

Review

Social behavior and climate change: how rising temperatures shape insect societies

Marko Bračić^{1,*,#}, Madeleine M Ostwald^{2,*,†} and Jelena Bujan^{1,‡}

Eusocial insects are master regulators of their thermal environments, using collective behaviors to thrive in diverse and extreme climates. Nevertheless, accelerating climate warming threatens to destabilize insect societies, due to the temperature-dependence of behaviors that underpin group organization and functioning. Here, we synthesize recent advances in our understanding of temperature effects on social behavior, focusing on the eusocial bees, wasps, ants, and termites, and scaling from individual behaviors to group-level processes. We highlight evidence that warming temperatures can disrupt foraging behavior, compromise the stability of social interactions, and exceed the capacity for nest thermoregulation. To find general patterns and understand evolutionary consequences of these effects, we need more long-term studies, research that incorporates fitness measurements, and a greater focus on tropical species, as well as understudied taxa like wasps and termites. Subtle behavioral shifts could unravel finely balanced social interactions, with cascading effects on colony fitness and ultimately ecosystem functioning.

Addresses

¹ Ruder Bošković Institute, Division for Marine and Environmental Research, Bijenička cesta 54, Zagreb 10000, Croatia² Queen Mary University of London, School of Biological and Behavioural Sciences, Mile End Road, London E1 4NS, UKCorresponding author: Bujan, Jelena (jelena.bujan@irb.hr)

* Contributed equally to this work.

<https://orcid.org/0000-0001-6528-3572>† <https://orcid.org/0000-0002-9869-8835>‡ <https://orcid.org/0000-0002-7938-0266>**Current Opinion in Insect Science** 2026, **76**:101525This review comes from a themed issue on **Social insects**Edited by **Hua Yan** and **Clint Penick**For complete overview about the section, refer “[Social insects \(2026\)](#)”

Available online 6 April 2026

<https://doi.org/10.1016/j.cois.2026.101525>2214–5745/© 2026 The Author(s). Published by Elsevier Inc. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>).

Introduction

Insect declines are prevalent across the globe, and climate change is a major cause of these declines [1]. While

social insects possess important adaptations for cooperatively buffering against short-term environmental change, these benefits might fail under prolonged or extreme thermal stress [2,3]. Unlike solitary insects, which can easily find a more favorable nesting site, established social insect colonies are semi-sessile, often persisting at a single location for many years and relying on central place foraging [4]. These traits that facilitate the growth of large colonies may also limit their ability to escape unfavorable environmental conditions under climate warming. How might social insect colonies mitigate these challenges as temperatures rise? Behavior is their first line of defense against thermal stress.

To understand how thermal stress affects social behavior, we review recent evidence from eusocial bees, wasps, ants, and termites (Table S1). Insect societies span a broad spectrum of obligate and facultative social behaviors [5]. In this review, we focus primarily on effects on the coordinated colony behaviors of the obligately eusocial insects, which represent the best studied insect social groups. The bulk of this research focuses on a small subset of well-studied, typically temperate-dwelling groups (especially honey bees and bumble bees), despite the high diversity of eusocial species in the tropics.

We examine how temperature influences different facets of social behavior, given its central role in regulating metabolic, developmental, and behavioral processes in ectotherms. Although our emphasis is on rising temperatures, temperature fluctuations, including unusually cool conditions, can also negatively affect eusocial insects [6]. We first consider the impact of high temperature on individual decision-making in terms of foraging activity and efficiency. We then scale up to group-level dynamics, including aggressive and protective behaviors, as well as communication among nestmates. Finally, we explore how temperature stress affects species’ extended phenotype — the nest — which can help buffer against environmental changes. We conclude by highlighting the broader consequences of these changes for the species and for species’ interactions, including mutualisms.

Foraging

Foraging is among the most thermally stressful tasks in social insects’ repertoires, during which they must leave thermoregulated nests and perform metabolically and cognitively demanding behaviors in ambient conditions.

Insects are therefore particularly vulnerable to the acute effects of heat stress while foraging, which can translate to colony-level impacts on resource intake. Warming temperatures can restrict, expand, and shift daily and seasonal foraging windows, with responses varying widely across species and local climates [7–10, but see 11].

Higher temperatures can accelerate foraging activity by increasing flight performance and food intake speed in bees and wasps [12–14], running speed in ants [15,16], and decomposition rates by termites [17]. Despite higher foraging speeds, there is growing evidence that heat stress reduces foraging effectiveness [18]. Although bumble bees flew faster under experimental heatwaves, they foraged for shorter durations, visited fewer flowers, and collected less pollen [14,19,20]. Larger pollen loads compound the effects of heat stress in bumble bees, increasing thoracic temperature by as much as 2°C across the range of pollen load sizes [21]. Some bees can partially mitigate this risk: nectar-loaded honey bees adjust their flight mechanics to reduce metabolic rate and increase evaporative cooling [22], while hovering bumble bees increase convective cooling by self-induced airflow [23]. Higher flight speed has also been suggested to facilitate heat loss through increased convection, although this mechanism has not been directly tested (reviewed in [6,24]). Still, overall foraging efficiency seems to be reduced under higher temperatures due to thermal stress. Consequently, in-nest recovery between foraging bouts may be an important mechanism of thermal recovery [25].

Acute cognitive impacts of heat stress may impair complex navigation and memory tasks associated with foraging. Heat-stressed bumble bees perform worse at associative learning and memory tasks [26] (Figure 1a). High temperatures impair homing behavior in foraging honey bees, reducing the probability of return to the hive [27], but seem to increase homing efficiency in the temperate ant *Formica cinerea* by increasing its running speed [16]. Simultaneously, climate warming can expose brood to stressful temperatures that impair development and adult cognition, with downstream effects on foraging efficiency. Honey bees and bumble bees reared at high temperatures show deficits in short-term learning and memory as adults [28,29]. To our knowledge, similar learning experiments have not yet been done in non-flying social insects.

Aggression, recognition, and protection

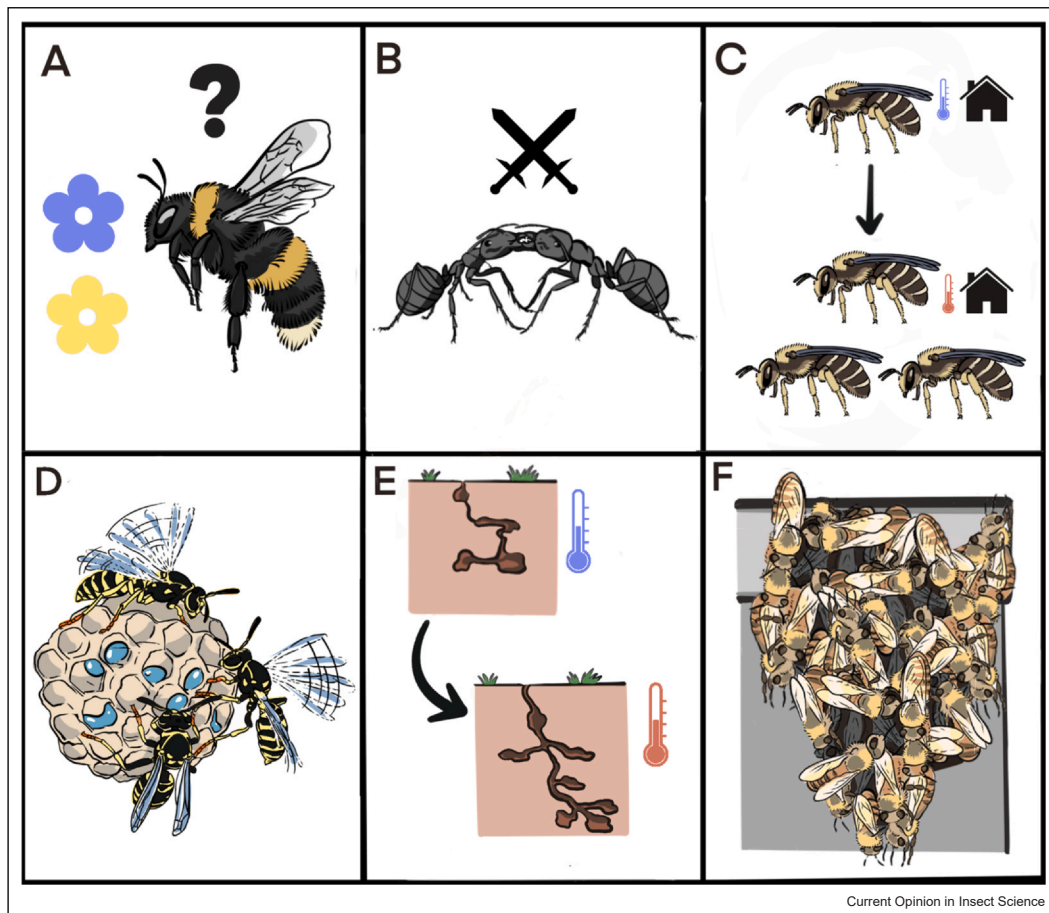
Close interactions between nestmates enable social insects to thrive across diverse habitats. By collectively directing aggression toward other colonies and species, social insects can dominate resources, raid nests, and defend their colonies. However, aggression is impacted by temperature, potentially disrupting social

organization. Elevated temperatures result in increased aggression of dominant paper wasp queens toward co-foundresses [30]. Similarly, wasps (*Polybia paulista*) and trap-jaw ants (*Odontomachus chelifer*) were aggressive toward nestmates when exposed to higher temperatures (Figure 1b), potentially due to disrupted nestmate recognition linked to changes in cuticular hydrocarbons (CHCs) [31,32]. Beyond nestmate recognition, CHCs also serve a role in cuticular waterproofing [33]. Their chemical composition can be altered by temperature and humidity in wasps, ants, and bumble bees ([34–37], for a small elevation effect in termites see [38]). In ants, CHC alterations under dry conditions are linked to higher drought resistance [35,39,40], which allows them to forage in drier conditions [41]. This potential tradeoff between beneficial physiological adaptations and impaired nestmate recognition due to rapid change in CHCs could negatively affect social interactions and aggression.

In addition to impacts on nestmate aggression, temperature influences aggression between colonies and species. In ants, aggression between colonies is higher in warmer populations of a high-elevation ant (*Tetramorium alpestre*), and in ants from warmer urban populations [42,43]. Within one ant community, temperature altered aggression toward both con- and allo-species, with some species becoming more and others less aggressive [44]. These results suggest that climate warming could reshape species interactions at the community level. However, other studies in ants and wasps have not detected an influence of temperature on aggression between colonies [45–47].

Temperature-sensitive aggression could also shape protective behavior in mutualistic relationships of social insects. For example, *Pseudomyrmex* ants defend host plants that provide extrafloral nectar and shelter, and many *Lasius* ants tend aphids that provide honeydew in exchange for protection. At higher temperatures, *Pseudomyrmex spinicola* ants were less effective at defending their bullhorn acacia host against encroaching plants [48], while four ant species were more effective at defending the whistling thorn acacia host against herbivory [49]. Under experimental warming, more ants visited pea plants but did not provide additional benefits for the plants [50]. At the community level, rising temperatures reduced the number of ant species interacting with each extrafloral-nectar-bearing plant [51]. In a cactus–ant system, species active at higher temperatures were more thermally tolerant but provided less protection, potentially reducing plant benefits and endangering these mutualisms [52]. A similar effect was observed in an ant–aphid mutualism: black garden ants (*Lasius niger*) had lower foraging efficiency, as they collected less honeydew at the highest temperature compared to the intermediate temperature, despite no decrease in ant

Figure 1



Current Opinion in Insect Science

Heat stress impacts individual- and colony-level behavior in social insects, which can result in (a) cognitive impairment [26], (b) increased aggression [42], and (c) increased cooperative behavior [68]. Social insects can mitigate heat stress through (d) evaporative cooling and fanning [56], (e) relocation of the nest to deeper, cooler soil layers [61], and (f) nest evacuation or bearding [65].

activity or aphid honeydew production [53]. These contrasting results indicate that the effect of warming could disrupt some mutualisms, but this will depend on the specific mutualistic system.

Social insects in mutualistic relationships can also suffer from the indirect effect of warming when their partner is heat-stressed and provides fewer resources. For example, plants produced fewer floral rewards at high temperatures, which reduced bee fecundity and survival [19,54]. Integrating information on thermal tolerances of mutualistic partners with the fitness costs imposed by rising temperatures would allow us to predict the consequences of disrupted mutualisms.

Colony thermoregulation and development

The semi-sessile nature of eusocial insects can be a disadvantage in a changing climate, as relocating an

entire colony is costly. Yet, despite these time and energy constraints, nest relocation is part of the life history of many social insects, and often occurs in response to seasonality or environmental stress [4]. Since colonies cannot always relocate quickly, they rely on behavioral and physiological strategies to reduce acute thermal stress. At the individual level, social insects, particularly the flying species, have an impressive repertoire of mechanisms to regulate their body temperature (reviewed in [6,24,55]). At the colony level, thermoregulation depends on coordinated social behavior. When faced with heat stress inside the nest, flying social insects will often respond with wing fanning behavior to promote convective cooling of the nest, sometimes combined with regurgitation of gut fluids and water collection to promote evaporative cooling [6,56] (Figure 1d). However, fanning becomes ineffective when ambient temperatures are too high [28], so a paper wasp species in a

warmer climate relied on it less than a species in a colder climate [56]. Evaporative cooling is a widespread thermoregulation strategy in bees and wasps ([56,57], reviewed in [28]), but there is no evidence for its use in non-flying social insects.

Colonies regulate nest temperature primarily to protect the most thermally sensitive members of the colony: its brood, including eggs, larvae, and pupae. Inability to protect brood from high temperatures can reduce egg survival, although moderate temperature increase generally speeds up development [58]. Acute heat stress might be less detrimental for ground nesters, as they can relocate brood to optimal-temperature nest areas, a task performed by specialized workers [59]. For example, during prolonged heat exposure ground-nesting ants will move the nest chambers deeper in the soil [60,61] (Figure 1e), while fungus-growing termites (genus *Macrotermes*) adjust nest architecture to meet the thermal needs of their symbiotic fungi [62]. Thus, nest-building behavior is plastic in both ants and termites, although direct evidence of thermally induced changes in nest architecture exists only for ants.

Although nests provide protection and allow for a degree of thermoregulation, the vulnerability and protective mechanisms depend on the nest type. While ground nesters are usually well buffered from climatic extremes, above-ground nests can be exposed to direct sunlight and experience much greater climatic variability [63]. Canopy nesters have limited ways to escape rising temperatures, and this is especially true for species that nest inside cavities where movement is restricted. One strategy is brood cooling. Automated tracking of European honey bees (cavity nesters) shows collective movement away from the brood area during heatwave conditions, increasing airflow to heat-sensitive brood [64]. Honey bees can also move a subset of workers outside the nest (bearding) in response to increased temperatures (Figure 1f), but prolonged bearding impairs brood development [65]. Some canopy ant species avoid detrimental temperatures by absconding the nest with brood [66]. Both bearding and absconding are energetically costly and may compromise brood development.

Cavity nesters also show plasticity in nest choice and organization. They might be able to choose a cavity that heats more slowly, such as a live twig over a dead twig, and in so doing reduce overheating. In some cavity nesters, namely ants, a single colony can occupy multiple separated nest sites. Although this polydomy of canopy ants likely evolved as an adaptation to scarce canopy nest sites [67], it might be beneficial in a warmer world.

In addition to affecting brood development and nest thermoregulation, rising temperatures may also influence the expression of sociality and ultimately fitness. In facultatively eusocial sweat bees, longer growing seasons permit the rearing of a worker brood prior to the reproductive brood, increasing the frequency of eusocial nesting [68] (Figure 1c). Direct fitness consequences of global warming are rarely investigated, but a recent study suggested that the projected increase in temperature by 2030 could decrease male production in tropical ant species by 55% [69], which directly reduces reproductive output.

Synthesis and outlook

Behavior is the first line of defense against thermal stress in social insects, with coordinated nest thermoregulatory behaviors offering a crucial lifeline against rising temperatures. Nevertheless, many essential colony behaviors are already disrupted under contemporary climate conditions. Foraging is limited under high temperatures due to reduced flight efficiency and cognitive impairment [26,27]. Warming can also unravel delicate social interactions, increasing nestmate aggression and reducing nestmate recognition, likely due to changes in CHC profiles that trade off with waterproofing adaptations [32,34]. Finally, warming can destabilize thermally-sensitive aggressive and mutualistic interactions between colonies and species [42–44,48–50]. However, mutualistic interactions remain understudied and seem to respond in a context-specific manner, making it difficult to predict how warming will affect these relationships. How and when these subtle alterations in social dynamics might precipitate larger changes in colony social cohesion remains a complex, outstanding question.

Major taxonomic and geographical gaps limit our understanding of the social effects of warming. Studies of temperate honey bees and bumble bees predominate, while the effects on stingless bees, wasps, and especially termites are still largely unknown and should be a priority for future research (Table S1). Additionally, responses of tropical social insects are underexplored even though the tropics hold the majority of global biodiversity, and the most vulnerable insect taxa [70].

Due to the urgency of these questions and practical constraints of studying warming effects across the full diversity of social insects, the focus on key behavioral traits offers a powerful framework for predicting vulnerability across species. For example, ground nesting species may be better insulated against high temperatures and can further alter the nest depth and structure (Figure 1e). In contrast, cavity or canopy nesters often respond by temporarily or permanently leaving the nest

(Figure 1f). Abscending from the nest with brood allows for exposure to predation, so arboreal nesting species might be at a higher risk, although this remains to be tested. Nest-cooling might be easier to perform for flying social insects, which use fanning and evaporative cooling strategies (Figure 1d), while non-flying ones rely more on other mechanisms, such as retreating to cooler microclimates. The energetic and fitness consequences of these differences have rarely been compared.

While evidence is growing for the acute effects of warming on individual behaviors, their colony-level impacts on survival, reproduction, and fitness are still unclear. Addressing these gaps requires long-term, multi-population studies that explicitly link behavioral responses to fitness consequences. Importantly, this research should determine how individual-level changes in behavior or interactions scale up to impact colony functioning. Microclimate availability and its use merit a stronger focus in future work, as an important strategy by which social insects can avoid overheating [63]. Further, while many studies have investigated the indirect impact of temperature, only a few have experimentally tested warming scenarios [19,25,64]. Manipulative studies that experimentally simulate warming offer powerful, underutilized tools for accelerating our understanding of these effects. More broadly, understanding how temperature reshapes the stability of eusocial societies while accounting for microclimate will be essential for predicting the persistence of these ecologically dominant groups in a rapidly warming world.

Funding

MB was supported by the European Union as part of the NextGenerationEU program: grant NPOO.C3.2.R2-I1.06.0034. JB was funded by an ERC Starting Grant (101161952 - IGNITE-ERC-2024-STG).

Data Availability

No data were used for the research described in the article.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

Acknowledgements

We thank Iva Čupić for creating accurate illustrations of social behaviors and Karl Roeder for his helpful comments on the manuscript.

Supporting information

Supplementary data associated with this article can be found in the online version at doi:10.1016/j.cois.2026.101525.

References and recommended reading

Papers of particular interest, published within the period of review, have been highlighted as:

- of special interest
- of outstanding interest

1. Wagner DL, Grames EM, Forister ML, Berenbaum MR, Stopak D: **Insect decline in the Anthropocene: death by a thousand cuts.** *Proc Natl Acad Sci* 2021, **118**:e2023989118, <https://doi.org/10.1073/pnas.2023989118>
2. Menzel F, Feldmeyer B: **How does climate change affect social insects?** *Curr Opin Insect Sci* 2021, **46**:10-15, <https://doi.org/10.1016/j.cois.2021.01.005>
3. Ostwald MM, Da Silva CRB, Seltmann KC: **How does climate change impact social bees and bee sociality?** *J Anim Ecol* 2024, **93**:1610-1621, <https://doi.org/10.1111/1365-2656.14160>
4. McGlynn TP: **The ecology of nest movement in social insects.** *Annu Rev Entomol* 2012, **57**:291-308, <https://doi.org/10.1146/annurev-ento-120710-100708>
5. Wilson EO: **The Insect Societies.** Belknap Press; 1976.
6. Perez R, Aron S: **Adaptations to thermal stress in social insects: recent advances and future directions.** *Biol Rev* 2020, **95**:1535-1553, <https://doi.org/10.1111/brv.12628>
7. Kasper ML, Reeson AF, Mackay DA, Austin AD: **Environmental factors influencing daily foraging activity of *Vespula germanica* (Hymenoptera, Vespidae) in Mediterranean Australia.** *Insectes Sociaux* 2008, **55**:288-295, <https://doi.org/10.1007/s00040-008-1004-7>
8. Wehner R, Wehner S: **Parallel evolution of thermophilia: daily and seasonal foraging patterns of heat-adapted desert ants: *Cataglyphis* and *Ocymyrmex* species.** *Physiol Entomol* 2011, **36**:271-281, <https://doi.org/10.1111/j.1365-3032.2011.00795.x>
9. Davies AB, Eggleton P, Van Rensburg BJ, Parr CL: **Seasonal activity patterns of African savanna termites vary across a rainfall gradient.** *Insectes Sociaux* 2015, **62**:157-165, <https://doi.org/10.1007/s00040-014-0386-y>
10. Jaboor SK, da Silva CRB, Kellermann V: **The effect of environmental temperature on bee activity at strawberry farms.** *Austral Ecol* 2022, **47**:1470-1479, <https://doi.org/10.1111/aec.13228>
11. Youngsteadt E, Prado SG, Keleher KJ, Kirchner M: **Can behaviour and physiology mitigate effects of warming on ectotherms? A test in urban ants.** *J Anim Ecol* 2023, **92**:568-579, <https://doi.org/10.1111/1365-2656.13860>
- The first study that measured temperature preferences of ants and combined them with field-measured operative temperatures to assess how urban and rural ant populations respond to warming.
12. Kovac H, Stabentheiner A, Brodschneider R: **Foraging strategy of wasps – optimisation of intake rate or energetic efficiency?** *J Exp Biol* 2018, **221**:jeb174169, <https://doi.org/10.1242/jeb.174169>
13. Kenna D, Pawar S, Gill RJ: **Thermal flight performance reveals impact of warming on bumblebee foraging potential.** *Funct Ecol* 2021, **35**:2508-2522, <https://doi.org/10.1111/1365-2435.13887>
14. Gérard M, Gardelin E, Lehmann P, Roberts KT, Sepúlveda-Rodríguez G, Sisquella C, Baird E: **Experimental elevated temperature affects bumblebee foraging and flight speed.** *Proc R Soc B Biol Sci* 2024, **291**:20241598, <https://doi.org/10.1098/rspb.2024.1598>
15. Roeder DV, Paraskevopoulos AW, Roeder KA: **Thermal tolerance regulates foraging behaviour of ants.** *Ecol Entomol* 2022, **47**:331-338, <https://doi.org/10.1111/een.13118>
16. Słipiński P, Cerdá X: **Higher soil temperatures cause faster running and more efficient homing in the temperate thermophilous ant *Formica cinerea* (Hymenoptera: Formicidae).** *Myrmecol News* 2022, **32**:149-158, https://doi.org/10.25849/MYRMECOL.NEWS_032:149
17. Zanne AE, Flores-Moreno H, Powell JR, Cornwell WK, Dalling JW, Austin AT, Classen AT, Eggleton P, Okada K, Parr CL, et al.:

- Termite sensitivity to temperature affects global wood decay rates.** *Science* 2022, **377**:1440-1444, <https://doi.org/10.1126/science.abo3856>
18. White SA, Dillon ME: **Climate warming and bumble bee declines: the need to consider sub-lethal heat, carry-over effects, and colony compensation.** *Front Physiol* 2023, **14**:1251235, <https://doi.org/10.3389/fphys.2023.1251235>.
This important review highlights the sublethal consequences of warming below insects' thermal limits. They discuss evidence for diverse colony and individual impacts of warming from the bumblebee literature.
19. Hemberger JA, Rosenberger NM, Williams NM: **Experimental heatwaves disrupt bumblebee foraging through direct heat effects and reduced nectar production.** *Funct Ecol* 2023, **37**:591-601, <https://doi.org/10.1111/1365-2435.14241>.
Experimental simulation of heatwaves for bumble bees (*Bombus impatiens*) foraging on oilseed rape in greenhouses. Heat-stressed bees took fewer, shorter foraging trips, which was also associated with reduced nectar production by heat-stressed plants.
20. Kuo Y, Lu Y-H, Lin Y-H, Lin Y-C, Wu Y-L: **Elevated temperature affects energy metabolism and behavior of bumblebees.** *Insect Biochem Mol Biol* 2023, **155**:103932, <https://doi.org/10.1016/j.ibmb.2023.103932>
21. Naumchik M, Youngsteadt E: **Larger pollen loads increase risk of heat stress in foraging bumblebees.** *Biol Lett* 2023, **19**:20220581, <https://doi.org/10.1098/rsbl.2022.0581>.
In field-caught foraging bumble bees, thoracic temperature increased with pollen load, rising by up to 2°C.
22. Glass JR, Burnett NP, Combes SA, Weisman E, Helbling A, Harrison JF: **Flying, nectar-loaded honey bees conserve water and improve heat tolerance by reducing wingbeat frequency and metabolic heat production.** *Proc Natl Acad Sci* 2024, **121**:e2311025121, <https://doi.org/10.1073/pnas.2311025121>
23. Glass JR, Petranek C, Dillon ME: **Induced airflow cools hovering bumble bees.** *Proc R Soc B Biol Sci* 2026, **293**:20252166, <https://doi.org/10.1098/rspb.2025.2166>
24. Glass JR, Johnson MG, Harrison JF: **When does temperature limit bee flight? Identifying the missing pieces of the puzzle.** *J Exp Biol* 2025, **228**:jeb250891, <https://doi.org/10.1242/jeb.250891>.
The review highlights key gaps in understanding how temperature limits bee flight and the thermoregulatory mechanisms individual bees use to cope with these constraints.
25. Quinlan GM, Feuerborn C, Hines HM, Grozinger CM: **Beat the heat: thermal respites and access to food associated with increased bumble bee heat tolerance.** *J Exp Biol* 2023, **226**:jeb245924, <https://doi.org/10.1242/jeb.245924>.
Experimental test of bumble bee exposure to alternating heat-stress and recovery phases. Prior heat exposure worsened bees' heat responses rather than improving them (i.e. no heat priming effect). Recovery period, duration, and food access were important for mitigating the effects of heat exposure.
26. Gérard M, Amiri A, Cariou B, Baird E: **Short-term exposure to heatwave-like temperatures affects learning and memory in bumblebees.** *Glob Change Biol* 2022, **28**:4251-4259, <https://doi.org/10.1111/gcb.16196>
27. DesJardins NS, Chester EK, Ozturk C, Lynch CM, Harrison JF, Smith BH: **Synergistic negative effects between a fungicide and high temperatures on homing behaviours in honeybees.** *Proc R Soc B Biol Sci* 2024, **291**:20240040, <https://doi.org/10.1098/rspb.2024.0040>
28. Jones JC, Oldroyd BP: **Nest thermoregulation in social insects.** *Advances in Insect Physiology*. Elsevier; 2006:153-191.
29. Perl CD, Johansen ZB, Moradinour Z, Guiraud M, Restrepo CE, Wen Jie V, Miettinen A, Baird E: **Heatwave-like events during development are sufficient to impair bumblebee worker responses to sensory stimuli.** *Front Ecol Evol* 2022, **9**:776830, <https://doi.org/10.3389/fevo.2021.776830>
30. Tibbetts EA, Reeve HK: **Two experimental tests of the relationship between group stability and aggressive conflict in Polistes wasps.** *Naturwissenschaften* 2008, **95**:383-389, <https://doi.org/10.1007/s00114-007-0336-x>
31. Michelutti KB, Batista NR, Lima-Junior SE, Cardoso CAL, Antonialli-Junior WF: **Temperature increase impairs recognition among nestmates in the social wasp *Polybia paulista* H. von Ihering, 1896 (Vespidae: Polistinae: Epiponini).** *Papéis Avulsos Zool* 2022, **62**:e202262059, <https://doi.org/10.11606/1807-0205/2022.62.059>
32. Silva KL, Batista NR, Antonialli-Junior WF: **Effect of temperature on nestmate recognition in the ant *Odontomachus chelifer*.** *Sociobiology* 2024, **71**:e9944, <https://doi.org/10.13102/sociobiology.v71i3.9944>
33. Menzel F, Morsbach S, Martens JH, Räder P, Hadjaje S, Poizat M, Abou B: **Communication versus waterproofing: the physics of insect cuticular hydrocarbons.** *J Exp Biol* 2019, **222**:jeb210807, <https://doi.org/10.1242/jeb.210807>
34. Michelutti KB, Soares ERP, Sguarizi-Antonio D, Piva RC, Suárez YR, Cardoso CAL, Antonialli-Junior WF: **Influence of temperature on survival and cuticular chemical profile of social wasps.** *J Therm Biol* 2018, **71**:221-231, <https://doi.org/10.1016/j.jtherbio.2017.11.019>
35. Baumgart L, Wittke M, Morsbach S, Abou B, Menzel F: **Why do ants differ in acclimatory ability? Biophysical mechanisms behind cuticular hydrocarbon acclimation across species.** *J Exp Biol* 2022, **225**:jeb243847, <https://doi.org/10.1242/jeb.243847>
36. Straub F, Kuppler J, Fellendorf M, Teuscher M, Vogt J, Ayasse M: **Land-use stress alters cuticular chemical surface profile and morphology in the bumble bee *Bombus lapidarius*.** *PLoS One* 2022, **17**:e0268474, <https://doi.org/10.1371/journal.pone.0268474>
37. Maihoff F, Sahler S, Schoger S, Brenzinger K, Kallnik K, Sauer N, Bofinger L, Schmitt T, Nooten SS, Classen A: **Cuticular hydrocarbons of alpine bumble bees (Hymenoptera: Bombus) are species-specific, but show little evidence of elevation-related climate adaptation.** *Front Ecol Evol* 2023, **11**:1082559, <https://doi.org/10.3389/fevo.2023.1082559>
38. Camarota F, Viana-Junior AB, Vidal DM, Zarbin PHG, Neves FS: **Changes in the cuticular profile but not aggression of termites along a tropical elevation gradient.** *Ecol Entomol* 2025, **50**:74-83, <https://doi.org/10.1111/een.13379>
39. Menzel F, Zumbusch M, Feldmeyer B: **How ants acclimate: impact of climatic conditions on the cuticular hydrocarbon profile.** *Funct Ecol* 2018, **32**:657-666, <https://doi.org/10.1111/1365-2435.13008>
40. Ostwald MM, Tretter S, Buellesbach J, Calixto JM, Fewell JH, Gadau J, Baudier KM: **Body mass and cuticular hydrocarbon profiles, but not queen number, underlie worker desiccation resistance in a facultatively polygynous harvester ant (*Pogonomyrmex californicus*).** *J Comp Physiol B* 2023, **193**:261-269, <https://doi.org/10.1007/s00360-023-01488-3>
41. Menzel F, Fiocca K, Steiner EB, Gordon DM: **Cuticular hydrocarbons and collective response to water stress in a desert ant.** *Funct Ecol* 2026, **40**:26-35, <https://doi.org/10.1111/1365-2435.70224>
42. Krapf P, Arthofer W, Ayasse M, Steiner FM, Schlick-Steiner BC: **Global change may make hostile – higher ambient temperature and nitrogen availability increase ant aggression.** *Sci Total Environ* 2023, **861**:160443, <https://doi.org/10.1016/j.scitotenv.2022.160443>.
Higher historical temperatures (and nitrogen content of workers) increased aggression toward other colonies, suggesting that global warming could intensify aggression within populations.
43. Harris BA, Stevens IADR, Mathis KA: **The effect of urbanization and temperature on thermal tolerance, foraging performance, and competition in cavity-dwelling ants.** *Ecol Evol* 2024, **14**:e10923, <https://doi.org/10.1002/ece3.10923>
44. Menges V, Rohovsky M, Rojas Feilke R, Menzel F: **Species-specific behavioural responses to environmental variation as a potential species coexistence mechanism in ants.** *Proc R Soc B Biol Sci* 2024, **291**:20240439, <https://doi.org/10.1098/rspb.2024.0439>
45. Segev U, Burkert L, Feldmeyer B, Foitzik S: **Pace-of-life in a social insect: behavioral syndromes in ants shift along a climatic**

- gradient. *Behav Ecol* 2017, **28**:1149-1159, <https://doi.org/10.1093/beheco/ax079>
46. Jandt JM, Detoni M, Loope KJ, Santoro D: **Vespula wasps show consistent differences in colony-level aggression over time and across contexts.** *Insectes Sociaux* 2020, **67**:367-381, <https://doi.org/10.1007/s00040-020-00768-3>
 47. Detoni M, Johnson SL, Adams CIM, Bengston S, Jandt JM: **Older, but not wiser: social wasp colony defensive behavior decreases with time, not experience.** *Insectes Sociaux* 2023, **70**:81-96, <https://doi.org/10.1007/s00040-022-00893-1>
 48. Shastri A, Smith A, Amador-Vargas S: **Experimentally elevated temperatures reduce activity and host defense in the acacia ant *Pseudomyrmex spinicola*.** *Behav Ecol Sociobiol* 2025, **79**:121, <https://doi.org/10.1007/s00265-025-03660-x>
 49. Tamashiro RA, Milligan PD, Palmer TM: **Left out in the cold: temperature-dependence of defense in an African ant-plant mutualism.** *Ecology* 2019, **100**:e02712, <https://doi.org/10.1002/ecy.2712>
 50. Magnoli SM, Keller KR, Lau JA: **Mutualisms in a warming world: how increased temperatures affect the outcomes of multi-mutualist interactions.** *Ecology* 2023, **104**:e3955, <https://doi.org/10.1002/ecy.3955>
 51. Câmara T, Cavalcante NT, Falcão HM, Santana E, dos Santos Silva Teixeira G, Arnan X: **Increased temperatures could heighten vulnerability of an ant-plant mutualism.** *Oecologia* 2024, **207**:8, <https://doi.org/10.1007/s00442-024-05646-4>
 52. Fitzpatrick G, Lanan MC, Bronstein JL: **Thermal tolerance affects mutualist attendance in an ant-plant protection mutualism.** *Oecologia* 2014, **176**:129-138, <https://doi.org/10.1007/s00442-014-3005-8>
 53. Blanchard S, Van Offelen J, Verheggen F, Detrain C: **Towards more intimacy: moderate elevation of temperature drives increases in foraging and mutualistic interactions between *Lasius niger* and *Aphis fabae*.** *Ecol Entomol* 2021, **46**:406-418, <https://doi.org/10.1111/een.12982>
 54. Walters J, Barlass M, Fisher R, Isaacs R: **Extreme heat exposure of host plants indirectly reduces solitary bee fecundity and survival.** *Proc R Soc B Biol Sci* 2024, **291**:20240714, <https://doi.org/10.1098/rspb.2024.0714>
 55. Lahondère C: **Recent advances in insect thermoregulation.** *J Exp Biol* 2023, **226**:jeb245751, <https://doi.org/10.1242/jeb.245751>
 56. Stabentheiner A, Nagy JM, Kovac H, Käfer H, Petrocelli I, Turillazzi S: **Effect of climate on strategies of nest and body temperature regulation in paper wasps, *Polistes biglumis* and *Polistes gallicus*.** *Sci Rep* 2022, **12**:3372, <https://doi.org/10.1038/s41598-022-07279-0>
 57. Ostwald MM, Smith ML, Seeley TD: **The behavioral regulation of thirst, water collection and water storage in honey bee colonies.** *J Exp Biol* 2016, **219**:2156-2165, <https://doi.org/10.1242/jeb.139824>
 58. Abril S, Oliveras J, Gómez C: **Effect of temperature on the development and survival of the Argentine ant, *Linepithema humile*.** *J Insect Sci* 2010, **10**:97, <https://doi.org/10.1673/031.010.9701>
 59. McGregor S, Uslu FE, Sakar MS, Keller L: **Targeted worker removal reveals a lack of flexibility in brood transport specialisation with no compensatory gain in efficiency.** *Sci Rep* 2024, **14**:4850, <https://doi.org/10.1038/s41598-024-55244-w>
 60. Sankovitz M, Purcell J: **Ant nest architecture is shaped by local adaptation and plastic response to temperature.** *Sci Rep* 2021, **11**:23053, <https://doi.org/10.1038/s41598-021-02491-w>
 61. García Ibarra F, Jouquet P, Bottinelli N, Bultelle A, Monnin T: **Experimental evidence that increased surface temperature affects bioturbation by ants.** *J Anim Ecol* 2024, **93**:319-332, <https://doi.org/10.1111/1365-2656.14040>
 62. Vesala R, Harjuntausta A, Hakkarainen A, Rönholm P, Pellikka P, Rikkinen J: **Termite mound architecture regulates nest temperature and correlates with species identities of symbiotic fungi.** *PeerJ* 2019, **6**:e6237, <https://doi.org/10.7717/peerj.6237>
 63. De Frenne P, Beugnon R, Klings D, Lenoir J, Niittynen P, Pincebourde S, Senior RA, Aalto J, Chytrý K, Gillingham PK, et al.: **Ten practical guidelines for microclimate research in terrestrial ecosystems.** *Methods Ecol Evol* 2025, **16**:269-294, <https://doi.org/10.1111/2041-210X.14476>
- Highlights the importance of microclimate in the context of the organism that is studied and recommends best practices to measure microclimates in various microhabitats.
64. Jhavar J, Davidson JD, Weidenmüller A, Wild B, Dormagen DM, Landgraf T, Couzin ID, Smith ML: **How honeybees respond to heat stress from the individual to colony level.** *J R Soc Interface* 2023, **20**:20230290, <https://doi.org/10.1098/rsif.2023.0290>
- Use of an automated tracking system to monitor the movements of thousands of individual honey bees during experimental heat stress periods. At the colony level, there was an increase in activity and collective movement away from the heat-sensitive brood.
65. Rowe EB, Prathibha P, Marting PR, Smith ML: **Honey bee beards: internal and external factors driving mass thermoregulatory evacuation.** *Insectes Sociaux* 2026, **73**:201-211, <https://doi.org/10.1007/s00040-025-01046-w>
 66. Bujan J, Yanoviak SP: **Behavioral response to heat stress of twig-nesting canopy ants.** *Oecologia* 2022, **198**:947-955, <https://doi.org/10.1007/s00442-022-05143-6>
 67. Adams BJ, Schnitzer SA, Yanoviak SP: **Connectivity explains local ant community structure in a Neotropical forest canopy: a large-scale experimental approach.** *Ecology* 2019, **100**:e02673, <https://doi.org/10.1002/ecy.2673>
 68. Schürch R, Accleto C, Field J: **Consequences of a warming climate for social organisation in sweat bees.** *Behav Ecol Sociobiol* 2016, **70**:1131-1139, <https://doi.org/10.1007/s00265-016-2118-y>
 69. Uquillas A, Bonilla N, Arizala S, Bassett Y, Barrios H, Donoso DA: **Climate drives the long-term ant male production in a tropical community.** *Sci Rep* 2025, **15**:428, <https://doi.org/10.1038/s41598-024-84789-z>
 70. Holzmann KL, Schmitzer T, Abels A, Čorkalo M, Mitesser O, Kortmann M, Alonso-Alonso P, Correa-Carmona Y, Pinos A, Yon F, et al.: **Limited thermal tolerance in tropical insects and its genomic signature.** *Nature* 2026, **651**:672-678, <https://doi.org/10.1038/s41586-026-10155-w>