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Env. Pollution



CAUSES OF THE UNREMITTING HIGH AMBIENT LEVELS OF PM_{10} IN A SUBURBAN BACKGROUND SITE IN NE SPAIN

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Study Area

Manlleu and the air quality-monitoring site (42°00'12.1"N 2°17'13.5"E).



Introduction

Long- and short-term exposure to PM_{2.5} and PM₁₀ is linked to increased respiratory and cardiovascular morbidity, premature mortality [1-3], and substantial health-related costs (4.5% of global GDP) [4]. Despite significant efforts to reduce PM through policies in Europe, 97% of the EU-27 urban population still faces PM_{2.5} concentrations above WHO guidelines (5 µg m⁻³) [5]. Secondary PM, especially organic aerosols, poses a challenge for abatement. In Spain, Manlleu is a notable PM pollution hotspot, largely due to biomass burning and intensive agriculture. This study analyses long-term PM₁₀ and BaP measurements, combined with speciation data, to assess PM sources and impacts, particularly biomass combustion, in urban Europe.

Figure 1.

a) Daily PM₁₀ concentrations and annual 90.4 percentile value (should not exceed 50 μ g m⁻³) for 2006-2024. b) Daily BaP concentrations and annual average (should not exceed 1 ng m^{-3}) for 2008-2024. Sampling campaigns and complete chemical analysis of PM_{10} in Blue squares. Monthly concentrations from November to March (NDJFM) of 2008-2023: c) PM_{10} , d) BaP, e) average 1/T and f) average BaP/PM_{10} ratios.

Results and conclusions

- PM₁₀ levels in Manlleu have exceeded the daily limit since 2006
- BaP concentrations exceeded the EU target and are 3 to 9 times higher than in Barcelona.
- Highest PM₁₀ and BaP concentrations are observed in winter months.
- BaP/PM₁₀ ratios indicate significant biomass burning, with a clear correlation between colder temperatures and higher pollutant levels.
- Biomass burning significantly contributes to PM₁₀ and BaP, with a reduction in BaP due to improved industrial practices after 2015.
- Biomass Burning Organic Aerosol (BBOA).
- SOA contributed to PM₁₀ mainly in summer months, while BBOA and MinInd were dominant during Winter.



Figure 3.

Contributions (in $\mu g m^{-3}$ and percentage) of the four main sources identified to the gravimetric PM₁₀ concentrations determined in the 2021–2022 period. a) Whole period from December 2021 to December 2022. b) Winter studied period, NDJFM.







• Chemical mass closure indicates high contributions from organic carbon, ammonium, and combustion-related pollutants, with trace metals elevated during heating periods. • PM₁₀ source apportionment revealed four key sources: Secondary Organic Aerosol (SOA), Mineral and Local Industry (MinInd), Agriculture and Fodder Production (AgFo), and



Figure 4.

a) Chemical profiles of the PMF analysis applied to PM_{10} speciation datasets in 2021–2022 (122 samples, 47 species). PM component concentrations of species (y1-axis, blue bars) and total variable percentages (y2-axis, red squares). b) Seasonal evolution of the contribution of main pollution sources.



• Non-refractory PM₁ source apportionment revealed biomass burning, particularly domestic, as the dominant source during winter, with significant contributions from traffic and secondary processes. • Despite reductions in industrial emissions, regional measures to address biomass burning and NH₃ emissions from agriculture have been ineffective, highlighting the need for stricter controls and continuous monitoring.



References

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Analysis of air quality data from the official surveillance network in Manlleu, which records hourly concentrations of pollutants like NO, NO₂, SO_2 , O_3 , and PM_{10} . The last offline PM_{10} speciation campaign was conducted from 2021 to 2022 to analyse inorganic and organic compounds, with techniques such as ICP and GC-MS. Additionally, an online PM₁ speciation campaign was performed from 2016-2017 to monitor hourly variations in pollutants, including black carbon and organic aerosols, focusing on biomass burning sources. This combination of offline and online methods enabled a comprehensive evaluation of PM origins and variations in Manlleu. The receptor modelling analysis was done by applying the Positive Matrix Factorization model, (US-EPA-PMF 5.0) to PM



Figure 2. Syringe quartz filters after filtering lixiviates.

Figure 5. Time series, weekly and diel PM₁ concentrations (a), PM₁ compounds (b, g, h, I, j), OA sources (c, d, e, f) and BC sources (k, l).

