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1 **Survive a Warming Climate: Insect Responses to Extreme High**  
2 **Temperatures**

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16

17 Running title: Insect Responses to Heat Extremes

18

19 **Abstract:**

20 Global change includes a substantial increase in the frequency and intensity of  
21 extreme high temperatures (EHTs) which influence insects at almost all levels. The  
22 number of studies showing the ecological importance of EHTs has risen in recent  
23 years but the knowledge is rather dispersed in the contemporary literature. Here we  
24 reviewed the biological and ecological effects of EHTs actually experienced in the  
25 field, i.e. when coupled to fluctuating thermal regimes. First, we characterized EHTs  
26 in the field. Then, we summarized the impacts of EHTs on insects at various levels  
27 and the processes allowing insects to buffer EHTs. Finally, we argue that the  
28 mechanisms leading to positive or negative impacts of EHTs on insects can only be  
29 resolved from integrative approaches considering natural thermal regimes. Thermal  
30 extremes, perhaps more than the gradual increase in mean temperature, drive insect  
31 responses to climate change, with crucial impact on pest management and biodiversity  
32 conservation.

33

34 **Keywords:**

35 climate change, extreme climate, life history trait, demographics, heat tolerance,  
36 thermoregulation

37

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39

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## 41 **1. INTRODUCTION**

42 Temperature varies across temporal and spatial scales in terrestrial and aquatic  
43 ecosystems (65, 117). Most insect species experience thermal fluctuations that can  
44 involve sublethal or lethal extreme high temperatures (EHTs) which affect  
45 physiological processes (39) and ecological systems (34). Frequent EHTs may even  
46 lead to adaptive evolution in insects (65). For decades, however, most studies  
47 concerning thermal effects on insects concentrated on the response to mean  
48 temperature or to fluctuations within the tolerable range while ignoring variations in  
49 the characteristics of EHTs (89, 105). Recently, researchers realized the biological and  
50 ecological importance of fluctuating temperature regimes but little attention is paid to  
51 the impacts of ecologically-relevant EHTs (17).

52 Global climate change includes a substantial increase in the occurrence of EHTs  
53 (1, 60), and this increase will continue in the near future (96). By contrast, the  
54 occurrence of cold extremes (cold spells, cold nights) is largely decreasing (1, 34).  
55 Thus, in a warming world, insects will be challenged mostly by elevated body  
56 temperatures (65). Neglecting daily temperature extremes generates inaccurate  
57 predictions when modeling the impacts of climate warming on populations (89). The  
58 key role of EHTs in the response of insects to warming is now attracting more  
59 attention and there is an urgent need to synthesize the disparate works. Here, we  
60 provide an overview of how natural EHTs impact insect performance at individual,  
61 population and community levels, by considering that EHTs are filtered by  
62 microhabitats and buffered by insects through behavioral thermoregulation,

63 phenotypic plasticity (e.g., acclimation), ontogeny variation, adaptive evolution and  
64 the resilience of their network (Figure 1). Our synthesis complements previous works  
65 that focused on the effects of low temperatures (125), or increase in mean temperature  
66 (119) and fluctuation regimes (26), to finally provide a comprehensive understanding  
67 of the impact of climate change on insects.

## 68 **2. HOW DO WE CHARACTERIZE THERMAL EXTREMES?**

### 69 **2.1. Definition and description of extreme high temperatures**

70 Until recently, no unified definition of EHTs existed in an environmental perspective.  
71 Meteorologists define EHTs as the temperatures above the 90th, 95th or 99th  
72 percentile of temperature distributions (1, 60), and similar indices are used by  
73 ecologists (17). The definition of EHTs can also be based on their societal impacts.  
74 The historical heat wave that hit Europe in 2003 is often used as a reference (9).  
75 Finally, another definition comes from biology, with environmental temperature  
76 exceeding upper physiological thresholds of focal organisms (96). Overall, a hybrid  
77 definition for EHTs corresponds to an “episode where climate or climate-driven  
78 conditions trigger a negative threshold-like (non-linear) biological response” (6).

79 The variables used to characterize EHTs are highly variable among studies,  
80 resulting in difficulties to compare directly their results. Most studies on EHTs are  
81 opportunistic and usually lacking replication (126). Controlled field experiments often  
82 simulate a single level of a given climate extreme imposed at a small spatial scale  
83 (126). The lack of a clear description of characteristics of EHTs limits our  
84 understanding of how such irregular EHTs affect biological systems (46, 87, 92, 122).

85 Thus, resolving the main components of EHTs becomes more meaningful than a  
86 general definition for ecological research (Figure 2). Based on field temperature data  
87 including EHTs (87), we identified several important characteristics: daily maximum  
88 and minimum temperatures which define the daily amplitude during the EHT events;  
89 the frequency of EHT events over long temporal scale (typically the number of hot  
90 days during a given period); the time elapsed between each subsequent EHT event,  
91 defining the temporal variance of extreme events. By manipulating values of each  
92 variable and their combinations, experimental designs in laboratory, mesocosms or in  
93 the field can quantify the impacts of EHTs with varying intensity, frequency, duration  
94 and temporal variance on fitness-related life history traits and demography (Figure 2).

95 In the field, insects experience temperature cycles with a single or several  
96 successive hot days alternating with a few mild days (87, 148). Diurnal cycles also  
97 alternate several hot hours around mid-day with mild hours (89, 147). Thus, the  
98 effects of EHTs depend largely on their characteristics which are complex to describe  
99 and to simulate in the lab or mesocosms due to the stochastic nature of EHTs.  
100 Intensity, duration and frequency are often used to link ecological performance and  
101 EHTs (46, 92, 122), despite the high uncertainty in these metrics. Simple statistical  
102 characteristics of EHTs were designed to explore the impact of different temperatures  
103 during daytime (89) and nighttime (147), different frequencies of hot days (86, 145)  
104 and different amplitude of fluctuation (140). During hot events, insects can be injured  
105 by EHTs approaching their physiological thresholds but the mild temperatures  
106 between hot events allow insects to repair (5, 87). The temporal sequence of hot and

107 mild phases is therefore an important characteristic of EHTs (87, 148). The impact of  
108 EHTs should be a subtle balance between heat injury and recovery (5, 87, 148).

## 109 **2.2. Thermal extreme vs. mean temperature**

110 Thermal performance curves (TPCs) are often used to describe the response of  
111 biological performance to temperature based on constant temperature treatments (124).  
112 TPCs are asymmetrical and nonlinear, thereby generating Jensen's inequality (120)  
113 which is used to describe the biological effects of environmental variability (31).  
114 TPCs, combined with Jensen's inequality, are frequently used to compute insect  
115 performance under fluctuating/changing temperatures, feeding predictions of climate  
116 change impacts (15, 72, 105, 124, 134). Based on Jensen's inequality, the effects of  
117 temperature variation would be positive, small and dramatically negative at low,  
118 intermediate and high temperatures, respectively (31, 76). The left-skewed TPCs  
119 show that the thermal optimum ( $T_{opt}$ ) is close to the critical thermal maximum ( $CT_{max}$ ),  
120 meaning that even a small variation in mean temperature above  $T_{opt}$  can push insects  
121 over  $CT_{max}$ , leading to dramatic decline in performance and fitness (105).

122 By causing an increase in thermal variance and in the strength of the Jensen's  
123 inequality, EHTs negatively impact insect fitness even if mean temperature remains  
124 near the optimal range (87, 89, 105). Indeed, the impact of increasing variance on key  
125 traits related to fitness can be larger than an increase in mean temperature (89, 105,  
126 140). Greater variance caused by EHTs can lead to decreased maximum performance,  
127 truncated thermal performance breadth and narrow thermal safety margin (105, 134).  
128 However, ambient temperature fluctuates usually with reasonable amplitude up to

129 sublethal daily maxima. Near-optimal temperature may occur during some parts of the  
130 day (e.g., morning, night) and thus provide an opportunity for insects to recover from  
131 heat injuries occurring around midday (87, 148). A better performance is therefore  
132 expected under variable regimes than under constant high temperatures. For example,  
133 temperature variation can improve rather than depress the development of aphids  
134 when the mean temperature is higher than the  $T_{opt}$  for development, thereby  
135 challenging the “Kaufmann effect” (147). Importantly, using TPCs derived from  
136 constant temperatures to predict insect fitness in field conditions contains the  
137 unrealistic assumption that performance is independent of thermal history, rates of  
138 temperature change and exposure times (124). More work on the thermal responses to  
139 the EHT characteristics will certainly help to design novel approaches to draw TPCs  
140 that could incorporate the influence of EHTs on insect performance and fitness.

### 141 **2.3. The seasonal and diurnal patterns of EHTs**

142 The amplitude of warming, including more frequent and intense extremes, is higher in  
143 spring, autumn and winter than in summer (1, 60). However, the biological impacts of  
144 EHTs in these seasons are relatively overlooked. EHTs in spring affect the  
145 phenological synchronization between plants and herbivore insects (77, 115) and  
146 parasitoids (37). EHTs in autumn may affect diapause induction, subsequent  
147 overwintering survival and spring population growth (132). EHTs in winter may  
148 influence survival during diapause and post-diapause development (11, 127).  
149 Individuals of univoltine species can experience EHTs across different seasons (118).  
150 Yet we have little information on seasonal stage-specific responses and their



151 consequences at the population level. Moreover, the amplitude of nighttime warming  
152 is higher than daytime warming (1, 60). Nighttime warming leads to contrasting  
153 effects on thermal performance and fitness, in contradiction with the predictions based  
154 on constant temperatures (147) or daytime warming (8, 89). To date, however, only  
155 few studies explored the specific effects of diurnal vs. nocturnal patterns of EHTs on  
156 insects (5, 8, 89, 147).

### 157 **3. HOW DO INSECTS RESPOND TO THERMAL EXTREMES?**

#### 158 **3.1. Physiological, biochemical and symbiont responses**

##### 159 **3.1.1. Mechanisms underlying heat damage**

160 Heat damage in insects involves a series of disorders at the molecular, biochemical  
161 and physiological levels (14, 25). Heat-induced changes in the fluidity of  
162 phospholipidic membranes and cellular homeostasis initiate heat damage (14). The  
163 OCLTT hypothesis (oxygen and capacity limitation of thermal tolerance) proposes  
164 that performance is constrained by the capacity for oxygen delivery relative to oxygen  
165 demand. The delivery of oxygen is the limiting factor during EHTs (35, 114). This  
166 hypothesis was confirmed in several aquatic species (135). However, it is not  
167 generally applicable to terrestrial insects since they are unlikely to become oxygen  
168 limited (14, 79), except if they live in close microenvironment such as plant galls  
169 (109). Finally, EHTs can disrupt the cellular ion balance (hyperkalemia), thereby  
170 impairing neurophysiological functions (103) and damaging mitochondria (14).

##### 171 **3.1.2. Physiological and molecular responses**

172 Insects can produce and accumulate particular molecules to prevent protein  
173 denaturation or cell inactivation during EHTs. Polyols, e.g. mannitol in *Aphis gossypii*  
174 and sorbitol in *Bemisia argentifolii*, are accumulated to cope with daytime EHTs (54).  
175 Mannitol and sorbitol quantities were higher at noon than during early morning.  
176 Heat-shock proteins (Hsps) (39, 85) and cuticle proteins (102) are induced and/or  
177 accumulated to deal with EHTs. Apple maggots express Hsp increasingly from  
178 midday to peak in late afternoon, paralleling the dynamics of air and fruit temperature  
179 in summer (85). Locusts are more heat-tolerant at low- than at high-latitude as a result  
180 of their expression pattern of Hsp70 and Hsp90 (22). The potato aphid increases  
181 concentration of several exoskeletal proteins during heat stress (102).

### 182 **3.1.3. Symbiont-mediated responses**

183 Bacterial endosymbionts provide essential nutrients to their insect host. EHTs depress  
184 the diversity, abundance and activity of endosymbionts, decreasing heat tolerance and  
185 fitness of the host insect (38, 70). For example, EHTs decrease the quantity of  
186 symbiotic bacteria, lowering heat tolerance in ants (38) and aphid species (142), and  
187 altering fitness-related traits in stinkbugs (70). Nevertheless, endosymbiosis may  
188 assist insect hosts to adapt to natural EHTs. In aphids, clones with heat tolerant  
189 symbionts develop faster than clones without under EHTs (52). Mutants of the  
190 symbiont *Buchnera aphidicola* with lower expression of heat-shock gene *ibpA*  
191 decrease heat tolerance of pea aphids (142). In addition, facultative symbionts, e.g.  
192 *Serratia symbiotica* and *Acetobacter* can protect *B. aphidicola* from short-term heat  
193 shock, improving aphid fecundity and offspring survival under EHTs (100).

#### 194 **3.1.4. Morphological responses**

195 EHTs can induce changes in body size of insects. High developmental temperature  
196 generally produces small individuals due to enhanced metabolism and fast  
197 development (4, 20, 131). This temperature-size rule arises due to phenotypic  
198 plasticity or selection for smaller body sizes (4, 41), although smaller individuals are  
199 more vulnerable to dehydration and overheating (44). However, small body size may  
200 facilitate heat damage in insect gut or gut symbionts, depriving nutrients and water  
201 (70). However, we know little about the growth trajectories under repeated EHTs and  
202 the underlying physiological mechanisms. For example, frequent heat exposure  
203 produces large individuals sometimes(123, 140). The selective advantage of smaller  
204 size might become disadvantageous under repeated EHTs (44).

#### 205 **3.2. Fitness-related life history traits and demography**

##### 206 **3.2.1. Stage-specific responses**

207 All stages of insects may experience EHTs. Meanwhile, different stages or instars  
208 often have different thermal sensitivities (13, 74, 146). Therefore, EHTs may produce  
209 stage-specific thermal responses. Generally, the thermal response of development,  
210 survival, reproduction and longevity varies with life stage (144) and adults are often  
211 more vulnerable to EHTs. For example, EHTs reduced more reproduction when they  
212 occurred closer to adulthood (141, 143). Both early and reproductive stages of  
213 *Sitobion avenae* (146) and *Plutella xylostella* (143) are the most sensitive to EHTs  
214 compared to mature larvae. Thus, the stage at which the insects are exposed to EHTs  
215 is an important component setting the impact of EHTs. Three potential mechanisms

216 may be involved in ontogenetic variations of thermal tolerance. 1) “Microclimate  
217 selection” hypothesis states that different stages live in and thus adapt to different  
218 thermal environments (74, 108, 139). 2) “Body size” hypothesis (75) explains why  
219 larger larvae are more heat tolerant than the smaller early life stages (86, 144, 146)  
220 which have faster increase of body temperature (3) and water loss (45), but it fails to  
221 explain the decreased tolerance at adult stage. 3) “Bogert effect” hypothesis suggests  
222 that the immobile stages (egg and pupa) should be more heat tolerant than mobile  
223 adult and larval stages due to their lower ability for behavioral thermoregulation (58),  
224 but this hypothesis has been rejected frequently (75).

### 225 **3.2.2. Trait-dependent responses**

226 Different life history traits differ in their thermal sensitivity (69). Among the main life  
227 history traits, reproduction has the narrowest thermal range, followed by development,  
228 while survival has the widest. Reproductive potential is expected to be the most  
229 vulnerable fitness-related life history traits to EHTs (136, 143). For example, a single  
230 hot event of few hours does not affect survival but decreases reproduction in the  
231 moths *Plutella xylostella* (145) and *Grapholitha molesta* (80). The optimal  
232 temperature for reproduction is lower than for development (89, 147), while the  
233 optimal temperature for immature survival is often a relatively wide range. When  
234 temperature increases continuously and surpasses the optimum, development would  
235 be restricted and soon mortality occurs (46, 63, 89), indicating slightly higher  
236 temperature thresholds for survival than for development. Thus, the critical upper  
237 thermal limits are: reproduction < development < survival (89, 147).

### 238 **3.2.3. Immediate, cross-stage and transgenerational responses**

239 The impact of EHTs occurs at various temporal scales across insect life cycles,  
240 leading to immediate impacts at the exposed stage (24, 86, 116, 140), permanent  
241 damage at later stages through carry-over effects (24, 75, 141, 144, 146), and  
242 transgenerational effects via parental effects (145). The immediate impacts incur all  
243 the molecular, physiological, and morphological mechanisms developed above and  
244 are reflected in heat death or delayed development during the events. EHTs in early  
245 life stages can influence performance of later stages or next generation depending on  
246 the intensity of EHTs, although this is still a matter of debate. Lifecycle modularity  
247 hypothesis and adaptive decoupling hypothesis (116) state that early life stress has no  
248 effects on adult performance, probably because the stressful temperatures are not  
249 extreme enough. However, there is evidence that extreme temperatures occurring in  
250 egg, larval or pupal stages can decrease survival (144) and fecundity (24, 75, 141, 146)  
251 of adults through carry-over effects, or even result in decreased offspring survival via  
252 transgenerational effects (145). Nevertheless, we still know little about the  
253 mechanisms underlying carry-over and transgenerational effects of extremes.

### 254 **3.3. Population dynamics**

#### 255 **3.3.1. Phenology and voltinism**

256 Frequent EHTs increase overall mean temperature, advancing spring and postponing  
257 autumn in the calendar, and causing shorter winters and longer growing seasons (99).  
258 For non-diapausing insects, frequent EHTs may advance phenology in spring and  
259 postpone phenology in autumn, increasing thereby the number of annual generations

260 (2, 99). However, for insects with obligate diapause, the combination of longer  
261 summers and shorter/warmer winters delays adult emergence due to insufficient  
262 winter chilling for diapause termination and to the reduced ability to regulate the  
263 onset and termination of diapause (20). For insects with facultative diapause, winter  
264 warming leads to earlier diapause termination (127). Nevertheless, EHTs do not  
265 always significantly increase the overall mean temperature and their effects on  
266 phenology and voltinism remain largely unknown for most insects. Few consecutive  
267 hot days occurring in autumn or spring would potentially disturb diapause induction  
268 or post-diapause development, thereby altering the phenology of insects.

### 269 **3.3.2. Abundance**

270 Repeated EHTs generally depress insect survival, fecundity and abundance (20, 87, 89,  
271 101, 122, 147). For example, the increase in daily maximum temperature dramatically  
272 reduces arthropod biomass in a rainforest (82). However, the effects of EHTs on  
273 insect abundance are complex due to the variations in the characteristics of EHTs,  
274 local baseline climates and habitat uses, species-specific thermal sensitivity of insects  
275 and the biotic interactions with other species. For instance, frequent EHTs depress  
276 abundance of heat sensitive species *Sitobion avenae* and *Schizaphis graminum*  
277 whereas they have null or even positive impacts on the heat tolerant species  
278 *Ropalosiphum padi* (92). Frequent EHTs reduce the abundance of *S. avenae* at the  
279 southern margin but not in the north (92). EHTs may also favor soil insects by  
280 enhancing their reproduction and abundance (81). Extremely warm winters occurring  
281 at high latitudes can increase insect abundance (48). However, EHTs in the Arctic

282 winter also accelerate snowmelt, decreasing indirectly insect abundance (11).

### 283 **3.4. Biotic interactions and community structure**

#### 284 **3.4.1. Changes in community composition and structure**

285 Differences in thermal tolerance of insects at the same trophic level can alter the  
286 relative dominance and interspecific competition among species (11, 21, 46, 92),  
287 leading to EHT-driven changes in the composition of communities (130). For example,  
288 more frequent EHTs alter the structure of cereal aphid communities across large  
289 temporal and geographical scales (92). In this case, EHTs alter the dominant pest  
290 species in winter wheat crops (92). EHT during summer affects interspecific  
291 competition in Mediterranean ant communities by changing the circadian rhythm of  
292 foraging in subdominant ants but not for dominant species (21). Insects are usually at  
293 the intermediate or lower trophic levels in communities and ecosystems. The  
294 EHT-induced changes in the composition and structure of insect communities could  
295 potentially influence top-down and bottom-up processes in plant-insect  
296 herbivore-predator systems.

#### 297 **3.4.2. Cascading effects across multitrophic systems**

298 Increasing frequency and intensity of EHTs enhances vulnerability of species at  
299 higher trophic levels (23, 101), although this rule cannot be generalized (42, 122).  
300 Thus, EHTs are likely to reduce predator diversity and disrupt ecosystem services  
301 such as natural biological control (7). Nevertheless, biodiverse multi-trophic  
302 assemblages are more resilient to EHTs (46, 61, 122). For simplified food-chains

303 including only two trophic levels such as plant-insect herbivore (46), prey-predator  
304 (101, 122) or parasitoid-hyperparasitoid systems (23), the species at higher trophic  
305 levels are more susceptible to EHTs. However, the overall effects of EHTs on  
306 tri-trophic systems such as plant-insect herbivore-predator/parasitoid are much less  
307 dramatic and often similar to patterns under normal climates (46, 122). Studies on the  
308 relationship between EHTs and the responses of each trophic level are helpful to  
309 clarify the resilience of ecological communities to changes in climate extremes.

### 310 **3.5. Geographical distributions**

311 Heat tolerance is weakly linked to latitude in insects, but tolerant species often are in  
312 warm microhabitats while heat vulnerable species remain mostly in cooler  
313 microhabitats (42, 110). Thermal safety margin, defined as the difference between the  
314  $CT_{max}$  of insects and the ambient maximum temperature of the hottest month (66,  
315 134), is frequently used to predict insect distribution following EHTs. Thermal  
316 extremes exceeding  $CT_{max}$  restrict the thermal performance range and influence the  
317 distribution across latitudes and altitudes and in heterogeneous landscapes (72, 104,  
318 134), shaping thereby the distribution of insects worldwide (66, 94). EHTs can shift  
319 distributions through local extinction induced by mass mortality due to fitness decline,  
320 to mismatch in the phenology of insects and host plants (20, 97) or to interaction with  
321 landscape fragmentation (106). However, EHTs in winter or nighttime may release the  
322 constraint of low temperatures, leading to positive effects such as increased winter  
323 survival and dispersal abilities in insects (9, 10), resulting in range expansion.  
324 Prolonged heat waves lead to an increase in number of nights above the flight



325 threshold temperature, promoting the take-off behavior in *Thaumetopoea pityocampa*  
326 and causing a rapid distribution shift to higher altitudes in the Alps (9). Some species  
327 with higher adaptive potential in heat tolerance can expand their range to lower  
328 latitudes (56).

## 329 **4. HOW DO INSECTS BUFFER AGAINST THERMAL** 330 **EXTREMES?**

### 331 **4.1. Microhabitats as filters of thermal extremes**

#### 332 **4.1.1. Mosaic of microclimates**

333 Forecasting biological impacts of EHTs requires general understanding how  
334 microhabitats filter environmental fluctuations and whether the heterogeneity will  
335 allow insects to escape heat extremes (65, 113, 129). Living and non-living objects in  
336 the environment produce a complex mosaic of microclimates which depends on the  
337 nature of the objects and how the microhabitat transfers atmospheric variations (138).  
338 Little is known, however, on the dynamics of these microclimatic mosaics during  
339 thermal extremes (111). In the apple tentiform leaf miner, the location within the tree  
340 canopy mosaic of optimal, sub-optimal and lethal micro-environments can be totally  
341 reversed during EHTs (112). The temperature gradient at micro-scale matters much  
342 for arthropods to escape EHTs. For example, in the flat-rock spider in Australia, few  
343 centimeters below hot rocks are enough to change the temperature by 9°C (107).  
344 Similarly, the temperature heterogeneity on surface of leaves can be 6 to 12°C (121).  
345 Spider mites can fully exploit this heterogeneity by moving over few centimeters but

346 this possibility is restricted during EHTs (19). Aphids are more constrained for their  
347 feeding behavior and cannot really exploit this heterogeneity (18). EHTs are expected  
348 to alter these fine-scale temperature gradients (19), depending on the baseline  
349 buffering property of the microhabitat (138).

#### 350 **4.1.2. Buffering ability of microhabitats**

351 EHTs are transmitted through microhabitat before they reach insects. Different  
352 microhabitats filter atmospheric conditions differently. For example, differences  
353 of  $>5^{\circ}\text{C}$  in the maximal microclimate temperature were measured when comparing  
354 different habitats e.g. grassland, heathland and deciduous woodland (128). The  
355 understory of forests is probably the strongest buffer of EHTs (28), which is almost  
356 independent of the dominant tree species and of latitude (29). Insects would benefit  
357 from moving to these microhabitats during EHTs although a direct effect remains to  
358 be demonstrated. Nevertheless, some insect species are specialized to microhabitats at  
359 the top canopy and the potential for vertical zonation of herbivore insects across forest  
360 canopies is important (78). The temperature of leaves hit by solar radiation is higher  
361 than ambient air temperature in temperate and boreal latitudes (113), thus the leaf  
362 surface magnifies the amplitude of thermal extremes (138). The thermal limit of small  
363 herbivore arthropods living at the leaf surface is related to leaf temperature during  
364 EHTs (110). Therefore, these herbivores are already vulnerable to contemporary  
365 thermal extremes. Similarly, endophagous insects such as leaf miners and  
366 gall-inducing organisms are living in their structures (mines, galls) where temperature  
367 increases more than ambient air during thermal extremes (108, 109). By contrast,

368 aquatic systems buffer atmospheric temperature variations due to the thermal inertia  
369 of water bodies and geophysical processes (138). In general, we lack data on the  
370 temperature of most microhabitats to provide an exhaustive ranking from the best to  
371 the worse buffer of EHTs.

## 372 **4.2. Thermoregulation**

373 Behavioral thermoregulation across the mosaic of microclimates plays an important  
374 role in avoiding EHTs (95). Thermoregulatory behavior can directly increase survival  
375 during EHTs (12, 65, 138). Nevertheless, the availability of suitable microclimates is  
376 needed for thermoregulation (138), determining the efficacy of insect buffering  
377 against climate warming (19, 65). However, it is predicted that warming will reduce  
378 seasonal and diurnal temperature ranges (60), causing decreases in temperature  
379 heterogeneity across temporal and spatial scales (65) including microhabitats (19).  
380 Thus, behavioral thermoregulation alone may not be sufficient to buffer ongoing  
381 climate warming (19).

382 Behavioral thermoregulation can also bring costs and lead to ecological  
383 consequences.. More frequent EHTs increase occurrence of thermoregulatory  
384 behaviors (90, 137) and maintenance energy costs (65), and in turn decrease foraging  
385 activity, energy efficiency and reproduction (65, 87). For example, aphids leave their  
386 host plants by dropping off or walking downward to seek cooler microhabitats to  
387 escape EHTs (86, 90, 91, 102). Therefore, they may have to endure food and water  
388 deprivation and spend time and energy to find another suitable feeding site, thereby  
389 reducing reproduction opportunities (88). Moreover, behavioral thermoregulation may

390 also alter the microhabitat use of insects (7, 88). Differences in behavioral regulations  
391 and microhabitat use between species can change interspecific interaction and food  
392 web structure (7). Furthermore, thermoregulatory behavior may limit evolutionary  
393 adaptation by reducing or even eliminating the selection pressure of EHTs (58).

### 394 **4.3. Ontogenetic variations and tradeoff between different traits**

395 Insects evolved complex life cycles with multiple developmental stages differing in  
396 thermal sensitivity (13). However, the role of ontogenetic variation of thermal  
397 sensitivity in buffering EHTs has received little attention. Different stages/age cohorts  
398 can overlap in field populations, resulting in the coexistence of multiple life  
399 stages/age cohorts during growth season. When the mixed-stage population  
400 experiences EHTs, the most sensitive stages would be injured or die whereas the most  
401 tolerant may survive and develop to adult stage for reproduction during the following  
402 mild phases, thereby contributing to population recovery. Comparing to pupal and  
403 adult stage, medium and late larval stage can particularly resist heat stress in both  
404 immediate survival and subsequent reproduction (145) Larval stages may acquire  
405 sufficient nutrients and water to both resist EHTs and provide enough time to recover  
406 from heat injury, reducing thereby the carry-over effects of EHTs on adult  
407 reproduction.

408 The intrinsic rate of increase, a metric for fitness, integrates survival, development  
409 time and reproduction (87, 89). Different responses to EHTs and even trade-off  
410 between these traits may also help buffer impacts of EHTs. Temperatures above the  
411 optimum for reproduction reduce fecundity (eg, number of offspring) but can also

412 accelerate maturation and reproduction processes (143, 147). Then, when temperature  
413 surpasses the optimum for development, insects tend to invest more energy in heat  
414 tolerance to improve survival, generating costs that reduce reproduction output (146)  
415 and delay development at least temporally (116, 140). Thus, variations in thermal  
416 sensitivity of different traits may contribute to buffer heat stress in insects.

#### 417 **4.4. Phenotypic plasticity**

418 Phenotypic plasticity is a non-genetic strategy to cope with environmental variation  
419 including thermal extremes (49, 57, 133). Most of the insect responses detailed above  
420 relate to phenotypic plasticity. Four types of plasticity in heat tolerance were  
421 identified based on the time scale and temperature intensity. “Rapid hardening/heat  
422 shock” corresponds to a short and harsh exposure to high temperature for a few  
423 minutes or hours (68, 133). The other types of thermal plasticity are all generated by  
424 exposure to moderately high temperatures for several hours to days for “acclimation”  
425 (laboratory) or “acclimatization” (field) (62, 68), days to weeks for “development  
426 acclimation/plasticity” (67, 68, 133) and weeks to months for “transgenerational  
427 acclimation” (62). In general, previous exposure to elevated temperatures increases  
428 heat tolerance both within and across generations.

429       Nevertheless, the potential to buffer against EHTs through plasticity may be  
430 limited. Quantitatively, there is limited scope for plasticity to increase heat tolerance  
431 significantly in many arthropod taxa (49, 57, 67, 133). For instance, the increases in  
432  $CT_{max}$  of some *Drosophila* species and the moth *Manduca sexta* are limited to only  
433 0.6~1.0 °C (67, 133) and 1~2 °C (73), respectively, via hardening and developmental

434 acclimation processes. In addition, increased heat tolerance brings a series of  
435 concurrent costs in other plastic traits that are related to fitness (36, 83, 133). These  
436 costs can generate considerable transgenerational consequences such as declined  
437 offspring size and survival (93).

#### 438 **4.5. Evolutionary adaptation**

439 Frequent EHTs could induce evolution of thermal tolerance to buffer against high  
440 temperature impacts (16, 57). Thermal evolution occurs in small insects during  
441 artificial selections (40, 51). Long-term thermal selection is commonly found in local  
442 adaption, e.g. drosophila and ants at different latitudes (32, 104), or beetles and  
443 butterflies at different altitudes (43, 55). Urban heat island also triggers thermal  
444 adaptation. Urban workers of ants evolved a  $CT_{max}$  1°C higher than the rural  
445 populations (32). Local adaptation often results from a subtle interaction between  
446 generation time, heritability, covariance between phenotypes, phylogenetic inertia and  
447 the strength of selection (66).

448       However, rapid evolution of thermal traits hardly occurs under natural selection in  
449 insects. Higher feeding rates evolved only after decades in nature under more frequent  
450 EHTs in *Colias eurytheme* (55). EHTs imbedded in fluctuating patterns may constrain  
451 the evolution of thermal traits due to lower heritability under ramping temperature  
452 than artificial static temperature (98), slowing down the speed at which insects can  
453 track climate warming (51). Thermal selection can also modify fitness-related life  
454 history traits (59, 71). Thermal tolerance often incurs fitness costs due to antagonistic  
455 genetic interactions between traits (84). Thermal tolerance could also be shaped by

456 phylogenetic inertia rather than thermal selection in nature (66). To some extent,  
457 thermal evolution can be a way to buffer EHTs, but its mechanisms are still evasive.

## 458 **5. KNOWLEDGE GAP AND FUTURE PERSPECTIVES**

### 459 **5.1. Ecological relevance of experimental designs**

460 The biological impacts of complex EHTs are difficult to investigate  
461 experimentally because of their stochastic nature. Open top chambers, infrared  
462 irradiation and electronic heating systems are widely used to simulate climate  
463 warming including EHTs in the field (50). However, results from these studies are  
464 difficult to apply broadly because the mechanisms are not fully understood. The  
465 different statistical moments of EHTs (intensity, duration, and frequency) may have  
466 biological impacts differing in their direction and amplitude, and their interactive  
467 effects could generate counter-intuitive responses in insects. Simulated regimes of  
468 EHTs are highly diverse in the literature, including constant temperature for given  
469 periods or fluctuations with fixed cycles. Currently, these regimes do not include  
470 alternative patterns of EHTs and mild periods which allow stress recovery (87, 147).  
471 Here we call for experimental designs with enhanced ecological relevance in future  
472 studies by incorporating the complete statistics of EHTs despite the inherent logistic  
473 complexity of factoring several variables. In addition, the simulated temperature  
474 regimes should be based on microclimatic rather than macroclimatic patterns to  
475 improve the ecological relevance. Such level of complexity is mandatory to help us to  
476 draw the landscape of insect performances under EHTs in natural thermal conditions.

477 **5.2. Enlarging our vision of extremes**

478 EHTs are by climatological definition relatively infrequent but intense (eg, heat  
479 waves). The biological perspective is more conservative in the sense that the thermal  
480 events challenging the physiology of insects are likely frequent. Environmental  
481 conditions may become suddenly challenging for organisms when several normal  
482 events co-occur leading to environmental extremes (30). This shift in perspective was  
483 only rarely applied to insects but the methodology and the concepts exist (33). Insects  
484 are constrained by several factors in their microhabitats. Dryness aggravates the  
485 physiological stress during hot periods. Wind decreases body temperature in the short  
486 term, but it incurs higher water loss. The same set of environmental conditions may be  
487 extreme for individuals deprived of appropriate endosymbionts but near optimal for  
488 those that host these microorganisms. Entomological sciences need to embrace more  
489 original and innovative characterizations of extreme events, to be combined with the  
490 standard climatological view.

491 **5.3. Rise of integrative approaches**

492 The ultimate goal of global change biologists is to anticipate the future trend of  
493 biodiversity. This goal can only be achieved by comprehensive understanding of the  
494 mechanisms for organism-environment interactions, and predicting population  
495 dynamics and community structure across time/generations. It seems crucial to link  
496 physiological mechanisms of thermal stress and the demographic variables.  
497 Physiological and genomic approaches have been successfully integrated in studies to  
498 link physiological impacts of environmental changes and field patterns of thermal



499 stress. Intertidal ecologists probably led this vision for decades (53), but  
500 entomologists now are aligned (47). The next challenge is to integrate the ‘field  
501 ecophysiology’ of thermal stress, especially during EHTs, and the population  
502 dynamics. Given the importance of individual processes (e.g., carry-over effects,  
503 hardening of thermal tolerance, and more generally phenotypic plasticity), approaches  
504 including individual-based models (27) are promising candidates to interconnect these  
505 processes and link individual and population scales. In addition, microclimate models  
506 are needed (64, 111, 112) to simulate the heterogeneous environmental arena within  
507 which individuals and species are interacting. Nevertheless, all these processes need  
508 to be fine-scaled to incorporate cross-seasonal impacts of EHTs at given moment in  
509 the life cycle of insects. Intense collaborations between physiologists, ecologists,  
510 climatologists and modelers are key for successful developments.

## 511 **6. CONCLUDING REMARKS**

512 Thermal extremes are complex events that combine different statistical moments.  
513 This complexity adds up the difficulty of getting a comprehensive understanding of  
514 their impacts on biological processes. While thermal extremes are defined as rare  
515 events by climatologists, ecologists realized that thermal extremes are much more  
516 frequent once they are defined relative to the thermal biology of organisms.  
517 Nevertheless, the impacts of ongoing climate change on these “biologically-relevant  
518 thermal extremes” remain to be identified. In this context, the most studied biological  
519 models are *Drosophila* species (39, 51, 66) and other species with conservation  
520 importance (61, 82). The thermal ecology of most species with economic importance

521 for agriculture, horticulture and forestry remains largely unknown. Extreme absence  
522 of knowledge is an important frontier limiting our ability to anticipate the ecological  
523 consequences of extreme heat events.

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## **SUMMARY POINTS**

1. Thermal extremes are complex, multicomposite events. Despite a lack of unified definition of thermal extremes due to this complexity, we can describe natural temperature extremes with several variables related to both the environment and the species thermal biology, defining together the intensity, frequency and temporal variance of extreme high temperature events.
2. Classic thermal performance curves (TPCs) derived from constant temperature experiments cannot be used to estimate the biological impacts of thermal extremes. More sophisticated TPCs are needed to include the influence of the all statistical moments of thermal extremes.
3. Extreme high temperatures often, but not always affect insects at various levels, ranging from molecular and physiological responses, at individual to population scales, to community structure and functioning.
4. Thermal extremes can affect instantaneously survival. They can also produce long-lasting impacts on life history traits and demographics of insects. The magnitude of these effects depends mainly on thermal history, traits and life-cycle stage. Collectively, all these effects translate into population dynamics.
5. Insects may buffer partly or completely the thermal extremes via behavioral thermoregulation, phenotypic plasticity, ontogeny variations, trade-off between different traits and evolutionary adaptation. However, most of these buffering strategies probably have limited potential quantitatively, and they may incur costs for other fitness-related traits.

## **FUTURE ISSUES**

1. Innovative, ecologically-relevant experimental designs are crucially needed to better understand the effects of realistic natural thermal extremes on insect populations, communities and ecosystems.
2. Beyond the instantaneous impacts of thermal extremes, the carry-over effects of heat stress and the stage-specific responses of insects deserve more comprehensive studies to better detail the consequences of thermal extremes.
3. We lack studies on the relative effects of the various strategies to buffer against thermal extremes, including physiological and behavioral thermoregulation, ontogenetic variation, phenotypic plasticity and adaptive evolution in insects. These buffering strategies may incur trade-offs with fitness-related traits, generating complex interactive effects that need to be identified to feed prediction models.
4. The mechanisms underlying heat injury and thermal recovery of insects at the molecular, physiological and ecological levels should be revealed to better comprehend the combined effects of alternating thermal extremes and mild periods.
5. The impacts of extreme high temperatures occurring in fall, winter and spring as well as at night on insect diapause, phenology, interspecific interactions and communities remain relatively understudied.
6. Microclimate models should be used more extensively to generate realistic thermal environments within which insects experience thermal extremes under field conditions in their microhabitats.
7. The thermal biology in context of thermal extremes of insect species of economic importance such as agricultural, horticultural and forestry pests, and their potential biological control agents, should be given priority to improve pest management and beneficial conservation worldwide.

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543 **LITERATURE CITED**

- 544 1. Alexander LV, Zhang X, Peterson TC, Caesar J, Gleason BE, et al. 2006. Global  
545 observed changes in daily climate extremes of temperature and precipitation. *J.*  
546 *Geophys. Res.* 111:1–22
- 547 2. Altermatt F. 2010. Climatic warming increases voltinism in European butterflies  
548 and moths. *Proc. R. Soc. B* 277:1281–87
- 549 3. Angilletta MJ. 2009. Thermal Adaptation: A Theoretical and Empirical Synthesis.  
550 Oxford, UK: Oxford Univ. Press
- 551 4. Atkinson D. 1994. Temperature and organism size: a biological law for  
552 ectotherms? *Adv. Ecol. Res.* 25:1–58
- 553 5. Bai CM, Ma G, Cai WZ, Ma CS. 2019. Independent and combined effects of  
554 daytime heat stress and nighttime recovery determine thermal performance. *Biol.*  
555 *Open* 8:bio038141
- 556 6. Bailey LD, van de Pol M. 2016. Tackling extremes: challenges for ecological and  
557 evolutionary research on extreme climatic events. *J. Anim. Ecol.* 85:85–96
- 558 7. Barton BT, Schmitz OJ. 2009. Experimental warming transforms multiple  
559 predator effects in a grassland food web. *Ecol. Lett.* 12:1317–25
- 560 8. Barton BT, Schmitz OJ. 2018. Opposite effects of daytime and nighttime  
561 warming on top-down control of plant diversity. *Ecology* 99:13–20
- 562 9. Battisti A, Stastny M, Buffo E, Larsson S. 2006. A rapid altitudinal range  
563 expansion in the pine processionary moth produced by the 2003 climatic anomaly.  
564 *Glob. Change Biol.* 12:662–71
- 565 10. Battisti A, Stastny M, Netherer S, Robinet C, Schopf A, Roques A, Larsson S.  
566 2005. Expansion of geographic range in the pine processionary moth caused by  
567 increased winter temperatures. *Ecol. Appl.* 15:2084–96
- 568 11. Bokhorst S, Phoenix GK, Bjerke JW, Callaghan TV, Huyerbrugman F, Berg MP.  
569 2012. Extreme winter warming events more negatively impact small rather than  
570 large soil fauna: shift in community composition explained by traits not taxa.  
571 *Glob. Change Biol.* 18:1152–62
- 572 12. Bonebrake TC, Boggs CL, Stamberger JA, Deutsch C, Ehrlich PR. 2014. From  
573 global change to a butterfly flapping: biophysics and behaviour affect tropical  
574 climate change impacts. *Proc. R. Soc. B* 281:20141264
- 575 13. Bowler K, Terblanche JS. 2008. Insect thermal tolerance: what is the role of  
576 ontogeny, ageing and senescence? *Biol. Rev.* 83:339–55
- 577 14. Bowler K. 2018. Heat death in poikilotherms: Is there a common cause? *J. Therm.*  
578 *Biol.* 76:77–79
- 579 15. Bozinovic F, Bastías D A, Boher F, Clavijobaquet S, Estay SA, Angilletta MJ.

- 580 2011. The mean and variance of environmental temperature interact to determine  
581 physiological tolerance and fitness. *Physiol. Biochem. Zool.* 84:543–52
- 582 16. Buckley LB, Huey RB. 2016. How extreme temperatures impact organisms and  
583 the evolution of their thermal tolerance. *Integr. Comp. Biol.* 56:98–109
- 584 17. Buckley LB, Huey RB. 2016. Temperature extremes: geographic patterns, recent  
585 changes, and implications for organismal vulnerabilities. *Glob. Change Biol.*  
586 22:3829–42
- 587 18. Cahon T, Caillon R, Pincebourde S. 2018. Do aphids alter leaf surface  
588 temperature patterns during early infestation? *Insects* 9:34
- 589 19. Caillon R, Suppo C, Casas J, Woods AH, Pincebourde S. 2014. Warming  
590 decreases thermal heterogeneity of leaf surfaces: implications for behavioural  
591 thermoregulation by arthropods. *Funct. Ecol.* 28:1449–58
- 592 20. CaraDonna PJ, Cunningham JL, Iler AM. 2018. Experimental warming in the  
593 field delays phenology and reduces body mass, fat content and survival:  
594 Implications for the persistence of a pollinator under climate change. *Funct. Ecol.*  
595 32:2345–56
- 596 21. Cerda X, Retana J, Manzaneda A. 1998. The role of competition by dominants  
597 and temperature in the foraging of subordinate species in Mediterranean ant  
598 communities. *Oecologia* 117:404–12
- 599 22. Chen B, Feder ME, Kang L. 2018. Evolution of heat-shock protein expression  
600 underlying adaptive responses to environmental stress. *Mol. Ecol.* 27:3040–54
- 601 23. Chen C, Gols R, Biere A, Harvey JA. 2019. Differential effects of climate  
602 warming on reproduction and functional responses on insects in the fourth trophic  
603 level. *Funct. Ecol.* 33:693–702
- 604 24. Chen YY, Zhang W, Ma G, Ma CS. 2019. More stressful event does not always  
605 depress subsequent life performance. *J. Integr. Agr.* 18:2321–29
- 606 25. Chown SL, Nicolson SW. 2004. *Insect Physiological Ecology: Mechanisms and*  
607 *Patterns*. Oxford, UK: Oxford Univ. Press
- 608 26. Colinet H, Sinclair BJ, Vernon P, Renault D. 2015. Insects in fluctuating thermal  
609 environments. *Annu. Rev. Entomol.* 60:123–40
- 610 27. Crespo-Pérez V, Regniere J, Chuine I, Rebaudo F, Dangles O. 2015. Changes in  
611 the distribution of multispecies pest assemblages affect levels of crop damage in  
612 warming tropical Andes. *Glob. Chang Biol.* 21:82–96
- 613 28. De Frenne P, Rodriguez-Sanchez F, Coomes DA, Baeten L, Verstraeten G, et al.  
614 2013. Microclimate moderates plant responses to macroclimate warming. *Proc.*  
615 *Natl. Acad. Sci. USA* 110:18561–65
- 616 29. De Frenne P, Zellweger F, Rodríguez-Sánchez F, Scheffers BR, Hylander K, et al.  
617 2019. Global buffering of temperatures under forest canopies. *Nat. Ecol. Evol.*

- 618 3:744–49
- 619 30. Denny M, Hunt L, Miller L, Harley C. 2009. On the prediction of extreme  
620 ecological events. *Ecol. Monogr.* 79:397–421
- 621 31. Denny M. 2017. The fallacy of the average: on the ubiquity, utility and continuing  
622 novelty of Jensen's inequality. *J. Exp. Biol.* 220:139–46
- 623 32. Diamond SE. 2017. Evolutionary potential of upper thermal tolerance:  
624 biogeographic patterns and expectations under climate change. *Ann. NY Acad. Sci.*  
625 1389:5–19
- 626 33. Dowd WW, Denny MW. 2020. A series of unfortunate events: characterizing the  
627 contingent nature of physiological extremes using long-term environmental  
628 records. *Proc. R. Soc. B* 287:20192333
- 629 34. Easterling DR, Meehl GA, Parmesan C, Changnon SA, Karl TR, Mearns LO.  
630 2000. Climate extremes: observations, modeling, and impacts. *Science*  
631 289:2068–74
- 632 35. Ern R. 2019. A mechanistic oxygen- and temperature-limited metabolic niche  
633 framework. *Philos. Trans. R. Soc. B* 374:20180540
- 634 36. Esperk T, Kjaersgaard A, Walters RJ, Berger D, Blanckenhorn WU. 2016. Plastic  
635 and evolutionary responses to heat stress in a temperate dung fly: negative  
636 correlation between basal and induced heat tolerance? *J. Evol. Biol.* 29:900–15
- 637 37. Evans EW, Carlile NR, Innes MB, Pitigala N. 2013. Warm springs reduce  
638 parasitism of the cereal leaf beetle through phenological mismatch. *J. Appl.*  
639 *Entomol.* 137:383–91
- 640 38. Fan YL, Wernegreen JJ. 2013. Can't take the heat: high temperature depletes  
641 bacterial endosymbionts of ants. *Microb. Ecol.* 66:727–33
- 642 39. Feder ME, Blair NT, Figueras H. 1997. Natural thermal stress and heat-shock  
643 protein expression in *Drosophila* larvae and pupae. *Funct. Ecol.* 11:90–100
- 644 40. Folk DG, Zwollo P, Rand DM, Gilchrist GW. 2006. Selection on knockdown  
645 performance in *Drosophila melanogaster* impacts thermotolerance and  
646 heat-shock response differently in females and males. *J. Exp. Biol.* 209:3964–73
- 647 41. Forster J, Hirst AG, Atkinson D. 2012. Warming-induced reductions in body size  
648 are greater in aquatic than terrestrial species. *Proc. Natl. Acad. Sci. USA*  
649 109:19310–14
- 650 42. Franken O, Huizinga M, Ellers J, Berg MP. 2018. Heated communities: Large  
651 inter- and intraspecific variation in heat tolerance across trophic levels of a soil  
652 arthropod community. *Oecologia* 186:311–22
- 653 43. García-Robledo C, Kuprewicz EK, Staines CL, Erwin TL, Kress WJ. 2016.  
654 Limited tolerance by insects to high temperatures across tropical elevational  
655 gradients and the implications of global warming for extinction. *Proc. Natl. Acad.*

- 656 *Sci. USA* 113:680–85
- 657 44. Gardner JL, Peters A, Kearney MR, Joseph L, Heinsohn R. 2011. Declining body  
658 size: A third universal response to warming? *Trends Ecol. Evol.* 26:285–91
- 659 45. Gibbs AG, Louie AK, Ayala JA. 1998. Effects of temperature on cuticular lipids  
660 and water balance in a desert *Drosophila*: is thermal acclimation beneficial? *J.*  
661 *Exp. Biol.* 201:71–80
- 662 46. Gillespie DR, Nasreen A, Moffat CE, Clarke P, Roitberg BD. 2012. Effects of  
663 simulated heat waves on an experimental community of pepper plants, green  
664 peach aphids and two parasitoid species. *Oikos* 121:149–59
- 665 47. Giron D, Dubreuil G, Bennett A, Dedeine F, Dicke M, et al. 2018. Promises and  
666 challenges in insect-plant interactions. *Entomol. Exp. Appl.* 166:319–43
- 667 48. Goodsmann DW, Grosklos G, Aukema BH, Whitehouse C, Bleiker KP, et al. 2018.  
668 The effect of warmer winters on the demography of an outbreak insect is hidden  
669 by intraspecific competition. *Glob. Change Biol.* 24:3620–28
- 670 49. Gunderson AR, Stillman JH. 2015. Plasticity in thermal tolerance has limited  
671 potential to buffer ectotherms from global warming. *Proc. R. Soc. B*  
672 282:20150401
- 673 50. Halbritter AH, De Boeck HJ, Eycott AE, Reinsch S, Robinson DA, et al. 2020.  
674 The handbook for standardized field and laboratory measurements in terrestrial  
675 climate change experiments and observational studies (ClimEx). *Methods Ecol.*  
676 *Evol.* 11:22–37
- 677 51. Hangartner S, Hoffmann AA. 2016. Evolutionary potential of multiple measures  
678 of upper thermal tolerance in *Drosophila melanogaster*. *Funct. Ecol.* 30:442–52
- 679 52. Harmon JP, Moran NA, Ives AR. 2009. Species response to environmental  
680 change: impacts of food web interactions and evolution. *Science* 323:1347–50
- 681 53. Helmuth B, Kingsolver JG, Carrington E. 2005. Biophysics, physiological  
682 ecology, and climate change: Does mechanism matter? *Annu. Rev. Physiol.*  
683 67:177–201
- 684 54. Hendrix DL, Salvucci ME. 1998. Polyol metabolism in homopterans at high  
685 temperatures: accumulation of mannitol in aphids (Aphididae: Homoptera) and  
686 sorbitol in whiteflies (Aleyrodidae: Homoptera). *Comp. Biochem. Physiol. A*  
687 *Comp. Physiol.* 120:487–94
- 688 55. Higgins JK, MacLean HJ, Buckley LB, Kingsolver JG. 2014. Geographic  
689 differences and microevolutionary changes in thermal sensitivity of butterfly  
690 larvae in response to climate. *Funct. Ecol.* 28:982–89
- 691 56. Hill MP, Chown SL, Hoffmann AA. 2013. A predicted niche shift corresponds  
692 with increased thermal resistance in an invasive mite, *Halotydeus destructor*.  
693 *Glob. Ecol. Biogeogr.* 22:942–51



- 694 57. Hoffmann AA, Chown SL, Clusella-Trullas S. 2013. Upper thermal limits in  
695 terrestrial ectotherms: how constrained are they? *Funct. Ecol.* 27:934–49
- 696 58. Huey RB, Hertz PE, Sinervo B. 2003. Behavioral drive versus behavioral inertia  
697 in evolution: a null model approach. *Am. Nat.* 161:357–66
- 698 59. Huey RB, Kingsolver JG. 1989. Evolution of thermal sensitivity of ectotherm  
699 performance. *Trends Ecol. Evol.* 4:131–35
- 700 60. IPCC. Summary for policymakers. Climate Change 2013: The Physical Science  
701 Basis Contribution of Working Group I to the Fifth Assessment Report of the  
702 Intergovernmental Panel on Climate Change. Cambridge, UK: Cambridge Univ.  
703 Press
- 704 61. Isbell F, Craven D, Connolly J, Loreau M, Schmid B, et al. 2015. Biodiversity  
705 increases the resistance of ecosystem productivity to climate extremes. *Nature*  
706 526:574–77
- 707 62. Jensen A, Alemu T, Alemneh T, Pertoldi C, Bahrndorff S, et al. 2019. Thermal  
708 acclimation and adaptation across populations in a broadly distributed soil  
709 arthropod. *Funct. Ecol.* 33:833–45
- 710 63. Karl I, Stoks R, De Block M, Janowitz SA, Fischer K. 2011. Temperature  
711 extremes and butterfly fitness: conflicting evidence from life history and immune  
712 function. *Glob. Change Biol.* 17:676–87
- 713 64. Kearney M, Porter WP. 2016. NicheMapR - an R package for biophysical  
714 modelling: the microclimate model. *Ecography* 40:664–74
- 715 65. Kearney M, Shine R, Porter WP. 2009. The potential for behavioral  
716 thermoregulation to buffer “cold-blooded” animals against climate warming. *Proc.*  
717 *Natl. Acad. Sci. USA* 106:3835–40
- 718 66. Kellermann V, Overgaard J, Hoffmann AA, Flojgaard C, Svenning JC, Loeschke  
719 V. 2012. Upper thermal limits of *Drosophila* are linked to species distributions  
720 and strongly constrained phylogenetically. *Proc. Natl. Acad. Sci. USA*  
721 109:16228–33
- 722 67. Kellermann V, Sgro CM. 2018. Evidence for lower plasticity in CTMAX at  
723 warmer developmental temperatures. *J. Evol. Biol.* 31:1300–12
- 724 68. Kellermann V, van Heerwaarden B, Sgro CM. 2017. How important is thermal  
725 history? Evidence for lasting effects of developmental temperature on upper  
726 thermal limits in *Drosophila melanogaster*. *Proc. R. Soc. B* 284:20170447
- 727 69. Kellermann V, van Heerwaarden B. 2019. Terrestrial insects and climate change:  
728 adaptive responses in key traits. *Physiol. Entomol.* 44:99–115
- 729 70. Kikuchi Y, Tada A, Musolin DL, Hari N, Hosokawa T, Fujisaki K, Fukatsu T.  
730 2016. Collapse of insect gut symbiosis under simulated climate change. *MBio*  
731 7:e01578-16

- 732 71. Kingsolver JG, Buckley LB. 2017. Quantifying thermal extremes and biological  
733 variation to predict evolutionary responses to changing climate. *Philos. Trans. R.  
734 Soc. B* 372:20160147.
- 735 72. Kingsolver JG, Diamond SE, Buckley LB. 2013. Heat stress and the fitness  
736 consequences of climate change for terrestrial ectotherms. *Funct. Ecol.*  
737 27:1415–23
- 738 73. Kingsolver JG, Maclean HJ, Goddin SB, Augustine KE. 2016. Plasticity of upper  
739 thermal limits to acute and chronic temperature variation in *Manduca sexta* larvae.  
740 *J. Exp. Biol.* 219:1290–94
- 741 74. Kingsolver JG, Woods A, Buckley LB, Potter KA, Maclean HJ, Higgins JK.  
742 2011 Complex life cycles and the responses of insects to climate change. *Integr.*  
743 *Comp. Biol.* 51:719–32
- 744 75. Klockmann M, Günter F, Fischer K. 2017. Heat resistance throughout ontogeny:  
745 body size constrains thermal tolerance. *Glob. Change Biol.* 23:686–96
- 746 76. Koussoroplis AM, Pincebourde S, Wacker A. 2017. Understanding and predicting  
747 physiological performance of organisms in fluctuating and multifactorial  
748 environments. *Ecol. Monogr.* 87:178–97
- 749 77. Kudo G, Ida TY. 2013. Early onset of spring increases the phenological mismatch  
750 between plants and pollinators. *Ecology* 94:2311–20
- 751 78. Lawton JH. 1983. Plant architecture and the diversity of phytophagous insects.  
752 *Annu. Rev. Entomol.* 28:23–29
- 753 79. Lehmann P, Javal M, Terblanche JS. 2019. Oxygen limitation is not the cause of  
754 death during lethal heat exposure in an insect. *Biol. Lett.* 15:20180701
- 755 80. Liang LN, Zhang W, Ma G, Hoffmann A, Ma CS. 2014. A single hot event  
756 stimulates adult performance but reduces egg survival in the oriental fruit moth,  
757 *Grapholitha molesta*. *PLoS ONE* 9:e116339
- 758 81. Lindo Z. 2015. Warming favours small-bodied organisms through enhanced  
759 reproduction and compositional shifts in belowground systems. *Soil Biol.*  
760 *Biochem.* 91:271–78
- 761 82. Lister BC, Garcia A. 2018. Climate-driven declines in arthropod abundance  
762 restructure a rainforest food web. *Proc. Natl. Acad. Sci. USA* 115:E10397–406
- 763 83. Loeschcke V, Hoffmann AA. 2007. Consequences of heat hardening on a field  
764 fitness component in *Drosophila* depend on environmental temperature. *Am. Nat.*  
765 169:175–83
- 766 84. Loeschcke V, Krebs RA. 1996. Selection for heat-shock resistance in larval and in  
767 adult *Drosophila buzzatii*: comparing direct and indirect responses. *Evolution*  
768 50:2354–59
- 769 85. Lopez-Martinez G, Denlinger DL. 2008. Regulation of heat shock proteins in the

- 770 apple maggot *Rhagoletis pomonella* during hot summer days and overwintering  
771 diapause. *Physiol. Entomol.* 33:346–52
- 772 86. Ma CS, Hau B, Poehling HM. 2004. The effect of heat stress on the survival of  
773 the rose grain aphid, *Metopolophium dirhodum* (Hemiptera: Aphididae). *Eur. J.*  
774 *Entomol.* 101:327–31
- 775 87. Ma CS, Wang L, Zhang W, Rudolf V. 2018. Resolving biological impacts of  
776 multiple heat waves: interaction of hot and recovery days. *Oikos* 127:622–33
- 777 88. Ma G, Bai CM, Wang XJ, Majeed MZ, Ma CS. 2018. Behavioural  
778 thermoregulation alters microhabitat utilization and demographic rates in  
779 ectothermic invertebrates. *Anim. Behav.* 142:49–57
- 780 89. Ma G, Hoffmann AA, Ma CS. 2015. Daily temperature extremes play an  
781 important role in predicting thermal effects. *J. Exp. Biol.* 218:2289–96
- 782 90. Ma G, Ma CS. 2012. Climate warming may increase aphids' dropping  
783 probabilities in response to high temperatures. *J. Insect Physiol.* 58:1456–62
- 784 91. Ma G, Ma CS. 2012. Effect of acclimation on heat-escape temperatures of two  
785 aphid species: implications for estimating behavioral response of insect to climate  
786 warming. *J. Insect Physiol.* 58:303–309
- 787 92. Ma G, Rudolf V, Ma CS. 2015. Extreme temperature events alter demographic  
788 rates, relative fitness, and community structure. *Glob. Change Biol.*  
789 21:1794–1808
- 790 93. Macagno ALM, Zattara EE, Ezeakudo O, Moczek AP, Ledonrettig C. 2018.  
791 Adaptive maternal behavioral plasticity and developmental programming mitigate  
792 the transgenerational effects of temperature in dung beetles. *Oikos* 127:1319–29
- 793 94. Martinet B, Lecocq T, Smet J, Rasmont P. 2015. A protocol to assess insect  
794 resistance to heat waves, applied to bumblebees (*Bombus Latreille*, 1802). *PLoS*  
795 *ONE* 10:e0118591
- 796 95. May ML. 1979. Insect thermoregulation. *Annu. Rev. Entomol.* 24:313–49
- 797 96. Meehl G A, Tebaldi C. 2004. More intense, more frequent, and longer lasting heat  
798 waves in the 21st century. *Science* 305:994–97
- 799 97. Merrill RM, Gutierrez D, Lewis OT, Gutierrez J, Diez SB, Wilson RJ. 2008.  
800 Combined effects of climate and biotic interactions on the elevational range of a  
801 phytophagous insect. *J. Anim. Ecol.* 77:145–55
- 802 98. Mitchell KA, Hoffmann AA. 2010. Thermal ramping rate influences evolutionary  
803 potential and species differences for upper thermal limits in *Drosophila*. *Funct.*  
804 *Ecol.* 24:694–700
- 805 99. Mitton JB, Ferrenberg SM. 2012. Mountain pine beetle develops an  
806 unprecedented summer generation in response to climate warming. *Am. Nat.*  
807 179:163–71

- 808 100.Montllor CB, Maxmen A, Purcell AH. 2002. Facultative bacterial endosymbionts  
809 benefit pea aphids *Acyrtosiphon pisum* under heat stress. *Ecol. Entomol.*  
810 27:189–95
- 811 101.Montserrat M, Guzman C, Sahun RM, Belda JE, Hormaza JI. 2013. Pollen supply  
812 promotes, but high temperatures demote, predatory mite abundance in avocado  
813 orchards. *Agric. Ecosyst. Environ.* 164:155–61
- 814 102.Nguyen TT, Michaud D, Cloutier C. 2009. A proteomic analysis of the aphid  
815 *Macrosiphum euphorbiae* under heat and radiation stress. *Insect Biochem. Mol.*  
816 *Biol.* 39:20–30
- 817 103.O’Sullivan J, Macmillan HA, Overgaard J. 2017. Heat stress is associated with  
818 disruption of ion balance in the migratory locust, *Locusta migratoria*. *J. Therm.*  
819 *Biol.* 68:177–85
- 820 104.Overgaard J, Kearney MR, Hoffmann AA. 2014. Sensitivity to thermal extremes  
821 in Australian *Drosophila* implies similar impacts of climate change on the  
822 distribution of widespread and tropical species. *Glob. Change Biol.* 20:1738–50
- 823 105.Paaijmans KP, Heinig RL, Seliga RA, Blanford JI, Blanford S, Murdock CC,  
824 Thomas MB. 2013. Temperature variation makes ectotherms more sensitive to  
825 climate change. *Glob. Change Biol.* 19:2373–80
- 826 106.Piessens K, Adriaens D, Jacquemyn H, Honnay O. 2009. Synergistic effects of an  
827 extreme weather event and habitat fragmentation on a specialised insect herbivore.  
828 *Oecologia* 159:117–26
- 829 107.Pike DA, Webb JK, Shine R. 2012. Hot mothers, cool eggs: nest-site selection by  
830 egg-guarding spiders accommodates conflicting thermal optima. *Funct. Ecol.*  
831 26:469–75
- 832 108.Pincebourde S, Casas J. 2015. Warming tolerance across insect ontogeny:  
833 influence of joint shifts in microclimates and thermal limits. *Ecology* 96:986–97
- 834 109.Pincebourde S, Casas J. 2016. Hypoxia and hypercarbia in endophagous insects:  
835 Larval position in the plant gas exchange network is key. *J. Insect Physiol.*  
836 84:137–53
- 837 110.Pincebourde S, Casas J. 2019. Narrow safety margin in the phyllosphere during  
838 thermal extremes. *Proc. Natl. Acad. Sci. USA* 116:5588–96
- 839 111.Pincebourde S, Murdock CC, Vickers M, Sears MW. 2016. Fine-scale  
840 microclimatic variation can shape the responses of organisms to global change in  
841 both natural and urban environments. *Integr. Comp. Biol.* 56:45–61
- 842 112.Pincebourde S, Sinoquet H, Combes D, Casas J. 2007. Regional climate  
843 modulates the canopy mosaic of favourable and risky microclimates for insects. *J.*  
844 *Anim. Ecol.* 76:424–38
- 845 113.Pincebourde S, Suppo C. 2016. The vulnerability of tropical ectotherms to

- 846 warming is modulated by the microclimatic heterogeneity. *Integr. Comp. Biol.*  
847 56:85–97
- 848 114.Portner H. 2001. Climate change and temperature-dependent biogeography:  
849 oxygen limitation of thermal tolerance in animals. *Naturwissenschaften*  
850 88:137–46
- 851 115.Posledovich D, Toftegaard T, Wiklund C, Ehrlén J, Gotthard K. 2018.  
852 Phenological synchrony between a butterfly and its host plants: Experimental test  
853 of effects of spring temperature. *J. Anim. Ecol.* 87:150–61
- 854 116.Potter KA, Davidowitz G, Woods HA. 2011. Cross-stage consequences of egg  
855 temperature in the insect *Manduca sexta*. *Funct. Ecol.* 25:548–56
- 856 117.Potter KA, Woods A, Pincebourde S. 2013. Microclimatic challenges in global  
857 change biology. *Glob. Chang Biol.* 19:2932–39
- 858 118.Radchuk V, Turlure C, Schtickzelle N. 2013. Each life stage matters: the  
859 importance of assessing the response to climate change over the complete life  
860 cycle in butterflies. *J. Anim. Ecol.* 82:275–85
- 861 119.Robinet C, Roques A. 2010. Direct impacts of recent climate warming on insect  
862 populations. *Integr. Zool.* 5:132–42
- 863 120.Ruel JJ, Ayres MP. 1999. Jensen’s inequality predicts effects of environmental  
864 variation. *Trends Ecol. Evol.* 14:361–66
- 865 121.Saudreau M, Ezanic A, Adam B, Caillon R, Walser P, Pincebourde S. 2017.  
866 Temperature heterogeneity over leaf surfaces: the contribution of the lamina  
867 microtopography. *Plant Cell Environ.* 40:2174–88
- 868 122.Sentis A, Hemptinne JL, Brodeur J. 2013. Effects of simulated heat waves on an  
869 experimental plant-herbivore-predator food chain. *Glob. Chang Biol.* 19:833–42
- 870 123.Sentis A, Hemptinne JL, Brodeur J. 2017. Non-additive effects of simulated heat  
871 waves and predators on prey phenotype and transgenerational phenotypic  
872 plasticity. *Glob. Chang Biol.* 23:4598–608
- 873 124.Sinclair BJ, Marshall KE, Sewell MA, Levesque DL, Willett CS, et al. 2016. Can  
874 we predict ectotherm responses to climate change using thermal performance  
875 curves and body temperatures? *Ecol. Lett.* 19:1372–85
- 876 125.Sinclair BJ, Vernon P, Klok CJ, Chown SL. 2003. Insects at low temperatures: an  
877 ecological perspective. *Trends Ecol. Evol.* 18:257–62
- 878 126.Smith MD. 2011. An ecological perspective on extreme climatic events: a  
879 synthetic definition and framework to guide future research. *J. Ecol.* 99:656–63
- 880 127.Stuhldreher G, Hermann G, Fartmann T. 2014. Cold-adapted species in a  
881 warming world—an explorative study on the impact of high winter temperatures  
882 on a continental butterfly. *Entomol. Exp. Appl.* 151:270–79
- 883 128.Suggitt AJ, Gillingham PK, Hill JK, Huntley B, Kunin WE, Roy DB, Thomas CD.

- 884 2011. Habitat microclimates drive fine-scale variation in extreme temperatures.  
885 *Oikos* 120:1–8
- 886 129.Sunday JM, Bates AE, Kearney MR, Colwell RK, Dulvy NK, Longino JT, Huey  
887 RB. 2014. Thermal-safety margins and the necessity of thermoregulatory  
888 behavior across latitude and elevation. *Proc. Natl. Acad. Sci. USA* 111:5610–15
- 889 130.Tougeron K, Damien M, Le Lann C, Brodeur J, van Baaren J. 2018. Rapid  
890 responses of winter aphid-parasitoid communities to climate warming. *Front.*  
891 *Ecol. Evol.* 6:173
- 892 131.Tseng M, Kaur K M, Pari SS, Sarai K, Chan D, et al. 2018. Decreases in beetle  
893 body size linked to climate change and warming temperatures. *J. Anim. Ecol.*  
894 87:647–59
- 895 132.Vallières R, Rochefort S, Berthiaume R, Hébert C, Bauce É. 2015. Effect of  
896 simulated fall heat waves on cold hardiness and winter survival of hemlock  
897 looper, *Lambdina fiscellaria* (Lepidoptera: Geometridae). *J. Insect Physiol.*  
898 73:60–69
- 899 133.van Heerwaarden B, Kellermann V, Sgro CM. 2016. Limited scope for plasticity  
900 to increase upper thermal limits. *Funct. Ecol.* 30:1947–56
- 901 134.Vasseur DA, DeLong JP, Gilbert B, Greig HS, Harley CDG, et al. 2014. Increased  
902 temperature variation poses a greater risk to species than climate warming. *Proc.*  
903 *R. Soc. B* 281:20132016
- 904 135.Verberk WC, Bilton DT. 2013. Respiratory control in aquatic insects dictates their  
905 vulnerability to global warming. *Biol. Lett.* 9:20130473
- 906 136.Walsh BS, Parratt SR, Hoffmann AA, Atkinson D, Snook RR, Bretman A, Price  
907 TAR. 2019. The impact of climate change on fertility. *Trends Ecol. Evol.*  
908 34:249–59
- 909 137.Wehner R, Wehner S. 2011. Parallel evolution of thermophilia: daily and seasonal  
910 foraging patterns of heat-adapted desert ants: *Cataglyphis* and *Ocymyrmex*  
911 species. *Physiol. Entomol.* 36:271–81
- 912 138.Woods HA, Dillon ME, Pincebourde S. 2015. The roles of microclimatic  
913 diversity and of behavior in mediating the responses of ectotherms to climate  
914 change. *J. Therm. Biol.* 54:86–97
- 915 139.Woods HA. 2013. Ontogenetic changes in the body temperature of an insect  
916 herbivore. *Funct. Ecol.* 27:1322–31
- 917 140.Xing K, Hoffmann A, Ma CS. 2014. Does thermal variability experienced at the  
918 egg stage influence life history traits across life cycle stages in a small  
919 invertebrate? *PLoS ONE* 9:e99500
- 920 141.Zani PA, Cohnstaedt LW, Corbin D, Bradshaw WE, Holzapfel CM. 2005.  
921 Reproductive value in a complex life cycle: heat tolerance of the pitcher-plant

- 922 mosquito, *Wyeomyia smithii*. *J. Evol. Biol.* 18:101–105
- 923 142.Zhang B, Leonard SP, Li Y, Moran NA. 2019. Obligate bacterial endosymbionts  
924 limit thermal tolerance of insect host species. *Proc. Natl. Acad. Sci. USA*  
925 116:24712–18
- 926 143.Zhang W, Chang XQ, Hoffmann AA, Zhang S, Ma CS. 2015. Impact of hot  
927 events at different developmental stages of a moth: the closer to adult stage, the  
928 less reproductive output. *Sci. Rep.* 5:10436
- 929 144.Zhang W, Rudolf V, Ma CS. 2015. Stage-specific heat effects: Timing and  
930 duration of heat waves alter demographic rates of a global insect pest. *Oecologia*  
931 179:947–57
- 932 145.Zhang W, Zhao F, Hoffmann AA, Ma CS. 2013. A single hot event that does not  
933 affect survival but decreases reproduction in the diamondback moth, *Plutella*  
934 *xylostella*. *PLoS ONE* 8:e75923
- 935 146.Zhao F, Hoffmann AA, Xing K, Ma CS. 2017. Life stages of an aphid living  
936 under similar thermal conditions differ in thermal performance. *J. Insect Physiol.*  
937 99:1–7
- 938 147.Zhao F, Zhang W, Hoffmann AA, Ma CS. 2014. Night warming on hot days  
939 produces novel impacts on development, survival and reproduction in a small  
940 arthropod. *J. Anim. Ecol.* 83:769–78
- 941 148.Zhu L, Wang L, Ma CS. 2019. Sporadic short temperature events cannot be  
942 neglected in predicting impacts of climate change on small insects. *J. Insect*  
943 *Physiol.* 112:48–56

944  
945

#### 946 **RELATED RESOURCES**

- 947 Bale JS, Masters GJ, Hodkinson ID, Awmack C, Bezemer TM, et al. 2002. Herbivory  
948 in global climate change research: direct effects of rising temperature on insect  
949 herbivores. *Glob. Change Biol.* 8:1–16
- 950 Grant PR, Grant BR, Huey RB, Johnson MTJ, Knoll AH, Schmitt J. 2017. Evolution  
951 caused by extreme events. *Philos. Trans. R. Soc. B* 372:20160146
- 952 Hance T, van Baaren J, Vernon P, Boivin G. 2007. Impact of extreme temperatures on  
953 parasitoids in a climate change perspective. *Annu. Rev. Entomol.* 52:107–26
- 954 Harris R, Beaumont LJ, Vance TR, Tozer CR, Remenyi TA, et al. 2018. Biological  
955 responses to the press and pulse of climate trends and extreme events. *Nat. Clim.*  
956 *Change* 8:579–87
- 957 Sgrò CM, Terblanche JS, Hoffmann AA. 2016. What can plasticity contribute to  
958 insect responses to climate change? *Annu. Rev. Entomol.* 61:433–51
- 959 Ummenhofer CC, Meehl GA. 2017. Extreme weather and climate events with

960 ecological relevance: a review. *Philos. Trans. R. Soc. B* 372:20160135  
961 van de Pol M, Jenouvrier S, Cornelissen JH, Visser ME. 2017. Behavioural,  
962 ecological and evolutionary responses to extreme climatic events: challenges and  
963 directions. *Philos. Trans. R. Soc. B* 372:20160134  
964 Vázquez DP, Gianoli E, Morris WF, Bozinovic F. 2017. Ecological and evolutionary  
965 impacts of changing climatic variability. *Biol. Rev.* 92:22–42

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985 **Figure captions**

986 **Figure 1.** Diagram illustrating how natural thermal extremes are filtered (often  
987 intensity is lowered) by microhabitats and buffered by insects through behavioral  
988 thermoregulation, phenotypic plasticity, ontogeny variations and adaptive evolution to  
989 ultimately increase survival to heat. Even after filtering and buffering, thermal  
990 extremes still impact insect performance at the individual, population and community  
991 levels within and across generations.

992

993 **Figure 2.** Characterizing Extreme High Temperatures (EHTs) in the field. (a) A  
994 temporal series of microclimatic temperature for a theoretical insect illustrates the  
995 different metrics used to describe the occurrence of EHTs (in *red*) and mild periods (in  
996 *blue*): their intensity (maximal and minimal temperatures of hot days ( $HT_{\max}$  and  
997  $HT_{\min}$ ) and mild days ( $MT_{\max}$  and  $MT_{\min}$ ), respectively); frequency (the number of  
998 consecutive hot days and the frequency at which hot periods occur); temporal  
999 variance (the timing of occurrence of hot and mild phases); the biological threshold  
1000 (temperature inducing thermal stress in a given organism, or  $CT_{\max}$ ). The dichotomy  
1001 between hot and mild events hides a large range of possibilities when designing  
1002 experiments to simulate natural thermal extremes. (b–c) In particular, the maximum  
1003 and minimum temperatures of mild and hot days vary within their own range. (d)  
1004 These complex thermal extremes with a daily sub-unit are embedded in naturally  
1005 fluctuating temperature regimes that can be described by their temporal variance of  
1006 EHT occurrence and their frequency (eg, the number of consecutive hot or mild days).

1007 Here, the temporal arrangement of hot and mild days varies across two axis defined  
1008 by number of hot days during a period of 10 days (frequency) and the variance in the  
1009 number of mild days between each hot day (temporal variance). Note that in panel *d*,  
1010 the global mean and variance of temperature remain the same across all  
1011 configurations at a given frequency of hot days.

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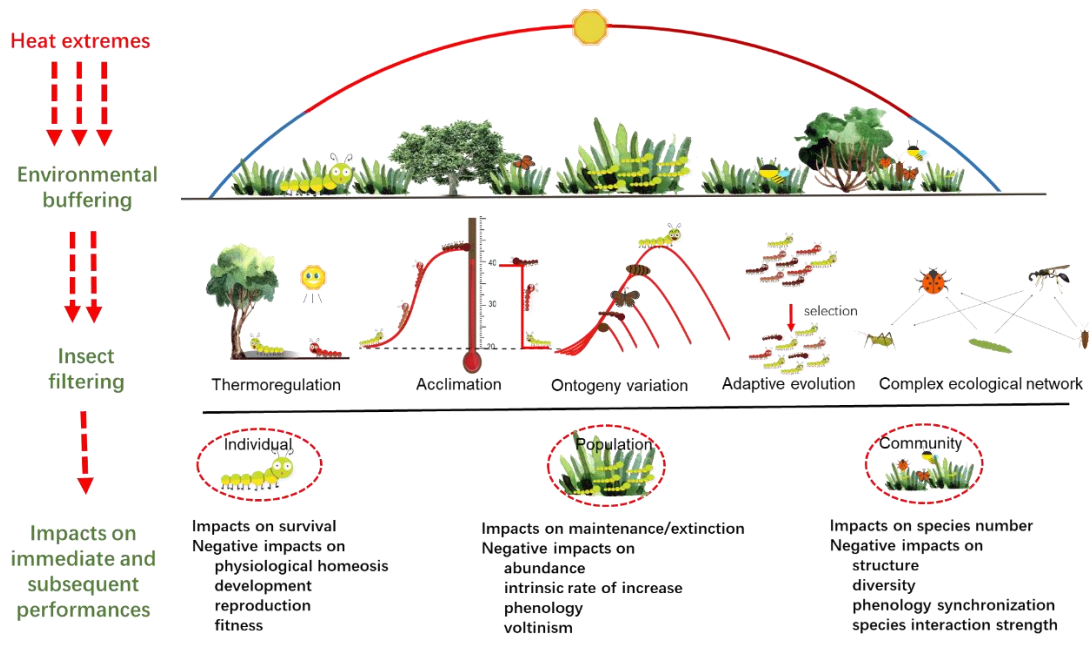
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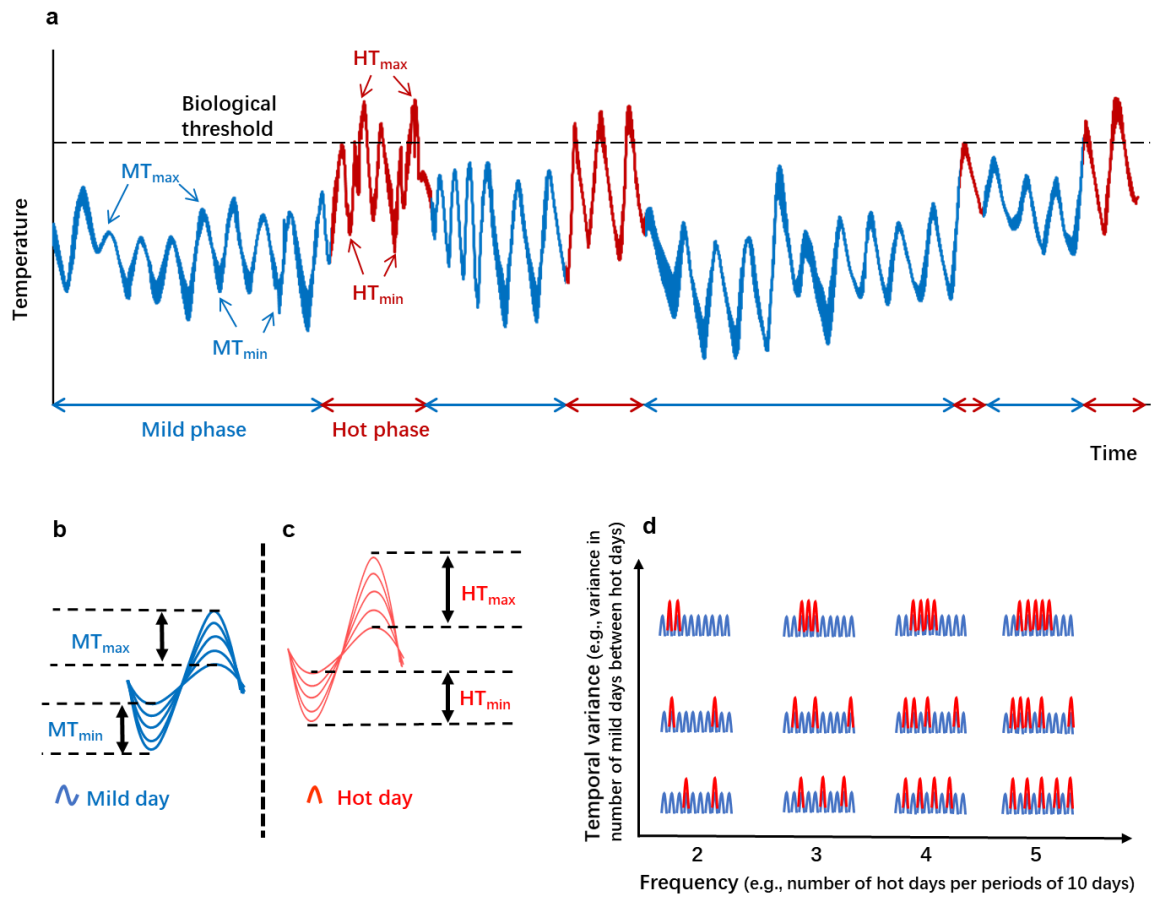
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Figure 1



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Figure 2