in the factors profile restricted by 4 allowed changes in the calculated Q (dQmax) and monitoring major factors swaps (Noris and Duvall, 2014).

295 Additionally, constraining the base run can improve the solution when data is limited by reducing the rotational space (Dai et al., 2020). The PMF software has the functions to "pull down maximally," "pull up maximally," "set to zero," and manually set the profile concentrations. While the first two options are soft constraints, the third and fourth are hard constraints and require a high level of confidence in the magnitude of the profile contributions. For this study, only soft contains are contemplated. The constraint increases in the final Q value, which should be less than 5% (i.e. %dQ < 5%), the recommended maximum change (Noris and Duvall, 2014). For this study, the %dQ was set by default <0.5%.

300 Due to missing data, all species except cations are included in the model. Only total OC and EC from carbonaceous matter are included for Duts-LRT and Volcanic-LRT events since comprising all carbon species could particularly weigh those and overshadow trace elements for the sources. For those LRT events, the potential run period is reduced until April 2021 since the tracer minerals Na, K, and Mg were not measured ~~afterward~~afterwards.

The factor interpretation of this study was primarily supported by the source characterization made by Gómez-Marín et al. (2021) for the studied station.

A final element comparison is made between the days of events with a positive relative contribution ($g_{i,k} > 0$) and the days categorized as before (DtE₋₁₅ to DtE₋₄) and after events (DtE₄ to DtE₁₅).

305 3 Results

This section is divided into four subsections. The first subsection shows possible aerosol events from BB-LRT, Dust-LRT, and Volcanic-LRT. The second is focused on the temporal and regional characterization of the events, including a description of

meteorological patterns and possible sources of aerosols. Finally, the third and fourth subsections locally assess the identified events using ground-level $PM_{2.5}$ concentration and chemical compositions, respectively.

310 3.1 Identification of events

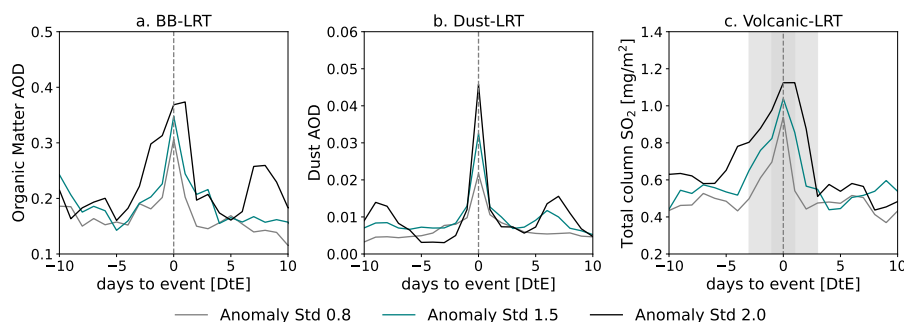


Figure 2. Average magnitudes of OM-AOD (a.), Du-AOD (b.), and $TCSO_2$ (c.) concentration around the events. Different colors represent different anomalies. The shadowed region delimits the event. Du-AOD is on a scale that is ten times lower than OM-AOD.

The products OM-AOD, Du-AOD, and $TCSO_2$ from the CAMS reanalysis were used to identify aerosol from BB-LRT, Dust-LRT, and Volcanic-LRT, respectively. Tab. 1 shows different statistics about the identification for each kind of event using the threshold of 0.8σ , 1.5σ , and 2.0σ . The intensest events reached maximum magnitudes of 0.66 for OM-AOD (March 30, 2020), 0.11 for Du-AOD (June 24, 2020), and $1.96 \text{ mg}/\text{m}^2$ for $TCSO_2$ (September 10, 2022). These specific days exceeded the average values for identified events more than twice for OM-AOD and $TCSO_2$ and more than five times for Du-AOD. Notably, OM-AOD reached 77% of the total AOD in the intensest event. Regarding the variability, all identified events showed high variability with a standard deviation around 42%, 90%, and 35% of the average for OM-AOD, Du-AOD, and $TCSO_2$, respectively.

Fig. 2 shows the average behavior of each variable centered around the event peak for the threshold of 0.8σ , 1.5σ , and 2.0σ . The events exhibit a clear peak centered on DtE_0 for most of the considered threshold. However, events identified with 2.0σ exhibit noisy behavior around the peak for BB-LT and volcanic-LRT. In particular, OM-AOD exhibits a second peak after the event peak. This could be caused by either the smaller sampling size or high aerosol loads that do not meet the duration criteria for the threshold. Besides, the strictest thresholds (2.0σ) result in a considerable reduction of studied events (see Tab. 1), which is critical for the assessment of $PM_{2.5}$ chemical composition.

Based on the study's objectives, only one identification threshold was selected for each event, allowing the investigation of more accurate identification indexes. Based on the results and literature review about the expected frequency of each type of event for the city (Mendez-Espinosa et al., 2019; SIATA, 2021), we selected events that exceed 0.8σ for BB-LRT and 1.5σ for Dust-LRT and Volcanic-LRT. Anomalies over 2.0σ significantly limited the number of days for the analysis, whereas 0.8σ probably oversamples volcanic and dust aerosol load events, whose affectation is recorded as infrequent. Moreover, the

330 selection ensures yearly representativity for each event (see supplementary Figure S2). BB-LRT, Dust-LRT, and Volcanic-LRT were represented during the complete study period, at least for three years each. **Notably, in 2020, these events covered 19.4%, 6.6%** These events cover 13%, 8%, and 12.013% of the days/year, respectively. With this selection, the sampling campaign can represent BB-LRT, Dust-LRT, and Volcanic-LRT events in the PMF with 31, 19, and 32 samples.

3.2 Regional analysis of events

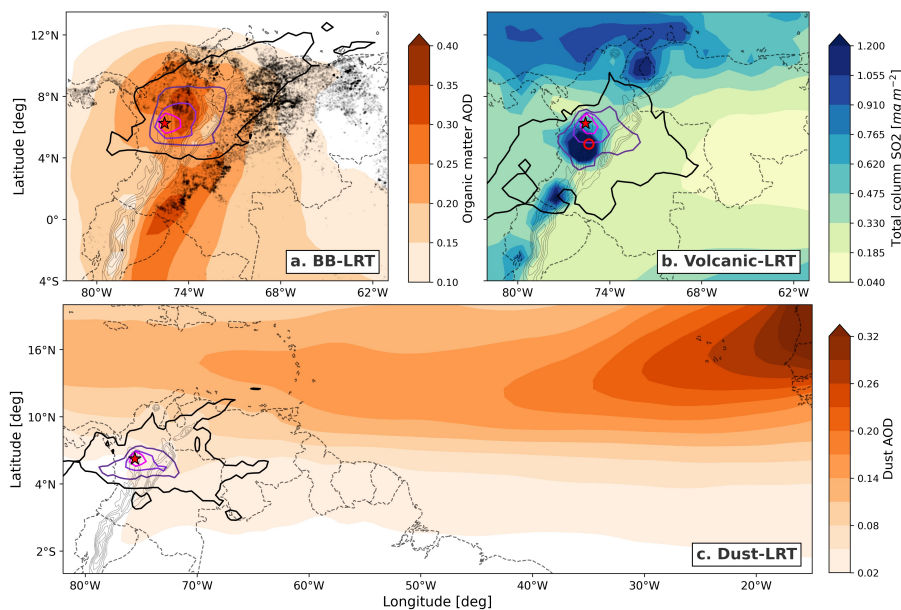


Figure 3. Mean spatial distribution of OM-AOD (a.), TCSO₂ (b.), and Du-AOD (c.) for days with each type of LRT event. Pink-to-black contours enclose the regions from which 20, 10, 5, and 1% of air masses arrive at the AV according to the back-trajectories. Gray dots in a. show MODIS-retrieved hotspots associated with fires with a 70% of confidence. The red star marks the location of the AV, while the red circle in the upper right panel is the location of the Nevado del Ruiz Volcano. The black contours show the terrain elevation from 1500 to 5000 masl.

335 Regional analysis was conducted to establish the meteorological conditions that may favor pollution LRT events and identify potential aerosol sources. The average spatial distribution of the identified AOD-OM, Du-AOD, and the TCSO₂ during the aerosol LRT events is shown in Fig. 3, with the integrated back trajectories of air masses arriving at the AV. OM-AOD values above 0.2 occur over most of Colombia during BB aerosol events (Fig. 3 a), consistent with a high number of fire hotspots in northern South America (Venezuela and Colombia). In particular, the back trajectories during these days suggest that a
 340 large part of the air masses arriving at the AV (10 to 20 %) come from the northeast, where OM-AOD exhibits its maximums (exceeding magnitudes of 0.35).

Volcanic-LRT (Fig. 3 b) shows high SO₂ values (over 1.2 mg m⁻²) in the atmospheric column above the Nevado del Ruiz volcano (red hollow circle). Additionally, another two spots are evident: the first one, toward southwestern Colombia, could be

related to another degassing volcano called the Galeras, whereas the other one is on Maracaibo Lake, with relatively high values possibly associated with oil extraction (see Fioletov et al. (2016)). According to the back-trajectories, during Volcanic-LRT events, around 10% of the air masses arriving in the AV come from the southeast, where the Nevado del Ruiz hotspot exists. A smaller fraction of air masses (1%) arrive from the hotspot in the country's southwest. Similarly, a clear dust propagation pattern is evident from the Sahara desert towards the Caribbean and, to a lesser extent, NSA (see Fig. 3 c). While air masses for BB-LRT and Volcanic-LRT have different directions, most air masses that reach the AV during Dust-LRT events come from the east, showing consistency in the analysis.

Despite the short study period, the identified events suggest a marked annual cycle (Fig. 4). During March, 50.0% of studied days represent the BB-LRT. April and February also showed high proportions of this event, with 43.3% and 24.8%. Concerning the dust events, two maximums are observed, April and July, with a proportion of 23.3% and 23.4%. The distribution for volcanic-LRT ranges from June to September, with a clear peak in August with a frequency of 49.2%. Additionally, as shown in Fig. 4, external events overlap in some cases. The higher intersections observed were for BB-LRT and Dust-LRT in April and for BB-LRT and volcanic-LRT in September. Here, where days were defined as both, e.g., BB-LRT and Dust-LRT, they were added to the joint classification (i.e., BB-LRT & Dust-LRT) and were not included in the respective singular classifications. Overall, the annual cycle of these frequencies and the local $PM_{2.5}$ concentration show the event's possible influence on $PM_{2.5}$ is non-linear.

Although LRT events ~~exhibited a marked season, some non-event days in the different months occur~~ display a marked seasonality, a significant percentage of days in each month have a negligible impact from LRT events (see Fig. 4), suggesting that intraseasonal variations are also relevant in explaining the occurrence of these events. To elucidate specific meteorological patterns during events days, Figs. 5 and 6 present composites of meteorological variables for the low (800 to 700 hPa) and mid (600 to 400 hPa) troposphere for the different kinds of events. The Andes mountain range generally causes a large spatial variability in the low-level wind field. The winds blowing from the east of the Andes cross the mountain range through two zones with relatively low altitudes (hereafter mountain passes), which are demarcated with white boxes (see Figs. 5 a, b, c).

Anomalous dry winds of around 1 m/s originate from the north during BB-LRT events (Fig. 5 d), connecting the AV with the high OM-AOD region. Additionally, winds blowing from Venezuela, where multiple BB hotspots were identified, reach northern Colombia through the mountain pass located north of the Andes, at the border between Colombia and Venezuela (see Fig. 5 a). Low-level winds during LRT events are in agreement with the back trajectories of Fig. 3 a. A reduction of 2 to 3 m/s in the intensity of mid-level winds (between 5°N and 8°N; see Figure 6 a, d) was also observed along with less rainy conditions in northern Colombia.

For Dust-LRT events, low-level winds flow across the two mountain passes in the Andes (Fig. 5 b) with increased easterly winds north of Colombia and northeasterly winds east of the Andes (Fig. 5 e), representing a dry flow from the Atlantic Ocean where Du-AOD is higher. Figs. 6 b and e, show that mid-level winds have notable eastward direction with anomalous winds coming from the Caribbean and less rainfall in NSA.

Finally, days with volcanic-LRT events are characterized by southeasterly winds blowing toward the AV (see Fig. 5 c) from the Nevado del Ruiz volcano region. These winds may be favored by anomalous southeasterly winds of more than 3 m/s

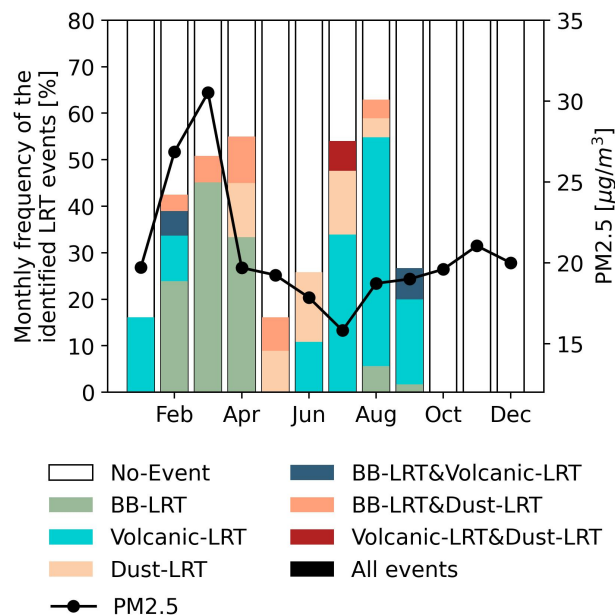


Figure 4. Monthly frequency of CAMS-based identified days with BB-LRT (green bars), Dust-LRT (wheat color bars), and volcanic-LRT (blue bars) events as identified from the CAMS reanalysis. Overlapped events are depicted in dark blue (BB and Volcanic), orange (BB and Dust), and red (Volcanic and Dust) bars, while since different LRT events can happen at the same time. White bars represent the period-frequency of no-identified days without LRT events, while the black line shows the monthly average PM_{2.5} concentration ($\mu\text{g}/\text{m}^3$) for the MED-BEME station.

crossing the southern mountain pass, as shown in Fig. 5 f, along with drier air reaching the AV. Less rainy conditions are also present in the Colombian Andes (Fig. 6 f), particularly between the AV and the Nevado del Ruíz volcano.

Noteworthy, the three types of events are accompanied by less rainy and drier atmospheric conditions in northern Colombia, Venezuela, and the Andes, respectively, with rainfall anomalies greater than -4 mm/d and specific humidity anomalies up to -1 gkg^{-1} . These results allow us to highlight the importance of precipitation to enhance or even trigger the aerosol LRT events to the AV.

3.3 PM_{2.5} concentrations change

After identifying possible LRT events and the meteorological conditions favoring them, a local analysis was performed to connect and validate the different events' influence on AV's local air quality. The daily concentration of PM_{2.5} reasonably responds to the BB-LRT identified by OM-AOD. Fig. 7 compares the PM_{2.5} concentrations before (DtE₋₁₅ to DtE₋₄), during (DtE₋₃ to DtE₃) and after (DtE₄ to DtE₁₅) each event. BB-LRT had the highest PM_{2.5} increments, especially at DtE₀. 86.7% of values during the peak day exceed the average PM_{2.5} for the study period (20.25 $\mu\text{g}/\text{m}^3$). Besides, over half of PM_{2.5} concentration during DtE₀ exceeds twice the WHO (2021) guidelines for daily concentrations (15 $\mu\text{g}/\text{m}^3$). On DtE₀, the

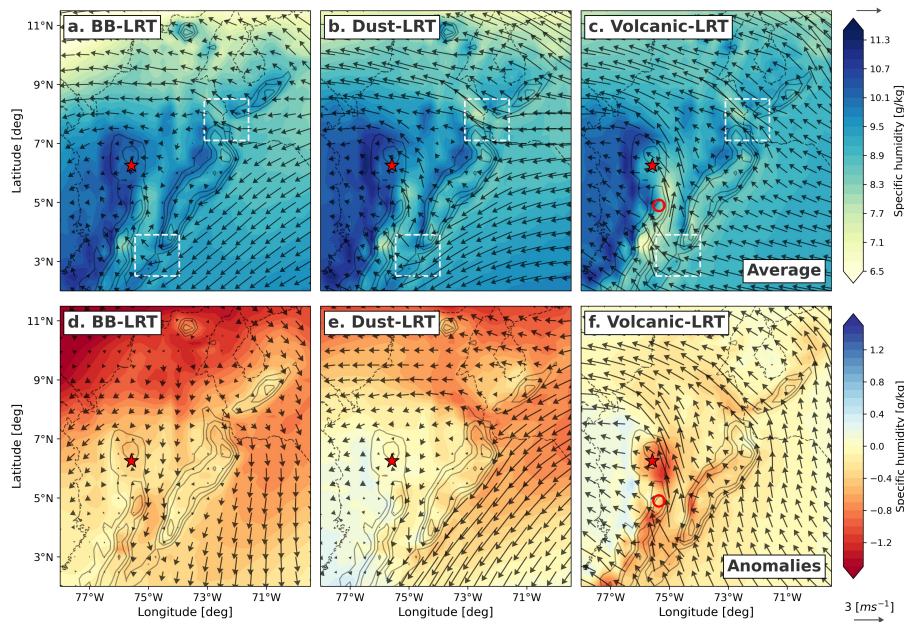


Figure 5. Meteorological composites for low-level (800 to 700 hPa) winds and specific humidity during days with (a.) BB-LRT, (b.) Dust-LRT, and (c.) volcanic-LRT events. d, e, f: as upper panels but for anomalies. A red star marks the Aburrá Valley, and the red circle in the upper right panel is the Nevado del Ruíz volcano. The white rectangles are two mountain passes in the Colombian Andes. The black contours show the terrain elevation from 1500 to 5000 masl.

PM_{2.5} average concentrations ($31.48 \mu\text{g}/\text{m}^3$) were significantly higher than the surrounding days ($p\text{-value} \leq 0.05$), according to the Mann-Whitney test. Similarly, the event days before the peak (Dte₋₃ to Dte₋₁) had significantly higher concentration compared to days before and after the identified BB-LRT events ($p\text{-value} \leq 0.05$). In contrast, concentrations following the event peak are similar to the average magnitude, indicating a sudden decrease.

As for BB-LRT, the peak of the events for Dust-LRT showed higher concentrations than days before the event ($p\text{-value} \leq 0.1$). On the other hand, the increase of PM_{2.5} due to Volcanic-LRT was consistent ~~throughout the event~~ at the beginning of the events, not just during the peak. After the events, the PM_{2.5} concentration for Dust-LRT remains at similar levels. However, outliers are observed ~~after the events~~, particularly in March 2020, a period strongly affected by BB-LRT. On the other hand, the concentration of PM_{2.5} right after the peak of Volcanic-LRT ~~significantly decreases. Although, forward-presented significantly lower concentrations (p-value < 0.1), contrasting with the subsequent~~ days (Dte₈ to Dte₁₅) ~~present an average magnitude similar to during the events~~. Overall, it is clear that the BB-LRT event has a higher impact on PM_{2.5} concentration to the AV than Dust-LRT and Volcanic-LRT.

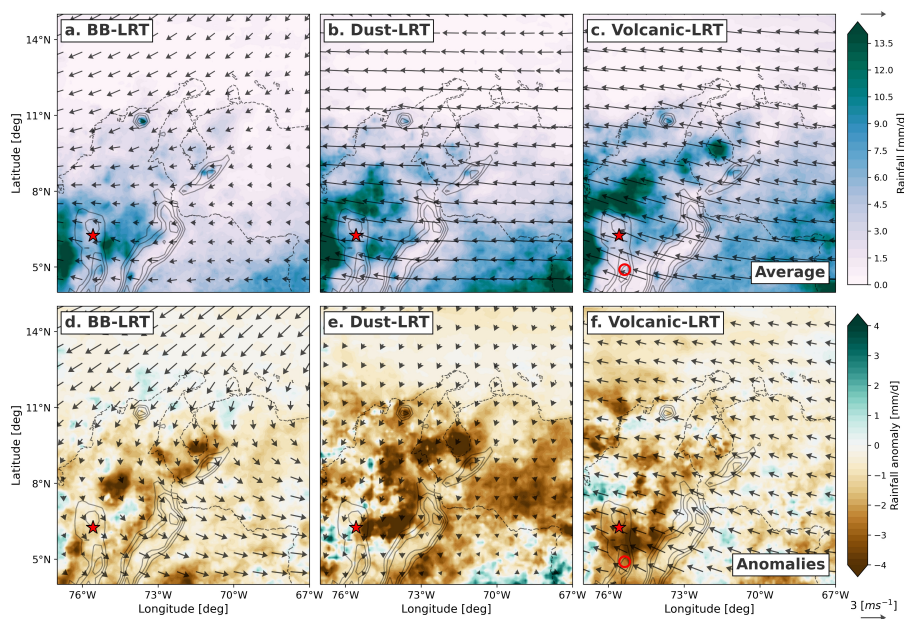


Figure 6. Meteorological composites for mid-level winds (600 to 400 hPa) and rainfall during days with (a) BB-LRT, (b) Dust-LRT, and (c) Volcanic-LRT events. d, e, f: as upper panels but for anomalies. The red star marks the Aburrá Valley, and the red circle in the upper right panel is the Nevado del Ruíz volcano. The black contours show the terrain elevation from 1500 to 5000 masl.

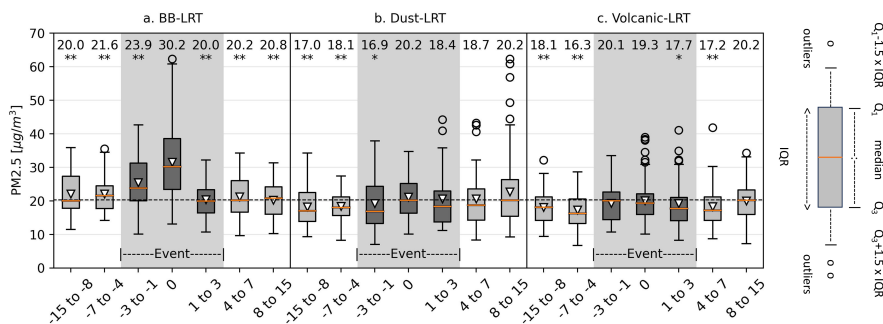


Figure 7. PM_{2.5} concentrations before, during and after (a.) BB, (b.) dust and (c.) volcanic aerosols events. The chart's top presents the median concentration of each dataset, along with the significance level compared with the peak of the data, ≤ 0.1 (*) or ≤ 0.05 (**). The shadowed darker boxes represent the events. The down triangles represent the average concentration for each day range, and the grey dashed line represents the total study period average. The boxes are limited by the first and third quartile, Q_1 and Q_2

3.4 PMF and chemical composition change

405 The PMF models of the BB-LRT and Volcanic-LRT presented low-residual-responses-to-no-outlier-values-For-a-good-fit,
and-all-residuals-are-normally-distributed;however,for the Dust-LRT, three-days-were-marked-as-outliers-and-removedsome

Table 2. Statistics performance of PMF for models during days affected by BB-LRT, Dust-LRT, and Volcanic-LRT

Properties	BB-LRT	Dust-LRT	Volcanic-LRT
<u>Model statistics</u>			
Samples	31	16	32
N. factors	6	6	6
Non-weak species	32	27	27
Bad species	OC5,EC6	Be	—
$Q/Q_{expected}$	0.99	1.00	0.93
Add. uncertainty[%]	16	11	12
<u>BS mapped factor [%]</u>	<u>82</u>	<u>82</u>	<u>88</u>
<u>DISP swaps</u>	<u>0</u>	<u>0</u>	<u>0</u>
<u>Results-Model statistics for PM_{2.5}</u>			
PM _{2.5} error [$\mu\text{g}/\text{m}^3$]	<u>2.50</u>	<u>1.84</u>	<u>2.60</u>
<u>R²</u>	<u>0.92</u>	<u>0.90</u>	<u>0.73</u>
<u>RM [%]</u>	<u>99.31</u>	<u>98.24</u>	<u>97.50</u>
<u>LRT factor statistics for PM_{2.5}</u>			
<u>PM_{2.5} [$\mu\text{g}/\text{m}^3$]</u>	11.14	6.77	6.46
PM _{2.5} [%]	37.61	33.93	30.85
PM _{2.5} error-BS 25th [$\mu\text{g}/\text{m}^3$]	<u>2.50</u> <u>9.17</u>	<u>1.84</u> <u>4.00</u>	<u>2.60</u> <u>5.66</u>
<u>R²PM_{2.5} BS 50th [$\mu\text{g}/\text{m}^3$]</u>	<u>0.92</u> <u>10.53</u>	<u>0.90</u> <u>5.63</u>	<u>0.77</u> <u>6.53</u>
<u>RM-PM_{2.5} BS 75th [%$\mu\text{g}/\text{m}^3$]</u>	<u>99.31</u> <u>11.48</u>	<u>98.24</u> <u>6.95</u>	<u>97.50</u> <u>8.47</u>

components residuals exceed the recommended range of -3 to 3 (Noris and Duvall, 2014), so three samples were excluded due to outlier data. In general, the three final selected PMF models showed a good performance, meeting acceptance criteria with a reconstructed mass (RM) in the range of 80 to 120 and an R² coefficient greater than 0.8 (Noris and Duvall, 2014). The $Q/Q_{expected}$ results show a good convergence, where the $Q/Q_{expected}$ ratio for the selected base models were close to 1, a determinant parameter for the number of factors and the component categorization. A summary of the models' statistics is given in Table 2.

In evaluating errors of the PMF, none of the three simulated scenarios showed significant rotational ambiguity, nor were substantial random errors in the dataset after running the DISP and BS methods. Results for the DISP method showed no factor swaps for all dQmax values. In the BS analysis, outputs were considered stable, yet not all base factors were mapped to the boot factors. On average, the percentage of factors correctly mapped was 84%, which is in line with (Noris and Duvall, 2014), where a minimum of 80% mapped factors are suggested for interpretability and to support the number of factors selected. The target BB-LRT, Dust-LRT, and Volcanic-LRT profiles for the models were mapped in 88%, 80%, and 98% of the runs, respectively. The 25th, 50th and 75th percentile of the PM_{2.5} contribution rates calculated for the target profiles in the BS

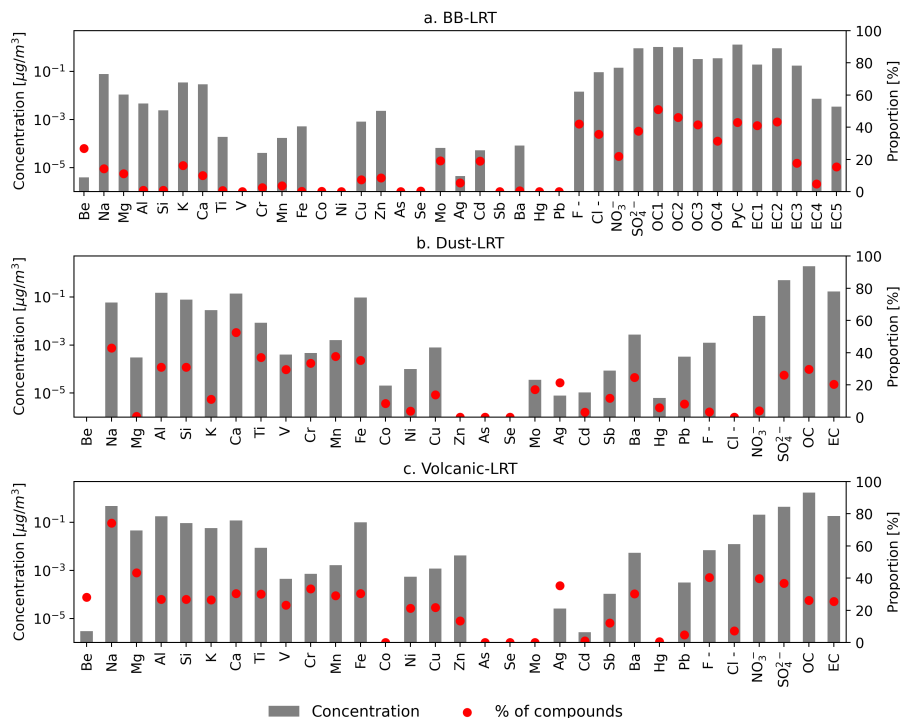


Figure 8. PMF output profiles for (a.) BB-LRT, (b.) Dust-LRT, and (c.) volcanic-LRT events. The bars represent the average concentration of the compounds, while the red points denote the compound's average percentage contribution to the whole element concentrations. Refer to Supplementary Figure S3 to see the contribution of all factors.

420 runs are presented in Table 2. The contribution rate variability represented by the BS runs is particularly important for small database models, where contribution rate results might be unstable compared to the more stable profile (Feng et al., 2023).

Final base models were constrained to improve the correspondence between the chemical profiles found by the PMF and the profiles expected based on the identified emission sources. Specific constraints were defined in the different modeling scenarios to refine the factor profiles. The model's factors are presented in Figure S3. For the BB-LRT model, three soft constraints were applied to the "coal boiler" factor to pull up the EC1, Se, and As concentrations. Conversely, the Ag, Se, and EC1 concentrations were pulled down in Factors "ceramic industry", "gasoline" and "Diesel". For the Dust-LRT model, four soft constraints were established: for "vehicular emissions", the concentrations of EC and Ni were maximally adjusted downward and upward, respectively; for "resuspended material," Ni was reduced to improve the fit; and for "biomass burning," OC was maximally increased. In the volcanic-LRT event scenario, two soft constraints were implemented. For volcanic-LRT, we pulled down the concentration of Cu and Mg, while Se was pulled up for "secondaries" and Cu was pulled down for "vehicular". For the models, the %dQ (i.e. the Q change because of the constraint) is < 1%.

The results from the PMF for BB-LRT ascribed concentration to six emissions sources (or factors): BB (37.6%), coal boiler (20.1%), ceramic industry (16.7%), gasoline (12.8%), Diesel (11.4%) and Incineration (1.4%). The factor of contribution

for BB-LRT in the sampling days varies from 0.0 to 4.5 (maximum identified on 25 March 2020), which means that not all
435 identified event days are backed by the PMF (The factor of contribution for the models ran are in Supplementary Figure S4).
Of the 31 samples identified, 27 days (equivalent to 87.0%) have positive contributions from BB abscribed. The PMF profile of
the BB factor, represented in Fig. 8a, is identified by the dominant contribution ($\geq 30\%$) of OC species, PyC and some anions.
The identified contribution of this factor in OC, but especially in OC1 (50.9%) and OC2 (46.1%), agrees with observations
made by Chow et al. (2004) for vegetation burning. Similarly, the high contribution of CyP (43.0%) back the profile since
440 BB produces $\sim 50\%$ of global CyP emissions (Santín et al., 2016). Regarding anions, the contribution to F^- (42.1%) head
contribution and support the identification and hypothesis of LRT as a trace element with high lifetime (Jayarathne et al.,
2014). The contribution to SO_4^{2-} (37.7%) and NO_3^- (22.0%) evidence secondary pollutants formation from the BB emissions
during daytime and nighttime, respectively (Rastogi et al., 2014). Moreover, the influence over Cl^- concentration (35.7%) may
evidence semiarid vegetation burning (Andreae et al., 1998). Other species as K (16.2%) traces BB emissions (Yu et al., 2018;
445 Rastogi et al., 2014).

For the Dust-LRT event days, the model identified the dust contribution (33.9%) and others from BB (27.8%), vehicular
(12.4%), resuspended material (6.5%), incineration (7.7%), and ceramic (11.7%). Due to only total OC and EC being con-
sidered, splitting the vehicular emissions between diesel and gasoline is difficult. The contribution factor for dust varies from
0.0 to 4.1 (maximum identified on 27 February 2020). For this event, 81.2% (13/16 samples) of the identified days exhibited
450 a positive contribution. The dust profile identification is supported by a high contribution ($>31.0\%$) of Al, Ca, Fe, and Si (\geq
 $0.1 \mu g/m^3$), as Fig. 8b illustrates. Similarly, Malaguti et al. (2015) highlights Fe, Al, Ca, Ti, and Mg in the fine Sahara's
dust aerosols. Ca and Ti are specially weighted for LRT from the Sahara desert since the other elements are more generic
crustal aerosol tracers (Martinez-Verduzco et al., 2023; Nicolás et al., 2008). The dust contributes to 52.6% and 37.0% of Ca
and Ti concentration, respectively. The profile also diverges from the general crustal material in the contribution of secondary
455 inorganic aerosol as SO_4^{2-} probably from $(NH_4)_2SO_4$ (Varrica et al., 2019; Malaguti et al., 2015), with 33.9% of average
concentration in the samples. Likewise, a particular mix between dust minerals and OC (29.7%) might also be a characteristic
to identify the dust profile (Aymoz et al., 2004; Malaguti et al., 2015).

For the days of Volcanic-LRT events, the model identified 30.8% of volcanic ~~aerosols~~ aerosol contribution. Other sources
were BB (29.1%), vehicular emissions (15.7%), resuspended dust and ceramic industry (11.0%), secondary aerosol formation
460 (9.1%), and incineration (4.3%). The volcanic factor presented contributions varying from 0.0 to 3.9 (maximum identified
on 6 February 2020). According to the model, the volcanic factor influences only 78.1% (25/32 samples) of identified days.
As expected Satsumabayashi et al. (2004), the volcanoes profile in Fig. 8c, shows a high contribution of SO_4^{2-} (36.9%). The
contribution but not the concentration of SO_4^{2-} was surpassed by F^- and NO_3^- . F^- from volcanic emissions frequently
concentrated onto the ash surface as CaF_2 , AlF_3 or Na_2SiF_6 (Bia et al., 2020; Delmelle et al., 2021); while NO_3^- may relate
465 with the oxidized trace species HNO_3 (Martin et al., 2012). Minerals with relatively large concentrations in the identified profile
include Na, Al, Ca, Fe, Si, K, and Mg in descendent order presented concentration $>0.04 \mu g/m^3$ (contribution of $>26.5\%$).
The finding of Trejos et al. (2021) and Vanegas et al. (2021) supported the volcanic profile, with Si, Al, Fe, Ca, K, Mg, and
Na as the main minerals for the Nevado del Ruiz Volcano ashes. Although the composition might vary between volcanoes, the

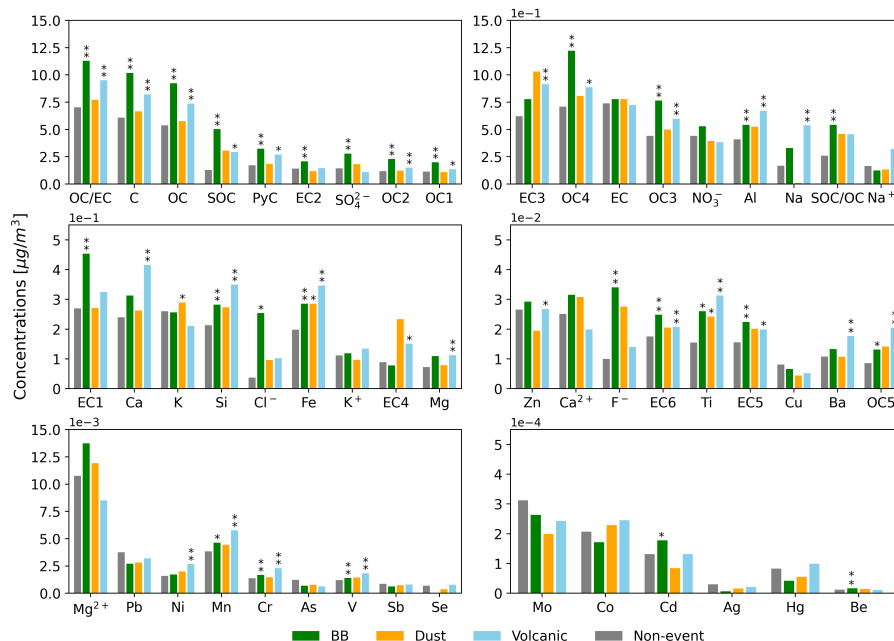


Figure 9. Campaign average concentration for non-event and positive-contribution days for BB-LRT, Dust-LRT, and Volcanic-LRT. The bars represent the concentration of the compounds along with the significance level compared with the days before and after the events, ≤ 0.1 (*) or ≤ 0.05 (**). The y-axes change for every figure to better identify concentrations. Supplementary Table S1-S2 presents these values and the mean concentrations of the compounds for days around (i.e., before and after) each event.

470 results for Sangay volcanic ashes in ~~the region~~ [South America](#) also back up the profile with similar main elements (Moran-Zuloaga et al., 2023). From these, Fe, Al, and Na are the most representative for the ashes diameter $< 2.5 \mu m$ (Mason et al., 2021); while Ca, Na, Si, and K might be particularly persistent after LRT (Ruggieri et al., 2012). [Cu and Zn are other tracers observed here and identified before for Colima Volcano in the southeast of the ring of fire in Mexico \(Miranda et al., 2004\).](#)

475 The total daily composition of $PM_{2.5}$ is also evidence of the impact of the LRT event. Fig. 9 shows the average daily concentration of the campaign-measured data for each kind of event and when no event is identified. This only considers days with a positive contribution factor from PMF results. Although the non-event average concentration is presented, the statistical comparison is made with the days before and after events to maintain similar meteorological conditions (i.e. the combined time-series of before (DtE₋₁₅ to DtE₋₄) and after event (DtE₊₄ to DtE₊₁₅) is statistically compared to the event time-series (DtE₋₃ to DtE₊₃) using the Mann Whitney test. Unlike the PMF model, ~~this comparison contains cations and all the comparison in Fig. 9 contains an analysis of the cations, the~~ carbon matter species ~~characterized for every event and widely-used ratios such as and the~~ OC/EC and SOC/OC ~~.The ratios for every type of event. Here,~~ the major elements generally have a more significant increment in LRT events. Some elements supported the model's fingerprint (Fig. 8), e.g., OC, OC1, OC2, SO_2^4 for BB; Fe, Al, and Ti for dust; and Si, Al, Fe, Ca, Mg, and Na for volcanic aerosols ~~.(Fig. 9).~~

480

significantly higher for BB, presenting a median OC/EC ratio of 11.3 that surpasses common urban combustion ratios like from fossil fuel (~ 4), combustion, and Diesel exhausted (<1) (Pani et al., 2019). Although OC/EC is more commonly used to identify sources of urban combustion and BB, some studies have shown its potential for determining the influence of volcanic activity (Pongpiachan et al., 2019). This is supported by evidence of carbon enrichment resulting from volcanic influence (Martinsson et al., 2009), which is also observed in this study, with consistently high concentrations in all OC species and almost all EC species (EC3, EC4, EC5, EC6). For volcanic aerosols, OC/EC had a median of 9.5. Likewise, the formation of SOC significantly increased for both events, with magnitudes of 5.0 and 2.9 for BB and volcanic aerosols, respectively. Heavy metals Cr (2.30 ngm^{-3}), Ni (2.67 ngm^{-3}), and Mn (5.76 ngm^{-3}) concentrations notably increased in event days for volcanic aerosols. Only the maximum concentration for Ni (25.05 ngm^{-3}) exceeds the concentration limit suggested for the annual average of the compound (20 ngm^{-3}) by European Commission (2019). Increments for Mn (4.64 ngm^{-3}) and Cr (1.67 ngm^{-3}), in addition to Cd (0.18 ngm^{-3}), were observed for BB. The (European Commission, 2019) set an annual limit of 5 ngm^{-3} for the Cd average.

The elevated concentrations of ions in the days of events (Fig. 9) also support the profiles selected (Fig. 8) and align with the literature. For the BB, in addition to the ions discussed observed in the PMF profile for the BB, the cations K^+ are representative ions (Rastogi et al., 2014; Moreno et al., 2023) that present significant increments for this type of event. Regarding the volcanic aerosol compositions (Fig. 9), the observed increment on Na^+ and K^+ also align with previous reports (Moreno et al., 2023; Mather et al., 2003; Roberts et al., 2018). On the other hand, although the PMF's fingerprint presented a high contribution of SO_2^{4-} and F^- , this was not enough for a significant rise in daily concentrations (Mather et al., 2003) shown in Fig. 9.

4 Discussion

Our analysis focused on events with high aerosol load and frequency, supported by a relatively long $\text{PM}_{2.5}$ chemical composition record. This approach avoided restricting the study to short periods (i.e., sporadic singular large-scale events) and made it possible to quantify longer-term and more frequent events (i.e., the long-term LRT transport of pollutants to the AV and not just the infrequent large-impact LRT events). We used satellite-derived data to identify regional-scale transboundary events and evaluate the effects on $\text{PM}_{2.5}$ concentration increments and composition.

Similar to other studies, thresholds were set for identifying events (Ridley et al., 2012; Carn et al., 2008); however, their selection in this study was representative of local events rather than high-polluted sporadic events. The OM-AOD threshold for BB-LRT events (0.2) agrees with previous results (0.2-0.3) (Kaiser et al., 2012; Misra et al., 2020). The Du-AOD (0.02) threshold was lower (i.e., 0.05) than suggested by Ridley et al. (2012) and that used by Achilleos et al. (2020) to track outstanding episodes but is of a comparable order of magnitude. In contrast, the TCSO_2 threshold in this study for volcanic degassing events differed from that discussed in the literature. The threshold used (0.87 mg/m^2) was lower more than ten times than the used for another degassing volcano (17.15 mg/m^2) (Carn et al., 2008).

515 The lower TCSO₂ threshold derived in this study is likely linked to the CAMS product we used. While we use the CAMS
reanalyses for OM-AOD, Dust-AOD, and TCSO₂ for consistency (i.e., tracers from the same model), the representation of
the actual magnitude of TCSO₂ in the reanalysis product is lower than the operational product. The reanalysis uses an older
climatological (2005-2010) emissions dataset (CAMS-GLOB-VOLC) for volcanic degassing (Granier et al., 2019) and only
520 assimilates satellite AOD. While satellite AOD is a useful product to help represent volcanic properties (e.g., volcanic plumes,
SO₂ concentrations), the operational product also assimilates satellite TCSO₂ (CAMS, 2023), providing more constraint on
the absolute magnitude of TCSO₂. However, regarding this study, the identification process is more important based on the
data variability rather than the overall magnitude.

The identified BB-LRT reproduced the seasonality described in the literature (Mendez-Espinosa et al., 2019; Hernandez
et al., 2019; Rodríguez-Gómez et al., 2022). Peak PM_{2.5} concentrations occurred in February and March for BB-LRT events.
525 In concordance with Mendez-Espinosa et al. (2019) and Henao et al. (2021), our findings show that the average conditions in
this period are characterized by wind transport from the northeast of the valley, covering part of the Orinoco and Caribbean
regions. BB aerosols from fires in the Orinoco can reach the valley through two passes in the Andes mountain range. The lim-
ited transport throughout this complex topography might promote the longer stance and accumulation of aerosols, increasing
their concentration in the AV (Ballesteros González, 2021). Furthermore, the back-trajectories warn about the Middle Mag-
530 dalena Valley, which suggests high BB emissions according to the average and the hotspots of OM-AOD. This area has been
previously considered critical for open fires in Colombia (Bolaño-Díaz et al., 2022; Ballesteros González, 2021). According to
Ballesteros González (2021), fires from this zone are more likely to affect the Andean cities than fires from the Orinoco due to
the mountain barriers.

The meteorology characterization for BB-LRT showed some conditions that might enhance the longer influence of the
535 identified events. The assessment highlights the dry conditions in NSA as promoters of both the occurrence of fires and LRT of
pollutants. In contrast, despite the high emissions of Amazon fire smoke in the southern hemisphere, these do not considerably
increase aerosols in the city due to the scavenging of particles in their transport (Hamburger et al., 2013). The BB-LRT around
August might be related to fires in that region (SIATA, 2021; Hamburger et al., 2013).

The estimated contribution to PM_{2.5} concentrations during the BB-LRT showed the highest magnitude for the studied events,
540 with 11.14 $\mu\text{g}/\text{m}^3$ (37.6%), [backed up by the comparison of PM_{2.5} concentration](#). According to Ballesteros-González et al.
(2020), the contribution from BB was calculated to be 4.7 $\mu\text{g}/\text{m}^3$ for the monitored city in February 2010 and 2018. This
finding is consistent with an average (maximum) estimation of 6.0 (12.5) $\mu\text{g}/\text{m}^3$ for samples taken in the same month in 2020.
Furthermore, the PMF results support the record of historical events, such as March 2020 at the beginning of the COVID-
19 pandemic. For this event, the PMF estimates a maximum contribution of 50.0 $\mu\text{g}/\text{m}^3$ out of 60.9 $\mu\text{g}/\text{m}^3$. The particular
545 event was characterized by local emissions suddenly decreasing and climbing in PM_{2.5} concentration (Henao et al., 2021;
Mendez-Espinosa et al., 2020).

Like BB-LRT, the monthly frequency of Dust-LRT events coincides with the annual cycle described by other studies. South
America experiences a dust reception from the Sahara desert mainly in two trimesters, from March to May and June to August
(Prospero et al., 2020). The seasonality of dust events partially coincides with the months of occurrence of BB-LRT events,

550 causing high overlapping between the identified days for these two types of events. Consequently, the identified weak impact on $PM_{2.5}$ by dust events controlled by Andes Mountain (Prospero et al., 2020) might be overshadowed by the most critical effect of BB-LRT. The only widely recognized event recorded for the study period occurred on June 24 and 25, 2020 (Mendez-Espinosa et al., 2020; SIATA, 2021). According to the PMF results, this exhibited almost two times higher contribution to $PM_{2.5}$ than the average event considered in this study. However, the PMF identified equivalent contributions in February 2020.
555 The overlapping between these events [and the small database size for the Dust-LRT model](#) might provoke impurities in the PMF factor related to Sahara dust. However, the dust profile aligns with the literature; the organic carbon contribution evidenced a likely mix with a combustion source.

In contrast to BB-LRT and Dust-LRT events, the impact of Volcanic-LRT is expected to be primarily determined by the direction of the wind rather than an actual seasonality in the emissions. Volcanic-LRT events occurred more frequently from
560 July to September, when low to mid-level winds blew from the Nevado del Ruíz volcano region in the southeast, facilitating the transport of emitted aerosols to the AV. According to the sulfur dioxide (SO_2) V2 catalog, available on <https://so2.gsfc.nasa.gov/measures.html> (Fioletov et al., 2023), the AV belongs to the influence area of the Nevado del Ruíz volcano. Indeed, this study identified a significant increment in $PM_{2.5}$ concentrations. According to the PMF results, on average, the contribution for $PM_{2.5}$ was $6.46 \mu gm^{-3}$. Likewise, Casallas et al. (2024) linked the increase in $PM_{2.5}$ concentration during the JJA trimester
565 with volcano activity in Cali, a relatively equidistant city located south of the volcano.

Although the Volcanic-LRT profile is predominantly made by SO_4^{2-} , the total concentration of the elements does not show a significant increment compared with surrounding days. This lack of significance in SO_4^{2-} has been observed in other cities where urban sources of SO_2 dominated (Miyakawa et al., 2007). However, some alternative hypotheses highlight regarding the magmatic gas state, pointing out that temperature before emissions might provoke low S oxidation and moderate SO_4^{2-}
570 formation (Mather et al., 2003) since SO_4^{2-} precursor formed at high temperature frequently close to the vent (Roberts et al., 2018). For instance, as evidenced in the Volcanic-LRT profile, Cl-poor plumes might reduce S oxidation and forward SO_4^{2-} (Mather et al., 2003).

The results have several implications for future research in the region. Modeling studies in the AV have only considered local emissions as inputs (e.g., Henao et al., 2020; Hernández et al., 2022), and it has been highlighted the need to include external
575 pollution sources as input in chemical transport models to have a more complete representation of air quality in the city. Further, this study suggests conducting shorter chemical campaigns and human health impact studies for target elements and sources, especially in the valley and upwind. For instance, special attention to the carbonaceous matter, F^- and Cd emitted from BB in open fires is needed for human and ecosystem health studies in Colombia (Tuomisto et al., 2008; Jayarathne et al., 2014), since this source is particularly important for the region. The infrequent contribution to Sahara dust is still significant, and there is
580 particular interest in the northern part of the country, where Sahara's most intense events significantly increase the mortality relative risk (Arregocés et al., 2023). On the other hand, Colombia has 12 active volcanoes, highlighting the importance of degassing activity in the country. Monitoring emissions and human health exposure is a primary need for municipalities, especially those close to active volcanoes such as Nevado del Ruíz. Both BB and Volcanic event contribution to heavy metals raise alerts around possible body metal accumulation and diseases such as thyroid cancer (Malandrino et al., 2020; Vigneri

585 et al., 2017). The inhalation intake of these heavy metals sums up to other exposures, such as ingestion and dermal route, which might also be higher near the source (e.g., from water and crop pollution). The increment in Mn and Cr might provoke $PM_{2.5}$ toxicity in lung epithelial cells (Yuan et al., 2019). A deep assessment of the Cr(VI) and Cr(III) needs to be done to evaluate the correct toxicity status of Cr.

Finally, this study advertises the need for cooperation among local, national, and international entities to manage complex
590 aerosol sources and human exposure.

5 Conclusions

This study evaluates the impact of BB-LRT, Dust-LRT, and Volcanic-LRT aerosols in the $PM_{2.5}$ concentration and chemical composition in the AV, a densely populated and mountainous region in the tropical Andes. Events for each aerosol type were identified using data with different anomalies from CAMS reanalysis products. Meteorological data from ERA5 and GPM,
595 along with a back-trajectories analysis, were used to link the aerosol events in the AV with the sources and to characterize the meteorological patterns that determine the LRT of aerosols. The impact of these regional events on the local $PM_{2.5}$ was finally evaluated with local concentration comparisons and characterized through PMF modeling. This study used a unique $PM_{2.5}$ composition campaign from April 2019 to October 2020.

The methodology allowed for identifying periods with LRT of pollutants, separating the days before, during, and after
600 the events. During event days, the back-trajectories of air masses arrive from zones previously reported as critical sources of aerosols in the region. This enables the connection between aerosol events in the AV and the sources. The sources were identified as regional fires for BB-LRT, Saharan sands for Dust-LRT, and emissions from the Nevado del Ruíz volcano for volcanic-LRT. The different types of events showed a very marked annual variability. BB-LRT events occurred mainly in February, March, and April, with a lower occurrence during August. Dust-LRT events showed a higher incidence from April to
605 August. Volcanic-LRT showed a high occurrence at the beginning of the second semester (from June to September) and during January and February. No events were identified during October, November, and December.

The weather patterns for BB-LRT events exhibited anomalous low-level northeasterly winds, favoring regional transport of aerosols. Additionally, drier conditions in northern Colombia and Venezuela promoted the occurrence and spread of fires in these regions. For Dust-LRT, more intense winds from the Caribbean at medium levels favor transport to the AV when the
610 aerosol load is high. Moreover, during Volcanic-LRT events, anomalous southeasterly winds at low and medium levels were present, along with low precipitation in the southeast of the AV, where the Nevado del Ruíz volcano is located, enabling the transport to the AV.

When evaluating the impact of LRT events on ground-level $PM_{2.5}$, BB-LRT events resulted in more significant increases in concentrations. The relative increment on $PM_{2.5}$ from Dust-LRT and Volcanic-LRT events was smaller but still significant.
615 Similarly, underpinned by literature-supported profiles, the PMF results evidenced the event's influence on the $PM_{2.5}$ chemical composition. OC species such as OC1 and OC2, anions as F^- and the traces of secondary pollutants SO_4^{2-} and NO_3^- primarily determine the profile identification for BB-LRT. On the other hand, general crustal minerals, together with Ca and Ti, define

the Sahara-influenced Dust-LRT profile. Regarding Volcanic-LRT, SO_4^{2-} and characteristic minerals Na, Al, Ca, Fe, Si, K, and Mg represented the profile. Further comparison in positive contributed event days supports the PMF results although identified
620 no substantial SO_4^{2-} rise during volcanic degassing events. This aligned with the observed low Cl^- contribution, supporting the reduced S oxidation hypothesis and forward production of SO_4^{2-} . Carbon and ion identification increments were mainly observed for BB-LRT and Volcanic-LRT. Lastly, the profile associated with significant ~~increment~~ increments in the heavy metals Cr, Mn, Cd, and Ni raises concerns. Future studies using chemical models, larger events datasets, and more detailed measurement and characterization techniques may offer further explanations for this ~~increment~~ increase.

625 The findings have several implications for future research. For instance, modeling studies in the region have only considered local emissions as inputs, and external pollution sources, especially volcanic degassing, need to be included as inputs in chemical transport models.

Code availability. A Python code for calculating the back-trajectories is available at https://github.com/MariaPa96/Basic_Trajectories_python.

630 *Data availability.* The CAMS reanalysis (Inness et al., 2019) data were downloaded from the Atmosphere Data Store at <https://ads.atmosphere.copernicus.eu/cdsapp#!/dataset/cams-global-reanalysis-eac4?tab=overview>. Meteorological data of the ERA5 reanalysis were downloaded from the Climate Data Store at <https://cds.climate.copernicus.eu/cdsapp#!/dataset/reanalysis-era5-pressure-levels> and GPM precipitation data was retrieved from the NASA Earthdata at https://disc.gsfc.nasa.gov/datasets/GPM_3IMERGHH_07/summary. The hourly $\text{PM}_{2.5}$ concentration were download from the official network platform at https://siata.gov.co/siata_nuevo/. The datasets of wind for the trajectories
635 calculation were downloaded from NCEP-NCAR Reanalysis 1 (<https://psl.noaa.gov>). The $\text{PM}_{2.5}$ chemical campaign datasets are available upon request to MGM (mgomez@elpoli.edu.co).

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