

# Temperature affects chemical defense in a mite-beetle predator-prey system

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## Supplementary Text S1 – detailed equations

The chemical defense dynamic and functional response models can be formalized (Heethoff and Rall 2015) by describing the number of prey individuals – either chemically defended or undefended – attacked (and eaten) by a predator as:

$$\frac{dN}{dt} = -F \quad (1a)$$

$$\frac{dS}{dt} = P_s - e_s \frac{F_S}{N} \quad (1b)$$

where  $N$  ( $N \text{ A}^{-1}_{\text{area}}$ ) is the change of prey individual density (eq. **1a**),  $S$  ( $\text{ng N}^{-1}$ ) is the change of per capita defensive secretions (eq. **1b**) over  $t$  (h) time,  $F$  ( $N \text{ h}^{-1}$ ) is the functional response of undefended and  $F_S$  ( $N_S \text{ h}^{-1}$ ) is the function response of defended mites. Furthermore  $e_s$  ( $\text{ng N}^{-1}$ ) and  $P_s$  ( $\text{ng N}^{-1}\text{h}^{-1}$ ) are factors describing the amount of defensive secretion lost per attack and the per capita secretion production over time, respectively.

To model the functional response, Heethoff and Rall (2015) chose a traditional type II functional response in habitat-free foraging space (Holling 1959; Vucic-Pestic et al. 2010) and extended the classic equation resulting in two new models;  $F$  for undefended mites (eq. **2a**) and  $F_S$  for defended (eq. **2b**):

$$F = \frac{\sigma\alpha(N - N_S)}{1 + \sigma\alpha T_h(N - N_S) + (1 - \sigma)\alpha T_U(N - N_S) + \alpha T_U N_S} \quad (2a)$$

$$F_S = \frac{\alpha N_S}{1 + \sigma\alpha T_h(N - N_S) + (1 - \sigma)\alpha T_U(N - N_S) + \alpha T_U N_S} \quad (2b)$$

where  $T_h$  ( $\text{h N}^{-1}$ ) is the handling time,  $\alpha$  ( $\text{A}_{\text{area}} \text{h}^{-1}$ ) is the attack rate,  $\sigma$  is the success rate and  $T_U$  ( $\text{h}$ ) is the time needed/wasted for failed attacks.

The gross change of the per capita secretion rate over time ( $S_t$ ) without any disturbance from predators (eq. **1b** with  $F_S = 0$ ) can be written as [eq. **3**, Heethoff and Rall (2015)]:

$$S_t = \frac{-K_S + K_S e^{\frac{R_S t}{K_S}} + S_{t=0}}{e^{\frac{R_S t}{K_S}}} \quad (3)$$

with  $K_S$  ( $\text{ng N}^{-1}$ ) as the reservoir size and  $R_S$  ( $\text{ng h}^{-1} \text{N}^{-1}$ ) as the regeneration rate of a subpopulation of defended prey. This assumes, that the regeneration of the defensive secretions starts immediately after a depletion event and until the maximum reservoir filling under given circumstances are reached. Both sides of eq. **3** can be  $\log_{10}$ -transformed to ensure normality of the model residuals. Consequently, eq. **3** can be used to fit experimentally measured quantitative secretion data over time to estimate  $K_S$  and  $R_S$  for different treatments after total depletion of the glandular contents (e.g. by artificial stimuli; Heethoff and Raspotnig (2012)). Both parameters are used to describe the *per capita* secretion production over time  $P_S$  (eq. **4**) to finally model the system with the two ordinary differential equations (eq. **1a** and eq. **1b**) but see Heethoff and Rall (2015) for further details.

$$P_S = R_S \left(1 - \frac{S}{K_S}\right) \quad (4)$$

To model the survival of prey,  $K_s$  ( $\text{ng N}^{-1}$ ) and  $R_s$  ( $\text{ng h}^{-1} \text{N}^{-1}$ ) were subsequently used together with the discovery rate  $\alpha$  ( $\text{A}_{\text{area}} \text{h}^{-1}$ ) and handling times  $T_h$  ( $\text{h N}^{-1}$ ) for model parametrization, thereby  $\alpha$  ( $\text{A}_{\text{area}} \text{h}^{-1}$ ) and  $T_h$  ( $\text{h N}^{-1}$ ) were calculated according to Binzer et al. (2012) and Rall et al. (2012) based on a predator mass of 3.5 mg (corresponding to the rove beetle *Stenus junco*, a model predator of oribatid mites, Heethoff et al. 2011) and measured prey body masses  $m_{\text{mite}}$  ( $\mu\text{g}$ ) (mean for each temperature, see supplementary material). The rate of successful attacks  $\sigma$  in case of undefended prey was set to 0.28 and the amount of defensive secretion lost per attack  $e_s$  to  $50 \text{ ng N}^{-1}$  based on empirical data from Heethoff et al. (2011) as well as Raspotnig (2006) and Heethoff (2012), respectively.

As we were interested in the number of prey specimens surviving because of chemical defense at a given temperature, we used eq. 5 (for more details see Brückner and Heethoff 2018) to extract this information from the simulated data. The calculated (eq. 5) effective chemical defense is therefore the number of surviving mite individuals caused by chemical defense  $\Delta_{chem}$  ( $\text{N A}_{\text{area}}^{-1}$ ) at a given time:

$$\Delta_{chem} = N_{S,t_n} - N_{t_n} \quad (5)$$

where  $N_{S,t_n}$  ( $\text{N A}_{\text{area}}^{-1}$ ) is the number of living, defended mites at a given time (i.e. model including chemical defense and other parameters like body mass) and  $N_{t_n}$  ( $\text{N A}_{\text{area}}^{-1}$ ) is the number of living, undefended mites at a given time (i.e. the null-model excluding chemical defense, but including other parameters e.g., body mass differences). Finally, we simulated both scenarios (eq. 1a and eq. 1b) and calculate the effective chemical defense  $\Delta_{chem}$  (eq. 5) over time at the different temperature regimes using the generic `Isoda()`-function in R (R Core

R\_Core\_Team 2019) with a total number of 100 prey individuals and a time step length of 0.1 h ( $n_{\text{steps}}= 1,000$ ).

## References

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