

Article Title: Monsoon meteorology and agricultural phenology structure atmospheric transport of aerial biomass across the Indian subcontinent

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5

6 **Abstract:** The movement of organisms through the atmosphere is a fundamental but under-observed
7 ecological process that redistributes biomass, genes, and ecological interactions across continents.
8 Aeroecology has revealed that such movements are strongly determined by synoptic and mesoscale
9 atmospheric dynamics, which jointly determine when, where, and at what altitudes animals initiate,
10 sustain, and terminate flight. However, current understanding is derived largely from temperate
11 climatologies, leaving tropical aeroecology, where atmospheric circulation regimes are
12 fundamentally different, poorly resolved. Here, I synthesize existing literature on radar observations,
13 migration studies, and atmospheric dynamics to examine how monsoon-dominated tropical
14 atmosphere structures aerial movement across the Indian subcontinent at inter-continental (>1000
15 km) and intra-peninsular (100 – 500 km) scales.

16 Across trophic levels, long-distance migratory movements are jointly organized by synoptic-scale
17 monsoon flow and its seasonal reversal, orographic channeling and uplift along the Himalayan
18 barrier, and spatial constriction at coastal bottlenecks along the western coast of the peninsula. These
19 processes generate a common set of migration corridors for both, birds and insects, including trans-
20 Himalayan, circum-Himalayan, Indo-African, and Indo-East/Southeast Asian pathways. At regional
21 scales, evidence from the limited but critical radar observations indicates that aerial dispersal is
22 tightly coupled to intraseasonal active–break phases of the monsoon, the stability and intermittency
23 of diel boundary-layer transitions (including the strength of nocturnal low-level jets), and –
24 particularly for insect pests – cropping cycles and agricultural phenology.

25 Despite these distinctive dynamics, radar aeroecology in the tropics remains sparse. India’s rapidly
26 expanding weather-radar network, especially its dual-polarization X- and C-Band coverage across the
27 western coast and the western Himalayan arc, provides an exceptional but underutilized platform for
28 strengthening tropical aeroecology. By explicitly integrating monsoon meteorology, ecology, and
29 radar aeroecology, this synthesis provides a framework for advancing a mechanistic, globally
30 representative understanding of tropical aeroecology.

31

32 **Keywords:** aeroecology, birds, bats, insects, migration, monsoon, weather radar

33

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34

35 1. Introduction

36 More than a million animal species enter the lower atmosphere at some point in their lives to
37 disperse, migrate, mate, or perform other essential life-history functions (Drake & Gatehouse 1995;
38 Alerstam 2009; Chapman et al. 2015). Aeroecology conceptualizes this aerosphere not as a passive
39 physical medium, but as an important ecological system structured by dynamic atmospheric
40 processes, within which animals must sense, navigate, and interact while occupying an energetically
41 costly niche (Kunz et al. 2008). Because wind velocity, turbulence, and convection govern the
42 energetic costs of flight and the displacement potential of airborne organisms, a mechanistic
43 understanding of aerial movement requires explicit consideration of atmospheric dynamics
44 (Shamoun-Baranes et al. 2010; Shamoun-Baranes et al. 2017; see **Box 1** for a glossary of key
45 atmospheric terms).

46

47 Broadly, aerial animal movement is structured by three interacting atmospheric controls operating
48 across spatio-temporal scales (**Figure 1**): (i) large-scale (synoptic) circulation that establishes the
49 background wind field, including pressure gradients and frontal dynamics (Shamoun-Baranes et al.
50 2017), (ii) mesoscale and boundary-layer processes that regulate the vertical structure and diel
51 evolution of the lower atmosphere, including boundary-layer growth and collapse, low-level jets
52 (LLJs), and convection (**Figure 2**; Reynolds et al. 2005; Wainwright et al. 2020), and (iii)
53 topographic forcing, whereby mountains and coastlines modify airflow through channeling or
54 orographic acceleration resulting in migration bottlenecks and corridors (Ranjan et al. 2023; Hawkes
55 et al. 2024; Ram et al. 2024). Together, these processes determine the speed, direction, stability, and
56 vertical stratification of winds, thereby shaping when, where, and at what altitudes animals initiate,
57 sustain, and terminate flight (Westbrook & Isard 1999; Shamoun-Baranes et al. 2010; Potter et al.
58 2018; Wainwright et al. 2020; Jentsch & Weidinger 2022; Mikkola et al. 2023).

59

60 Modern radar systems – especially Doppler Weather Radars (DWRs) – have emerged as powerful
61 tools in aeroecology, uniquely capable of simultaneously observing atmospheric structures and the
62 movement of vast assemblages of flying animals embedded within them (Holleman et al. 2008;
63 Dokter et al. 2011; Bauer et al. 2019). However, despite the advantages, they remain underutilized in
64 ecological research across most tropical countries (but see Drucker et al. 2025). Unlike dedicated
65 biological radars, DWRs are designed for meteorological monitoring rather than animal detection¹

1 ¹Purpose-built biological radars are typically deployed as single, study-specific systems (e.g., entomological Vertical Looking Radars
2 or Robin and MERLIN™ bird radar systems), whereas Doppler Weather Radars (DWRs) operate as extensive national-scale networks.
3 Crucially, DWRs are government-operated infrastructures that substantially reduce the financial barriers associated with acquiring,
4 deploying, and maintaining dedicated biological radars (Drake et al. 2020; Bauer et al. 2024). Given that a modern biological radar unit
5 can cost USD \$150,000–\$250,000, with substantial annual operating expenses, access to national DWR networks represents a
6 particularly important opportunity for researchers in the Global South.

66 (Chapman et al. 2003; Chilson et al. 2012; Chilson et al. 2017; Drake et al. 2020; Long et al. 2020;
67 see also Bauer et al. 2024). Their scanning strategies, widening beam geometry with distance, and the
68 frequent overlap between biological and meteorological echoes complicate ecological interpretation
69 (**Figure 3**; Gauthreaux et al. 2020). Interpreting biological backscatter often requires independent
70 validation through aerial sampling using drones (Löcken et al. 2020), helikites, aerial nets, or suction
71 traps (Chapman et al. 2002; Chapman et al. 2004; Mungee et al. 2025), ecological expertise on the
72 known occurrence or emergence of animals (Frick et al. 2012; Stepanian et al. 2020; Anjita et al.
73 2024), or the application of advanced post-processing and machine-learning techniques (Lin et al.
74 2019). While hundreds of studies in Europe and North America have leveraged operational DWR
75 networks to quantify migration magnitude, vertical stratification, and wind coupling (e.g., Dokter et
76 al. 2011; Hu et al. 2016a; Hu et al. 2016b; Shamoun-Baranes et al. 2017), comparatively few have
77 examined tropical systems (Rogers et al. 2020; but see Kearsley et al. 2022; Anjita et al. 2024;
78 Drucker et al. 2025). As a result, much of modern aeroecological inference has been shaped by
79 atmospheric conditions typical of mid-and high-latitude regions (Gauthreaux et al. 2003; Shamoun-
80 Baranes et al. 2010; Dokter et al. 2011; Shamoun-Baranes et al. 2017).

81
82 However, atmospheric structure varies systematically with latitude (Rife et al. 2010; Sandeep et al.
83 2015; Luiz et al. 2024). Differences in seasonality and circulation regimes can fundamentally alter
84 the ecological context in which aerial movement evolves and operates (Drucker et al. 2025). For
85 instance, shorebird departure timing and migration trajectories were closely associated with
86 seasonally reversing monsoon circulation along the East Asia–Australasian Flyway (Lei et al. 2025),
87 while the bird migration in Colombia was temporally more diffuse and uniform across the season as
88 compared to the strongly pulse-driven migration of temperate North America (Drucker et al. 2025).
89 Although still relatively few, these studies suggest that differences in atmospheric structure between
90 tropical and temperate climates, often interacting with regional topography, can generate distinct
91 aeroecological dynamics (Rogers et al. 2020).

92
93 Despite these indications, empirical studies explicitly examining aerial movement within tropical
94 atmospheric systems remain scarce, leaving a major gap in our understanding of how aerial
95 movement ecology. Here, I synthesize existing evidence to examine how monsoon-driven tropical
96 atmospheric processes structure aerial movement, using India as a focal case. The Indian
97 subcontinent provides a particularly compelling system for examining these dynamics at lower
98 latitudes. The region lies at the intersection of a seasonally reversing monsoon circulation (**Figure 4**),
99 major orographic barriers, and multiple large-scale migratory networks, including the Central Asian
100 Flyway of birds and the Himalayan insect flyway (Miyabayashi & Mundkur 1999; Boere & Stroud
101 2006; Hawkes et al. 2025). The steep elevational gradients of the Himalayas intersect the monsoon
102 circulation to channel, lift, trap, or accelerate aerial fauna, producing movement dynamics that differ
103 fundamentally from those described in temperate systems. Yet these complementary sources of
104 information – ecological observations, atmospheric climatology, and weather-radar infrastructure –
105 remain largely unintegrated.

106

107 To address this gap, I pursue three objectives. First, I review the development of radar-based research
108 on aerial fauna in India – from early entomological radar campaigns to recent DWR applications –
109 focusing on how these observations reveal coupling between aerial organisms and synoptic and
110 mesoscale atmospheric structure. Second, I synthesize the country’s principal migratory corridors
111 and aeroecological hotspots across birds and insects, examining how aerial movement is organized
112 by three interacting atmospheric controls: large-scale monsoon circulation, mesoscale and boundary-
113 layer dynamics, and topographic forcing across major mountain systems and coastlines. Third,
114 building on this synthesis, I identify key gaps in our understanding of how aerial fauna interact with
115 monsoon-driven atmospheric structure across scales, and outline how these gaps can be addressed
116 using the spatial coverage, temporal continuity, and observational capabilities of India’s rapidly
117 expanding DWR network.

118

119

120 **2. Methods**

121 I conducted a structured literature search to synthesize existing knowledge on radar-based
122 aeroecology and atmospheric drivers of aerial fauna movement in India. Literature searches were
123 performed in the Web of Science Core Collection database without temporal restriction using two
124 main search strings: (i) Radar-based studies: TS=((radar* OR "weather radar*" OR "vertical looking
125 radar*" OR "entomological radar*") AND (bird* OR avian OR insect* OR arthropod* OR bat* OR
126 chiroptera* OR "biological echo*" OR bioscatter*)) AND (India OR "Indian subcontinent")), and (ii)
127 Atmospheric determinants of aerial movement and migration: TS=((migrat* OR dispers* OR
128 flyway* OR "windborne movement" OR emergence) AND (bird* OR avian OR insect* OR
129 arthropod* OR locust* OR aphid* OR planthopper* OR bat* OR chiroptera*)) AND (atmospher* OR
130 monsoon OR "boundary layer" OR "low-level jet" OR "synoptic wind*") AND (India OR "Indian
131 subcontinent")) NOT TS=("emerging disease" OR zoonos* OR virus* OR rabies OR "nipah" OR
132 pathogen*).

133

134 Titles and abstracts were screened for relevance to radar observations, aerial movement, and
135 atmospheric drivers of aerial fauna in the Indian subcontinent. Because several foundational radar-
136 entomology studies conducted in India between 1960 and 1990 are incompletely indexed in
137 international databases, supplementary manual searches were undertaken to retrieve archival
138 publications, institutional reports, and regionally published journals. Citation chaining was used to
139 identify additional relevant sources.

140

141 **3. Radar Observations of Aerial Fauna in India**

142 Radar-based studies of aerial fauna in India remain limited in number and historically sporadic. The
143 available literature is characterized by campaigns linked to technological experiments (Ramana
144 Murty et al., 1964; Mazumdar et al., 1965; Riley & Reynolds 1997), opportunistic observation of
145 bioscatterers by meteorologists (Suresh et al. 2005), or to specific agricultural pest outbreaks (Riley
146 & Reynolds 1997; Amarjyothi et al. 2021; Anjita et al. 2024), rather than as a sustained research
147 discipline at the intersection of ecology and meteorology.

148

149 The earliest documented radar study of aerial fauna in India used a 3.2-cm (X-Band; **Figure 3**)
150 meteorological radar to track Desert Locust (*Schistocerca gregaria*) around New Delhi,
151 demonstrating predominantly wind-driven transport within the lower boundary layer (~ 1 to 1.5 km)
152 and a clear diel cycle of convective ascent and evening descent (Ramana Murty et al., 1964;
153 Mazumdar et al., 1965; **Figure 2**; also see **Box 1**). Crucially, the swarm arrival and activity was
154 tightly linked to the phase structure of the southwest monsoon; invasion followed an anomalously
155 wet 15-day period, but peak aerial activity occurred during a subsequent temporary break in monsoon
156 conditions, when rainfall subsided and lower-tropospheric winds were well organized. With the
157 return of active monsoon conditions and a shift in wind direction, swarm activity rapidly diminished
158 and locusts disappeared from the region. These early observations already illustrated how insect
159 movement in the region is strongly coupled to the alternating active and break phases of the monsoon
160 circulation (**Box 1**).

161
162 A major phase of development occurred between 1985 and 1995, when the UK Overseas
163 Development Administration tested prototypes of the first nutating vertical-looking entomological
164 radar near Hyderabad, southern India, in collaboration with the International Crops Research Institute
165 for the Semi-Arid Tropics (ICRISAT), as part of a programme on the ecology and management of
166 the cotton bollworm (*Helicoverpa armigera*; King et al. 1990; Riley et al. 1990; Riley et al. 1992;
167 Reynolds et al. 1999). This represented the first sustained effort in India to quantify aerial insect
168 assemblages using coordinated radar observations, mark–recapture, and aerial sampling using
169 helikites. The studies showed that adult bollworms emerged at dusk (~20 min after sunset), with peak
170 densities about one hour later, followed by declining but sustained nocturnal activity. Early peaks
171 were dominated by ovipositing females, with mate-searching males appearing later in the night. Most
172 individuals remained within ~10 m of the surface, and no mass high-altitude ascents were observed,
173 indicating predominantly local dispersal rather than long-distance migration (Riley et al. 1992).

174
175 From the same campaign, Reynolds & Wilson (1989) showed that during the winter agricultural
176 season (early *rabi*; November–December), large numbers of Hemiptera undertook nightly windborne
177 flights at ~100 m above ground. Aerial catches were dominated by rice planthoppers, leafhoppers,
178 and aphids, alongside high densities of predatory mirids. Radar observations documented mass dusk
179 take-off events and secondary nocturnal density peaks later in the night after 23:00 IST, potentially
180 representing long-distance migrants. Back-trajectory wind-field analyses suggested source regions
181 100–200 km away, often from irrigated rice landscapes along the Krishna River basin. In some cases,
182 insect invasions coincided with synoptic disturbances that strengthened and veered low-level winds,
183 including cyclonic systems moving inland from the Arabian Sea.

184
185 Riley et al. (2013) extended this work to eastern India, using elevated aerial nets – rather than radar
186 per say – in West Bengal to examine long-distance migration of aphids and associated small insects.
187 Samples were similarly dominated by planthoppers, leafhoppers, aphids, and predatory mirids. In
188 contrast to the ICRISAT study, substantial activity occurred during both day and night; however,
189 peak activity and movements still occurred after dusk. Wind-field analyses indicated displacement
190 distances of 300–500 km overnight, with migrants typically originating in Bangladesh or northern

191 West Bengal and advected southwest under post-monsoon north-easterlies (**Figure 4**). The study also
192 highlighted the role of nocturnal temperature inversions in sustaining passive flight by enabling
193 insects to remain within stable, fast-moving nocturnal wind layers, including LLJ-like flows
194 developing above the inversion. These atmospheric conditions substantially increased potential
195 displacement distances during warm tropical nights.

196
197 Because birds produce stronger radar returns than most insects, they are generally easier to detect
198 with traditional S- and C-band DWR (**Figure 3**). Nevertheless, the first explicitly bird-focused radar
199 study in India appeared only later. Suresh et al. (2005) used the S-band DWR at the Cyclone
200 Detection Radar Station (Chennai, South India) to investigate strong reflectivity returns (28–38 dBZ)
201 observed during morning and evening twilight between December and April. After ruling out clouds,
202 particulates, and refractive turbulence, the authors attributed these signals to mass departures and
203 arrivals of wetland birds commuting between coastal roosts and foraging areas. Several lines of
204 evidence supported this interpretation: reflectivity enhancements occurred exclusively over land,
205 peaked precisely at civil dawn and dusk, formed expanding and contracting horizontal “biological
206 fronts” between ~200–600 m a.g.l., and exhibited radial velocities exceeding background winds by
207 up to $\pm 10 \text{ m s}^{-1}$, consistent with powered flight rather than passive drift. Thus, despite their capacity
208 for powered and directional flight, vertebrate commuting behavior remained structured by boundary-
209 layer transitions and low-level wind organization. Importantly, Suresh et al. (2005) provided one of
210 the only quantitative demonstrations in India of using DWR reflectivity to estimate bird densities:
211 $\sim 2,400 \text{ birds km}^{-3}$, implying 8,000–13,000 individuals within the sampled $5 \times 7 \text{ km}$ region, values
212 broadly consistent with independent roost counts. To the best of my knowledge, till date this remains
213 the only study in India to explicitly quantify aerial vertebrate densities using DWR.

214
215 The 2020 desert locust upsurge across West and Northwest India marked a renewed application of
216 the national DWR network for biological monitoring, representing the first coordinated use of DWR
217 for this purpose since the early opportunistic studies of Ramana Murty et al. (1964) and Mazumdar et
218 al. (1965). Rao et al. (2020) used data from radars in Jaipur (S-Band) and New Delhi (C-Band) to
219 resolve fine-scale swarm morphology, including swarm fronts, internal density gradients, and
220 leading-edge movement, enabling reliable mapping of swarm progression across heterogeneous
221 terrain. Subsequent studies demonstrated that locust swarms produce distinct reflectivity signatures
222 (10–30 dBZ) and Doppler velocity fields closely aligned with near-surface winds (Amarjyoti et al.
223 2021). Anjita et al. (2024) further showed that swarms can be detected at distances of 40–80 km as
224 coherent non-meteorological echo patches forming shallow layers within the lowest 1–1.5 km of the
225 atmosphere, and are often embedded within LLJ regimes associated with monsoon onset or dryline
226 boundaries.

227
228 Despite these advances, radar-based investigations of aerial fauna in India remain remarkably sparse
229 and episodic, spanning only a handful of campaigns over more than five decades. However, a
230 substantial body of ecological and observational research – largely independent of radar – already
231 documents major migration routes, dispersal pathways, and seasonal aggregation zones for alate
232 organisms across the Indian subcontinent, often in close association with prevailing wind systems

233 and atmospheric structure. Synthesizing these patterns provides the much needed framework for
234 understanding how atmospheric processes organize aerial movement across the region and for
235 advancing radar aeroecology.

236

237 **4. Atmospheric Controls of Aerial Movement Across the Indian Subcontinent**

238 Birds, as powered fliers, exercise strong control over heading, altitude, and airspeed, and can often
239 compensate for moderate to strong crosswinds (Sinelschikova et al. 2020). Their navigation
240 integrates multiple sensory systems including magnetoreception, solar and stellar compasses, and
241 orographic landmarks (Emlen 1975a, 1975b; Pennycuick 1975; Alerstam 1979a; Alerstam 1979b;
242 Schmidt-Koenig et al. 1991; Green 2003; Newton 2008; Pennycuick 2008; Goto et al. 2017;
243 Aschwanden et al. 2019; Wiltschko & Wiltschko 2019). In contrast, most migratory insects are
244 windborne and rely primarily on synoptic-scale wind fields, boundary-layer structure (including
245 LLJs), and temperature thresholds to determine activity and movement (Schaefer 1976; Westbrook &
246 Isard 1999; Reynolds et al. 2005). While directional orientation occurs in many taxa, long-distance
247 movement is achieved predominantly through selective ascent into favorable wind layers rather than
248 sustained course correction (Achtemeier 1991; Chapman et al. 2010; Chapman et al. 2015; Hu et al.
249 2016a; Hu et al. 2016b; but see Menz et al. 2011). Because these two groups differ fundamentally in
250 their degree of aerodynamic control and atmospheric coupling, their interactions with monsoon-
251 driven wind regimes and boundary-layer dynamics also differ, and so I consider them separately in
252 the following subsections.

253

254 **4.1 Avian Movement Systems**

255 Migratory birds provide the clearest framework for understanding how large-scale biological
256 movements interact with atmospheric circulation and regional topography across the Indian
257 subcontinent. The majority of long-distance migrants in India are embedded within the Central Asian
258 Flyway (CAF), with fewer species across the East Asia–Australasian Flyway (EAAF) and the West
259 Asia–East African Flyway (WAEAF) (Miyabayashi & Mundkur 1999; Boere & Stroud 2006; **Figure**
260 **5**).

261 CAF links high-latitude breeding areas across northern Eurasia – including Arctic Russia and Siberia
262 – with wintering regions across the Indian subcontinent and few parts of South Asia and Middle East.
263 Southbound autumn migration coincides with the development of the Siberian High – a large cold-
264 season continental anticyclone that intensifies the Asian winter monsoon, generating persistent
265 northerly low-level flow from the Eurasian interior toward South Asia and Indian Ocean (Zhang et al.
266 2025). In spring, the mid-latitude westerlies and the Western Disturbances interact with the
267 Himalayas to provide tailwinds and uplift that help migratory birds gain altitude to cross the
268 mountains more efficiently (Lang & Barros 2004; Dimri & Dash 2012; Dimri et al. 2015; Koll et al.
269 2015; Puranik & Karekar 2017; Hunt et al. 2018). While the Asian monsoon system establishes the
270 large-scale atmospheric context for seasonal migration (**Figure 4 & 5**), the specific routes and
271 movement dynamics of migrants are ultimately shaped by regional mesoscale processes and complex
272 topography that modulate wind assistance and barrier permeability at finer spatial scales (Kumar et
273 al. 2020a). Accordingly, CAF is increasingly viewed as an umbrella system comprising distinct, non-

274 overlapping corridors, each characterized by different atmospheric, orographic, and ecological
275 conditions (Kumar 2019; Kumar et al. 2020a).

276

277 The Western circum-Himalayan corridor is used predominantly by species capable of obligate or
278 facultative soaring flight (e.g. Black Kites, Greater and Indian Spotted Eagles, Tawny Eagles,
279 Western Marsh Harriers, Montagu's Harriers, Pallid Harriers, and the Common and Demoiselle
280 Cranes; Terraube et al. 2012; Kumar et al. 2020a; Baraiya et al. 2022; Mi et al. 2022; Ram et al.
281 2024; Baraiya et al. 2025; Kannan et al. 2025). This association suggests that thermal uplift likely
282 plays an important role in structuring this route, as the arid and semi-arid landscapes of Central Asia
283 and Pakistan generate strong daytime thermal convection that can facilitate energetically efficient
284 soaring flight. The Trans-Himalayan corridor within the CAF is used by a smaller number of species
285 with specialized, high-altitude powered flight (e.g. Bar-headed Goose, Brown-headed Gull, Eurasian
286 Wigeon, Northern Pintail, Garganey, and Ruddy Shelduck; Bishop et al. 2015; Palm et al. 2015;
287 Prins & Namgail 2017; Li et al. 2020; Kannan et al. 2025; Panagoda et al. 2025), and finally the
288 Eastern circum-Himalayan route supports a few circum-navigating species such as the Himalayan
289 Vultures and Northern Shovelers (Palm et al. 2015; Ummee et al. 2025) and partially overlaps with
290 the EAAF (but see Palm et al. 2015).

291

292 Some species, however, change corridors, e.g. Black Kites (Literák et al. 2022) and Himalayan
293 Vultures (Ummee et al. 2025), although the drivers of such switching remain poorly understood, and
294 are likely influenced, at least in part, by atmospheric conditions. Ruddy Shelducks, Common
295 Cuckoo, and Siberian Rubythroat circumnavigate the Himalayas along either the western or the
296 eastern corridors (Miyabashi & Mundurkar 1999; Trierweiler et al. 2014; Palm et al. 2015; Cheng et
297 al. 2024; Zhao et al., 2024), while Demoiselle Cranes take the western detour in spring but cross the
298 high-altitude Himalayas and Qinghai-Tibet Plateau during autumn (Galtbalt et al. 2022; Mi et al.
299 2022). The circum-Himalayan detours can be up to 36% longer than the shortest trans-Himalayan
300 trajectory (Baraiya et al. 2025), and thus most likely reflect the trade-off between wind assistance and
301 migratory distance and/or time (Khacher 1978; Hawkes et al. 2011, Hawkes et al. 2013; Delany &
302 Namgail 2017; Parr et al. 2017; Prins & Namgail 2017).

303

304 In contrast to the Himalayan-centered migration systems of the CAF, large-scale movements across
305 eastern Asia occur within a more open coastal atmospheric corridor associated with the EAAF, which
306 similarly links high-latitude breeding areas across Arctic and temperate East Asia with wintering
307 regions extending through Southeast Asia, Indonesia, and Australasia (Yong et al. 2015; **Figure 5**).
308 CAF and EAAF also overlap meteorologically in that both are embedded within the seasonal
309 monsoon circulation (Yong et al. 2015). However, the dominant physical controls shaping migration
310 differ between the two systems: with stronger roles of topographic barriers and uplift dynamics of the
311 Himalayan-Tibetan region along the CAF and predominance of coastal wind systems and maritime
312 atmospheric circulation across the EAAF. Crucially, along the EAAF, tailwind support associated
313 with the strengthening East Asian winter monsoon circulation during autumn appears particularly

314 important for southbound migration, whereas temperature gradients, emerging productivity and
315 resource availability along the flyway, play a greater role during spring movements (Zhang et al.
316 2025).

317

318 The western margin of the Asian continent supports a third major migration system that intersects the
319 Indian subcontinent: the WAEAF (Newton 2008). WAEAF is among the most poorly documented
320 flyways in the world and is frequently omitted from global syntheses due to the paucity of tracking
321 data (Guilherme et al. 2023). Amur Falcons and Common Cuckoos represent the best-studied
322 migrants along this route (Ganpule 2011; Cheng et al. 2024; Kaur et al. 2024), migrating from
323 Northeast Asia to South and East Africa and using Northeast India as a major staging region where
324 large congregations refuel before undertaking transoceanic flights across the Arabian Sea and
325 western Indian Ocean (**Figure 5**). These movements suggest that migration along WAEAF is closely
326 linked to seasonal transitions in the Asian monsoon system and the displacement of the Inter-Tropical
327 Convergence Zone (ITCZ; **Figure 4**). During autumn, migrants arriving in Northeast India typically
328 stage during October–November before departing toward Africa under the post-monsoon circulation
329 regime. At this time, the establishment of continental anticyclonic circulation across West Asia and
330 the Arabian Peninsula generates coherent easterly flow across the northern Arabian Sea, providing
331 favorable tailwind support for long-distance transoceanic flights toward East Africa. In contrast,
332 spring return movements occur during the pre-monsoon transition, when atmospheric circulation over
333 the Arabian Sea becomes increasingly influenced by strengthening southwesterly flow associated
334 with the developing Somali Jet and the northward displacement of the ITCZ. Importantly, many
335 migrants appear to time their movements to precede the peak Somali Jet phase, which typically
336 develops during the mature southwest monsoon (June–August).

337 Across all three flyways, autumn migration is closely aligned with the development of continental
338 high-pressure belt (ITCZ) and the strengthening of monsoon outflow, which generate persistent
339 tailwinds: northerlies along the CAF, easterlies across the Arabian Sea along the WAEAF, and
340 coastal northeasterlies along the EAAF, and facilitate southward displacement. In contrast, spring
341 migration occurs under transitional circulation regimes, where westerlies, pre-monsoon flows, and
342 Western Disturbances interact with the Himalayan barrier to wind assistance and access to high-
343 altitude crossing routes. The Himalayas act as both a major barrier and a dynamic filter, structuring
344 corridor selection through a combination of orographic uplift, channeling, and energetic constraints,
345 while coastal and oceanic margins enable alternative, wind-mediated pathways, particularly along the
346 eastern and western flanks of the subcontinent. This framework also provides important context for
347 understanding the movement ecology of aerial insect migrants in the region.

348

349 **4.2 Insect Movement Systems**

350 Insect migration typically occurs as multi-generational, wind-assisted movement across regional and
351 continental scales, in which successive generations track shifting seasonal resources rather than
352 completing closed individual migratory circuits (Brown & Chippendale 1974; Drake & Gatehouse
353 1995; Chapman et al. 2015; Satterfield et al. 2020). Unlike vertebrate migrants, these movements are

354 often mediated by developmental and physiological adaptations such as diapause and migratory
355 morphs that synchronize dispersal with seasonal environmental conditions. Because each generation
356 relies heavily on atmospheric transport and makes independent movement decisions, migratory
357 connectivity between breeding and non-breeding populations is generally weak, resulting in
358 extensive population mixing across large spatial scales (Drake & Gatehouse 1995; Chapman et al.
359 2013; Gao et al. 2020).

360
361 Despite their ecological importance, substantial knowledge gaps persist and the diversity of
362 migratory taxa and the spatial extent of their flyways remain largely unknown (Prakash et al. 2022).
363 Across the globe, only a small fraction of insect taxa, e.g. brown planthopper (*Nilaparvata lugens*),
364 globe skimmer dragonfly (*Pantala flavescens*), monarch butterfly (*Danaus plexippus*), Silver-Y moth
365 (*Autographa gamma*), have been studied in detail (Drake & Reynolds 2012; Chapman et al. 2015;
366 Wu et al. 2019; Hu et al. 2024). Unlike birds, for which complete migratory routes have been
367 reconstructed through satellite and GPS tracking, no insect species has yet been tracked along its full
368 migration pathway (but see Menz et al. (2011) for a partial route analysis of the Death's-head
369 hawkmoth *Acherontia atropos*). Therefore, the coupling between insect migration and atmospheric
370 circulation must largely be inferred indirectly from radar observations, wind trajectory analyses, and
371 population-level genetic evidence. Below, I synthesize the best-resolved insect migratory systems
372 within the Indian subcontinent and their atmospheric coupling.

373
374 The Himalayan insect flyway, analogous to the CAF, was recently formalized by Hawkes et al.
375 (2025) based primarily on evidence from migratory Diptera. However, field evidence for traffic along
376 this flyway is extensive even beyond Diptera. Gatter (1980) recorded continuous northward spring
377 migrations of butterflies, noctuid moths, syrphid flies, and bumblebees at elevations >5,000 m, with
378 hoverfly fluxes reaching ~19 individuals per minute at Thorong La in Nepal (5,416 m). These rates
379 are comparable to migratory traffic recorded at substantially lower temperate mountain passes, such
380 as the Pyrenees (2,263 m; Hawkes et al. 2024), highlighting the remarkable vertical extent of aerial
381 migration in lower-latitude systems. Observations from Nepal also corroborate high-altitude spring
382 crossings (~3,600 m) by hoverflies and butterflies (Westmacott & Williams 1954; Reynolds et al.
383 2024). Isotopic analyses of the globe skimmer dragonfly (*Pantala flavescens*) reaching the Maldives
384 indicate origins in Nepal, implying long-distance, potentially trans-Himalayan trajectories consistent
385 with flight records up to 6,300 m (Hobson et al. 2012). More recently, Chandra et al. (2019)
386 documented 2,261 insect species in the Indian Trans-Himalaya, including recognized migrants from
387 Odonata, Lepidoptera, Hemiptera, Diptera, and Hymenoptera.

388
389 Collectively, these observations indicate that the Indian Himalayas support a recurrent,
390 atmospherically structured migratory corridor. The broad seasonal direction of autumn and spring
391 movements suggests a role for monsoon-driven synoptic circulation analogous to that documented
392 for migratory birds, although the mesoscale, vertically stratified atmospheric mechanisms shaping
393 these migrations remain unexplored due to the absence of complete-route tracking data for insects.
394 Nevertheless, earlier entomological studies have noted that Himalayan insects frequently exploit diel

395 wind systems of the valley–pass topographic accelerators (along-valley wind circulation regimes;
396 **Box 1**; Westmacott & Williams 1954; Mani 1968; Gatter 1980; Reynolds et al. 2024).

397
398 A second major insect movement system linked to the Indian subcontinent appears to occur along the
399 Afro–Asian corridor analogous to the WAEAF for birds. The proposed trans-Himalayan migratory
400 pathway of the globe skimmer dragonfly, considered the longest regular non-stop migration relative
401 to body size (Hedlund et al. 2021), extends from northern India across the subcontinent and onward
402 toward East Africa (Anderson 2009; Hobson et al. 2012; Ranjan et al. 2023). Offshore records of the
403 painted lady (*Vanessa cardui*) across the northern Indian Ocean (Ecuador 1992), together with its
404 recently demonstrated 4,200 km trans-Atlantic migration (Suchan et al. 2024), indicate that large
405 Lepidoptera are capable of monsoon-assisted Indo–African crossings, even though population-level
406 connectivity remains unresolved. The fall armyworm (*Spodoptera frugiperda*) has also been
407 proposed to have reached the Indian subcontinent from Africa (Johnson 1987). The same
408 atmospheric corridor has been confirmed for desert locusts with rapid trans-Indian Ocean transport
409 found tightly coupled to monsoon dynamics and Indian Ocean Dipole–driven rainfall anomalies
410 (Feng et al. 2025). Energetic modeling and wind trajectory analyses indicate that such transoceanic
411 movements depend critically on favorable tailwinds and are unlikely to be achieved through self-
412 powered flight alone (Hedlund et al. 2021). Smaller insects, which often rely more heavily on passive
413 atmospheric transport, may be capable of similar long-distance dispersal across this corridor,
414 although the mechanisms remain poorly understood. Genetic studies support an Indian origin for
415 several Drosophilidae and Tephritidae species now established in East Africa, implicating seasonal
416 wind-driven dispersal as the underlying mechanism (Tsacas 1984; Jacquard et al. 2013; also see
417 Johnson 1987). Large offshore arrivals of blowflies in the Maldives also support mass transoceanic
418 transport (Hawkes et al. 2024). Large-scale wind assisted migrations of Culicoides are also known to
419 occur on the movements of the ITCZ (Sellers 1980). Taken together, these observations indicate a
420 recurring monsoon-mediated aerial corridor between India and East Africa, with insects transported
421 toward India by south-westerly monsoon winds and potentially redistributed westward or southward
422 during the winter monsoon, likely via the Maldives and Seychelles (Anderson 2009; Hobson et al.
423 2012). These bidirectional exchanges are driven by seasonal shifts in the ITCZ and associated wind-
424 field reversals, which establish synoptic-scale pathways linking South Asia, the Arabian Peninsula,
425 East Africa, and the central Indian Ocean.

426
427 Another large-scale migratory system links Southeast and East Asia and partially intersects the
428 Indian subcontinent, broadly coinciding with the EAAF for birds. Populations of brown planthoppers
429 from India (primarily Assam and West Bengal) and the Malay Archipelago act as donor populations
430 for the larger Southeast and East Asian cluster, with net gene flow directed toward Southeast Asia
431 (Wu et al. 2019; Hu et al. 2024). The Bay of Bengal branch of the southwest monsoon transports
432 large swarms from Northern Indochina into East India at altitudes of ~500–1,500 m, where
433 boundary-layer convergence can concentrate migrants before westward dispersal across the Indo-
434 Gangetic Plain. These winds transport insects across hundreds to over a thousand kilometers in a
435 single night (Drake & Reynolds 2012; Otuka et al. 2012; Chapman et al. 2015; Hu et al. 2024).
436 Autumn migration occurs during the weakening monsoon and the establishment of continental

437 northerly flow, enabling southward return transport toward subtropical wintering regions. Trajectory
438 modeling and long-term monitoring indicate that the common cutworm (*Spodoptera litura*)
439 undertakes seasonal cross-border migrations along the southern Himalayan margin, moving from
440 Northeast India, Bangladesh, and Northern Myanmar into Southwestern China during spring and
441 returning toward South Asia in autumn, with these movements facilitated by monsoon-driven airflow
442 across the region (Song et al. 2024). Although India lies just outside the formal East Asian insect
443 flyway (Hu et al. 2025), many agricultural pests remain genetically and demographically linked to
444 Indochinese source populations, making India integral to the wider Asian migratory system.

445
446 Beyond trans-boundary flyways, windborne insect movement within India at shorter scales (~100–
447 500 km) is increasingly recognized as an important ecological phenomenon (Ramesh et al. 2012;
448 Jyothi et al. 2021; Vinayan et al. 2023). In the Indian subcontinent these pathways frequently
449 coincide with the inland penetration of the Arabian Sea monsoon LLJ, which extends across
450 peninsular and central India during the southwest monsoon (Viswanadhapalli et al. 2020; **Figure 4**).
451 Monsoon LLJ is often associated with a strongly stratified nocturnal boundary layer that can
452 concentrate aerial organisms within relatively thin atmospheric layers (Desamsetti et al. 2010;
453 Dwivedi et al. 2020), creating favorable conditions for wind-assisted drift and the formation of
454 coherent insect layers detectable by radar. Recent studies indicate that migratory insect abundance
455 can be comparable to or even greater during daytime (e.g. windborne dispersal of mosquitoes in
456 northeast India; Reynolds et al. 1996) than nighttime, particularly under warmer atmospheric
457 conditions, suggesting that daytime boundary-layer convection and thermal uplift may also contribute
458 to large-scale insect transport across India (Gao et al. 2025).

459
460 Several experimental and observational studies support the plausibility of wind-assisted dispersal
461 across the Indian subcontinent. Flight-mill experiments show that *H. armigera* can sustain flights
462 exceeding eight hours under still-air conditions, suggesting that moderate wind assistance could
463 result in substantial downwind displacement (Pedgley et al. 1987; Vaishampayan & Singh 1996;
464 Jyothi et al. 2021). Field observations have linked sudden increases in moth abundance to synoptic
465 wind regimes, especially southeasterly flows during February–April and occasionally during
466 October–December when cyclonic systems develop over the Arabian Sea (Pedgley et al. 1987). In
467 the teak defoliator (*Hyblaea puera*), dispersal occurs primarily during April–July when high
468 population densities trigger migration from early-flushing to later-flushing teak stands, producing
469 short-range wind-assisted movements of ~5–10 km and occasionally longer atmospheric transport
470 (Nair & Sudheendrakumar 1986). Seasonal wind reversals also structure aphid dispersal across the
471 Indo-Gangetic Plain, where mustard aphids ascend hundreds of metres into the boundary layer and
472 can be transported tens to hundreds of kilometres within a single night (Ghosh et al. 2019). Flight
473 activity of red flour beetle (*Tribolium castaneum*) and lesser grain borer (*Rhyzopertha dominica*)
474 peaks during the warm monsoon and post-monsoon months but declines sharply during cooler
475 periods (Rajan et al. 2018), highlighting the role of temperature thresholds along with monsoon
476 cycles. Collectively, these observations indicate that regions lying beneath the inland extension of the
477 monsoon LLJ, particularly across Maharashtra, Madhya Pradesh, and Rajasthan, may function as
478 major corridors for drift-dominated insect transport during stable monsoon nights.

479

480 Species capable of sustained powered flight migrate primarily in response to ecological cues while
481 still interacting with atmospheric structure. Butterfly migrations provide the most well-documented
482 example. Large swarms dominated by *Catopsilia* and *Papilio* species have been recorded along the
483 Kalpakkam coastline of South India, moving predominantly northward during October and
484 southward during July (Ramesh et al. 2012). These movements frequently occur across or even
485 against prevailing winds, indicating that migration is driven largely by seasonal rainfall patterns,
486 host-plant phenology, and reproductive cycles rather than passive atmospheric advection alone.
487 Nevertheless, migration timing remains closely coupled to diel atmospheric conditions. Peak flight
488 activity occurs during periods of strong solar heating but relatively weak convective turbulence.
489 Large butterfly movements often occur near transitions in monsoon rainfall regimes, implying
490 sensitivity to intraseasonal variability such as active–break phases of the monsoon (Ramesh et al.
491 2012). Similar atmospheric coupling has also been demonstrated in the transcontinental spread of
492 desert locust swarms, whose rapid redistribution across the Indian Ocean region has been shown to
493 be tightly phase-locked to intraseasonal variability in the monsoon circulation (Feng et al. 2025).
494 Danaine butterflies likewise undertake large-scale seasonal migrations across Southern India, moving
495 westward toward the Western Ghats following the southwest monsoon (October–November) and
496 returning toward the eastern plains during the pre-monsoon period (April–May; Vinayan et al. 2023).
497 Evidence further suggests that populations breed alternately between the Western Ghats and the
498 eastern plains, implying a multi-generational migration cycle in which successive generations
499 undertake the reverse movement (Kunte 2005; Bhakare & Ogale 2018). Together, these observations
500 indicate that even actively flying migrants remain strongly constrained by atmospheric structure, with
501 ecological drivers interacting with diel atmospheric and synoptic monsoon dynamics to shape
502 migration timing and direction.

503

504 **5. Tropical Aeroecology: Knowledge Gaps and Observational Opportunities**

505 Although the available literature remains sparse compared with better-studied regions such as Europe
506 and North America, and many migratory pathways within the Indian subcontinent remain poorly
507 resolved (Tyagi et al. 2022), it nevertheless provides strong evidence for a convergence of large-
508 scale, intercontinental migratory routes across taxa. Both birds and insects utilize the same three
509 principal corridors intersecting the Indian subcontinent (Trierweiler et al. 2014; Chapman et al. 2015;
510 Hu et al. 2016a; Anderson 2009). This indicates that these international flyways are not taxon-
511 specific but are instead emergent properties of the underlying atmospheric system and synoptic
512 monsoon circulation: both groups exploit favorable tailwind assistance at large scales, even though
513 the mechanisms by which they maintain heading and regulate movement differ (Alerstam 1979a;
514 Chapman et al. 2010; Chapman et al. 2015; Hu et al. 2016a).

515

516 Additionally, intra-peninsular movement is tightly coupled to agricultural phenology² and
517 spatiotemporal shifts in resource availability. Seasonal cropping cycles, host-plant dynamics, and
518 resource availability associated with monsoon progression structure the timing, magnitude, and
519 direction of dispersal in many taxa, including major agricultural pests and migratory butterflies
520 (Kunte 2005; Ramesh et al. 2012; Rajan et al. 2018; Ghosh et al. 2019; Vinayan et al. 2023). Aerial
521 transport operates on populations whose emergence, abundance, and distribution are themselves
522 seasonally structured, linking atmospheric transport processes with ecological and phenological
523 dynamics. This coupling is especially important within the Indian subcontinent, where the inland
524 penetration of the monsoon LLJ gives rise to recurrent transport corridors that redistribute these
525 seasonally generated populations. From an observational perspective also, this presents a critical
526 opportunity: the relative predictability of agricultural phenology, together with extensive agronomic
527 datasets, provides a powerful but underutilized framework for interpreting radar signals in terms of
528 species composition and ecological function, potentially making tropical aeroecological systems
529 more tractable than their temperate counterparts. The sparsity of literature therefore reflects limited
530 integration of meteorological and atmospheric processes with ecological research, rather than the
531 absence of large-scale insect movement systems, suggesting that India likely hosts far more extensive
532 and structured migratory dynamics than currently documented. In the sections that follow, I first
533 synthesize the key distinctions that emerge between temperate and tropical aeroecological systems,
534 and then outline how these differences can be explicitly investigated using the expanding network of
535 Doppler weather radars.

536

537 **5.1 Tropical vs. Temperate Aeroecology: Emerging Contrasts**

538 In temperate regions, aerial migration is typically organized around transient synoptic events,
539 producing discrete, pulse-like movements that are tightly coupled to passing weather systems
540 (Gauthreaux & Belser 2003; Liechti 2006; Van Doren & Horton 2018). In contrast, tropical monsoon
541 systems are defined by persistent but seasonally reversing circulation. Over the Indian subcontinent,
542 strong and spatially coherent southwesterly low-level jets (LLJs) during the summer monsoon
543 transition to weaker, more heterogeneous northeasterly flows in the post-monsoon period (Gadgil
544 2003; Goswami 2005; Roxy et al. 2015). This fundamental difference shifts aerial movement from
545 episodic transport windows to sustained, seasonally structured displacement where the same
546 atmospheric corridors can be reused under reversing wind regimes. During the summer monsoon,
547 LLJs can facilitate rapid, long-distance transport, with insects such as planthoppers routinely
548 advected over >1000 km within a single night within highly stratified flow (Otuka et al. 2012; Hu et
549 al. 2024). By comparison, post-monsoon movement becomes weaker and more intermittent, often
550 relying on localized synoptic enhancement and favorable nocturnal boundary-layer conditions
551 (Reynolds & Wilson 1989; Riley et al. 2013).

552 A second key contrast lies in the diel structure of the atmospheric boundary layer (ABL), which
553 governs the formation and accessibility of transport layers. Much of temperate aeroecology is

7 ²In the Indian subcontinent, agricultural phenology is structured around three primary cropping seasons: the monsoon-
8 driven *khari* season (June–October; e.g. rice, maize, cotton), the winter *rabi* season (October–March; e.g. wheat,
9 mustard, pulses), and the short pre-monsoon *zaid* season (March–June; e.g. cucurbits, fodder crops, and some pulses)

554 interpreted within a relatively canonical ABL cycle: a deep daytime convective boundary layer
555 (CBL; ~1–3 km) transitions rapidly after sunset into a shallow, strongly stratified stable boundary
556 layer (SBL; typically <0.5–0.6 km). This predictable collapse provides a consistent temporal window
557 at dusk, gating take-off, ascent, and altitude selection, and enabling migrants to exploit coherent,
558 low-turbulence nocturnal layers for efficient transport (Hu et al. 2016). In monsoon-dominated
559 tropical systems, however, this structure is far less consistent. Although the CBL similarly deepens
560 during the day, nocturnal stabilization is frequently delayed or weakened. Cloud cover, wet surfaces,
561 and persistently warm nighttime temperatures suppress radiative cooling, often delaying surface
562 decoupling by several hours after sunset (Sandeep et al. 2015; Mehta et al. 2017a,b). As a result, a
563 fully developed SBL is not a reliable nocturnal state, and the lower atmosphere may remain weakly
564 mixed or only partially stratified well into the night. Consequently, access to these transport layers is
565 contingent rather than guaranteed, and coherent vertical layering may only emerge under specific
566 combinations of stability and wind shear (Sandeep et al. 2015; Mehta et al. 2017; Algarra et al.
567 2019). This results in greater spatio-temporal variability in aerial density, vertical structuring, and
568 radar detectability across both diel and synoptic scales. These differences fundamentally alter both
569 the predictability and vertical organization of aerial movement.

570 A third distinction, largely absent from temperate systems, is the strong influence of intraseasonal
571 variability. The Monsoon Intraseasonal Oscillation (MISO), expressed through active–break cycles,
572 imposes a dominant ~10–20 day modulation on atmospheric structure. During active phases,
573 enhanced convection, deep cloud cover, and precipitation disrupt boundary-layer stratification,
574 increase turbulence, and suppress organized aerial movement. In contrast, break phases are
575 characterized by reduced convection, clearer skies, and the development of more stable and vertically
576 stratified nocturnal conditions, favoring the formation of concentrated bioscatter layers and
577 enhancing radar detectability. These cycles also modulate LLJ strength and inland penetration, as
578 well as boundary-layer depth and stability (Krishnamurti & Bhalme 1976; Sikka 2006), thereby
579 structuring migration across taxa, from locust redistribution under strengthened flow to episodic
580 hemipteran transport across the Indo-Gangetic Plain (Reynolds & Wilson 1989; Amarjyothi et al.
581 2021). Unlike temperate systems, where migration is primarily organized around short-lived synoptic
582 events, tropical movement dynamics are therefore embedded within a hierarchy of interacting
583 timescales, with intraseasonal variability exerting a dominant control on when and where movement
584 can occur.

585 Overall, these atmospheric differences translate into a stronger coupling between aerial movement
586 and ecological production in tropical systems. Tropical aerial transport operates on populations
587 whose emergence, abundance, and spatial distribution are themselves structured by monsoon-driven
588 resource dynamics. In intensively cultivated tropical landscapes, agricultural phenology, particularly
589 the timing of sowing, crop growth, and harvest cycles, further amplifies this coupling by
590 synchronizing pulses of aerial biomass with favorable transport conditions. As a result, atmospheric
591 flows do not simply redistribute organisms, but repeatedly connect temporally shifting source
592 populations with distant sink regions across the subcontinent.

593

594 5.2 India's Weather Radar Network: Observational Capacity and Research Frontiers

595 Globally, advances in dual-polarization and signal processing now enable routine separation of
596 biological and meteorological scatterers and, increasingly, discrimination among biological targets
597 (Stepanian et al. 2016; Dokter et al. 2019; Matthews et al. 2025). Opportunistic observations have
598 been largely replaced by mechanistic inferences such as relating migration intensity to boundary-
599 layer stabilization, quantifying concentration of migrants within LLJs, and forecasting migration
600 timing and routes (Shamoun-Baranes et al. 2010; Chilson et al. 2017; Van Doren & Horton 2018;
601 Horton et al. 2020). National mosaics of density, migration intensity, ground speed and flight
602 direction are available for birds and insects (Dokter et al., 2011; Chilson et al. 2017; Nilsson et al.
603 2019; Horton et al. 2020; Mungee et al. 2025). Standardized products such as vertically integrated
604 reflectivity and migration traffic rate have enabled cross-site and cross-season synthesis (Shamoun-
605 Baranes et al., 2014; Dokter et al. 2019; Lin et al. 2019; Nilsson et al., 2019).

606 India is uniquely positioned to support comparable advances. The India Meteorological Department
607 (IMD) currently operates a network of 51 weather radars, comprising 22 S-band, 19 X-band, 9 C-
608 band, and 1 Ka-band systems, of which 33 are dual-polarization (**Figure 6**). The network spans
609 almost the full climatic breadth of India, and the coverage is particularly dense along the western
610 coastline (to monitor the inland penetration of Arabian Sea monsoon flow), and across the western
611 Himalayan arc (to resolve high-elevation atmospheric processes). These regions coincide with major
612 migratory corridors and with strong gradients in elevation, habitat, and agricultural phenology,
613 creating unusual opportunities to examine how atmospheric transport, topographic forcing, and
614 source-sink population dynamics interact across both plains and complex terrain. The network is
615 expanding rapidly under the Government of India's *Mission Mausam* initiative, which aims to
616 increase aerospace coverage to >95% through ~120–130 radars by 2026–2027.

617

618 The chain of dual-polarized X-band radars along the western Himalayan arc is particularly well
619 suited to resolving insect movement within complex terrain, which currently remains almost entirely
620 uncharacterized in the region. Owing to their shorter wavelength and higher sensitivity to small
621 targets, X-band systems can detect dense fluxes of even small-bodied insects, offering a largely
622 untapped opportunity to examine windborne insect transport across mountain passes, valleys, and
623 orographically structured airflow from Jammu and Kashmir through Ladakh to Uttarakhand. By
624 contrast, avian migration across the Himalaya has been relatively well studied, particularly with
625 respect to large-scale crossings and flight strategies. However, the organization of movement within
626 the Indian subcontinent, especially post-entry redistribution across the peninsula (e.g., the contrasting
627 east- and west-coast routes in Brown-headed Gulls (Panagoda et al. 2025)), remains poorly resolved.
628 Existing studies linking movement to habitat, temperature, and precipitation (Urfi 2011; Komal et al.
629 2017; Gupta 2023; Bhagwat et al. 2025) rarely incorporate atmospheric structure explicitly, limiting
630 inference about regional-scale redistribution and its drivers. The dense network of C- and S-band
631 radars along the coastlines and major wetland systems (**Figure 6**) offers a clear pathway to resolving
632 these dynamics, building on early demonstrations of radar-based migration detection (Suresh et al.
633 2005).

634

635 These intra-peninsular radars also provide a critical opportunity to resolve the coupling between
636 atmospheric transport and ecological production, particularly the role of agricultural phenology in
637 structuring the timing, composition, and magnitude of aerial biomass entering monsoon-driven flow.
638 Integrating radar observations with trajectory modelling, ground-based sampling, and emerging tools
639 such as genomics and isotopic tracing can enable the identification of source populations, sink
640 regions, and recurrent migration corridors across the subcontinent, providing a mechanistic basis for
641 understanding redistribution processes across both ecological and agricultural systems. These
642 challenges extend across taxa and highlight the need for systems that allow direct calibration of
643 radar-derived biological signals.

644

645 Another underexplored opportunity lies in integrating bat movement ecology with India's radar
646 network. Although bats constitute a substantial component of aerial biomass, they remain
647 underrepresented in radar-based studies in the tropics, where long-distance migration is less prevalent
648 than in birds or insects (Frick et al. 2012a,b; Moussy et al. 2013; Sangavi et al. 2021; Murugavel et
649 al. 2023). However, bat emergences from large colonies generate some of the most intense, spatially
650 coherent, and temporally predictable vertebrate fluxes within the nocturnal boundary layer (Horn and
651 Kunz 2008; Chilson et al. 2012; Meade et al. 2019). These tightly constrained events produce
652 compact, high-density reflectivity plumes that are readily distinguishable from birds, insects, and
653 precipitation in weather radar data (McCracken et al. 2008; Frick et al. 2012; Vanitharani et al. 2013;
654 Werber et al. 2023). Globally, such systems have been central to calibrating and validating radar-
655 based aeroecological inference (Kunz et al. 2008; Chilson et al. 2012; Horn & Kunz 2008). In India,
656 several well-documented colonies fall within the footprint of the expanding IMD radar network,
657 presenting immediate opportunities to integrate radar observations with telemetry and colony-based
658 counts to quantify diel and seasonal variability in vertebrate aerial flux.

659

660 Despite this expanding infrastructure, a critical bottleneck remains access to and standardization of
661 radar data. Volumetric data are not consistently archived, and access to raw scans is often restricted,
662 limiting biological applications (Bhat et al. 2025). In the United States, the expansion of
663 aeroecological research followed the public availability of long-term Next Generation Weather Radar
664 (NEXRAD) archives, while in Europe the European Network for the Radar Surveillance of Animal
665 Movement (ENRAM) has enabled continental-scale analyses of major flyways. Similar progress has
666 recently emerged in Australia following open access to operational radar archives. India currently
667 lacks a centralized, accessible repository of long-term datasets, and integration with auxiliary data
668 (e.g., reanalysis winds, radiosonde profiles, ecological observations) remains limited.

669 India lacks a centralized, accessible repository of long-term raw datasets. In addition, there is limited
670 integration between radar data and auxiliary datasets such as reanalysis winds, radiosonde profiles,
671 and ecological observations, constraining mechanistic interpretation. Addressing these limitations
672 will require coordinated efforts to (i) establish open-access radar archives with standardized formats,
673 (ii) develop reproducible workflows for biological signal extraction and classification, (iii) integrate

674 radar data with atmospheric and ecological datasets, and (iv) build interdisciplinary capacity linking
675 meteorology and ecology. Given the scale and diversity of India's radar network, such developments
676 would enable unprecedented insight into the organization of aerial movement across monsoon-
677 dominated tropical systems.

678

679 **6. Conclusion**

680 By synthesizing multiple lines of evidence, this study shows that aerial movement across the Indian
681 subcontinent is structured by monsoon-driven atmospheric dynamics operating across scales. Aerial
682 movement across taxa is governed by the interaction of the same three globally fundamental sets of
683 processes: large-scale circulation, boundary-layer dynamics, and topographic forcing (Kunz et al.
684 2008; Shamoun-Baranes et al. 2010). However in monsoon-dominated tropical systems, their
685 meteorological and hence ecological expression differs in important ways. Seasonal wind reversals,
686 strong intraseasonal variability, and pronounced diel boundary-layer transitions create a distinct set of
687 ecological opportunities and constraints for dispersal, migration, and biomass transport. Recognizing
688 these differences is essential for developing a truly global understanding of aerial life. India's
689 aeroecology is fragmented across radar studies, natural history, agriculture, ornithology, entomology,
690 and meteorology. A national weather-radar-based aeroecology system would integrate these
691 disjointed bodies of knowledge and transform our understanding of aerial biodiversity. For ecologists
692 and meteorologists seeking to interpret biological signatures in Indian radar data, the current
693 literature therefore provides a useful spatiotemporal scaffold, identifying key corridors, broad
694 seasonal windows, and likely diel structure.

695

696 **Conflict of Interest**

697 The author declares no conflicts of interest.

698

699 **Author Contributions**

700 M.M. conceived the study, carried out the analyses and literature synthesis, interpreted the results,
701 wrote the manuscript, and takes responsibility for the content of the work.

702 **Funding**

703 The author acknowledges institutional support from Azim Premji University for the time and
704 resources that enabled this work.

705 **Acknowledgments**

706 I thank Dr. Subrata Kumar Das (Indian Institute of Tropical Meteorology, Pune) for generously
707 sharing dual-polarized C-band radar data from Silkheda, which will form the basis of a follow-up
708 study (in preparation). Preliminary exploration of these data was instrumental in shaping the ideas
709 presented here and motivated the need for this synthesis. I am grateful to Bill Kunin, Ryan Neely III,
710 and Christopher Hassall for introducing me to the field of aeroecology, and to Christopher Hassall
711 and Ramana Athreya for inspiring me to pursue its development in India. I also acknowledge Azim
712 Premji University, especially Gautam Pandey, Amman Madan, and Nisha Butolia, for providing
713 institutional support, time, and resources that enabled this work.

714

715 **Data Availability Statement**

716 This article is a synthesis of previously published literature and publicly available datasets. No new
717 data were generated or analyzed for this study. The locations of Doppler Weather Radars (DWRs)
718 used to illustrate the Indian radar network were obtained from the India Meteorological Department
719 (IMD) website (<https://mausam.imd.gov.in/>) through an RTI Application. All other sources used in
720 the preparation of this manuscript are cited in the reference list.

721

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Box 1. Glossary of atmospheric terms

Along-valley wind circulation: A diurnal mountain-flow system with daytime upslope and up-valley winds and nighttime downslope and down-valley drainage flows, which can align with or oppose aerial movement.

Active–break monsoon phases: Intraseasonal (~10–20 day) shifts in convection and rainfall associated with MISO/BSISO variability:

- **Active phase:** Enhanced convection and widespread rainfall over the Indian subcontinent.
- **Break phase:** Suppressed convection, clearer skies, and more coherent low-level winds.

Atmospheric boundary layer (ABL): Lowest layer of the troposphere directly influenced by surface fluxes of heat, moisture, and momentum on timescales of hours or less.

- **Convective boundary layer (CBL):** Daytime ABL driven by surface heating, characterized by buoyant turbulence and strong vertical mixing.
- **Stable boundary layer (SBL):** Nocturnal ABL driven by radiative cooling, characterized by weak turbulence, strong stratification, and limited vertical mixing.

Dryline boundary: A mesoscale boundary separating moist and dry air masses, characterized by sharp humidity gradients and often acting as a zone of convergence and uplift.

Indian Ocean Dipole (IOD): A coupled ocean–atmosphere mode defined by an east–west sea surface temperature gradient across the tropical Indian Ocean, modulating monsoon circulation, winds, and rainfall.

Inter-Tropical Convergence Zone (ITCZ): A seasonally migrating zone of low pressure and deep convection where trade winds converge; shifts northward over India during summer and southward during autumn.

Low-level jet (LLJ): A horizontally coherent wind maximum in the lowest ~1–2 km of the atmosphere (typically ~10–25 m s⁻¹ in the monsoon region), enabling rapid and concentrated atmospheric transport.

Mid-latitude westerlies: Prevailing west-to-east winds in mid-latitudes associated with synoptic disturbances and large-scale atmospheric transport.

Monsoon Intraseasonal Oscillation / Boreal Summer Intraseasonal Oscillation (MISO/BSISO): Northward-propagating (~10–30 day) intraseasonal variability in tropical convection and circulation that modulates monsoon intensity, low-level jet strength, precipitation patterns, and boundary-layer structure across the Indian subcontinent.

Nocturnal monsoon inversion: A stable temperature inversion that develops within the nocturnal boundary layer during monsoon conditions, often co-occurring with the LLJ and suppressing vertical mixing, thereby maintaining coherent transport layers.

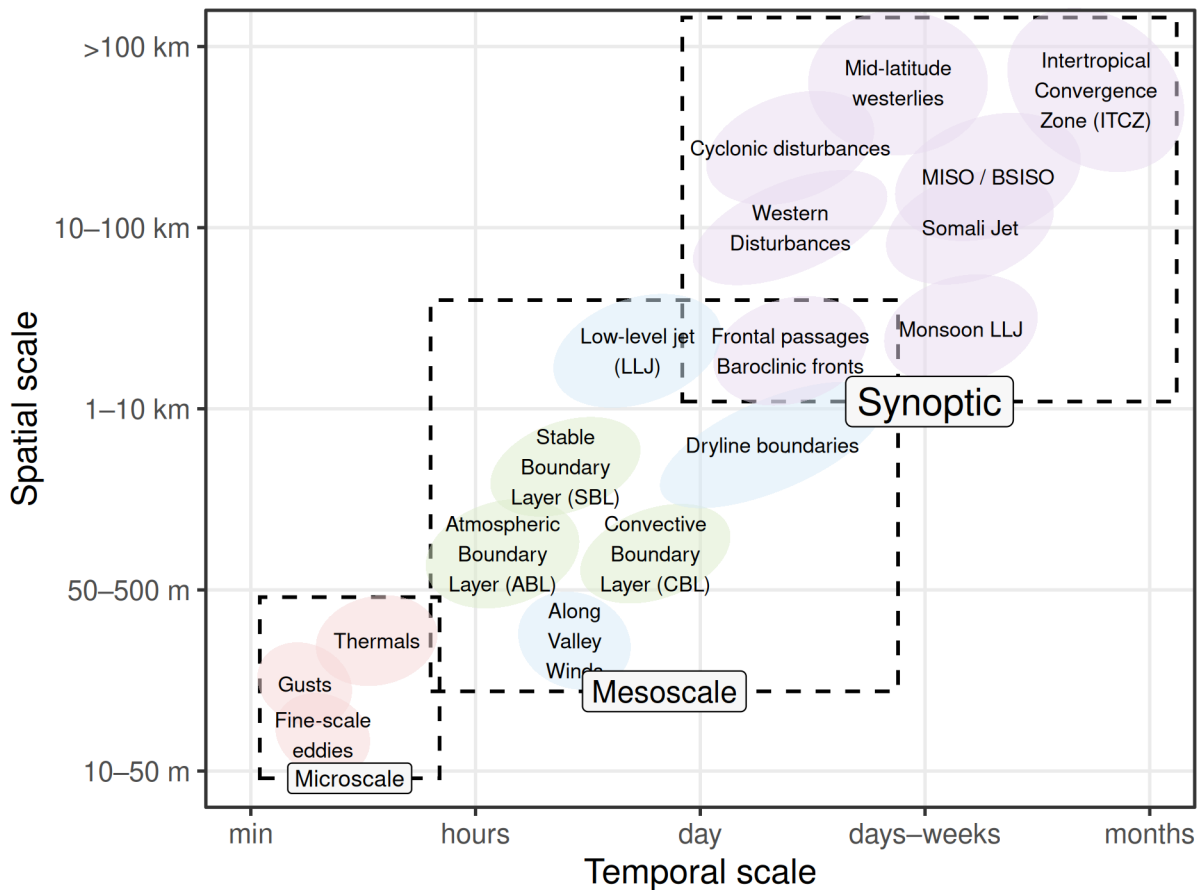
Northeast monsoon (October–November): Post-monsoon reversal to weaker northeasterly flow, restructuring windborne transport pathways across the subcontinent.

Somali Jet (Findlater Jet): A strong cross-equatorial low-level jet over the western Indian Ocean that drives southwesterly monsoon flow into India.

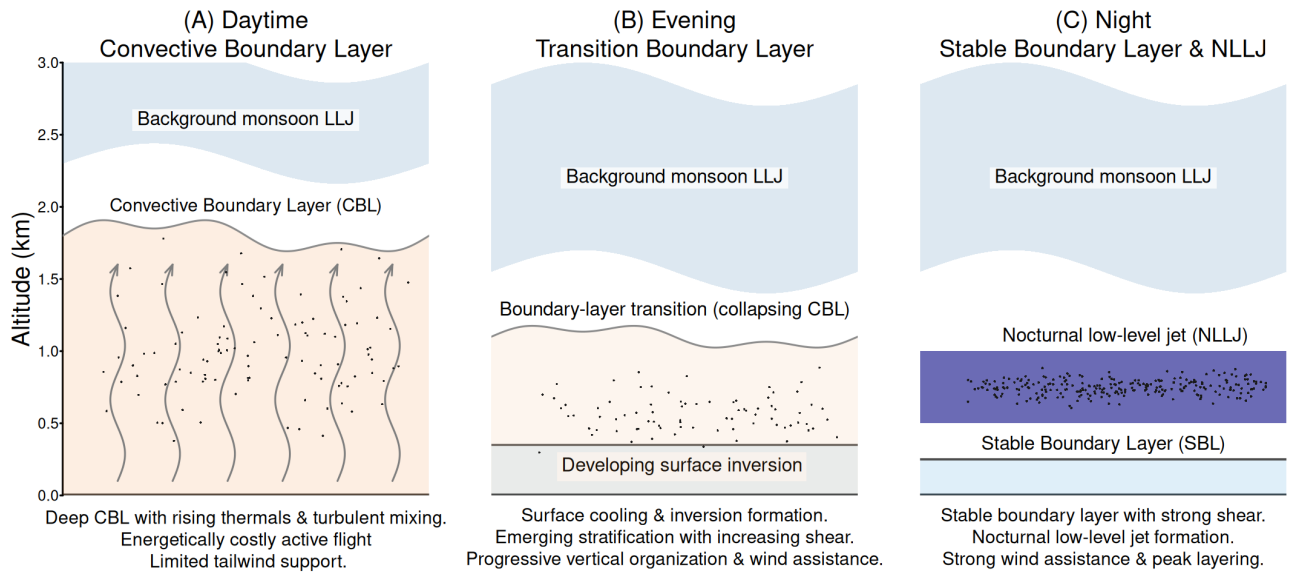
Southwest monsoon (June–September): Season of strong, persistent southwesterly low-level flow over India, often organized into high-speed transport corridors associated with the monsoon circulation.

Western Disturbances (WD): Eastward-propagating mid-latitude low-pressure systems embedded in the westerlies, originating over the Mediterranean region and influencing winter precipitation and synoptic flow over northern India.

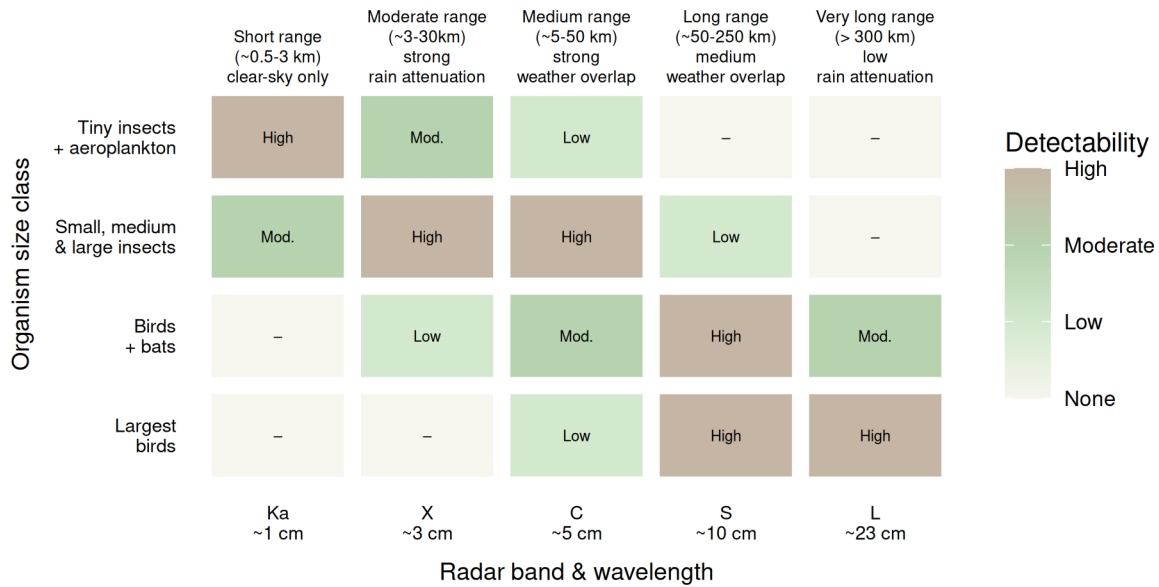
Vertical wind shear: Change in wind speed and/or direction with height; in stable conditions, shear is associated with the formation and maintenance of jets and thin, stratified transport layers.



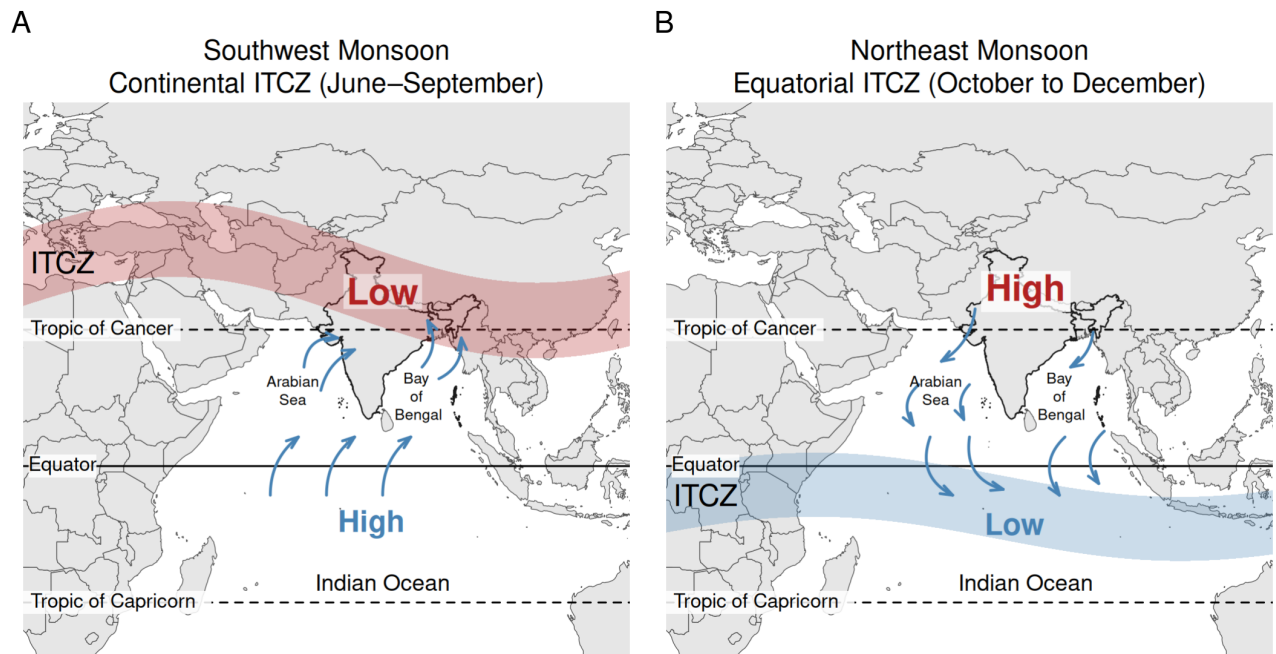
1255 **Figure 1. Spatio-temporal organization of atmospheric processes relevant to aerial movement**
 1256 **of animals.** Ellipses represent the typical temporal (x-axis) and spatial (y-axis) scales of key
 1257 atmospheric processes pertaining to aerial movement of animals. Colors distinguish four conceptual
 1258 groupings of processes (turbulence, boundary-layer, convective / shear systems, and large-scale
 1259 circulation), while dashed boxes highlight three scale regimes: microscale, mesoscale, and synoptic.
 1260 Mesoscale processes play a central role in governing take-off, departure, landing, and vertical
 1261 stratification (flight altitude), whereas large-scale migratory routes are primarily structured by
 1262 synoptic circulation. Microscale processes determine fine-scale behaviours such as foraging and local
 1263 predator–prey interactions.



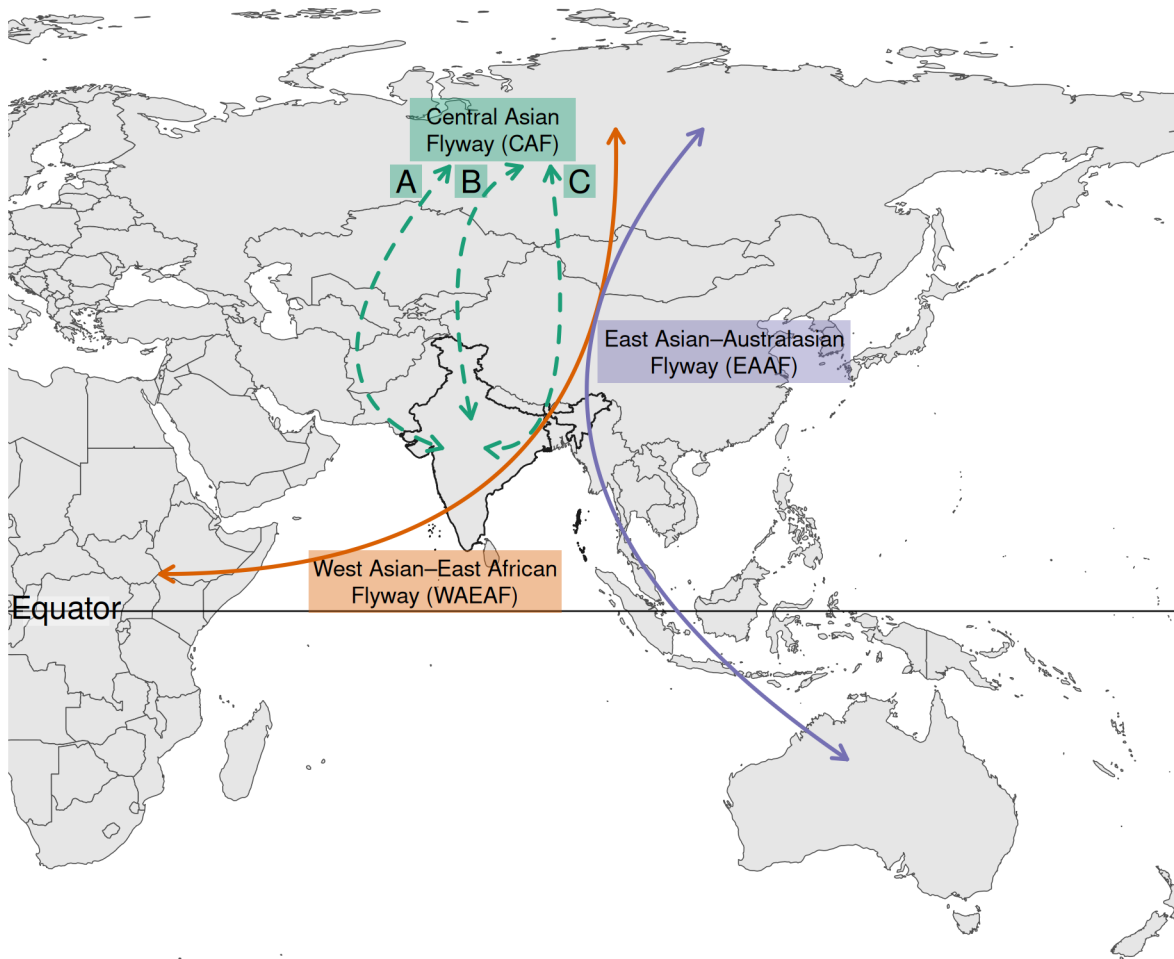
1264 **Figure 2. Diel reorganization of the troposphere and its aeroecological consequences.** The
 1265 structure of the lower atmosphere reorganizes predictably across the 24-hour cycle, generating
 1266 distinct transport regimes for aerial fauna. **(A)** During daytime, a deep convective boundary layer
 1267 (CBL) driven by surface heating produces strong turbulent mixing and weak vertical stratification.
 1268 Under these conditions, aerial organisms are broadly distributed, wind assistance is diffuse, and radar
 1269 returns are vertically widespread. **(B)** During the evening transition, surface cooling initiates
 1270 inversion formation and boundary-layer collapse, increasing vertical shear and promoting the
 1271 emergence of organized transport layers. **(C)** At night, a shallow stable boundary layer (SBL)
 1272 forms near the surface, capped by strong shear and, frequently, a nocturnal low-level jet (LLJ). This regime
 1273 generates coherent, wind-assisted transport layers above the inversion, leading to vertical
 1274 stratification of migrants and the strongest radar-defined bioscatter signatures of the diel cycle. Diel
 1275 transitions illustrate how boundary-layer physics governs not only flight energetics and directional
 1276 alignment, but also the radar-detectability and vertical structure of aerial biomass.



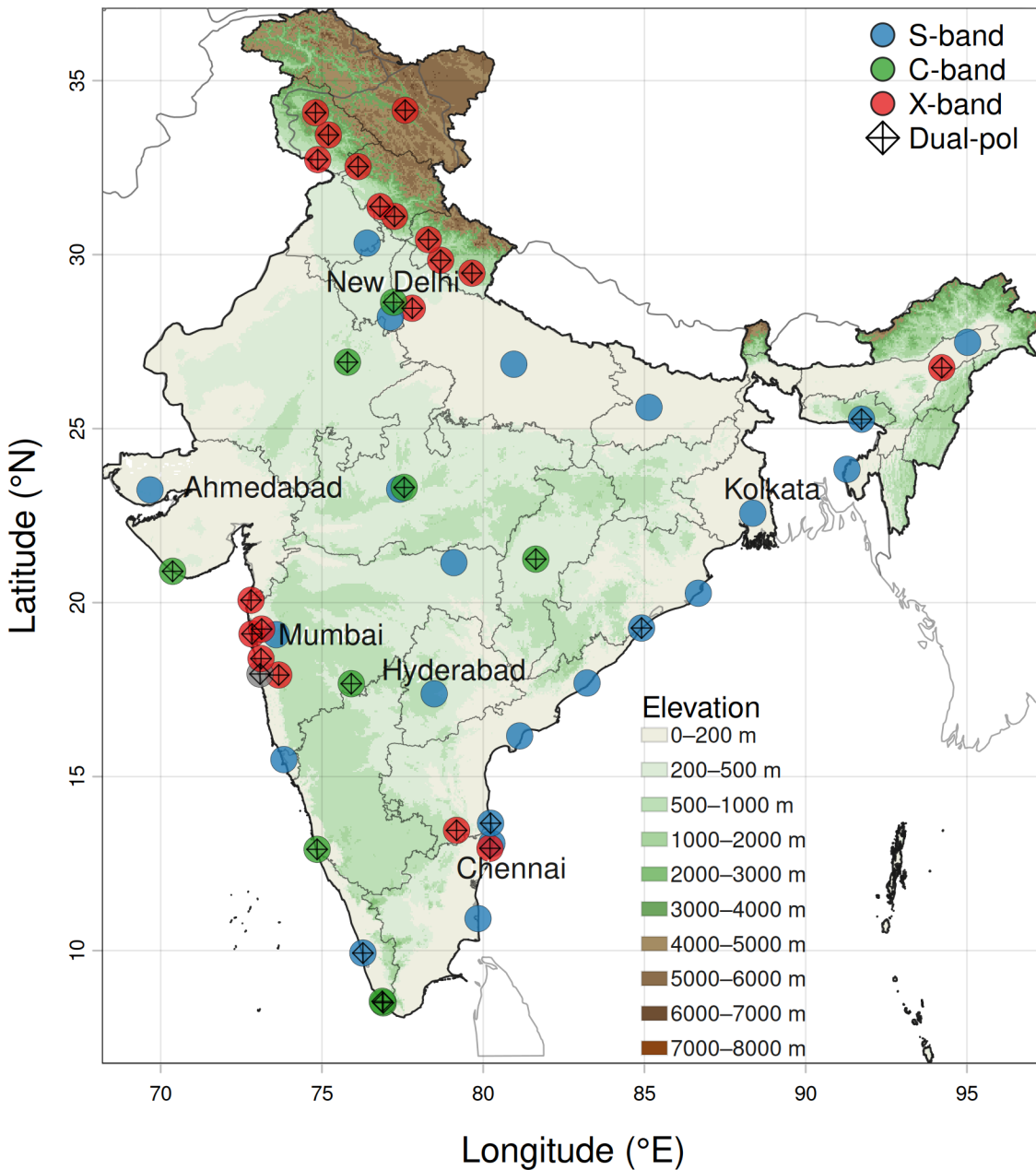
1277 **Figure 3. Radar wavelength and detectability of aerial organisms across major radar bands.**
 1278 Typical relationship between common weather radar wavelengths (x-axis) and the approximate size
 1279 of biological targets detectable in the atmosphere (y-axis). Short-wavelength radars (Ka- and X-band;
 1280 ~1–3 cm) are highly sensitive to small bioscatterers but operate over relatively short ranges and
 1281 experience strong attenuation in precipitation. Intermediate wavelengths (C-band; ~5 cm) represent a
 1282 trade-off between sensitivity and range, allowing detection of mixed bioscatter (insects, birds, and
 1283 bats) while overlapping strongly with meteorological echoes. Longer wavelengths (S- and L-band;
 1284 ~10–23 cm) experience little attenuation and enable long-range detection, but are primarily sensitive
 1285 to larger birds (see Gauthreaux Jr et al. 2019), with weak to no sensitivity to small insects. Detection
 1286 ranges shown are approximate and depend on radar power, beam geometry, atmospheric conditions,
 1287 and target density.



1288 **Figure 4. Seasonal reorganization of the Indian monsoon system. (A)** During the Southwest
 1289 Monsoon (June–September), strong cross-equatorial flow from the southern Indian Ocean intensifies
 1290 into southwesterly low-level winds over the Arabian Sea and penetrates the subcontinent. The Inter-
 1291 Tropical Convergence Zone (ITCZ) shifts northward over northern India, establishing a continental
 1292 low-pressure belt (monsoon trough) across the Indo-Gangetic Plain. This configuration promotes
 1293 sustained low-level transport pathways, particularly along the Arabian Sea and Bay of Bengal
 1294 branches, and can concentrate aerial fauna within shear zones and monsoon low-level jets. **(B)**
 1295 During the Northeast Monsoon (October–December), the circulation reverses. The ITCZ retreats
 1296 southward toward the Indian Ocean, high pressure dominates over the subcontinent, and
 1297 northeasterly surface winds prevail. This seasonal reversal alters both the direction and persistence of
 1298 wind assistance for aerial transport, restructuring potential dispersal corridors for aerial movement of
 1299 animals.



1301 **Figure 5. Monsoon-structured aerial migration corridors intersecting the Indian subcontinent.**
 1302 Three major avian flyways intersect the Indian subcontinent. The Central Asian Flyway (CAF)
 1303 comprises three principal corridors: (A) the western circum-Himalayan route, associated with soaring
 1304 migrants exploiting thermal uplift and arid convection; (B) the trans-Himalayan route, used by high-
 1305 altitude powered migrants crossing the Tibetan Plateau under favorable wind and uplift conditions;
 1306 and (C) the eastern circum-Himalayan route, linking South and Southeast Asia and partially
 1307 overlapping with the East Asia–Australasian Flyway (EAAF). Autumn movements are facilitated by
 1308 northerly and easterly flow associated with the winter monsoon, while spring movements are
 1309 supported by westerlies and pre-monsoon circulation that enable uplift and tailwind assistance across
 1310 the Himalayas. The EAAF is similarly shaped by the East Asian monsoon, whereas the West Asian–
 1311 East African Flyway (WAEAF) reflects transoceanic connections between South Asia and East
 1312 Africa mediated by post-monsoon easterlies and pre-monsoon southwesterlies linked to the Somali
 1313 Jet and the seasonal displacement of the Intertropical Convergence Zone (ITCZ). Arrows indicate
 1314 approximate migration corridors within broader flyway belts. Importantly, emerging evidence
 1315 presented in this synthesis suggests that these same atmospheric pathways also structure insect
 1316 migration and dispersal, pointing to a shared aeroecological framework across taxa.



1317 **Figure 6. Spatial distribution of India’s Doppler Weather Radar (DWR) network.** Radar sites
 1318 are coloured by frequency band (S-, C-, X-, and Ka-band), with dual-polarization systems indicated
 1319 by additional central markers. The current network comprises 51 radars, including 22 S-band, 19 X-
 1320 band, 9 C-band, and 1 Ka-band systems, of which 33 are dual-polarization. Coverage is particularly
 1321 dense along the western coastline, especially across Maharashtra, and across the western Himalayan
 1322 arc (Jammu & Kashmir, Ladakh, Himachal Pradesh, and Uttarakhand), where a chain of dual-
 1323 polarized X-band radars provides unprecedented coverage of high-altitude passes and orographically
 1324 structured airflow. These regions coincide with major migratory corridors and zones of strong
 1325 atmospheric forcing, making them especially suitable for studying aerial movement. [Ministry of
 1326 Earth Sciences, Indian Meteorological Department. (2026, March 5). *Details of operational DWR*
 1327 *network across India* (RTI Application No. IMETD/R/E/26/00078). Central Public Information
 1328 Officer, New Delhi.]