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Integrated dataset of atmospheric bioaerosols over east Asia

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Abstract. Bioaerosols are one of the main types of aerosols originating from the Earth's biosphere and are widely found in the atmosphere. They possess both biological attributes and aerosol characteristics, thereby exerting significant influences on climate, the environment, ecosystems, and public health. However, their regional-scale distribution, influencing factors, climatic and environmental impacts remain unclear due to the scarcity of observational data. This study firstly establishes an integrated bioaerosol dataset based on a large-scale dust–bioaerosol field campaign conducted across East Asia using unified sampling and analytical methods. The dataset systematically integrates atmospheric bioaerosol number concentrations and bacterial community structure at multiple taxonomic levels across 45 sites in China, Japan, South Korea, and Mongolia. In addition, meteorological variables (e.g., air temperature, relative humidity, wind speed and direction), air quality parameters (e.g., PM_{10} and $PM_{2.5}$), and Normalized Difference Vegetation Index (NDVI) data during the sampling period were incorporated from multiple sources. Further analysis of this integrated dataset indicates that bioaerosol number concentrations are negatively correlated with local NDVI. Moreover, there is a clear relationship between bioaerosol number concentration and air temperature, with a peak observed at approximately 10–15 °C. A pronounced diurnal variation in bioaerosol concentrations is also found, which is strongly associated with Aerosol Optical Depth (AOD) and particulate matter concentrations. In addition, substantial differences in community structure were observed across different underlying surface types, and the α-diversity indices (Shannon and Chao1 indices) were negatively correlated with NDVI. This dataset provides a robust foundation for advancing research on atmospheric bioaerosol processes, as well as their implications for climate, the environment, public health, and interdisciplinary studies. The dataset generated in this study is openly available via Zenodo (https://zenodo.org/records/19605145)(Huang et al., 2026).

1 Introduction

Bioaerosols are defined as living aerosol particles, biologically active components, and metabolic products of organisms, with particle diameters ranging from 0.001 to 100 μm (Fröhlich-Nowoisky et al., 2016; Després et al., 2012). For coarse-mode particles with diameters greater than 1 μm, these particles typically account for approximately 30% of both number and mass

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concentrations in urban and rural atmospheres, and their contribution can reach up to 80% in pristine rainforest regions (Fröhlich-Nowoisky et al., 2016). Bioaerosols are transported through the air, due to their small size and low density, can be readily dispersed across different environments (Van Leuken et al., 2016). Because they exhibit both physical and biological properties, they play important roles in public health, climate processes, and ecosystem functioning (Du et al., 2018; Huang et al., 2024a).

Most bioaerosols fall within the respirable size range (Bulski, 2020; Estillore et al., 2016), which allows their entry into the human body via the respiratory tract or penetrate compromised skin and mucous membranes, thereby posing potential health risks (Bulski, 2020). Deposition in different depths within the respiratory system may induce allergic or toxic responses in both humans and animals (Hofmann, 2011; Eriksen et al., 2023). Bioaerosols also play critical roles in modulating climate change. As airborne particulate matter, they absorb and scatter solar and terrestrial longwave radiation, thereby exerting direct regional and global radiative forcing (Huang et al., 2024a). Furthermore, bioaerosols can act as nuclei for cloud condensation, ice crystals, and precipitation, thereby influencing hydrological cycles and climate systems (Fröhlich-Nowoisky et al., 2016; Huang et al., 2025). A unique property of airborne biological components is their ability to produce ice-nucleating proteins (INPs), which can promote ice developing when subzero temperatures are relatively high (Hoose and Möhler, 2012). These biological INPs are active above −15 °C, whereas mineral dust particles typically are not (Morris et al., 2013). In addition, atmospheric microorganisms can influence the photochemical and chemical reactions of aerosols, thereby altering their composition (Péguilhan et al., 2025), with consequent feedbacks on climate processes (Meinander et al., 2022). Both viable and inactivated microbial cells exhibit comparable oxidative potential and may even enhance the oxidizing power of chemical species in the atmosphere (Samake et al., 2017).

Bioaerosol sampling has been conducted worldwide, yielding data on their distribution, composition, sources, and atmospheric roles, thereby advancing our understanding of their behavior and influencing factors (Huang et al., 2024b; Zhang et al., 2022b; Petersson Sjögren et al., 2023; Jabeen et al., 2023). However, existing studies lack standardized sampling and analytical protocols. Even when operated simultaneously, different sampling instruments may produce varying bioaerosol signals and exhibit differences in collection efficiency for different types of microorganisms (Mainelis, 2019; Mescioglu et al., 2021; Mbareche et al., 2018). The pore size and material of filter membranes can influence results of airborne microbiome studies (Guo et al., 2024; Jeong and Kim, 2021). Differences in sampling flow rate can also affect results, as high flow rates increase the risk of adverse effects (Boifot et al., 2024). Following collection, temperature conditions during sample transport and short- or long-term storage, can influence DNA degradation and alter sample composition (Clasen et al., 2020). These factors collectively reduce sample comparability and affect downstream analyses. During molecular analysis, significant differences in taxonomic richness, community composition, and diversity may arise from the selection of different 16S rRNA variable regions (Larsen et al., 2015; Lin and Ju, 2023). The application of different reference databases for taxonomic classification can also yield divergent results due to inherent database discrepancies (Balvočiūtė and Huson, 2017; Ramakodi, 2022). Together, these factors complicate cross-study comparisons. Although many studies aim to link environmental, meteorological, and chemical factors with bioaerosols, variations in sampling and analytical methods may lead to contradictory conclusions

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and hinder the development of robust findings, as it is often unclear whether discrepancies arise from environmental variability or methodological heterogeneity (Sajjad et al., 2023; Mainelis, 2019; Pogner et al., 2024).

At present, the spatial and temporal coverage of bioaerosol sampling remains sparse (Zawadowicz et al., 2019). Many observations are restricted to isolated field campaigns or single-site stations, with particularly limited coverage in arid regions and developing countries (Shammi et al., 2021). In addition, long-term continuous records are scarce, and standardized time series spanning multiple years and sites are largely lacking. This limitation not only constrains the detection of long-term trends and interannual variability but also hampers the use of observations for model calibration and the assessment of climate-scale impacts (Huang et al., 2024a; Safatov et al., 2022). Furthermore, methodological heterogeneity across studies complicates data integration and limits large-scale analyses (Alfaro-Perez et al., 2025). Given the highly dynamic nature of the atmosphere, bioaerosol concentrations exhibit substantial spatial and temporal variability (Šantl-Temkiv et al., 2019), and conclusions derived from studies with limited spatial or temporal coverage may therefore be inconsistent (Jiang et al., 2022). Together, these factors further reduce the robustness of integrated assessments of bioaerosol impacts on climate, the environment, ecosystems, and human health (Gashi et al., 2025).

To accurately quantify bioaerosol-related effects, there is an urgent need to expand observational network coverage, establish long-term continuous monitoring records, and advance methodological standardization to enable regional and transregional data integration and sharing (Huang et al., 2024a). Therefore, establishing a bioaerosol dataset with broad geographic coverage and unified sampling and analytical methodologies is of great importance for advancing scientific research and applications (Sajjad et al., 2023; Gashi et al., 2025; Boifot et al., 2024; Jiang et al., 2022). Such a dataset would enable improved characterization of the geographic distribution, spatiotemporal variability, and environmental and health impacts of bioaerosols. Under ongoing climate change, drought conditions are projected to intensify (Araujo et al., 2025; Xu et al., 2023), with East Asia expected to be strongly affected (Qi et al., 2024). Dust generated during sandstorm can promote the long-range transport of bioaerosols (Maki et al., 2019; Zhou et al., 2024). Considering the extensive arid and semi-arid regions of East Asia and the large populations residing downwind (Yu et al., 2023; Jin et al., 2022), the potential risks associated with bioaerosols in this region are relatively high. Establishing such a dataset will support improved assessment of public health risks under climate change and provide a scientific basis for mitigation and policy development.

2 Sampling sites and methods

2.1 Sampling sites

The DuBi (Dust–Bioaerosol) field sampling campaign was conducted over an extended period and across a broad spatial domain. From 2011 to 2021, sampling was performed at multiple sites across East Asia, including China, Japan, South Korea, and Mongolia (see Table 1 for details). These sites encompass a wide range of surface types and climatic conditions, providing strong regional representativeness and ensuring comparability among observations (Fig. 1).

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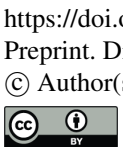

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This field observation campaign substantially enhanced the temporal and spatial coverage of bioaerosol measurements in East Asia. As a result, the compiled dataset enables a more comprehensive characterization of bioaerosol distributions across diverse environmental settings, thereby improving our understanding of the regional features and evolution patterns of bioaerosols in East Asia.

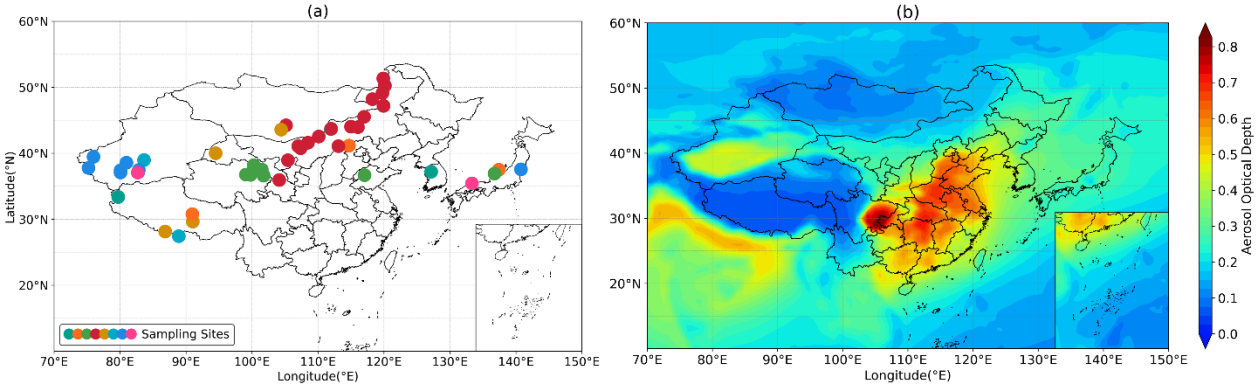

Figure 1: (a) Spatial distribution of sampling sites from 2011 to 2021; (b) spatial distribution of the mean aerosol optical depth (AOD) from 2000 to 2020.

Table 1: Summary of land surface characteristics, number of samples, locations, sampling period (local standard time, LST), and sampling duration throughout the DuBi field campaign conducted between 2011 and 2021.

Sampling Site	Land Surface	number of samples	Latitude & Longitude	Sampling Period	Sampling Duration
Dunhuang	-	13	（94.5° E, 40.1° N）	2011/9/9-2011/9/10 2021/7/11-2021/7/19	1-12h
Kanazawa	-	47	（136.7° E, 36.5° N）	2011/3/22-2011/5/1 2017/4/21-2017/5/15 2017/9/5-2017/9/28	6-15h
Hakui	-	2	（136.8° E, 36.9° N）	2013/4/28 2014/3/28	1h
Tsogt-Ovoo	-	14	（105.2° E, 44.2° N）	2014/3/16-2014/3/18 2017/4/29-2017/5/4	2-18h
Namie	-	59	（140.8° E, 37.6° N）	2015/8/19-2018/3/22	-
Erlianhot	Sandy land	111	（111.9° E, 43.7° N）	2016/3/30-2016/5/25 2017/8/17-2017/8/19	2-24h
Zhangbei	Sparse grassland	140	（114.7° E, 41.2° N）	2016/3/29-2016/5/31	2-15h
Jinan	Urban	121	（117.1° E, 36.7° N）	2016/3/23-2016/6/4	7-34h

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Dalanzadgad	Road	5	（104.4° E, 43.6° N）	2017/5/1-2017/5/3	-
Bayannur	Urban	2	（107.4° E, 40.8° N）	2017/8/4	3-6h
Ulanqab	Sparse grassland	2	（113.1° E,41.1° N）	2017/8/4-2017/8/5	3-8h
Xilinhot	Urban	1	（116.1° E,44.0° N）	2017/8/5-2017/8/6	8h
Arxan	Urban	4	（119.9° E,47.2° N）	2017/8/6-2017/8/7 2017/8/14	3-9h
Ewenki Autonomous Banner	Urban	7	（119.8° E,49.1° N）	2017/8/7-2017/8/8	4-11h
Ergun	Urban	3	（119.9° E,51.3° N）	2017/8/9-2017/8/10	3-9h
Labudalin	Wetland	2	（120.1° E,50.2° N）	2017/8/10-2017/8/11	4-10h
New Barag Left Banner	Sparse grassland	3	（118.3° E,48.2° N）	2017/8/13-2017/8/14	4-9h
East Ujimqin Banner	Sparse grassland	2	（117.0° E,45.5° N）	2017/8/15-2017/8/16	4-9h
Abag Banner	Sparse grassland	2	（115.0° E,44.0° N）	2017/8/16-2017/8/17	4-9h
Baotou	Sparse grassland	2	（110.1° E,42.5° N）	2017/8/19-20174/8/20	4-11h
Urad Middle Banner	Sparse grassland	2	（108.5° E,41.6° N）	2017/8/20-2017/8/21	4-9h
Urad Rear Banner	River dam	2	（107.0° E,41.1° N）	2017/8/21-2017/8/22	4-11h
Alxa Left Banner	Grassland	2	（105.5° E,38.9° N）	2017/8/22-2017/8/23	4-10h
Yongin	-	68	（127.16° E,37.2° N）	2018/3/9-2018/7/4	-
Lhasa	-	11	（91.0° E,29.6° N）	2018/5/7-2018/5/8 2018/5/27-2018/5/31	9-13h
Everest Base Camp, Xigaze	-	28	（86.9° E,28.1° N）	2018/5/11-2018/5/24	7-14h
Tazhong	Desert	40	（83.7° E,39.0° N）	2019/7/7-2019/7/30	9-22h
Minfeng	Grassland	31	（82.7° E,37.1° N）	2019/7/11-2019/7/31	11-12h
Tazhong–Minfeng route	Desert Cropland Desert	8	（82.9° E,37.5° N） （80.1° E,37.1° N） （80.1° E,37.4° N）	2019/7/23-2019/8/1	2-3h

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	Desert Grassland Grassland Grassland		（81° E,38.6° N） （75.2° E,37.8° N） （76° E,39.5° N） （82.8° E,37.1° N）		
Ngari		50	（79.7° E,33.4° N）	2019/7/7-2019/7/31	11-12h
Namtso	Alpine meadow	80	（91.0° E,30.8° N）	2020/8/3-2020/8/24	9-13h
Qinghai	Urban Alpine meadow Sparse grassland Urban Cropland and urban	8	（101.8° E,36.6° N） （99.8° E,36.7° N） （99.1° E,36.8° N） （100.1° E,37.3° N） （100.3° E,38.2° N） （101.5° E,37.5° N）	2020/7/16-2020/7/20	2-12h
Yuzhong	Cropland Grassland	62	（104.1° E,36.0° N）	2020/10/31-2020/11/19	2-6h
Yadong	Forest	141	（88.9° E,27.4° N）	2021/6/22-2021/8/30	11-13h

2.2 Sampling methods

A custom-designed bioaerosol sampler was employed for field collection to ensure operational stability and methodological consistency across sites. Instruments of this type have been widely used in bioaerosol research (Maki et al., 2010; Qi et al., 2023). To construct the sampler, a 13-mm polycarbonate membrane with 0.2 μm pore size (Whatman 111106 and GTTP01300) was inserted inside a 13-mm Swinnex filter holder (Millipore SX0001300) fitted. Polycarbonate membranes offer advantages such as high chemical resistance, high thermal stability, and a low tendency to adsorb proteins or extractable substances (Ferguson et al., 2019). Their optical transparency also enables accurate microscopic detection. A micro air pump (AS ONE MAS-1, Japan) with a standard flow rate of 12 L min^{-1} was connected to the sampler to draw atmospheric microorganisms onto the filter membrane (Fig. 2a, b).

Prior to sampling, both the filters and filter holders were sterilized. The assemblies were first rinsed with 75% ethanol, followed by ultrapure water to remove residual detergent. Subsequent processing of the filters and holders was conducted in a laminar-flow hood, where they were immersed in absolute ethanol and ultrapure water to further remove adhered organic residues. The dried filters and holders were then assembled and sterilized by autoclaving at 121 °C for 20 min, followed by ultraviolet irradiation for 30 min (Xue et al., 2024; Qi et al., 2021). The sterilized components were sealed in pre-sterilized collection tubes and sterile bags until use. During sampling, operators wore masks and sterile gloves, and the samplers were mounted at

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a height of 1.5 m above ground level. Real-time flowmeter readings obtained immediately before the micro-pump was turned on and immediately before it was turned off were recorded as the initial and final sampling flow rates, respectively. After sampling, filters were labeled with the sampling location and time, stored at −20 °C during transport, and subsequently transferred to −80 °C freezers for long-term preservation.

2.3 Fluorescence microscopy analysis

Bioaerosol particle concentrations were determined using DAPI staining (4,6-diamidino-2-phenylindole; D9542, Sigma), followed by fluorescence microscopy. 4% paraformaldehyde (250 μL) was introduced to each filter holder containing the membrane to fix the samples for 1–2 h. The filters were then rinsed with sterile ultrapure water to remove residual fixative. Under light-protected conditions, DAPI solution (10 μg mL^{-1}, 250 μL) was introduced to stain the samples for approximately 15 min (Maki et al., 2014), followed by a final rinse with sterile water.

During slide preparation, the membrane was treated with non-fluorescent immersion oil (IMMOIL-F30CC, Olympus) for wetting. The coverslip was then pressed firmly, and excess immersion oil was removed using a dust-free wipe. Samples were examined using a fluorescence microscope (BX53 and DP72, Olympus) with an excitation wavelength range of 340–390 nm A fluorescence microscope (BX53 and DP72, Olympus) providing a 340–390 nm excitation wavelength range was employed to examine the samples. (Tang et al., 2018).

As shown in Fig. 2e–g, four categories of particles were distinguishable under the fluorescence microscope based on their emitted colors: blue, yellow, white, and black. DAPI-stained microbial cells produced strong blue or blue-green fluorescence, whereas mineral particles appeared white (Maki et al., 2013). Black particles were attributed to black carbon (Liu et al., 2023). Yellow particles were generally interpreted as non-DNA-containing particles (Kepner and Pratt, 1994). However, other studies have shown that water-soluble salts (Liu et al., 2023) and organic matter (Mostajir et al., 1995) can also exhibit yellow fluorescence. In this study, most yellow-green particles were associated with mineral dust and were therefore considered to be mineral particles enriched in water-soluble salts and organic material. In this study, bioaerosols were defined as particles exhibiting cellular morphology and emitting blue-green fluorescence after DAPI staining. For each filter membrane, 20 randomly selected microscopic fields were imaged, and all particles belonging to the four categories were manually counted using the Cell Counter plugin in ImageJ.

Bioaerosol number concentrations, C (particles m^{-3}) were calculated using Eq. (1):

$$C = \frac{S_1 \times N_0}{S_0 \times V}, \quad (1)$$

where S_1 is the filtration area of the filter membrane (μm²), S_0 is the field-of-view area of the microscope (μm²), V is the sampled air volume (m^3), and N_0 is the average number of microbial particles observed per microscopic field of view (Maki et al., 2019).

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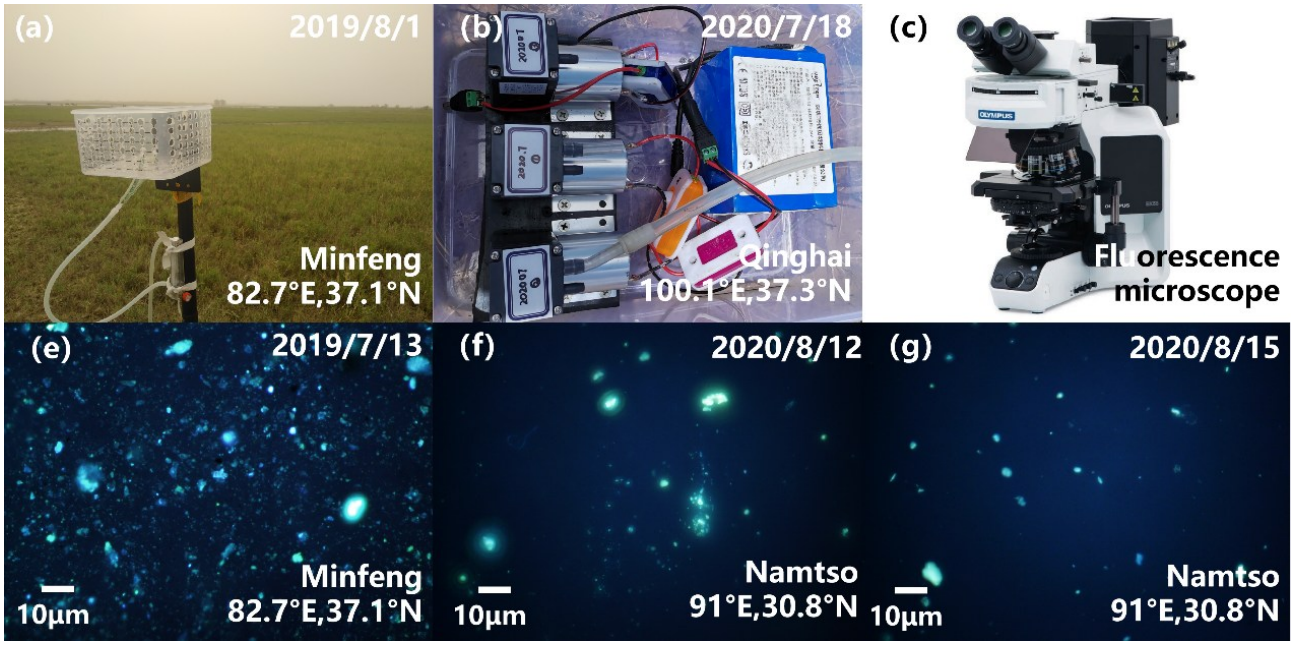

Figure 2: (a) Bioaerosols sampler used during the DuBi field sampling campaign; (b) power supply and sampling pump; (c) fluorescence microscope; (e–g) fluorescence micrographs of samples after DAPI staining.

2.4 16S rRNA gene sequencing

The 16S rRNA gene has been widely used to characterize microbial community composition and diversity (Giles et al., 2023; Nagarajan et al., 2023). The UltraClean Soil DNA Kit (MoBio, San Diego, CA) was employed to extract bacterial DNA in this study. Sequencing was performed on Illumina NovaSeq 6000 and Illumina MiSeq platforms in PE250, PE150, or PE300 modes. The bacterial 16S rRNA gene V4–V5 hypervariable region was amplified using the 515F/907R primer pair (forward: GTGCCAGCMGCCGCGGTAA; reverse: CCGTCAATTCMTTTRAGTTT) (Fang et al., 2017). Paired-end sequencing produced reads exceeding 250 bp for each sample, which were subsequently sorted according to their unique barcode sequences. Blank controls were processed alongside the samples during both DNA extraction and PCR amplification. No PCR amplicons were detected in these controls, demonstrating that the entire experimental workflow was free from detectable contamination (Qi et al., 2023; Karstens et al., 2019).

The Quantitative Insights into Microbial Ecology pipeline (QIIME2; version 2024.2) was used to assign taxonomic identities to the raw sequences (Bolyen et al., 2019). High-quality sequences were denoised and resolved into amplicon sequence variants (ASVs) at single-nucleotide resolution. Taxonomic classification was assigned using the SILVA v138 reference database (Pin et al., 2021). ASVs classified as chloroplasts, mitochondria, archaea, eukaryotes, or unassigned were removed prior to downstream ecological analyses.

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In this study, the Shannon and Chao1 indices were used to quantify the alpha diversity of microbial communities. These indices were calculated using the vegan package in R (v4.4.4). Differences in alpha diversity among surface types were assessed using the Wilcoxon rank-sum test to evaluate statistical significance (Loos et al., 2024).

2.5 Meteorological and air quality Data

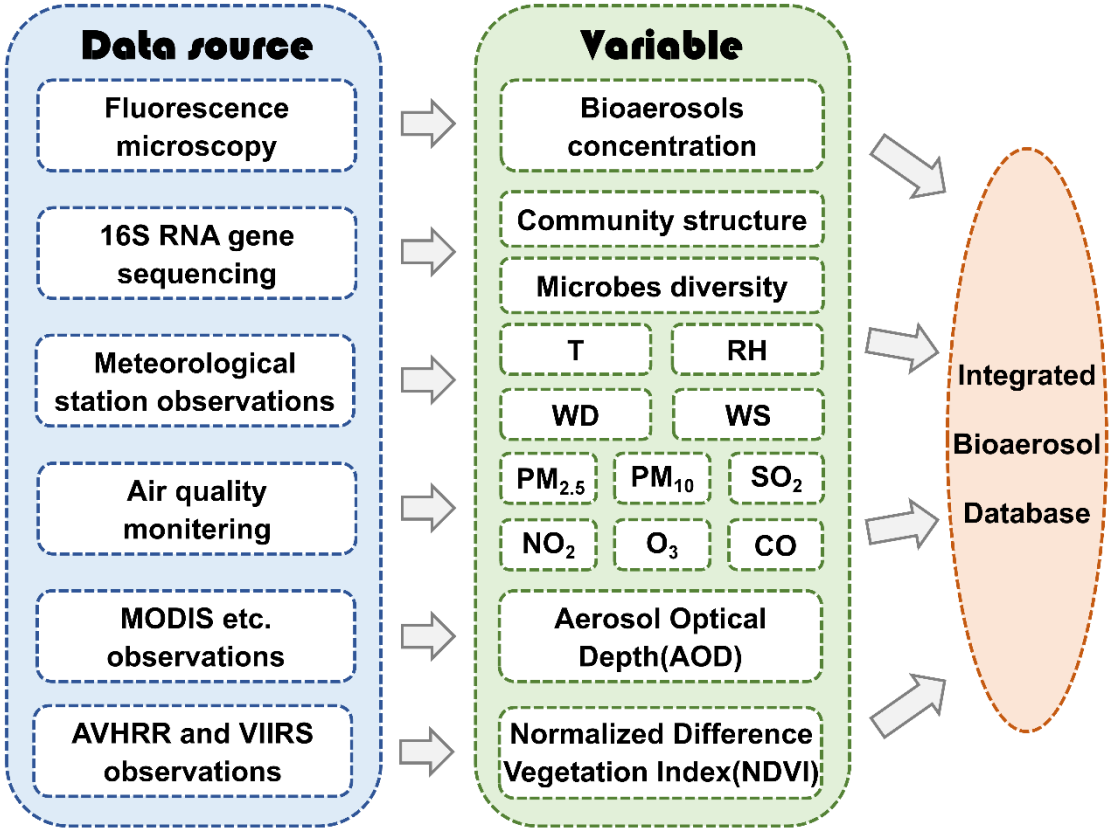

Figure 3: Schematic overview of the integrated database construction workflow and data sources.

The integrated dataset compiled in this study includes the following data products (Fig. 3):

Meteorological variables, such as air temperature, dew point, wind speed, wind direction, were obtained from surface meteorological observation stations. Because relative humidity (RH) is not directly reported at these stations, RH was derived using the Magnus formula based on temperature and dew-point measurements. (China stations: https://www.ncei.noaa.gov/pub/data/noaa/isd-lite/； stations outside China: https://www.ncei.noaa.gov/data/global-summary-of-the-day/access/)；

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Air-quality data were obtained from automatic monitoring networks. For China, mass concentrations were retrieved from the China National Urban Air Quality Real-Time Publishing Platform (https://air.cnemc.cn:18007). For Japan and South Korea, data were obtained from https://soramame.env.go.jp and https://www.airkorea.or.kr/;

AOD data were obtained from two satellite-derived products. The M2IMNXGAS dataset, based on MODIS and AVHRR retrievals, provides monthly AOD at a spatial resolution of 0.5° × 0.625°(Gelaro et al., 2017) (https://disc.gsfc.nasa.gov/datasets/M2IMNXGAS_5.12.4/summary?keywords=M2IMNXGAS_5.12.4). The MAIAC Land AOD Level-2 product (MCD19A2, Version 6.1) provides daily aerosol optical depth (AOD) data at a spatial resolution of 1 km × 1 km and is generated by applying the Multi-angle Implementation of Atmospheric Correction algorithm to observations acquired from the MODIS Terra and Aqua sensors. (https://www.earthdata.nasa.gov/data/catalog/lpcloud-mcd19a2-061);

NDVI data were obtained from the CDR product generated from AVHRR, VIIRS, and related sensors. Data are provided on a 0.05° × 0.05° spatial grid with daily temporal coverage. (Vermote and Program, 2019) (https://www.ncei.noaa.gov/data/land-normalized-difference-vegetation-index/access/).

3 Results and discussion

3.1 Correlation of bioaerosol concentrations and NDVI & air temperature

We analysed the relationships between bioaerosol concentration, NDVI, and air temperature across East Asia at a large spatial scale under non-special weather conditions (e.g., dust events). As shown in Fig. 4a, bioaerosol concentrations exhibit a significant negative correlation with NDVI. In regions with low NDVI, sparse vegetation cover and dry, loose soils facilitate the entrainment and emission of microorganisms into the atmosphere under wind forcing (Qi et al., 2023). In contrast, regions with high NDVI are characterized by stronger vegetation-mediated particle interception, which enhances dry deposition and consequently leads to lower atmospheric microbial concentrations (Zhai et al., 2022).

Figure 4b indicates that the relationship between temperature and bioaerosol concentration is non-linear. When temperatures are below approximately 15 °C, bioaerosol concentrations show a positive correlation with temperature. However, as temperature increases further, microbial concentrations decline markedly. A possible explanation is that, within an optimal temperature range, warming enhances atmospheric mixing and air movement, thereby promoting the dispersal of microorganisms into the atmosphere and resulting in higher concentrations (Cavicchioli et al., 2019). At excessively high temperatures, however, conditions become unfavorable for microbial growth and reproduction, and the abundance of microorganisms from certain sources may decrease accordingly (Zhang et al., 2022b).

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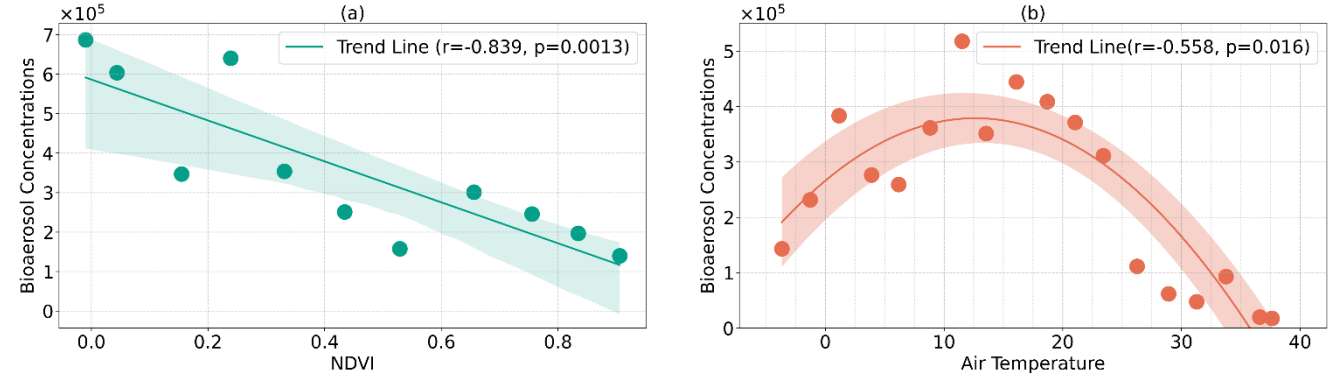

Figure 4: (a) Relationship between bioaerosol concentration and NDVI during the DuBi field sampling campaign. The green points represent mean bioaerosol concentrations calculated by binning NDVI into intervals of 0.1 and averaging all samples within each interval. (b) Relationship between bioaerosol concentration and air temperature. The orange points represent mean bioaerosol concentrations calculated by binning temperature into intervals of 2.5 °C and averaging all samples within each interval. The shaded areas indicate the 95 % confidence intervals.

3.2 Diurnal variation of bioaerosol concentration

Figure 5 shows the temporal variations in bioaerosol concentrations in Yuzhong, China, on 6 and 15 November 2020. As illustrated, bioaerosol concentrations on both days exhibit pronounced diurnal patterns rather than random fluctuations. Concentrations are lowest in the early morning (07:00–09:00), followed by a gradual increase, reaching a peak in the afternoon (15:00–17:00), and then decreasing thereafter. This pattern suggests that bioaerosol concentrations are governed by well-defined diurnal processes.

The concentrations of NO_2 and CO also display clear temporal variations with similar overall trends, suggesting that these two trace gases may share common sources. However, the temporal variations of NO_2 and CO on the two sampling days are not fully consistent with those of bioaerosol concentrations. This discrepancy implies that short-term variations in bioaerosol concentrations during the sampling period were not primarily controlled by anthropogenic emissions or changes in the atmospheric boundary layer height (Li et al., 2022). Instead, they are more likely associated with surface biological emissions and atmospheric dynamical processes. These processes include daytime increases in surface temperature that enhance microbial activity, intensified release from soil and vegetation surfaces, and strengthened surface turbulence that facilitates the entrainment of particles into the atmosphere (Gashi et al., 2025; Sharma et al., 2022; Zhang et al., 2022a; Kulmala et al., 2023).

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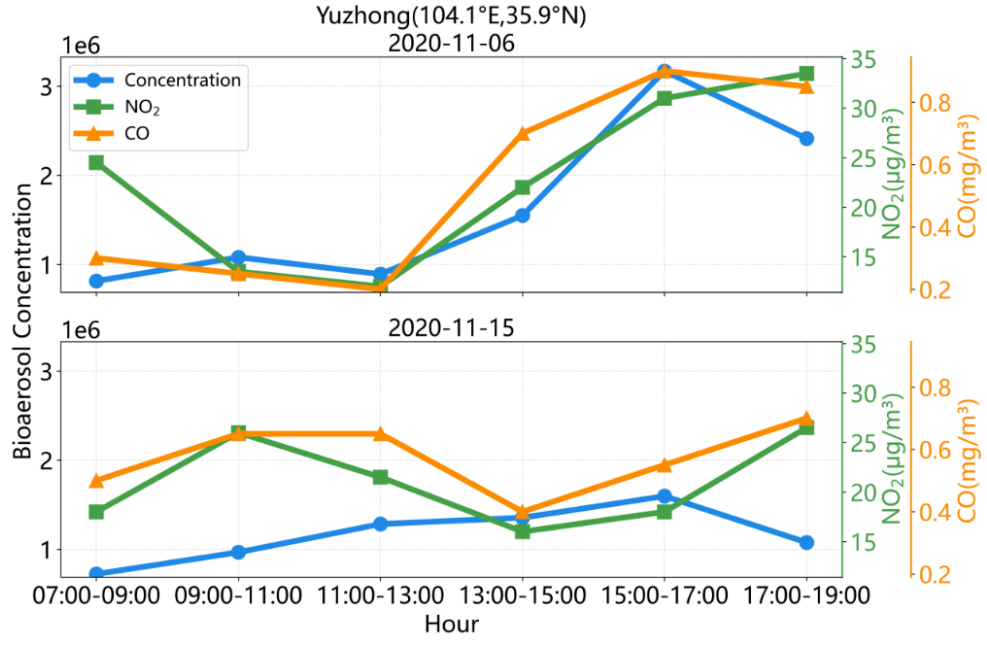

Figure 5: Temporal variations in bioaerosol concentration, NO_2, and CO in Yuzhong on 6 and 15 November 2020.

Figure 6 illustrates diurnal-scale variations in bioaerosol concentrations and related physical parameters in Yonago, Japan, from March to June 2018. As shown, bioaerosol concentrations exhibit pronounced high-frequency fluctuations on the diurnal timescale, indicating strong modulation by short-timescale processes. Within each month, concentrations consistently increase and then decrease, forming a recurrent "single-peak, within-month" pattern throughout March–June. This suggests that bioaerosols do not behave as random background noise but instead represent an aerosol component characterized by a well-defined temporal structure.

During the sampling period, a prominent concentration peak occurred in mid-April, substantially exceeding levels observed in other months and persisting for several consecutive days rather than representing a single-day anomaly. Given that no corresponding extreme peaks are observed in the other variables shown in the figure, this event is unlikely to have been driven solely by particulate matter (PM) or gaseous pollution. Instead, it more plausibly reflects a regional-scale, concentrated release from biological sources.

AOD at 550 nm reaches near-maximum values during the major bioaerosol concentration peaks and varies largely in phase with bioaerosol concentrations. The temporal variations in PM_{10} and $PM_{2.5}$ concentrations are broadly consistent with those of bioaerosol concentrations, with $PM_{2.5}$ exhibiting a slight lag. This indicates that, during this event, bioaerosols and non-biological particles were influenced by shared controlling processes rather than varying independently.

In contrast, the variations in NO_2 and O_3 concentrations are substantially smaller than those of PM and bioaerosols and do not show pronounced increases or decreases concurrent with bioaerosol peaks. This indicates that the observed bioaerosol concentration variations during the sampling period were not primarily controlled by gaseous pollutant emissions.

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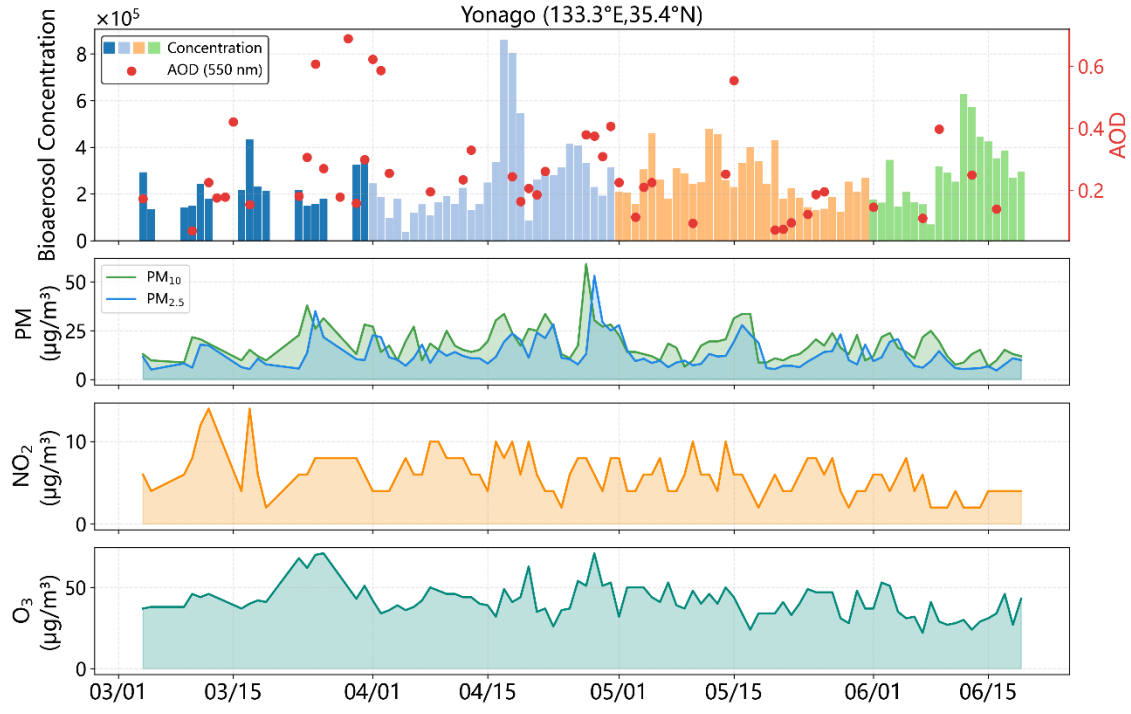

Figure 6: Daily variations in bioaerosol concentration, aerosol optical depth (AOD), PM2.5, PM10, NO_2, and O_3 in Yonago, Japan, during March–June 2018.

3.3 Bacterial community structure

To investigate the dominant taxa at different hierarchical levels across varying underlying surface types, this study selected the five taxa with the highest relative abundances for each surface type for analysis. The underlying surface was classified into six categories based on NDVI: NDVI < 0.1 was defined as very low vegetation cover; 0.1 ≤ NDVI < 0.2 as low vegetation cover; 0.2 ≤ NDVI < 0.4 as medium–low vegetation cover; 0.4 ≤ NDVI < 0.6 as medium vegetation cover; and 0.6 ≤ NDVI ≤ 0.8 as high vegetation cover. In addition, because the sampling campaign included aircraft measurements, an additional category representing the upper atmosphere was included.

Figure 7a presents the bacterial community composition at the phylum level. Proteobacteria dominate the community, with a relative abundance of 60.1%, followed by Firmicutes (24.8%). Among the identified phyla, Chloroflexi are primarily associated with areas of low vegetation cover (43.0%) and very low vegetation cover (33.2%). Actinobacteria and Acidobacteriota exhibit similar distribution patterns, occurring mainly in low-vegetation (33.9% and 35.3%, respectively) and very low vegetation cover regions (31.3% and 28.0%, respectively). These three phyla represent dominant bacterial groups in soils under varying vegetation conditions (Zhou and Wang, 2023). As soil and vegetation are important sources of airborne microorganisms, contributions from soil sources to atmospheric microbial populations become more pronounced under conditions of low vegetation cover (Archer et al., 2023; Xie et al., 2020).

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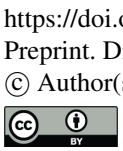

Figure 7b shows the bacterial community composition at the genus level. The dominant taxon at the genus level is *Lactococcus*, with a relative abundance of 23.4%. Among the identified genera, *Staphylococcus* (87.8%), *Alcaligenes* (67.3%), and *Enterococcus* (49.6%) are predominantly distributed in the upper atmosphere. *Paraburkholderia* (61.6%) is mainly distributed in areas with high vegetation cover. The pronounced enrichment of *Staphylococcus*, *Alcaligenes*, and *Enterococcus* in upper-atmosphere samples may be attributed to their strong environmental tolerance and diverse source characteristics, which facilitate survival during atmospheric vertical transport and under selective filtering processes in upper-atmosphere environments (García-Solache and Rice, 2019; Onyango and Alreshidi, 2018; Machado et al., 2023; Pedrosa-Silva and Venancio, 2023). *Paraburkholderia*, a genus typically associated with plants, receives sustained and stable biological inputs in regions with high vegetation cover, making it more likely to accumulate in near-surface air over densely vegetated areas (Esmaeel et al., 2018; Eberl and Vandamme, 2016).

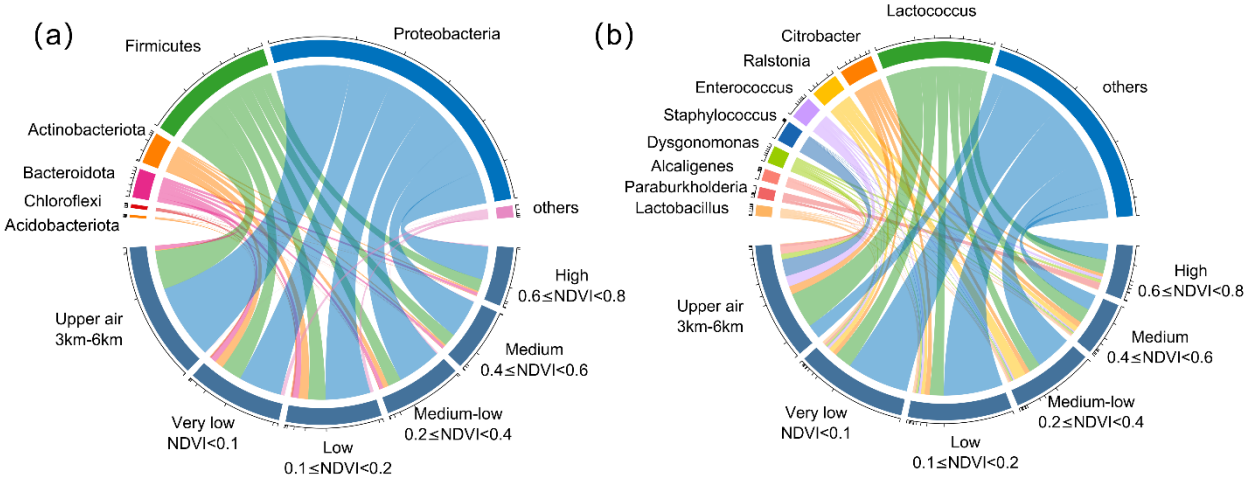

Figure 7: (a) Distribution of dominant taxa at the phylum level across different underlying surface types; (b) distribution of dominant taxa at the genus level across different underlying surface types.

3.4 Bacterial diversity

As shown in Fig. 8a and Fig. 8c, the mean values of both the Shannon and Chao1 indices across different underlying surface types follow the same decreasing order: low vegetation cover > very low vegetation cover > medium–low vegetation cover > high vegetation cover > medium vegetation cover. This pattern indicates that α diversity and species richness are highest in areas with low vegetation cover, whereas areas with medium vegetation cover exhibit the lowest diversity. Variations in α diversity across vegetation cover gradients reflect the combined effects of source inputs, multi-source mixing, and ecological filtering processes (Mantoani et al., 2024; Lu et al., 2024; Gashi et al., 2025). Low vegetation cover areas integrate both soil- and vegetation-derived biological sources and are typically associated with stronger surface disturbance, resulting in the highest species richness and diversity (Nie et al., 2024; Mu et al., 2020; Archer et al., 2023). In contrast, medium vegetation cover regions suppress soil resuspension while not yet developing the complex phyllosphere-associated microbial sources

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characteristic of highly vegetated areas. Consequently, these regions experience fewer source inputs and stronger environmental filtering, leading to the lowest α diversity (Zhai et al., 2022; Robinson et al., 2021).

Figures 8b and 8d further show that both the Shannon index ($r = -0.68$, $P = 0.015$) and the Chao1 index ($r = -0.818$, $P = 0.0011$) are negatively correlated with NDVI, with the Chao1 index exhibiting a stronger correlation. Based on the differing sensitivities of these two indices, it can be inferred that, under low-NDVI conditions, atmospheric microbial communities are more strongly influenced by external source inputs, leading to increased species richness. However, many of the newly introduced taxa occur at low relative abundances and therefore have a limited effect on community evenness, resulting in a smaller magnitude of change in the Shannon index compared to the Chao1 index.

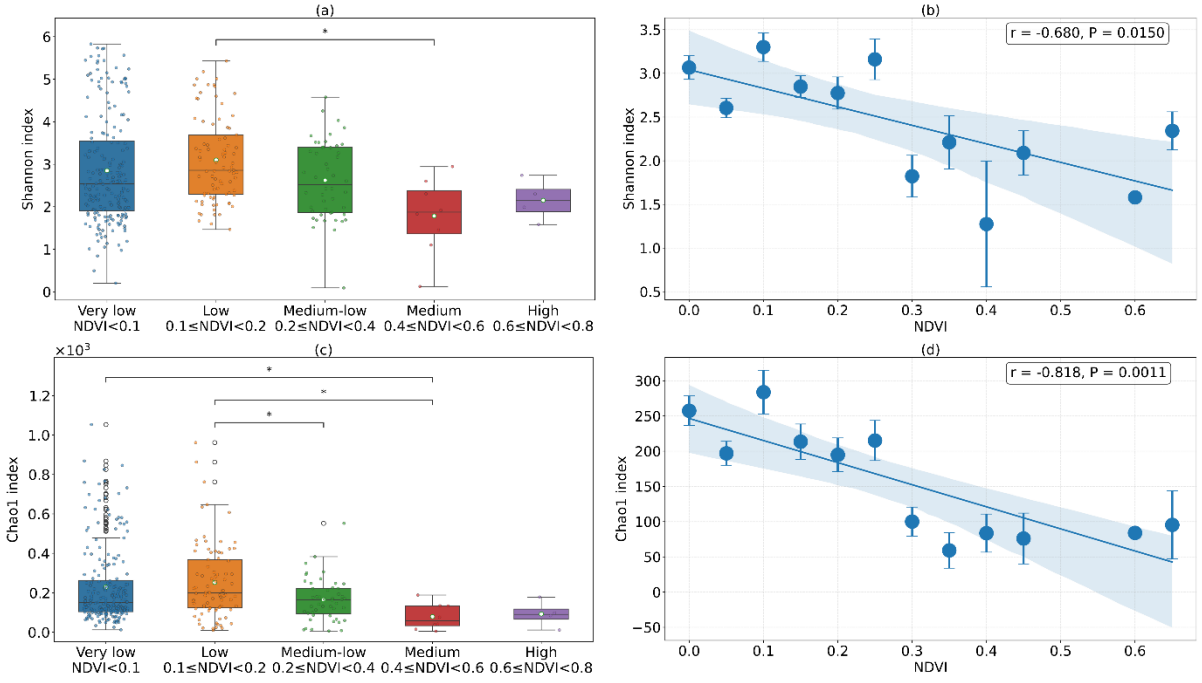

Figure 8: (a) Relationship between underlying surface types and the Shannon index; (b) relationship between NDVI and the Shannon index; (c) relationship between underlying surface types and the Chao1 index; (d) relationship between NDVI and the Chao1 index. Note: In panels (a) and (c), asterisks indicate statistically significant differences in diversity indices between sample groups based on the Wilcoxon rank-sum test, whereas the absence of asterisks indicates no significant difference. One to four asterisks denote $P < 0.05$, $P < 0.01$, $P < 0.001$, and $P < 0.0001$, respectively. In panels (b) and (d), blue dots represent mean diversity indices calculated by binning NDVI into intervals of 0.05 and averaging all samples within each bin; error bars indicate the standard error of the mean for each bin.

Data availability

Interested researchers can download the integrated dataset from https://zenodo.org/records/19605145(Huang et al., 2026). The non-biological aerosol particle number concentration data and the bacterial community structure data at the domain, class,

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order, family, and species levels during the sampling period are not currently publicly available. These data are available from the corresponding author upon reasonable request.

Conclusions

This study reports an integrated dataset of atmospheric bioaerosols obtained from a large-scale dust–bioaerosol field observation campaign conducted across East Asia using a unified sampling and analytical protocol. The dataset includes number concentrations of both biological and non-biological aerosols, bacterial community structure at multiple taxonomic levels, together with concurrent environmental variables and air quality parameters. The database enables the systematic analysis of correlations between regional-scale vegetation cover, temperature, and airborne microbial concentrations and community structure. The results indicate that bioaerosol concentrations exhibit a significant negative correlation with NDVI. Bioaerosol concentrations increase with air temperature below approximately 15 °C, whereas a decline is observed at higher temperatures. Temporal-scale analysis demonstrates that the diurnal variation in bioaerosol concentration exhibits a discernible intraday structure rather than random fluctuations. In addition, the daily evolution of bioaerosol concentrations shows a relatively high degree of synchrony with AOD and particulate matter concentrations. Community structure and diversity analyses further demonstrate pronounced ecological differentiation among regions with varying vegetation cover, with α-diversity showing a significant negative correlation with NDVI. By integrating concentration data, community structure, and multifactor environmental information within a unified methodological framework, this database constitutes a consistent and comprehensive foundation for investigating land–atmosphere interface ecological processes and the spatiotemporal evolution of regional bioaerosols, with clear substantial scientific and applied value.

Author contributions

JH and ZH conceived and designed the database framework, established the unified sampling and analytical protocols, and supervised the overall project, including data validation and quality control. TM, JS and JB contributed to methodological development, provided technical support, and assisted in data validation and manuscript revision. QD, FX, YW, YJ, XH, DL, DW and WZ conducted field sampling and laboratory analyses and curated the dataset. WZ and QD performed the formal data analysis and prepared the figures. WZ prepared the manuscript with contributions from all co-authors. All authors reviewed and approved the final manuscript.

Competing interests

The authors declare no conflicts of interest.

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