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34 Introduction

35 Pollen is one of the leading causes of allergic diseases worldwide (World Allergy
36 Organization, 2013), affecting an estimated 15-40 % of the European population (Rojo et
37 al., 2024). The prevalence of allergic rhinitis and other pollen-related conditions
38 continues to increase globally (Adams-Groom et al., 2022; Beggs et al., 2023; D'Amato et
39 al., 2007). The severity of allergic responses depends on several factors, including pollen
40 concentration, species-specific sensitization thresholds and individual susceptibility
41 (Rapiejko et al., 2007; Rodríguez-Rajo et al., 2010).

42 In recent decades, climate change has become a key factor influencing the dynamics of
43 pollen production and dispersion (Anderegg et al., 2021; Lake et al., 2017). Rising
44 temperatures and elevated atmospheric CO₂ levels contribute to increased pollen
45 production and cause earlier, longer and more intense pollen seasons for many allergenic
46 taxa. As a result, exposure periods are becoming more variable and less predictable, with
47 the timing of peak pollen concentrations often shifting from year to year, which
48 complicates forecasts of allergy risk (Pacheco et al., 2021; Paudel et al., 2021; Tomczyk
49 et al., 2025; Ziska et al., 2019).

50 Given these shifts, continuous and precise monitoring of airborne pollen has become
51 crucial for understanding exposure patterns and supporting public health management,
52 allergy forecasting and early-warning systems (Grewling et al., 2023; Maya-Manzano et
53 al., 2023). Pollen monitoring networks were initially established to support allergy
54 diagnosis and treatment but are now also used to study climate change impacts, track
55 invasive species and monitor airborne biological particles (Galan et al., 2014)

56 Most monitoring sites still rely on manual sampling methods, such as the Hirst-type
57 pollen trap (Hirst, 1952), which, despite being standardized and quality-controlled
58 through aerobiological networks (CEN/TS 16868, 2015), have several limitations. These
59 include delays of 1-9 days before data are available, low temporal resolution and
60 uncertainties related to airflow variations, counting methodology, adhesives and
61 observer differences (Adamov et al., 2024; Triviño Triviño et al., 2023). However, Hirst-
62 type traps remain the reference against which other pollen and fungal spore detection
63 methods are evaluated (Tummon et al., 2024). At the same time, the development and
64 validation of low-cost sensors are essential for expanding monitoring networks, as they
65 can help identify optimal locations for the subsequent deployment of automatic
66 samplers and improve overall spatial coverage. However, Hirst-type traps remain the
67 reference against which other pollen and fungal spore detection methods are evaluated
68 (Tummon et al., 2021).

69 In recent years, technological advances have enabled real-time detection of pollen and
70 other bioaerosols, using diverse approaches and instruments (Buters et al., 2024; Maya-
71 Manzano et al., 2023). These systems allow continuous monitoring, higher temporal
72 resolution and improved responsiveness for public health and research applications. The

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73 rapid evolution of these technologies is transforming pollen monitoring, providing critical
74 real-time information for allergy management, climate studies and environmental
75 research (Sauvageat et al., 2020). One of the devices is the Swisens Poleno Jupiter, an
76 ~~advanced~~ instrument designed for real-time bioaerosol monitoring, capable of detecting
77 and identifying airborne particles such as pollen. It combines digital holography with
78 fluorescence measurements, providing detailed information on both the morphology and
79 composition of particles to support the development of automatic pollen detection
80 systems (Erb et al., 2025). The recent studies highlight that automatic pollen detection
81 even with ~~advanced~~ machine learning, has not achieved full accuracy or maturity and
82 further work is needed to generalize such models across more pollen species and
83 conditions. They emphasize that significant improvements in algorithms, training data and
84 validation strategies are still required before automated pollen identification can match
85 expert human analysis (Farooq et al., 2025; Gimenez et al., 2024; Shamrat et al., 2024).
86 ~~We have undertaken some of these gaps by the extending and refining the reference~~
87 ~~pollen database for the Swisens Poleno Jupiter automatic monitoring system. The~~
88 ~~reference pollen database provided with the Swisens Poleno System comprises 14~~
89 ~~taxonomic classes and an additional class for water droplets. It was originally developed~~
90 ~~based on data collected at the site where the detector was built in Switzerland, without~~
91 ~~adaptation to new locations. In this study, the reference database used in the Swisens~~
92 ~~Poleno Jupiter system was further developed through extension and refinement in order~~
93 ~~to improve detection performance.~~

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95 ~~For this purpose, during the pollination seasons, local pollen samples were collected and~~
96 ~~used to retrain the neural network model. The new data enabled the incorporation of~~
97 ~~fluorescence measurements, whereas the initial model was based solely on holography~~
98 ~~prior to retraining. We hypothesised that the retrained model would provide more~~
99 ~~accurate and timely predictions of airborne pollen concentrations compared to models~~
100 ~~trained exclusively on foreign datasets, as well as improved performance due to the~~
101 ~~addition of fluorescence data.~~
102 ~~For this purpose, during the pollination seasons, local~~
103 ~~pollen samples were collected and used to retrain the neural network model. We~~
104 ~~hypothesised that the retrained model would provide more accurate and timely~~
105 ~~predictions of airborne pollen concentrations compared to models trained solely on~~
106 ~~foreign datasets.~~
107 The study describes the measurement campaigns carried out as part of
108 the development of automatic pollen detection methods. To date, there are no studies
109 explicitly emphasizing the importance of retraining neural network models using locally
110 collected pollen datasets that reflect real environmental conditions. Secondly, we
111 analysed the similarities and differences in the diurnal concentration patterns of five
112 pollen taxa — *Alnus* (alder), *Betula* (birch), *Fraxinus* (ash), *Pinus* (pine) and *Quercus* (oak).
These tree ~~species-genera~~ are known for their allergenic potential, particularly *Alnus* and
Betula from the Betulaceae family (Biedermann et al., 2019). Owing to their high pollen

productivity, airborne concentrations of these taxa can fluctuate substantially (Malkiewicz et al., 2016). We hypothesised that diurnal variability would show comparable patterns among taxa with similar pollen release mechanisms, while differing among ~~species-taxa~~ with distinct phenological or aerodynamic characteristics. Additionally, the availability of high-resolution hourly data enabled us to assess the influence of meteorological conditions on short-term variability in pollen concentrations. We used standard meteorological parameters, including temperature, relative humidity, precipitation, wind and sunshine duration, to obtain a comprehensive understanding of the factors influencing pollen concentrations. However, in order to obtain a broad view of the factors influencing pollen concentrations, we also included the planetary boundary layer height among the analysed parameters. ~~Although frequently mentioned in aerobiology, its influence on hourly pollen variability derived from automatic measurements has rarely been investigated compared to standard local meteorological drivers, with the first attempt in this direction made by Chappuis et al. (2020). Although frequently mentioned in aerobiology, its influence on hourly pollen variability has rarely been investigated compared to standard local meteorological drivers.~~

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This study provides two years analysis of diurnal airborne pollen dynamics using data from an automatic real-time monitoring system. It is the first to employ the Swisens Poleno Jupiter detector for continuous, high-resolution assessment of diurnal variability across multiple pollen seasons. In contrast to previous studies based on traditional Hirst-type volumetric traps (Clot, 2001; Kapyla, 1984; Kasprzyk et al., 2001; Scevkova et al., 2015), this approach allows for a more detailed and consistent evaluation of hourly fluctuations and interannual trends in pollen concentrations.

2. Data and methods

2.1 Automatic detection

Location and time frame

~~The Swisens Poleno Jupiter detector is installed in Wroclaw (southwestern Poland), on the roof of the Department of Climatology and Atmosphere Protection, University of Wroclaw, at an elevation of 20 meters above ground level, with no other high-rise structures in the immediate surroundings. The detector Swisens Poleno Jupiter is installed in Wroclaw (southwestern Poland), on the roof of the Department of Climatology and Atmosphere Protection, University of Wroclaw, at an elevation of 20 meters above ground level.~~ The instrument is co-located with a standard Hirst-type volumetric trap, allowing for direct comparison between automatic and manual monitoring methods. The measurements cover the periods of 2024 and 2025, encompassing two complete pollen seasons.

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Station location in Wrocław city



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Figure 1. Location of the Swisens Poleno Jupiter detector in Wrocław, Poland (red dot); the inset shows the location of Poland within Europe. The European map is based on Natural Earth data, while the city map uses data from OpenStreetMap contributors.

Description of the detector

The Swisens Poleno Jupiter (Swisens AG) is an ~~airborne-cytometer~~ [airflow cytometer](#) designed for real-time analysis of individual particles ranging from 0.5 to 300 μm , operating at a sampling flow rate of 40 liters per minute. As particles pass through the instrument, they are concentrated within the measurement chamber, where two orthogonally positioned lasers detect each particle, providing estimates of size and velocity based on light scattering. Once detected, the particle is captured by two greyscale holographic cameras positioned at 90° angles to each other and perpendicular to the airflow. The holographic images are reconstructed numerically to a resolution of 200×200 pixels, with each pixel representing $0.595 \times 0.595 \mu\text{m}$ in the physical domain. These images enable precise characterization of particle morphology. Immediately after imaging, laser-induced fluorescence (LIF) is measured. Particles are sequentially excited by three laser sources (280, 365 and 405 nm) and the resulting fluorescence is recorded across five spectral channels (333-381, 411-459, 465-501, 539-585 and 658-694 nm),

hereafter referred to by their central wavelengths: 357, 435, 483, 562 and 676 nm. Fluorescence emission is captured by silicon photomultipliers (SiPMs), producing up to 15 intensity measurements per particle. Due to saturation and single-photon excitation effects, the effective number of usable fluorescence channels is 13. The fluorescence data undergo additional preprocessing to enhance usability and robustness. In this work, analysis focuses on particle morphology (from holographic images) and chemical composition (from fluorescence intensities). The instrument also performs polarized-scattered-light measurements, which are not used here (Erb et al., 2024; Erb et al., 2025; Sauvageat et al., 2020).

Model architecture

The pollen classification model was based on a customised convolutional neural network architecture specifically optimised for processing greyscale images generated by the SwisensPoleno digital holography module. Unlike standard pre-trained architectures such as VGG16, which are primarily designed for RGB image recognition tasks, the applied network was adapted to the characteristics of monochromatic holographic data, reducing unnecessary model complexity and improving compatibility with the input format. The model processed two orthogonal pollen images separately before combining the extracted features through fully connected layers. Additionally, fluorescence measurements could be incorporated through a dedicated network branch, enabling classifications both with and without fluorescence input (Crouzy et al., 2025; Erb et al., 2025; Sauvageat et al., 2020).

Non-biological particles were removed prior to classification using a deterministic morphological filter described by Sauvageat et al. (2020). Water droplet training datasets were created from operational measurements collected during fog and rain events and manually cleaned to remove artefacts such as aggregates and debris. The model was trained entirely from scratch without the use of pre-trained neural networks. As emphasised by Erb et al. (2025), reliable pollen classification requires training datasets that reflect the full variability of each pollen taxon, including samples collected from different plants, locations, and meteorological conditions. This underlines the importance of the present study. The detailed network architecture is available on GitHub (MeteoSwiss Biometeorology Team, 2025)."

Measurement campaigns

The measurement campaigns were designed to collect high-quality reference data for retraining and validating the automatic pollen detection model. A total of five taxa- *Alnus*, *Betula*, *Fraxinus*, *Quercus* and *Pinus* - were included in the experiments, with the number of campaigns, recorded events, measurement duration and amount of pollen adjusted to

209 the availability and characteristics of each species (see Table 1). The total number of
210 recorded events ranged from 3 622 for *Fraxinus* to 7 233 for *Betula*, with measurement
211 durations between 19 min and 68 min and pollen quantities from 0.5 ml to 1.25 ml.

212 Each campaign consisted of a series of controlled laboratory experiments aimed at
213 introducing locally collected pollen into the automatic detector under standardized
214 conditions. The main goal was to extend the pollen database with accurately identified
215 local taxa and to ensure the consistency and reliability of data used for model
216 development.

217 The campaigns were carried out using the Swisens Poleno Atomizer, a device that enables
218 controlled aerosolization of pollen grains directly from a test tube into the Swisens Poleno
219 Jupiter detector. Swisens Atomizer delivers the particles directly to the Swisens Poleno
220 detector under controlled conditions. This system disperses selected particles at a stable
221 rate by combining vibration and airflow. The vibration frequency, amplitude and blower
222 speed can be precisely adjusted. The Swisens Atomizer is positioned directly above the
223 SwisensPoleno Jupiter so that the sampled air contains the target pollen particles. This
224 setup allows simultaneous measurement and labeling of all collected data with the
225 corresponding particle type (Erb et al., 2025). The entire process is managed via an
226 integrated computer system, which provides feedback on instrument performance and
227 enables data recording when the conditions are optimal.

228 Each campaign followed a structured workflow:

- 229 1. Pollen preparation - Inflorescences of *Alnus*, *Betula*, *Fraxinus*, *Quercus* and *Pinus*
230 were collected during their natural pollination periods and processed to obtain
231 clean pollen grains. It is important to monitor meteorological conditions and
232 collect catkins when they open naturally (during sunny weather and temperatures
233 above 10 °C) or to collect inflorescences and let them open in warm, dry
234 conditions.
- 235 2. Measurement phase - The Atomizer introduced the pollen into the detector, with
236 continuous monitoring to ensure high-quality data.
- 237 3. Data validation and selection - Collected signals were reviewed to remove artifacts
238 or non-pollen particles.
- 239 4. Dataset compilation - Verified data were organized using *Swisens Data Explorer*
240 software, forming the basis for neural network retraining.
- 241 5. Model retraining and testing - The new model was trained on a subset of the
242 dataset and evaluated on an independent portion to assess detection
243 performance.

244 The campaigns were repeated for each taxon to ensure representative coverage of the
245 dominant allergenic species in the region. Each dataset contained thousands of

individual particle measurements, reflecting both the natural variability of pollen availability and the specific conditions of each measurement session.

Table 1. Summary of total dataset sizes for each pollen taxon

Taxa	Total number of recorded events	Number of campaigns	Duration of the measurement phases	Amount of pollen per campaig
<i>Alnus</i>	6 409	5	50 min	1.25 ml
<i>Betula</i>	7 233	2	68 min	1.25 ml
<i>Fraxinus</i>	3 622	1	19 min	0.5 ml
<i>Quercus</i>	5 975	2	52 min	1 ml
<i>Pinus</i>	6 572	1	45 min	1 ml

For reliable particle identification, input data must contain only the target pollen type. Environmental samples often include unwanted particles such as dust, other pollen, or plant debris, necessitating a data cleaning step. Using the SwisensDataExplorer tool, expert annotators manually removed non-target particles, producing “clean” datasets for training. Typical exclusions included double particles, irregular shapes and out-of-range sizes, while blurred images were retained to preserve natural variability. Although manual cleaning is subjective and not an absolute ground truth, it ensures that essential particle characteristics are maintained. These cleaned datasets were then used to retrain a neural network and provide a new detection model.

2.2. Station surroundings

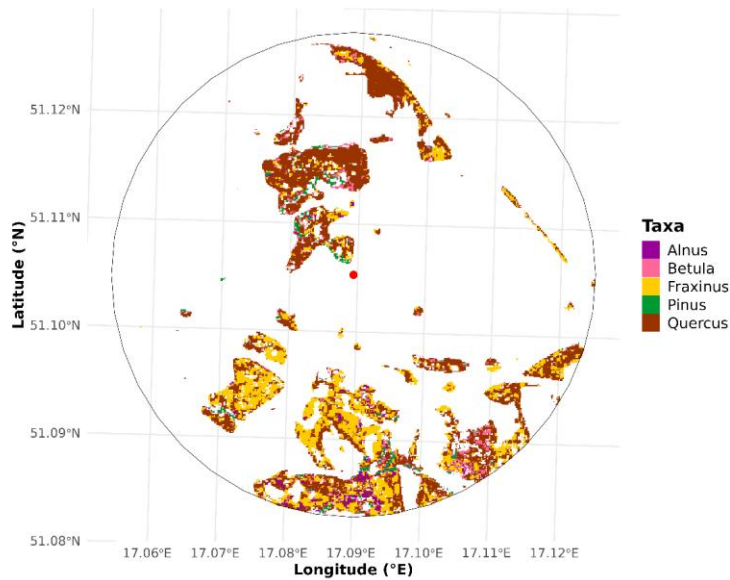


Figure 2. Station surroundings based on data from Grabska-Szwagrzyk et al. (2024); map prepared by the author.

Within a 2.5 km buffer around the monitoring station, the analyzed tree species are predominantly located to the north and south of the site. The northern area, corresponding to Park Szczytnicki, is mainly dominated by *Quercus*, *Pinus* and *Betula*. In contrast, the southern sector, located near the Odra River, contains the highest density of trees, particularly *Fraxinus* and *Quercus*, with additional occurrences of *Betula* and *Alnus* along the riverbanks. The western and eastern directions are sparsely vegetated, containing only isolated individual trees rather than larger clusters.

2.3. Hirst trap measurements

For validation purposes, airborne pollen samples were collected using a volumetric 7-day Hirst-type trap (Hirst, 1952), co-located with the Swisens Poleno detector at the same height on the roof of the Institute of Climatology and Atmospheric Protection. The Hirst trap continuously draws in air and pollen grains, along with other particles, adhere to a prepared sticky tape. After one week, the tape is replaced and laboratory slides are prepared, from which pollen grains are counted following European standards (Galán et al., 2014), i.e., from four continuous horizontal bands. These Hirst-type measurements served as reference data to validate the results obtained from the Swisens Poleno instrument and the neural network model.

2.5. Meteorological data

Meteorological data were obtained from automatic measurements at the Department of Climatology and Atmospheric Protection, where the detector and Hirst pollen trap are located. The dataset includes mean hourly records of temperature at 2m [°C], wind speed [m/s] and direction [°] at 10m, relative humidity at 2m [%], sum of precipitation [mm] and sunshine duration [hours]. Additionally, the WRF model was used to determine the hourly variability of planetary boundary layer (PBL) height [m]; however, only data for 2024 are presented, as simulations for the current year are still ongoing.

2.6. Method of analysis

Evaluation of the old and new model results

The pollen season was determined separately for both the automatic real-time detector (Swisens Poleno Jupiter) and the reference Hirst-type sampler using the 90% method (Nilsson and Persson, 1981). The start of the pollen season was defined as the day when the cumulative pollen count exceeded 5% of the annual total and the end as the day when it surpassed 95%. These season limits were then applied consistently for validation of the automatic measurements and for the subsequent statistical and diurnal variability analyses.

To evaluate temporal consistency, daily concentrations from two consecutive pollen seasons were compared for both the old and new automatic models (*hereafter referred to as “old” and “new”, respectively*) against Hirst reference data. The baseline model was originally developed by MeteoSwiss and trained on datasets obtained from the MeteoSwiss pollen monitoring network in Switzerland. The official GitHub repository provides access to *the old model – MCH 2022 model* (*the MCH 2022 model*) (MeteoSwiss Biometeorology Team, 2025MeteoSwiss, 2022). The current model (*new*) was retrained using locally collected pollen data from Wrocław, Poland, for the same taxa, with the aim of improving predictive performance under local environmental conditions. While the *oldprevious* model relied solely on holographic images, the new model incorporates both holography and fluorescence spectral information. Crouzytozy et al. (2025) demonstrated that including additional information improves detection performance; however, studies on the use of locally collected pollen to improve model performance for specific taxa remain scarce., *which is the primary focus of the present work.*

Agreement between the automatic method and the Hirst was quantified using the coefficient of determination (R^2) and the root mean square error (RMSE). R^2 assessed the proportion of variance explained, while RMSE provided an absolute measure of average daily deviation.

For evaluation by exposure level, hourly concentrations were binned into four classes (low, medium, high, very high). Thresholds were defined individually per taxon based on observed distributions. As an example, *Betula* (birch) classes were:

326 Low: ≤ 30 pollen grains m^{-3}

327 Medium: 31-80 pollen grains m^{-3}

328 High: 81-150 pollen grains m^{-3}

329 Very high: > 150 pollen grains m^{-3}

330 (Analogous taxon-specific thresholds were applied for *Alnus*, *Fraxinus*, *Pinus* and
331 *Quercus*.)

332 Model outputs (old vs. new) were evaluated for each taxon and concentration class using
333 contingency and skill metrics computed from counts of True Positives (TP), False Positives
334 (FP) and False Negatives (FN). Derived metrics included: Probability of Detection ($\text{POD} =$
335 $\text{TP}/(\text{TP}+\text{FN})$), False Alarm Ratio ($\text{FAR} = \text{FP}/(\text{TP}+\text{FP})$), Critical Success Index ($\text{CSI} =$
336 $\text{TP}/(\text{TP}+\text{FP}+\text{FN})$) and Success Rate ($\text{SR} = (\text{TP}+\text{TN})/\text{total}$, where TN is True Negatives). These
337 metrics were reported separately for each taxon and concentration level to characterize
338 model behavior across exposure ranges.

339

340 Diurnal variability of pollen concentrations

341 Hourly concentrations from the Swisens Poleno Jupiter (new model) were analyzed to
342 characterize diurnal patterns for each taxon. All timestamps were recorded in
343 Coordinated Universal Time (UTC). Differences in hourly distributions between the two
344 seasons were quantified and visualized to identify peak exposure hours.

345 Diurnal variability of meteorological parameters and pollen data

346 Finally, an overall correlation matrix was constructed to assess the relationships between
347 hourly meteorological variables (temperature, wind speed and direction, relative
348 humidity, precipitation and sunshine duration) and pollen concentrations obtained from
349 the Swisens Poleno Jupiter (new model). Subsequently, for each taxon, Spearman rank
350 correlations were calculated between pollen concentration and each meteorological
351 variable separately for each month within the respective flowering season, in order to
352 capture the seasonal and hourly variability of these relationships. Previous research has
353 shown that meteorological conditions strongly influence airborne pollen concentrations
354 (Dąbrowska-Zapart et al., 2020; Kluska et al., 2020; Malkiewicz et al., 2016; Tomczyk et
355 al., 2025), justifying the use of correlation analyses to assess their effect on hourly pollen
356 variability. Statistical significance was assessed at three levels ($p < 0.05$, $p < 0.01$ and $p <$
357 0.001) and correlation magnitudes and significance patterns were reported and mapped
358 to hourly contexts.

359 The hourly planetary boundary layer (PBL) height was determined using the WRF
360 numerical meteorological model, with model setup and parameterizations following
361 Skamarock et al. (2021). Spearman's rank correlations were then calculated between PBL

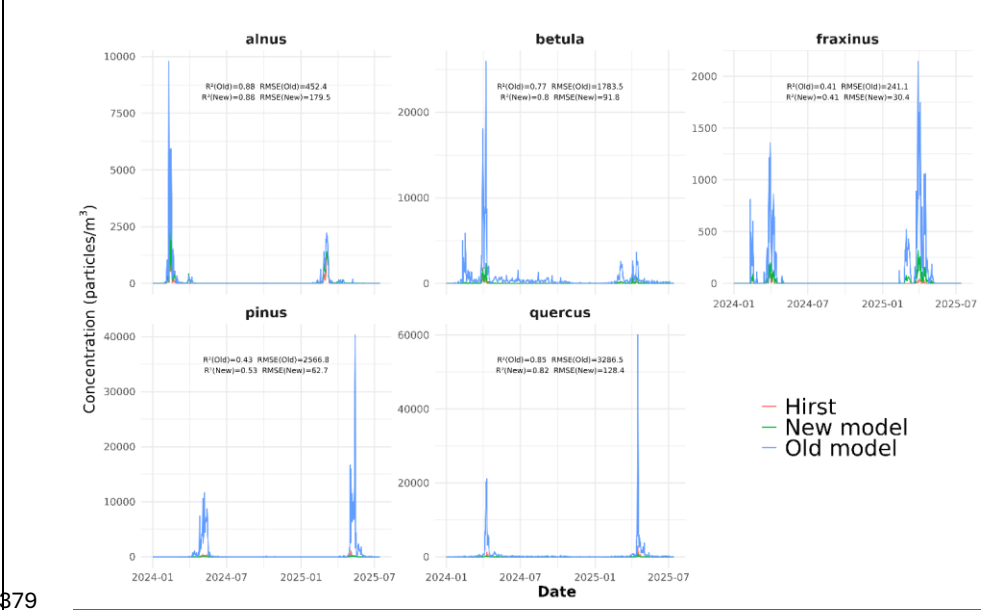
height and pollen concentrations to assess how boundary layer dynamics influence short-term variations in airborne pollen levels. Correlations for all variables were adjusted for multiple comparisons using the Bonferroni correction to account for the large number of observations.

A wind-rose analysis was performed using concurrent hourly wind direction and speed records to identify prevailing wind sectors associated with the highest pollen concentrations. These wind sector concentration patterns were compared spatially with land-cover/land-use maps (see Figure 2) to infer potential source areas and directional contributions to measured pollen loads. Figure 2 was created based on detailed tree distribution maps developed by Grabska-Szwagrzyk (2024), who classified 16 dominant tree species and genera across Poland using time series of Sentinel-2 imagery.

3. Results

3.1 Pollen concentrations from the Swisens Polleno Jupiter and Hirst

This presents a comparison between the machine learning model trained on Swiss pollen data (old model) and the newly retrained model using local taxa (new model) against Hirst measurements, highlighting differences in predicted concentrations, temporal patterns and deviations from observed values.



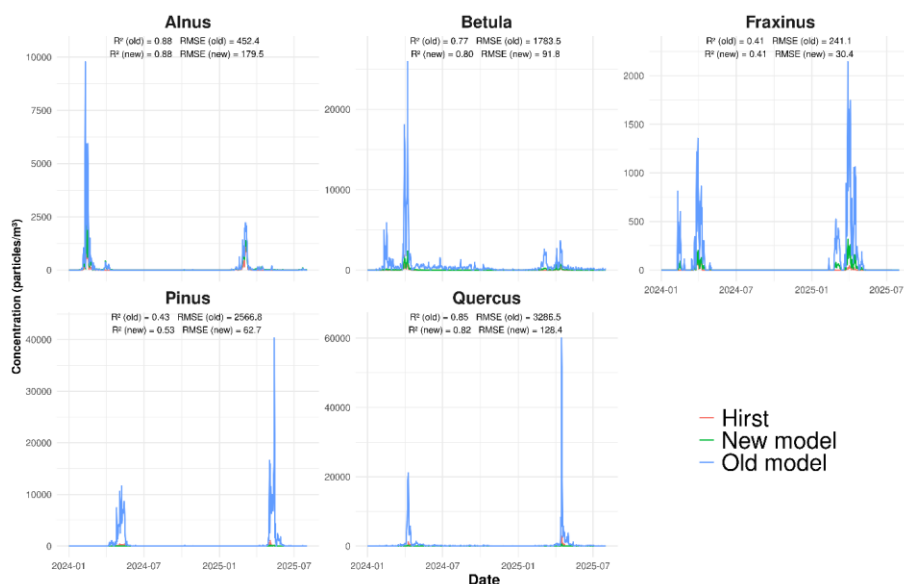


Figure 3. Daily pollen concentrations from old and new ML models compared with Hirst reference data.

For *Alnus* and *Betula*, both models capture the main seasonal peaks well, with *Alnus* showing a pronounced early spring maximum. The new model closely follows the Hirst observations in both shape and timing of peaks, while the old model tends to overestimate concentrations. For *Alnus*, both models achieve the same $R^2 = 0.88$, but the new model shows a much lower RMSE (179.5 vs 452.4, for old and new model respectively - here and thereafter), indicating improved accuracy. Similarly, for *Betula*, the new model slightly improves the determination coefficient ($R^2 = 0.80$ vs 0.77) and reduces RMSE substantially (91.8 vs 1783.5).

For *Fraxinus*, concentrations remain relatively low, with both models capturing the timing of peaks. The new model performs markedly better, maintaining the same $R^2 = 0.41$ but achieving a much lower RMSE (30.4 vs 241.1), reflecting greater precision in reproducing small variations.

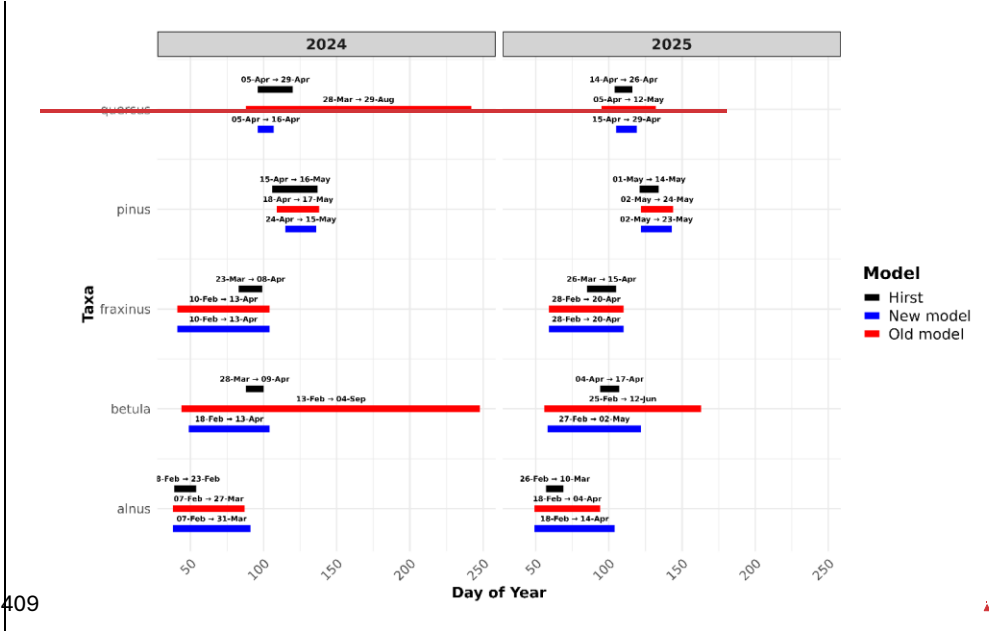
In the case of *Pinus*, the new model better reproduces both low and moderate concentration periods, while the old model significantly overestimates peak magnitudes. This improvement is confirmed by a higher R^2 (0.53 vs 0.43) and a significantly reduced RMSE (62.7 vs 2566.8).

401 For *Quercus*, a distinct late spring peak is observed. Both models correctly identify the
402 timing of high pollen events, but the new model more accurately represents the peak
403 structure and intensity, with a slightly lower R^2 (0.82 vs 0.85) and a drastically smaller
404 RMSE (128.4 vs 3286.5), highlighting a much-improved fit to observed concentrations.

405

406 **3.2 Comparison of Models - Pollen Season Duration**

407 This examines how each model predicts the start and end of the pollen season, analyzing
408 differences in season length and timing in comparison with observed data.



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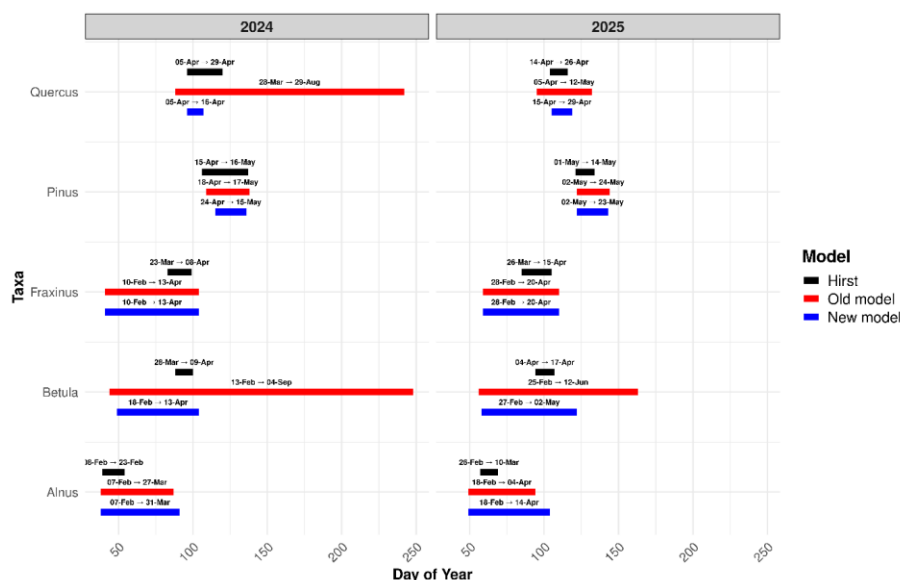


Figure 4. Start and end dates of pollen seasons for the studied taxa.

For *Alnus*, both years show that the new model reproduces the Hirst-derived start dates well, whereas the end dates are captured less accurately (2024: 7 Feb-27 Mar vs 8 Feb-23 Feb; 2025: 18 Feb-4 Apr vs 26 Feb-10 Mar), while the old model predicts a longer season (until the end of March or mid-April).

For *Betula*, the old model largely overestimates the pollen season duration (2024: 13 Feb-4 Sep; 2025: 25 Feb-12 Jun), while the new model (2024: 18 Feb-13 Apr; 2025: 27 Feb-2 May) better captures the Hirst timing (2024: 28 Mar-9 Apr; 2025: 4 Apr-17 Apr).

For *Fraxinus*, both models identify the onset of the season earlier than Hirst in 2024 (10 Feb-13 Apr vs 23 Mar-8 Apr), but the new model better reproduces the duration. A similar pattern appears in 2025, where the new model (28 Feb-20 Apr) aligns with Hirst (26 Mar-15 Apr) more accurately than the old model, which extends the season.

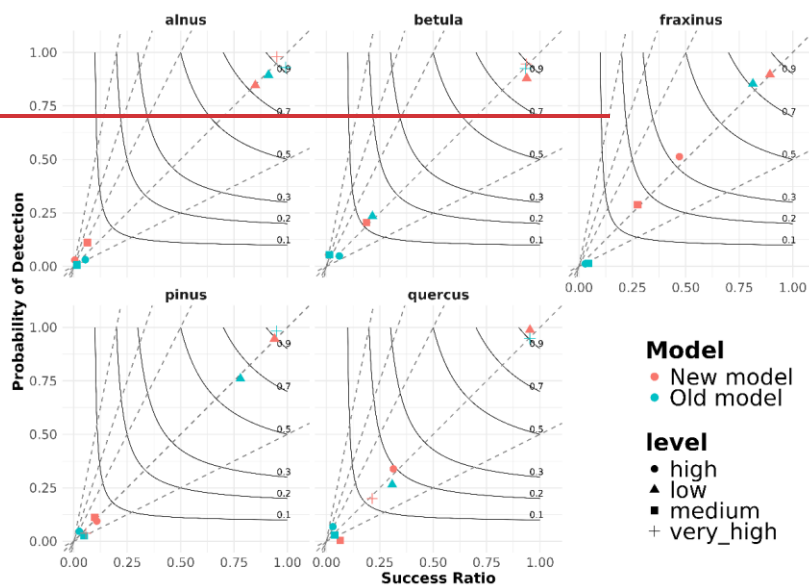
For *Pinus*, both models capture the general spring timing, but the new model aligns more closely with observations (2024: 18 Apr-17 May vs 15 Apr-16 May; 2025: 2 May-23 May vs 1 May-14 May), while the old model slightly underestimates the duration.

For *Quercus*, the new model accurately reproduces the observed periods (2024: 5 Apr-16 Apr vs 5 Apr-29 Apr; 2025: 15 Apr-29 Apr vs 14 Apr-26 Apr), whereas the old model

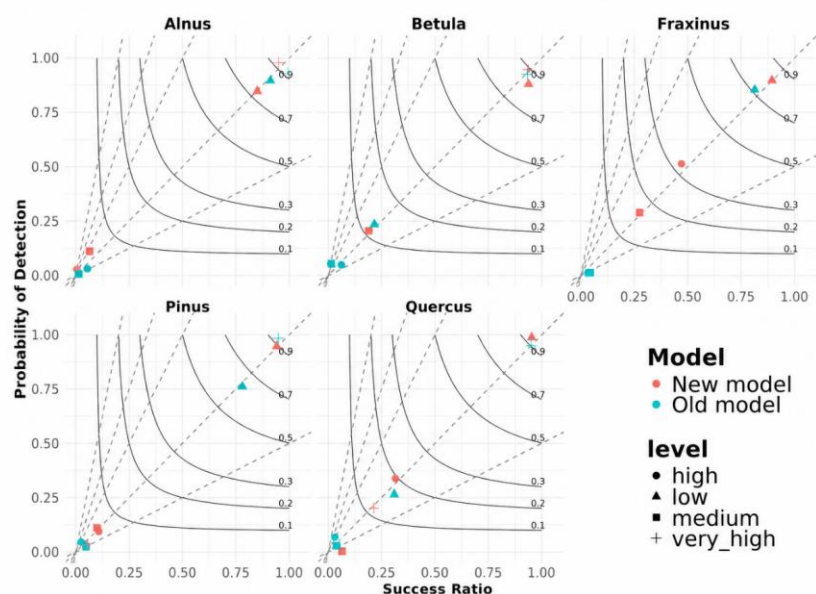
430 significantly extends the season (2024: 28 Mar-29 Aug; 2025: 5 Apr-12 May), indicating
431 overprediction of long-term pollen presence.

432 **3.3 Comparison of Models - Performance Diagram**

433 Model performance is evaluated using metrics such as accuracy, precision, recall and F1-
434 score. Performance diagrams and statistical analyses are provided to illustrate model
435 strengths and weaknesses.



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Figure 5. Performance of automatic ML models compared with Hirst reference data, shown as a performance diagram. The diagram is based on daily data from the 2024–2025 pollen season. The metrics include success ratio (SR), probability of detection (POD), false alarm ratio (FAR), and critical success index (CSI).

Figure 5. Performance of automatic ML models compared with Hirst reference data.

Detection of *Alnus* pollen remained challenging for both models at medium and high concentrations, resulting in low POD (<0.25) and success ratios below 0.2. The new model shows noticeable improvement at low concentrations (POD ≈ 0.75 , SR ≈ 0.7), indicating better sensitivity to early pollen occurrence, while both models perform perfectly at very high concentrations (POD ≈ 1.0 , SR ≈ 1.0), confirming reliability during peak events.

For *Betula*, the new model clearly outperforms the old one at low concentrations (POD ≈ 0.9 vs 0.6 ; SR ≈ 0.8 vs 0.6), while maintaining similar performance at higher levels (POD ≈ 1.0 , SR ≈ 1.0). Detection at medium levels remains limited for both models, though the new model achieves higher accuracy and fewer false alarms overall.

For *Fraxinus*, improvements are mainly visible at medium and high concentrations (old model: POD ≈ 0.1 , SR ≈ 0.1 ; new model: medium 0.3 , 0.3 and high 0.5 , 0.5), reflecting better seasonal detection capability. Very high concentrations are successfully detected

458 by both models ($POD \approx 1.0$, $SR \approx 1.0$), while at low concentrations the new model shows
459 moderate improvement over the old model.

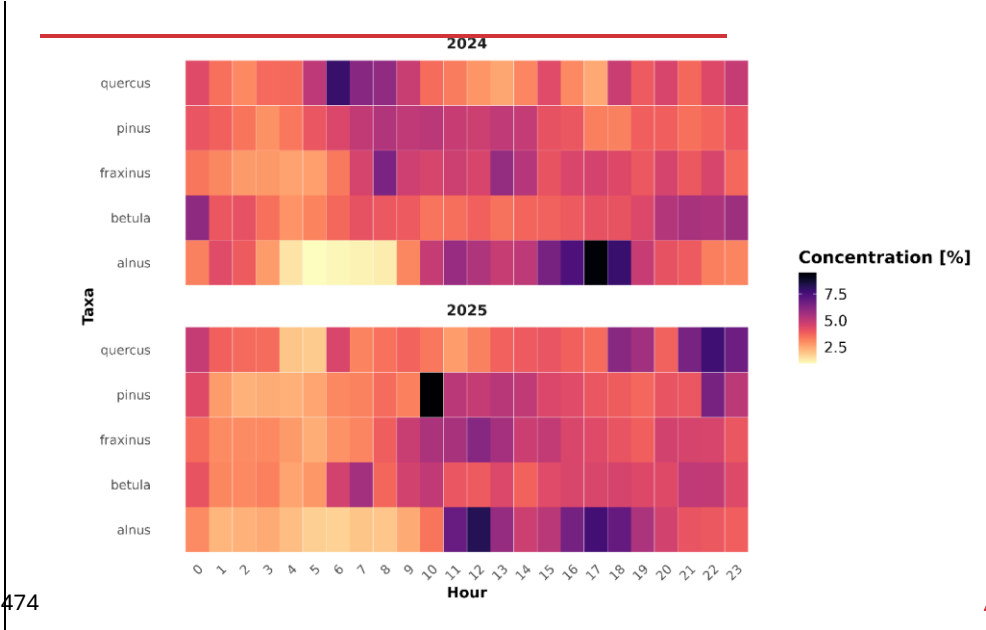
460 In the case of *Pinus*, the new model achieves strong detection performance at low
461 concentrations ($POD \approx 0.95$, $SR \approx 0.9$), outperforming the old model ($POD \approx 0.7$, $SR \approx 0.6$).
462 Both models detect very high concentrations perfectly ($POD \approx 1.0$, $SR \approx 1.0$), though
463 detection at medium and high levels remains less reliable.

464 For Quercus, the new model substantially improves detection at low concentrations (POD
465 ≈ 0.97 , $SR \approx 0.9$), outperforming the old model. Detection at medium concentrations
466 remains very low ($POD \approx 0.1$, $SR \approx 0.1$), while at high concentrations the new model
467 performs better than the old one. Very high concentrations are better detected by the old
468 model.

469

470 **3.4 Hourly Concentration Patterns over Two Years**

471 Hourly Swisens Polleno Jupiter (the new model) pollen concentration profiles for two
472 consecutive years are presented, showing diurnal trends, seasonal variations and year-
473 to-year differences.



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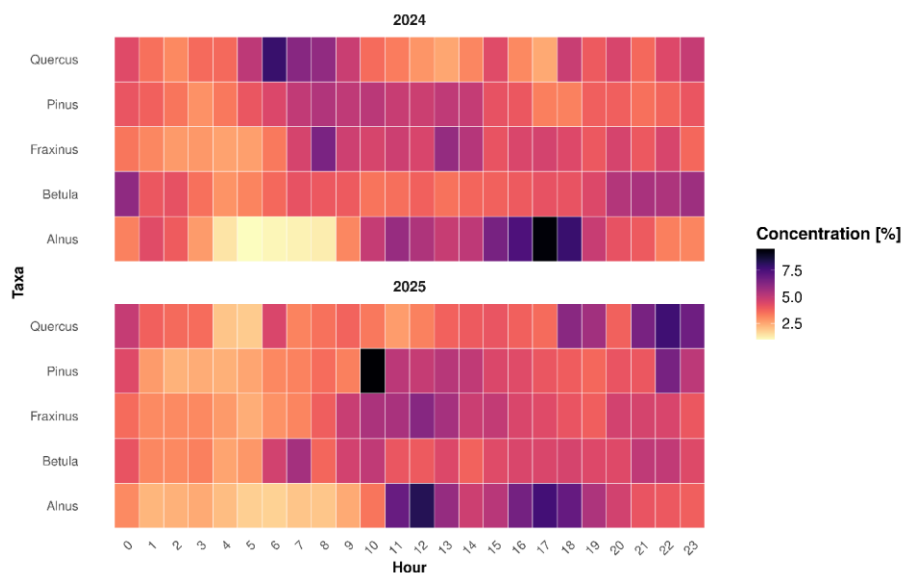


Figure 6. Hourly pollen profiles from the Swisens Poleno Jupiter for two consecutive years.

For *Alnus*, the highest contributions to the seasonal pollen load were consistently observed during the late afternoon hours, specifically between 16:00 and 18:00. In 2024, hourly contributions for 16:00, 17:00 and 18:00 reached 9.5%, 8% and 5% of the total seasonal load, while in 2025 they were slightly lower but still substantial, at 7.5%, 7% and 5%, respectively. This pattern reflects a typical daily trend, with the lowest pollen concentrations in the morning, higher levels around midday and a peak in the late afternoon, highlighting this period as critical for *Alnus* pollen dispersal and its overall impact on seasonal exposure.

Betula displayed a distinct temporal pattern, with peak pollen concentrations occurring in the evening between 20:00 and 00:00, as well as in the morning between 06:00 and 10:00. During these periods, hourly contributions ranged from 4% to 5% of the seasonal total, indicating that both morning and evening represent windows of elevated *Betula* pollen concentrations, with potential implications for allergy sufferers during these times.

For *Fraxinus*, pollen concentrations gradually increased in the morning, reaching a peak around midday (11:00–14:00), with hourly contributions ranging from 5% to 6%. Additional moderate peaks were observed in the late morning, early afternoon and evening, reflecting a more extended period of pollen availability compared to *Alnus* and

495 *Betula*. This pattern suggests that *Fraxinus* contributes to sustained exposure throughout
496 the central part of the day.

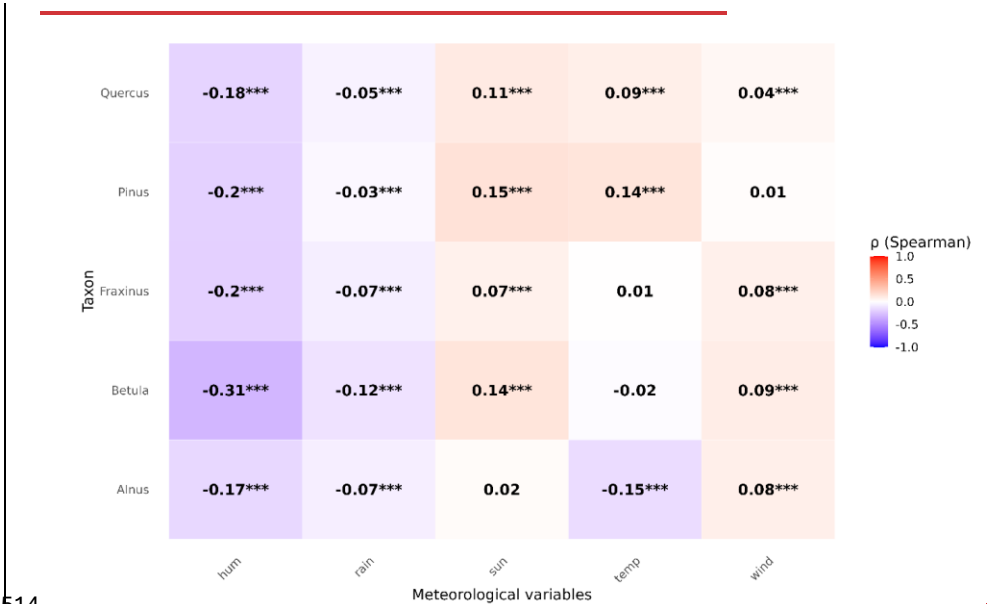
497 *Pinus* exhibited pronounced morning and midday peaks, with the highest hourly
498 contribution reaching 10% at 10:00 in 2025. Overall, pollen levels remained consistently
499 elevated throughout the day, ranging from 3% to 6%, with the lowest values observed
500 between 03:00 and 05:00. This extended period of substantial emission reflects both high
501 variability and prolonged daytime dispersal, making *Pinus* a major contributor to airborne
502 pollen over a broad temporal window.

503 *Quercus* exhibited a bimodal pattern of pollen concentrations, with morning peaks
504 occurring between 06:00 and 09:00 and a second, more pronounced peak in the evening
505 between 18:00 and 00:00. Hourly contributions during these periods reached up to 8% of
506 the seasonal load, indicating that both early morning and evening represent critical
507 windows for *Quercus* pollen exposure.

508

509 **3.5 Meteorological Parameters and Pollen Concentrations**

510 The Spearman correlation matrix (Fig. 7) provides an overview of the associations
511 between hourly meteorological parameters and pollen concentrations for multiple tree
512 taxa, thereby identifying the principal drivers of pollen variability and informing their
513 predictive potential.



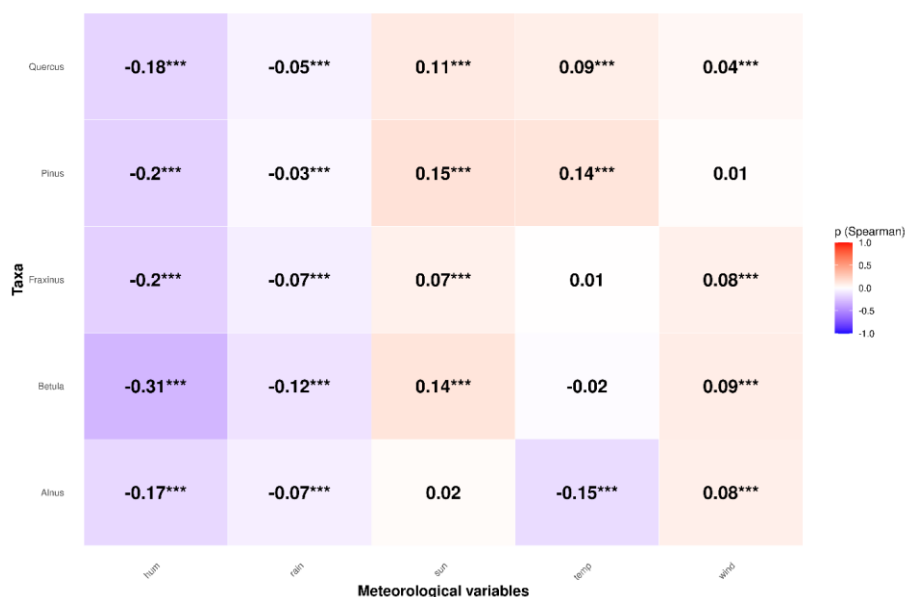


Figure 7. Spearman correlation matrix between meteorological variables and pollen concentrations. The correlation matrix is based on daily pollen season data (2024–2025) and corresponding meteorological data for the same period. Abbreviations: hum – relative humidity, rain – precipitation sum, sun – sunshine duration, temp – air temperature at 2 m, wind – wind speed.

Figure 7. Overall Spearman correlation matrix between meteorological variables and pollen concentrations.

In the case of *Alnus*, pollen concentration showed a clear negative correlation with relative humidity ($\rho = -0.17$, $p < 0.001$) and temperature ($\rho = -0.15$, $p < 0.001$). In contrast, the influence of wind and sunshine duration was relatively weak ($\rho = 0.08$ and 0.02 , respectively).

Betula exhibited the strongest negative correlation with relative humidity ($\rho = -0.12$, $p < 0.001$), whereas the effect of temperature was minimal ($\rho = -0.02$, ns) and solar radiation showed a moderately positive effect ($\rho = 0.14$, $p < 0.001$).

For *Fraxinus*, pollen concentration correlated moderately positively with sunshine duration ($\rho = 0.07$, $p < 0.001$) and wind ($\rho = 0.08$, $p < 0.001$), while humidity ($\rho = -0.07$, $p < 0.001$) and temperature ($\rho = -0.20$, $p < 0.001$) had weaker negative effects.

535 *Pinus* showed the highest positive correlations with sunshine duration ($\rho = 0.15$, $p < 0.001$)
536 and temperature ($\rho = 0.14$, $p < 0.001$), whereas humidity had a slight negative influence (ρ
537 $= -0.03$, $p < 0.001$) and wind was nearly neutral ($\rho = 0.01$, ns).

538 In the case of *Quercus*, pollen exhibited a moderate positive relationship with
539 temperature ($\rho = 0.09$, $p < 0.001$) and weaker positive correlations with sunshine duration
540 ($\rho = 0.11$, $p < 0.001$) and wind ($\rho = 0.04$, $p < 0.001$), while humidity showed a slight negative
541 effect ($\rho = -0.05$, $p < 0.001$).

542 The following analysis (Fig. 1S) focuses on the hourly correlations between these
543 variables, highlighting how their relationships change throughout the day.

544

545 Figure 1S. The hourly Spearman correlation matrix between meteorological variables and
546 *Alnus* pollen concentrations for the 2024–2025 season. hum – relative humidity, rain –
547 precipitation sum, sun – sunshine duration, temp – air temperature at 2 m, wind – wind
548 speed.

549 Figure 2S. The hourly Spearman correlation matrix between meteorological variables and
550 *Betula* pollen concentrations for the 2024–2025 season. hum – relative humidity, rain –
551 precipitation sum, sun – sunshine duration, temp – air temperature at 2 m, wind – wind
552 speed.

553 Figure 3S. The hourly Spearman correlation matrix between meteorological variables and
554 *Fraxinus* pollen concentrations for the 2024–2025 season. hum – relative humidity, rain –
555 precipitation sum, sun – sunshine duration, temp – air temperature at 2 m, wind – wind
556 speed.

557 Figure 4S. The hourly Spearman correlation matrix between meteorological variables and
558 *Pinus* pollen concentrations for the 2024–2025 season. hum – relative humidity, rain –
559 precipitation sum, sun – sunshine duration, temp – air temperature at 2 m, wind – wind
560 speed.

561 Figure 5S. The hourly Spearman correlation matrix between meteorological variables
562 and *Quercus* pollen concentrations for the 2024–2025 season. hum – relative humidity,
563 rain – precipitation sum, sun – sunshine duration, temp – air temperature at 2 m, wind –
564 wind speed.

565

566 ~~Figure 1S. Hourly Spearman correlation matrix between meteorological variables and~~
567 ~~*Alnus* pollen concentrations.~~

568 ~~Figure 2S. Hourly Spearman correlation matrix between meteorological variables and~~
569 ~~*Betula* pollen concentrations.~~

Figure 3S. Hourly Spearman correlation matrix between meteorological variables and *Fraxinus* pollen concentrations.

Figure 4S. Hourly Spearman correlation matrix between meteorological variables and *Pinus* pollen concentrations.

Figure 5S. Hourly Spearman correlation matrix between meteorological variables and *Quercus* pollen concentrations.

For *Alnus*, the Spearman results indicate a clear diurnal/seasonal pattern. The air temperature, in the pollen season, shows a strong positive correlation with pollen counts, particularly during midday-afternoon. Relative humidity is negatively correlated with pollen counts. The signal is the strongest in March, in early morning hours (around 08:00) and night/late-evening (19-21:00). When comparing February and March, temperature emerges as the dominant driver in February, while in March humidity is the meteorological variable most consistently associated with pollen concentrations. ~~while in March humidity becomes the most influential meteorological factor shaping pollen concentrations.~~

For *Betula*, Spearman correlations revealed strong positive associations between pollen concentrations and air temperature, particularly during daytime hours, with peak correlations reaching $r \approx 0.65-0.70$ ($p < 0.01$). Relative humidity was consistently negatively correlated ($r \approx -0.45$ to -0.60 , $p < 0.05-0.01$), while sunshine duration showed moderate positive effects in the morning. Rainfall effects were mostly non-significant. Seasonal differences indicated that in March, temperature and humidity effects were concentrated during the day, whereas in April, their influence extended more evenly into the evening and night, with wind becoming a more important factor.

For *Fraxinus*, air temperature positively influenced pollen concentrations, with correlations reaching $r \approx 0.60-0.68$ ($p < 0.01$), particularly during daytime hours in March and during the afternoon, evening and night in April. Relative humidity showed a negative relationship ($r \approx -0.40$ to -0.55 , $p < 0.05-0.01$), especially in the early and late hours of March and during the night in April. Sunshine duration, rainfall and wind effects were generally non-significant.

For *Pinus*, temperature effects were weaker compared to other taxa, showing moderate negative correlations during early morning and late-night hours in May ($r \approx -0.30$ to -0.40 , $p < 0.05$). Relative humidity consistently exhibited strong negative correlations ($r \approx -0.50$ to -0.65 , $p < 0.01$), particularly in May, while sunshine duration showed occasional positive associations during the afternoon. Wind and rainfall effects were negligible.

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605 For *Quercus*, pollen concentrations showed strong positive correlations with air
606 temperature throughout the entire day in April ($r \approx 0.60\text{--}0.72$, $p < 0.01$), while other
607 meteorological factors had no significant influence.

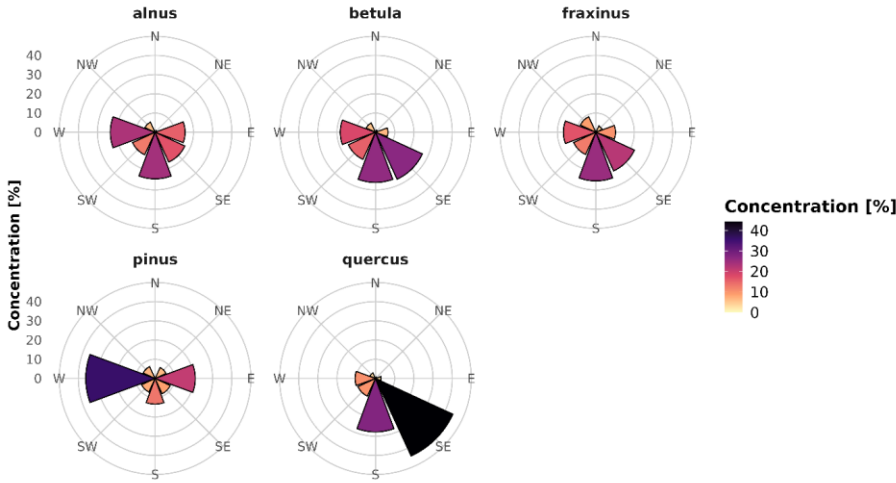
608
609 Table 2. Spearman’s correlations between hourly pollen concentrations and PBL height
610 in Wrocław (2024–2025).

Taxa	<i>Alnus</i>	<i>Betula</i>	<i>Fraxinus</i>	<i>Pinus</i>	<i>Quercus</i>
PBL	0.13 ^{***}	0.13 ^{***}	0.12 ^{***}	0.08 ^{***}	0.11 ^{***}

611
612 Spearman’s rank correlations between hourly pollen concentrations and planetary
613 boundary layer (PBL) height were positive for all taxa, indicating that higher PBL heights
614 are generally associated with increased pollen concentrations. The strongest correlations
615 were observed for *Betula* and *Alnus*, while *Pinus* showed the weakest relationship. All
616 correlations were statistically significant ($p < 0.001$), indicating a consistent link between
617 boundary layer dynamics and short-term variations in pollen concentrations and
618 highlighting the role of atmospheric mixing in pollen dispersion across different
619 *taxaspecies*.

620

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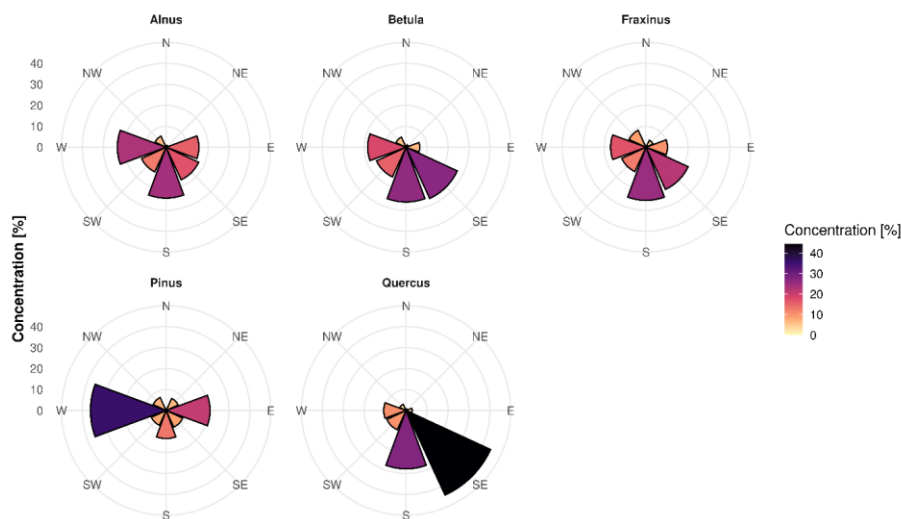


Figure 8. Influence of wind direction on pollen concentrations by taxa.

The relationship between pollen concentrations and and wind direction is presented in Fig. 8. For *Alnus*, *Betula* and *Fraxinus* the highest pollen concentrations are from the southern sector, which contributes about 25%. High contributions (20-25%) to total count is from the SE for *Betula* and *Fraxinus* and W for *Alnus*. The results are significantly different for *Pinus* and *Quercus*. *Pinus* pollen showed a clear west-east transport pattern, with the highest contributions from western (36.1%) and eastern (21%) winds, while *Quercus* pollen was mainly associated with southeastern (44.4%) and southern (27.8%) winds. Northern winds contributed minimally (<5%) to the dispersal of all taxa, with the lowest influence from NW, N and NE directions.

4. Discussion

Erb et al. (2025) discussed how to design measurement campaigns and mentioned that local data collection is **essential** for accurate model calibration. Our study confirms that the newly developed model, trained on locally collected pollen data, outperforms the model retrained on data from a different region. However, it is important to note that the new model was also trained using fluorescence information, which could

improve detection efficiency (Crouzy et al., 2025). The performance of the old model, which is based solely on holography, may vary over time depending on the presence of false signals, i.e. particles resembling specific pollen taxa, which can occur in the atmosphere and potentially interfere with the classification. New model more accurately capturing both the seasonal timing and the daily variability. In contrast, the previous model tends to overestimate concentration magnitudes and fails to represent temporal variations with sufficient precision.

The enhanced detection efficiency observed in this study aligns with the study of Maya-Manzano et al. (2023) for Munich (southern Germany), who reported comparable R^2 values for *Quercus* and higher values for *Fraxinus*, although lower for *Betula*. Their work also highlighted the weaker correspondence between real-time monitoring devices and Hirst-type measurements, underscoring the ongoing challenges in reconciling automated and traditional methods. While Hirst-type traps remain the gold standard in aerobiology, they have inherent limitations, such as low temporal resolution and dependence on manual counting. This raises important considerations when comparing real-time data with Hirst measurements (Tummon et al., 2024).

In our work, the locally trained model consistently outperformed the reference model, particularly at low and medium pollen concentrations, showing enhanced sensitivity and overall predictive accuracy. Both models achieved similar performance at high concentrations, indicating reliable detection under extreme pollen loads. This indicates that if the primary goal is to issue warnings for high pollen levels, models retrained using data from other regions may still perform adequately. However, the improved sensitivity of the locally retrained model at lower concentrations is particularly important for operational air-quality monitoring and early warning systems, as it allows for more timely identification of pollen events relevant to public health. Similar challenges related to underestimation at low concentrations were also reported by Tummon et al. (2024), further supporting the need for region-specific model retraining.

This study provides a detailed description of the measurement campaign and its methodology. The lowest amount of pollen was collected from *Fraxinus*, which was influenced by the low number of training events and consequently, *Fraxinus* exhibited the lowest R^2 among all taxa. This underscores the importance of properly extracting sufficient amounts of pollen from catkins. *Pinus* also showed a relatively low R^2 compared to *Alnus*, *Betula* and *Quercus*, which is likely due to interference from morphologically similar pollen grains, such as spruce or larch (Szczepanek et al., 2017). Therefore, one possible solution could be to aggregate data within the Pinaceae group.

Analysis of hourly pollen percentages revealed distinct diurnal patterns consistent with daily meteorological rhythms, as also reported by Chappuis et al. (2020).~~Analysis of hourly pollen percentages revealed distinct diurnal patterns consistent with daily meteorological rhythms.~~ *Alnus* peaked in the late afternoon (16:00–18:00) as the first

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major pollen-releasing taxon of the season. *Betula* and *Quercus* exhibited bimodal patterns, with morning and evening peaks. In contrast, *Fraxinus* and *Pinus* maintained relatively high pollen concentrations throughout the day. These results align with previous observations in Europe, including Ščevková et al. (2015) and Toth et al. (2011), who noted daytime peaks for *Fraxinus*, *Alnus*, *Corylus* and *Pinus*, as well as afternoon and night-time peaks for *Betula* and Cupressaceae-Taxaceae. Similar bimodal patterns have been reported by Dąbrowska-Zapart et al. (2020).

Our findings regarding evening peaks suggest that nocturnal maxima are influenced not only by local pollen release but also by the secondary deposition of previously emitted or transported pollen (Grewling et al., 2016; Ščevková et al., 2015). For instance, *Betula* and *Fraxinus* produce small pollen grains with low settling velocities, making them prone to long-distance transport and prolonged suspension within the atmospheric boundary layer. During daytime heating, thermal convection lifts pollen grains to higher altitudes; once convection weakens in the evening, these particles gradually descend, resulting in elevated nighttime concentrations (Clot, 2001; Kasprzyk et al., 2001; Kämpylä, 1984).

The local urban effects may also enhance nocturnal concentrations. The heating of anthropogenic surfaces (e.g., buildings and roads) during the day and subsequent nighttime radiative cooling can modify near-surface circulation, promoting weak turbulence and the resuspension of previously settled pollen grains. Such mechanisms may partly explain the persistence of elevated evening levels, particularly under calm and dry conditions. Given the low gravitational settling velocity of these taxa, deposition from the boundary layer (~1 km) may take more than 24 hours (Ščevková et al., 2015), further supporting the hypothesis that the observed evening and nocturnal peaks reflect both delayed sedimentation and localized resuspension.

Night-time increases were also noted in Kluska et al. (2020), who reported that low relative humidity and calm winds favor elevated concentrations of *Betula*, *Pinus* and Poaceae pollen at night. Resuspension of previously settled pollen appears to play a minor role in our dataset, as evening conditions, characterized by higher humidity, lower temperatures and moderate wind speeds, were generally unfavorable for significant re-entrainment, which is consistent with findings of Ščevková et al. (2015).

Meteorological factors were key drivers of hourly pollen variability. Our findings confirm the well-known pattern of increased pollen concentrations during warm and sunny days and decreased concentrations under high humidity or rainy conditions (Malkiewicz et al., 2016; Ščevková et al. 2015; Tomczyk et al. 2025; Ziska et al., 2019). However, rainfall was not always statistically significant, highlighting the complex nature of its influence on pollen concentrations, as some studies even report increases in concentrations after precipitation events (Kluska et al., 2020). The effect varies with rainfall intensity and type, as precipitation processes are associated with pollen wash-out. Kluska et al. (2020) showed that under low-intensity rainfall (<1.0–2.5 mm·h⁻¹), no decreases and even slight

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increases in pollen concentrations can occur. Interestingly, early-season pollen, such as that from *Alnus*, showed no significant correlation with sunlight and even a negative association with temperature. This contrasts with the findings of Borycka and Kasprzyk (2018), who reported that *Betula* did not exhibit significant correlations with meteorological variables, while *Alnus* showed strong relationships. In the present study, *Fraxinus* and *Betula* also displayed different patterns, with no consistent correlation with temperature, whereas later-season trees, such as *Pinus* and *Quercus*, exhibited positive associations with temperature. Wind was generally an important factor influencing pollen dispersal and increasing concentrations; however, this was not observed for *Pinus*. This is likely related to the morphological features of *Pinus* pollen, which has air sacs that enhance its dispersal, combined with the high pollen output of the species, making wind less critical for effective transport (Szczepanek et al., 2017).

Early-spring taxa, *Alnus* and *Betula*, displayed strong positive correlations with air temperature and negative correlations with relative humidity, particularly between 12:00 and 18:00, when concentrations typically increased. This pattern supports the hypothesis that warm and dry afternoon conditions promote pollen release, consistent with observations by Dąbrowska-Zapart et al. (2020) and Kluska et al. (2020), who also found that high temperature and solar radiation act as primary triggers of emission. For *Betula*, the most pronounced temperature effect occurred in the late afternoon and evening (14:00-22:00), coinciding with the timing of observed concentration peaks. However, it differs between months, which suggests that the phase of pollen release is likely correlated with meteorological conditions, so the early part of the pollination period in March shows different sensitivity compared to April. It is also well known that for early-flowering plant taxa, the timing of the pollen season can vary considerably from year to year (Malkiewicz et al., 2016). Interestingly, sunshine duration showed a relationship with both taxa only in March, mainly during the morning hours. Interestingly, sunshine duration was crucial only in March for both taxa and practically only during the morning hours.

Many studies emphasize the importance of temperature for the development of inflorescences; however, it is possible that pollen emission and flower opening are primarily triggered by sunshine duration. Favorable thermal conditions may then mainly support continued emission and the persistence of pollen in the air, which could explain the increased association during later hours which could explain why temperature becomes more influential during the later hours of the day (Malkiewicz et al., 2016; Nowosad et al., 2018; Picornell et al., 2019; Werner et al., 2021). Among all taxa, only *Betula* exhibited a correlation with wind around midday in April. Wind was previously identified as an important factor by Puc et al. (2015) and Puc (2006), whereas Tomczyk et al. (2025) reported that wind does not play a significant role. Nevertheless, it should be noted that their findings were based on daily data, while the present analysis is conducted on an hourly scale.

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Fraxinus exhibited a similar dependence on temperature and sunshine duration, with maximum correlations observed between 10:00 and 16:00. It is also evident that the daily pattern differs between months: in March, temperature was the most important factor during the day and evening, ~~while relative humidity showed higher correlation in the evening and night~~ ~~while relative humidity had a stronger effect in the evening and night~~. In April, temperature played a key role in the morning, evening and night, ~~whereas relative humidity showed the highest correlation at night~~ ~~whereas relative humidity was most influential at night~~. Kubik-Komar et al. (2018) also identified *Fraxinus* as a taxon whose pollen concentration is particularly related to temperature, noting that the timing of the season may vary depending on meteorological conditions during the first part of the year. Therefore, the observed correlations may also differ between months, depending on the prevailing weather conditions.

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Pinus showed negative correlations with temperature (morning and night) and with humidity (except in the morning), which is opposite to the results presented in the overall analysis (Fig. 7), where early-pollinating taxa showed negative correlations with temperature and *Pinus* showed positive correlations. This highlights the complex nature of the relationship between pollen concentration and meteorological conditions. It should be underlined that the aerodynamic properties of *Pinus* pollen allow it to remain airborne for a long time, which may explain the lack of correlation with current meteorological conditions, as this pollen could have been released earlier (Szczepanek et al., 2017).

Quercus, which flowers later in the season, was linked to temperature throughout the day. ~~*Quercus*, which flowers later in the season, was strongly responsive to temperature throughout the day~~, exhibiting a bimodal pollen release pattern with peaks in the morning (06:00–09:00) and evening (18:00–22:00), likely reflecting local meteorological control. Interestingly, temperature appeared to be important throughout the entire day only in April, probably because approximately 75% of the seasonal pollen amount occurs during the first two weeks of the season, as noted by Grewling et al. (2014).

The positive and statistically significant correlations between planetary boundary layer height and hourly pollen concentrations suggest that atmospheric mixing plays an important role in the short-term variability of pollen levels, as confirmed by Andújar-Maqueda et al. (2025). Stronger correlations observed for *Betula* and *Alnus* compared to *Pinus* may reflect differences in pollen release patterns and dispersal mechanisms. In particular, *Pinus* pollen, due to its aerodynamic shape and morphology, may be dispersed through mechanisms less directly linked to planetary boundary layer height. This is supported by the lack of a significant correlation with wind speed, ~~suggesting that other factors, such as local turbulence or canopy-level processes, could play a more important role in its short-term variability. It is reflected in planetary boundary layer (PBL) processes, where studies suggest that katabatic flows can enhance pollen concentrations at night~~.

while daytime convective turbulence promotes dispersion, contributing to variability near the surface. More generally, differences between pollen types arise from interacting factors such as source distribution, meteorological conditions, PBL structure, pollen properties, flowering phenology, and other environmental influences (Andújar-Maqueda et al., 2025), suggesting that other factors, such as local turbulence or canopy-level processes, could play a more important role in its short-term variability.

Wind direction was significantly associated with hourly pollen levels. Wind direction also played a significant role in shaping hourly pollen levels. The highest concentrations were typically linked to airflows from vegetated areas upwind of the monitoring site. This finding is consistent with Dąbrowska-Zapart et al. (2020), who also demonstrated that wind direction significantly influences short-term (hourly) variations in pollen and spore concentrations.

Overall, our results highlight the need to consider taxon-specific diurnal patterns, meteorological drivers and local environmental context when interpreting pollen measurements and developing predictive models. The combination of high temporal resolution monitoring and incorporation of local taxa into model training substantially improves the ability to capture real-time pollen fluctuations, peak exposure periods and species-specific variability, which are critical for accurate forecasting and public health applications.

6. Conclusion

Our results showed that retraining the neural network model with locally collected pollen data improves predictive accuracy for *Alnus*, *Betula*, *Fraxinus*, *Pinus* and *Quercus*, especially at low and medium pollen concentrations, compared to models trained on data from other regions. The retrained model correlates more closely with Hirst measurements in terms of the start and end of the season, as well as the concentration levels and can provide valuable information for pollen information system or pollen forecasting.

Based on two years of automatic pollen concentration measurements, we present diurnal pollen variability, which exhibits distinct patterns depending on the taxon. People with allergies should avoid pollen peak hours. These are:

- *Alnus*: late afternoon peak (16:00-18:00)
- *Betula*: evening peak (18:00-22:00)
- *Fraxinus*: midday peak (12:00-13:00), additional moderate afternoon peaks
- *Pinus*: high levels throughout the day, morning and midday peaks (10:00)

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- 834 • *Quercus*: bimodal pattern - morning (06:00-09:00) and evening (18:00-22:00)
835 peaks

836 Temperature and low relative humidity are the main drivers of pollen release, while
837 sunshine duration enhances emission for *Fraxinus*, *Pinus* and *Quercus*. Hourly analyses
838 reveal that the strength of correlations between pollen concentrations and
839 meteorological factors varies throughout the day and depends on the phase of the
840 flowering season or month.

841 The planetary boundary layer height has only a limited direct influence on hourly pollen
842 concentration variability, with particularly lower correlations observed for *Pinus*.

843 Wind direction affects pollen dispersal, with southern and southeastern winds
844 contributing most, highlighting the influence of local land cover and the surroundings of
845 the monitoring station.

846

847 **Code availability**

848 Code will be made available on request.

849

850 **Data availability**

851 ~~The training datasets are available at: link Data will be made available on request.~~

852

853 **Author contribution**

854 Tomczyk Szymon: Conceptualization, Methodology, Investigation, Formal analysis,
855 Writing - Original Draft, Writing – review and editing, Visualization.

856 Małgorzata Werner: Conceptualization, Supervision, Funding acquisition, Writing -
857 Review & Editing, Investigation.

858 Małgorzata Malkiewicz: Supervision, Writing - Review & Editing, Investigation.

859 Karol Bubel: Writing - Review & Editing, Investigation.

860

861 **Competing interests**

862 The authors declare that they have no conflict of interest.

863

864 **Acknowledgement**

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867

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871 Access, the authors have applied a CC-BY public copyright licence to any Author
872 Accepted Manuscript version arising from this submission.

873

874

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