

Host-parasitoid models: The interplay of density-dependent host growth and functional response of parasitoids

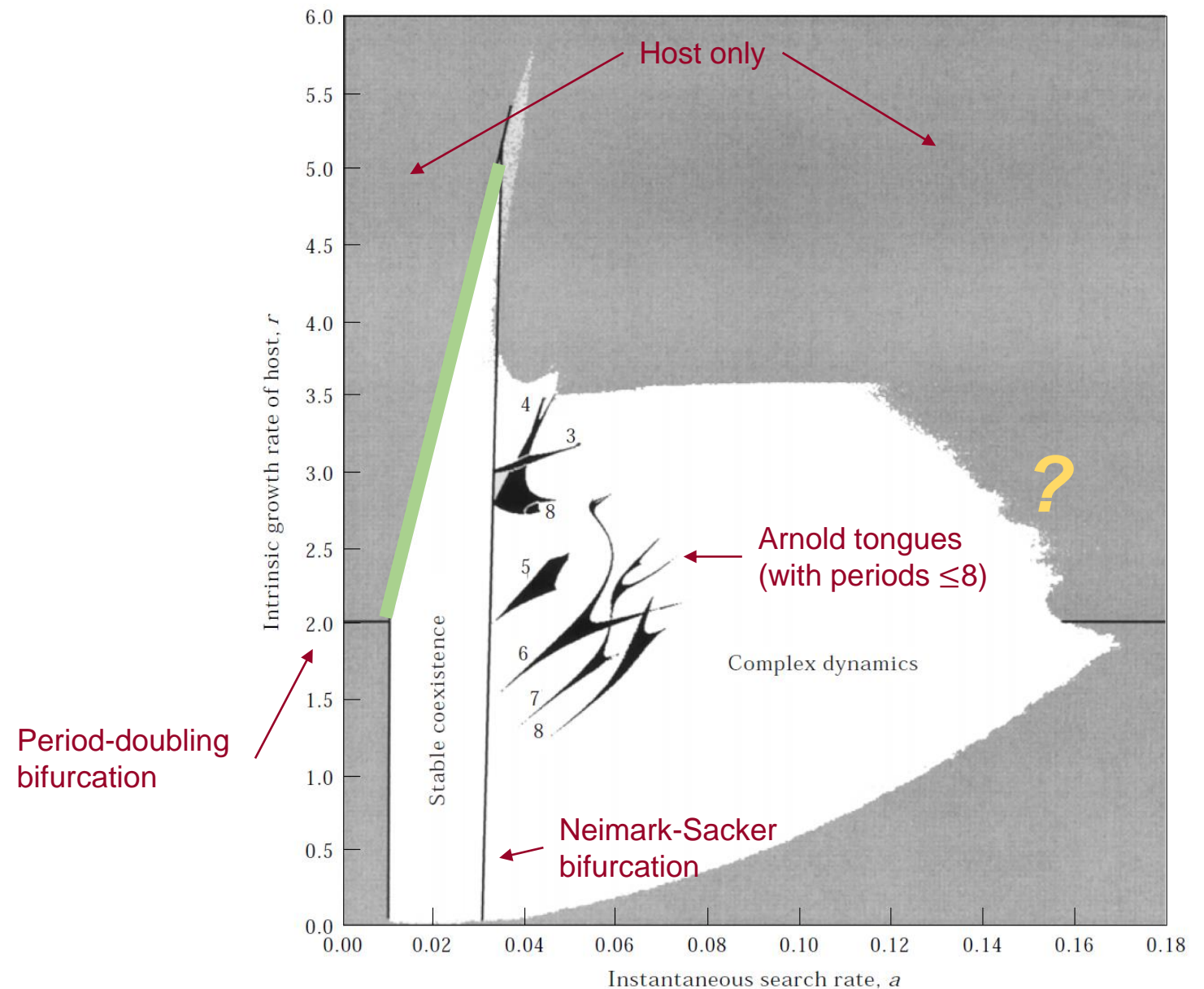
Frank M. Hilker

Ricker + type II

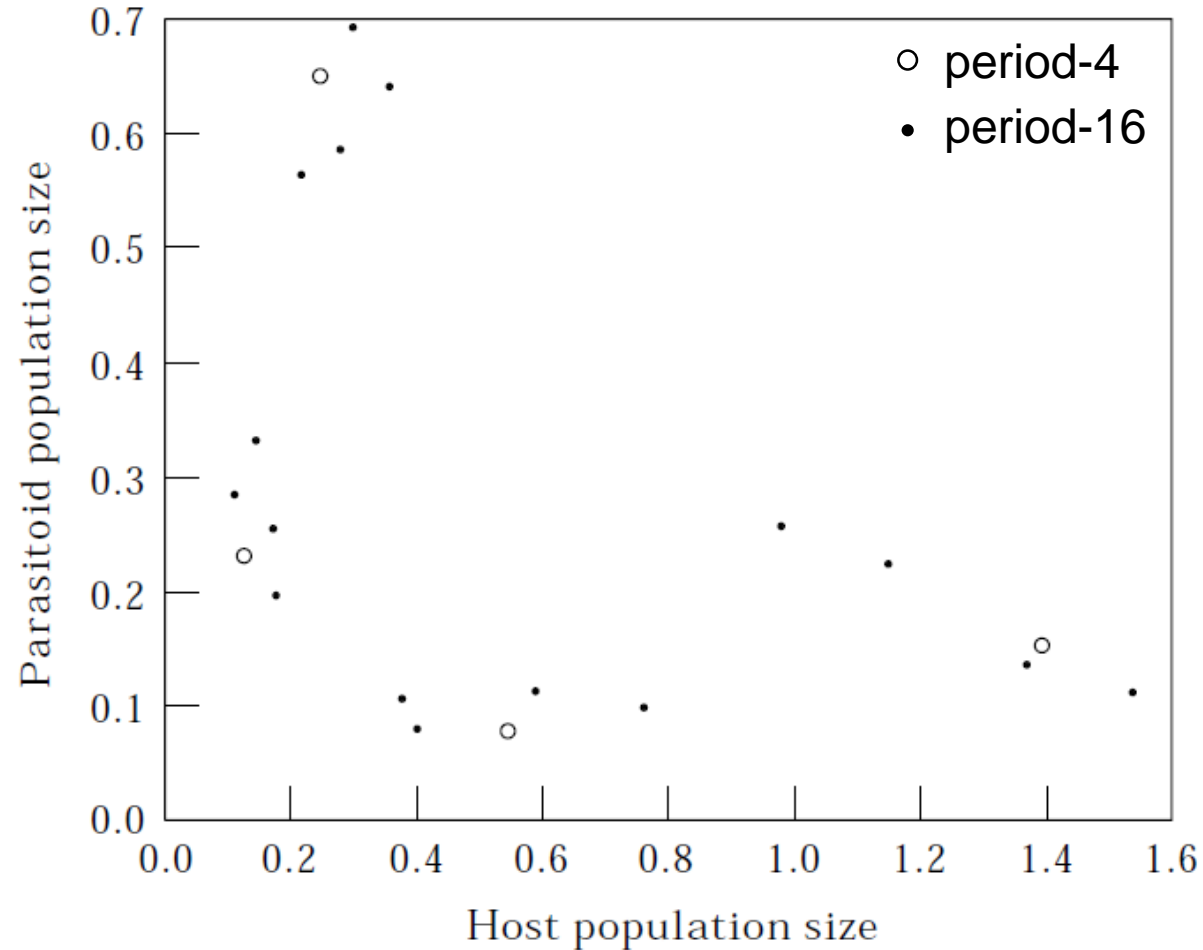
$$H_{t+1} = H_t e^{r(1-H_t) - \frac{aTP_t}{1+aT_hH_t}}$$

$$P_{t+1} = H_t \left(1 - e^{\frac{-aTP_t}{1+aT_hH_t}} \right)$$

$$T = 100, T_h = 1$$



Coexistence of two periodic attractors



$$r = 2.8, a = 0.043, T = 100, T_h = 1$$

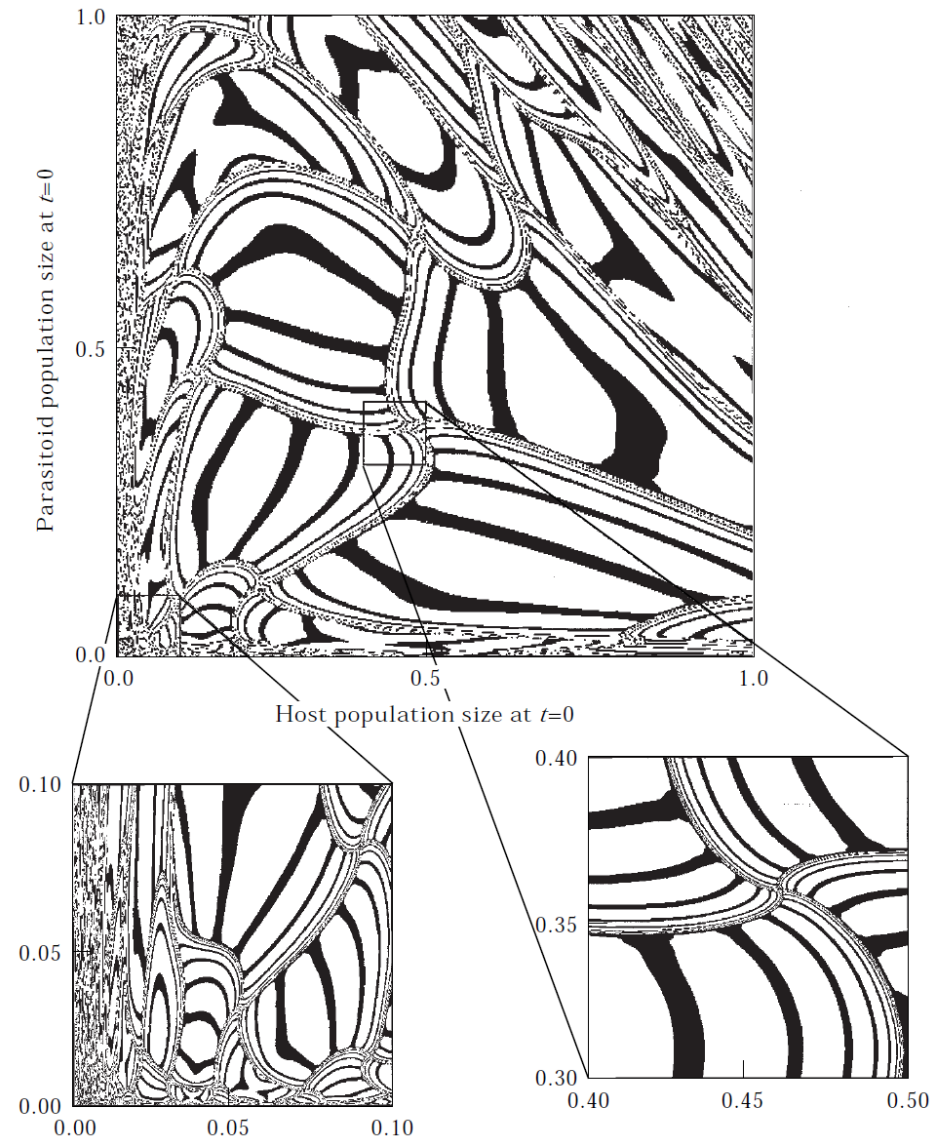
Other examples reported

- period-8 and period-15
- period-16 and period-45
- chaotic and six-piece chaotic

However, ***no report of more than two*** coexisting attractors.

There are fractal basins of attraction

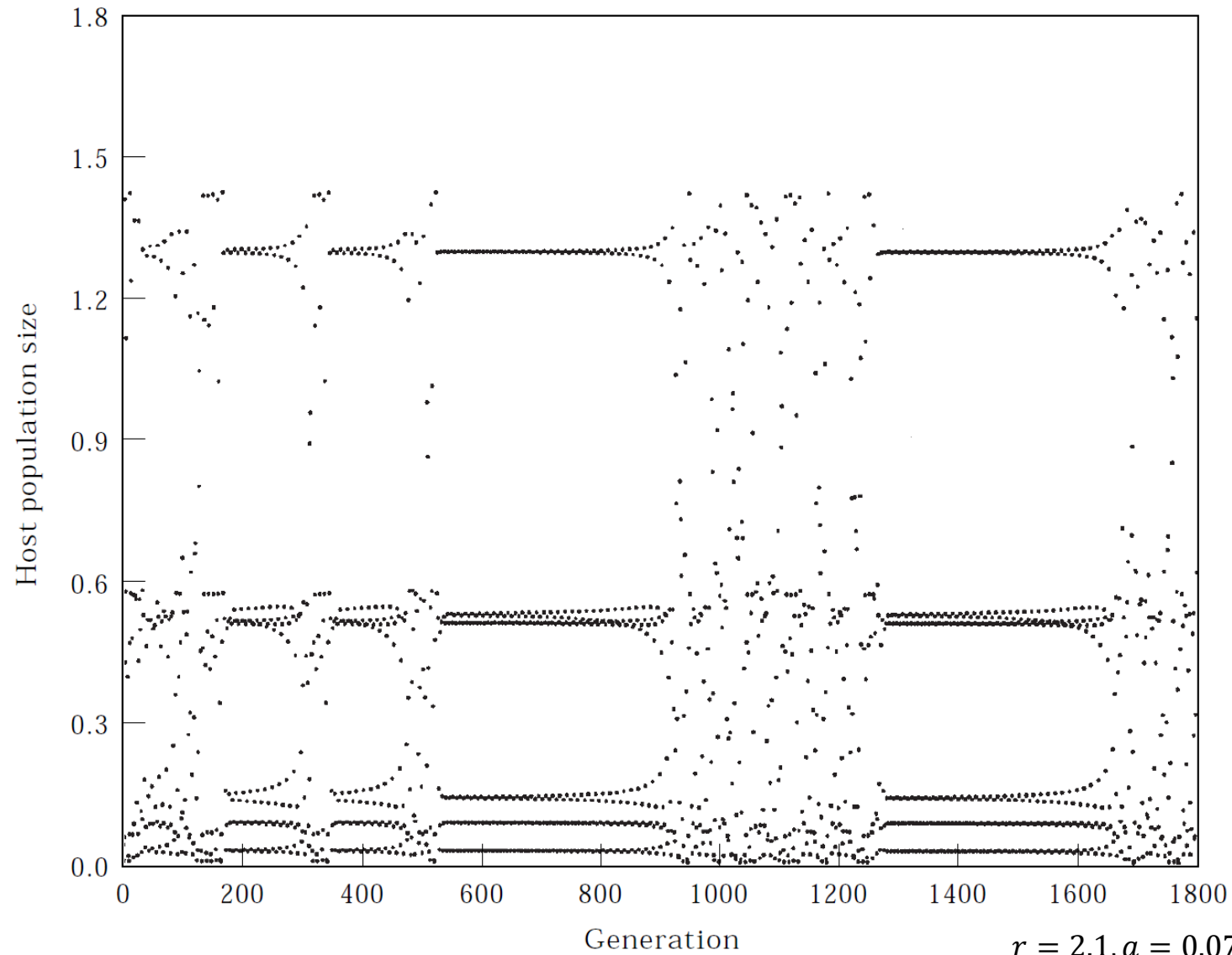
Here, for the 4- and 16-cycle



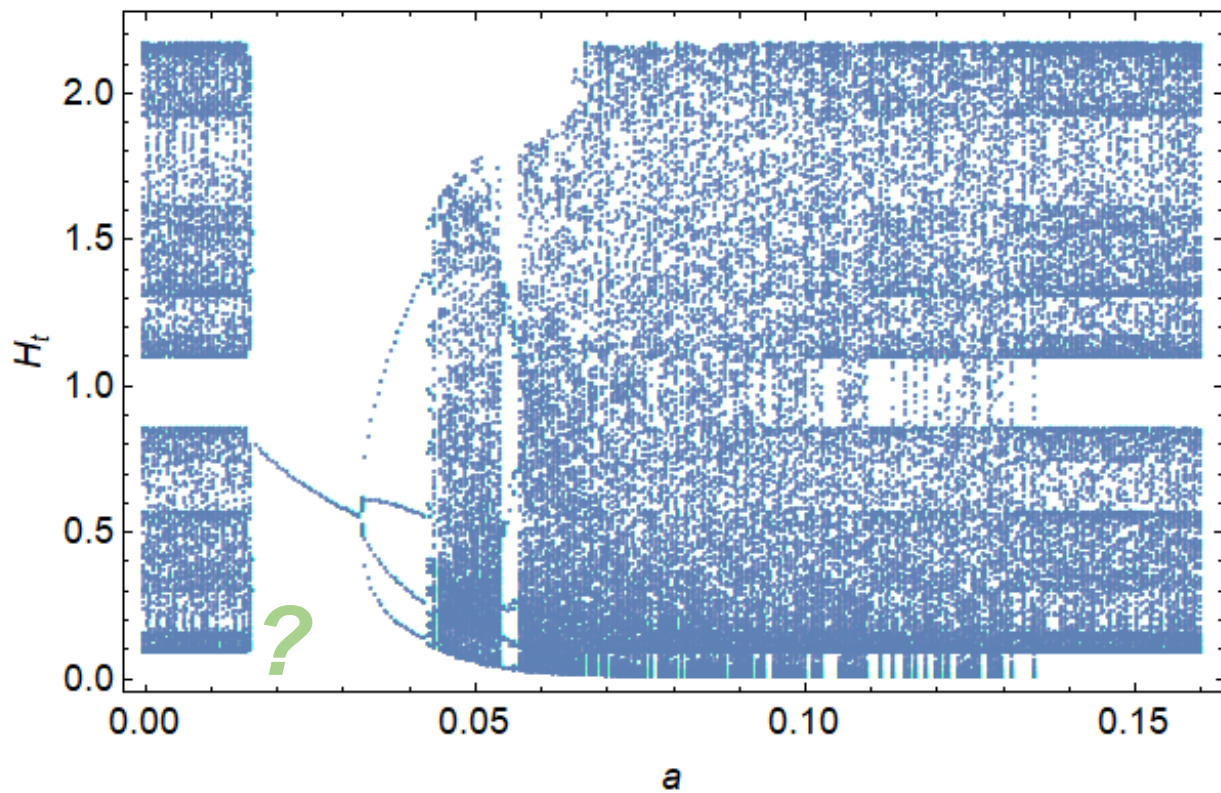
$$r = 2.8, a = 0.043, T = 100, T_h = 1$$

There is chaos

Here, intermittent

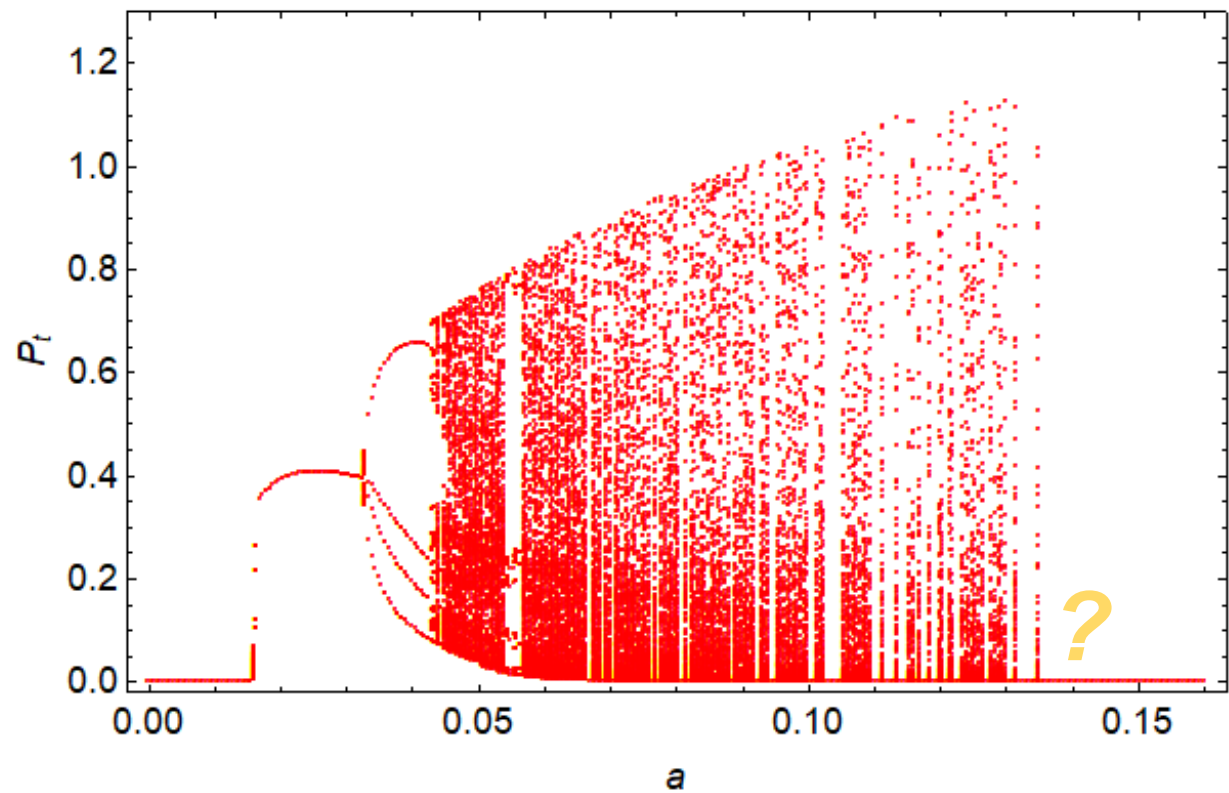


$$r = 2.1, a = 0.0717, T = 100, T_h = 1, H_0 = P_0 = 0.5$$



Parasitoids stabilize the unstable
host dynamics – how?

$r = 2.8, T = 100, T_h = 1$, random ICs

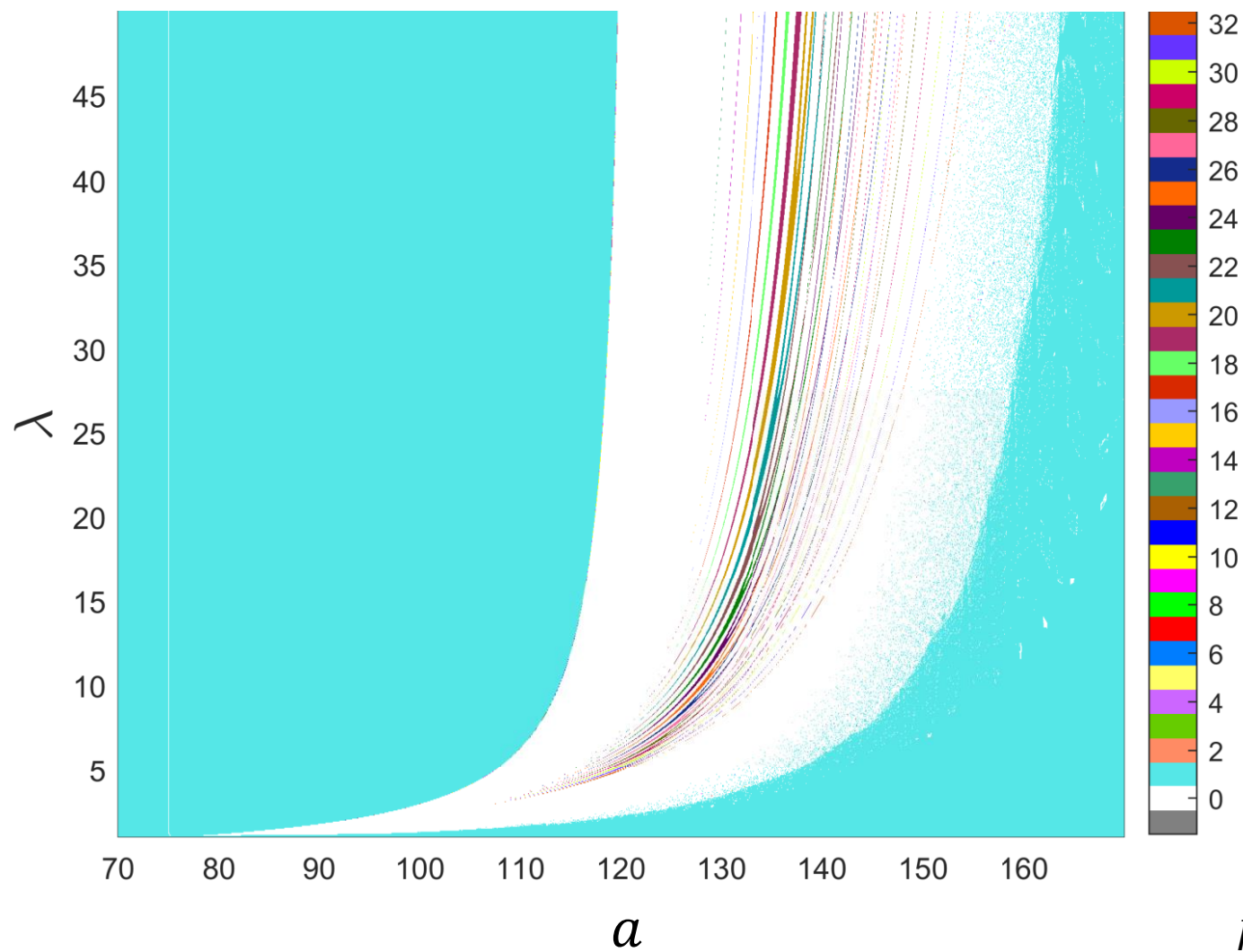


Why do the (quasi-)periodic cycles disappear?

Beverton-Holt + type II

$$H_{t+1} = \frac{\lambda H_t}{1 + (\lambda - 1)H_t} e^{\frac{-aP_t}{1+hH_t}}$$
$$P_{t+1} = H_t \left(1 - e^{\frac{-aP_t}{1+hH_t}}\right)$$

No Arnold tongues,
“just” periodic windows

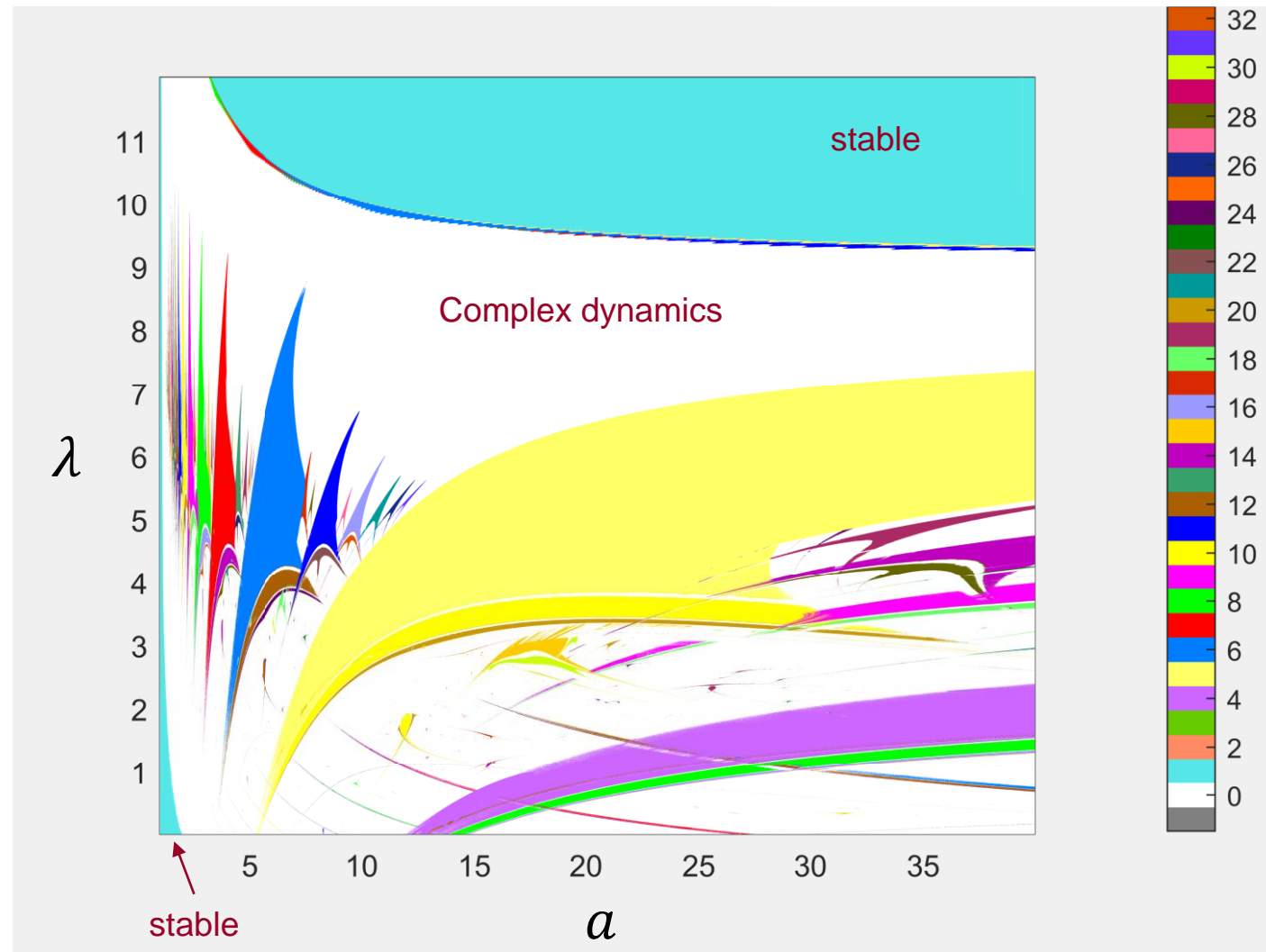
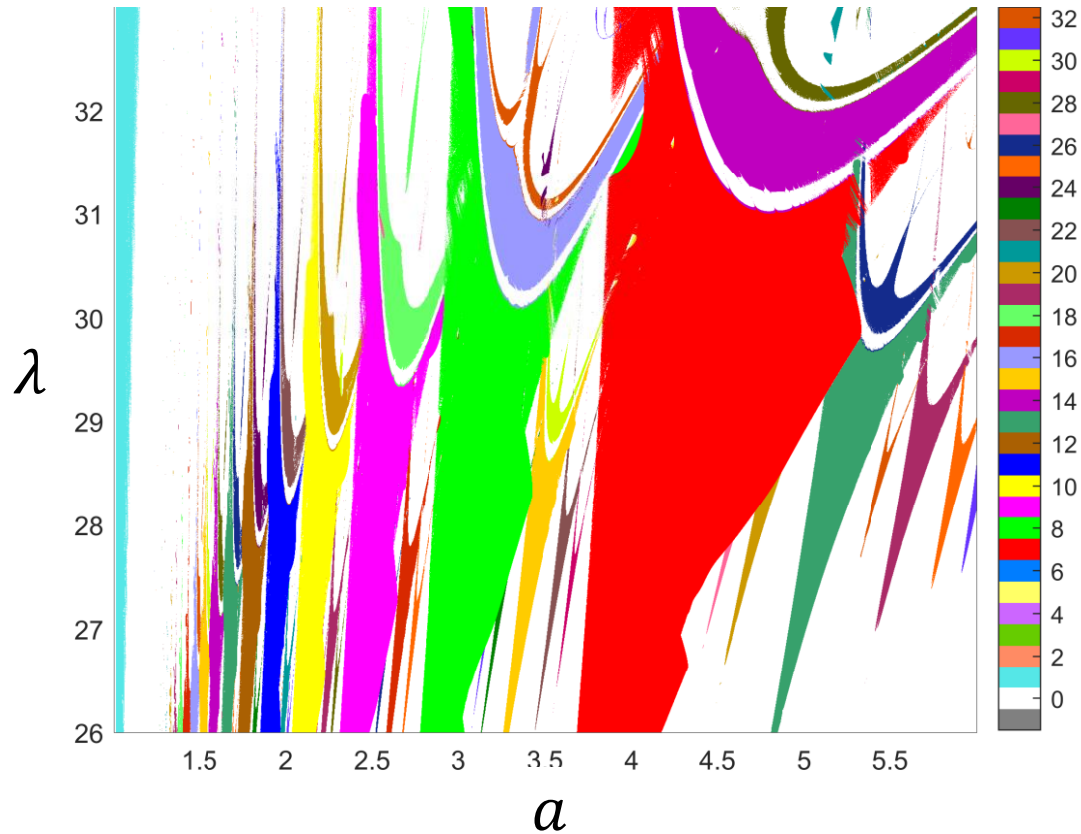


Beverton-Holt + type III

$$H_{t+1} = \frac{\lambda H_t}{1 + (\lambda - 1)H_t} e^{\frac{-aH_t P_t}{1+hH_t^2}}$$

$$P_{t+1} = H_t \left(1 - e^{\frac{-aH_t P_t}{1+hH_t^2}}\right)$$

Zoom



$$h = 7.4$$

Ricker + type III

$$\begin{aligned}H_{t+1} &= H_t e^{r(1-H_t/K) - \frac{aTH_tP_t}{1+cH_t+aT_hH_t^2}} \\P_{t+1} &= H_t \left(1 - e^{\frac{-aTH_tP_t}{1+cH_t+aT_hH_t^2}}\right)\end{aligned}$$

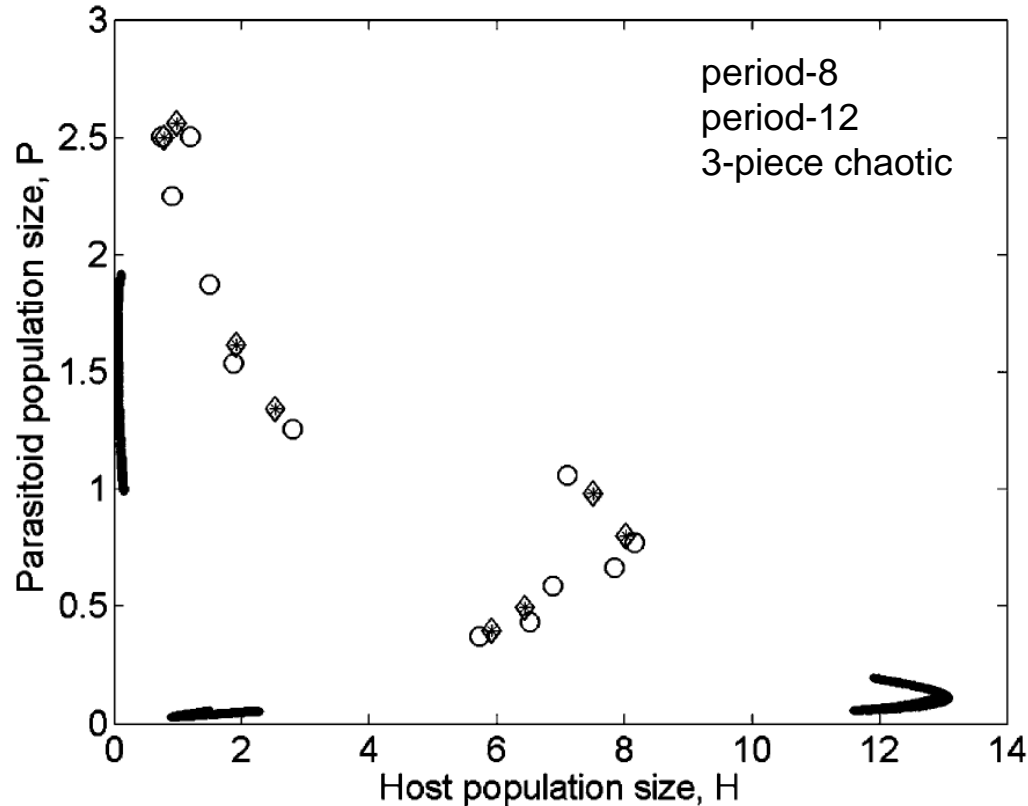
...at first glance, no particular novel dynamic behavior reported
(but more careful reading required)

Coexistence of three attractors

Ricker + parasitoid interference

$$H_{t+1} = H_t e^{r(1-H_t/K)-aP_t^{-m+1}}$$

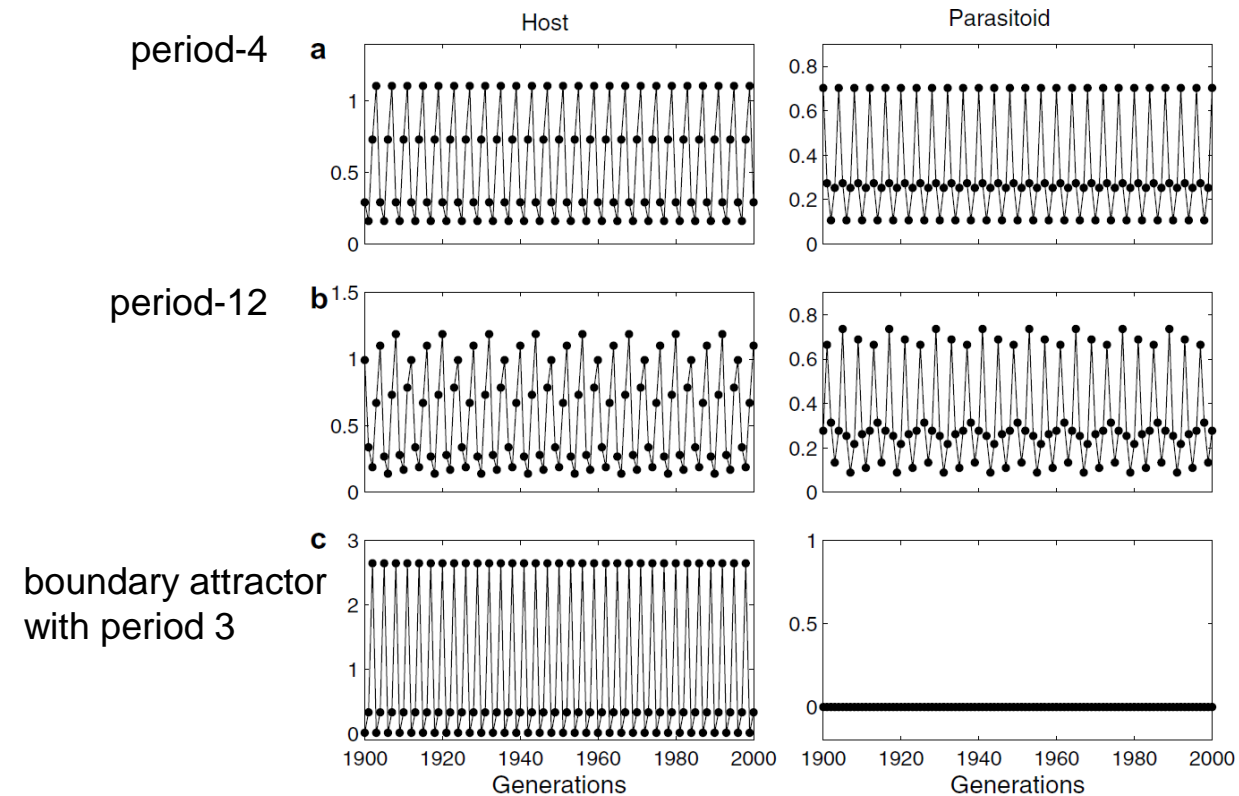
$$P_{t+1} = N_t \left(1 - e^{-aP_t^{-m+1}}\right)$$



Ricker + intergenerational parasitoid survival

$$H_{t+1} = H_t e^{r(1-H_t/K)-aP_t}$$

$$P_{t+1} = N_t(1 - e^{-aP_t}) + \delta P_t$$



Generalized host growth

Idea: Choose a general host growth that can be undercompensatory (like Beverton-Holt) or overcompensatory (like Ricker) depending on a parameter

$$\begin{aligned}H_{t+1} &= g(H_t) f(H_t, P_t) \\P_{t+1} &= N_t(1 - f(H_t, P_t))\end{aligned}$$

$$g(H) = \frac{\lambda H}{(1 + cH)^b}$$

Hassell (1976) map

$\lambda > 0$ intrinsic growth rate

$c > 0$ measure of intraspecific competition

$b \geq 0$ measure of compensation

$b = 0$ geometric growth

$0 < b < 1$ undercompensation

$b = 1$ Beverton-Holt

$b > 1$ overcompensation

Hassell + something else...

$$H_{t+1} = \frac{\lambda H_t}{\left(1 + aH_t e^{\frac{-aTP_t}{1+aT_hH_t}}\right)^b} e^{\frac{-aTP_t}{1+aT_hH_t}}$$

$$P_{t+1} = H_t \left(1 - e^{\frac{-aTP_t}{1+aT_hH_t}}\right)$$

“something else”

$$H_{t+1} = \frac{\lambda H_t}{(1 + aH_t e^{-AP_t})^b} e^{-AP_t}$$

$$P_{t+1} = H_t(1 - e^{-AP_t})$$

$b = 1$ Beverton-Holt

$b > 0$ Hassell

Liu et al. (22008)

Liu (2009)

$A = \text{constant or}$

$A = aP^{-m}$ parasitoid interference (Hassel-Varley)

Liu et al. (2008) Computer simulation of the dynamic complexity of a host-parasitoid model. International Conference on Computer Science and Information Technology

Liu, Hua (22009) Intra-specific density dependent effect of a host-parasitoid interaction model.

Zhu and Zhao (2009, *Chaos, Solitons and Fractals*)

- Coexistence of non-equilibrium attractors
 - not more than two in Kaitala et al. (1999) reported
 - Is $r > 2$ needed, i.e. instability in the host dynamics alone?
 - With Beverton-Holt host growth, is there also chaos possible with type II or III functional response?
- Bifurcation questions
 - How do parasitoids go extinct for large attack rates a ? (e.g., attractor crisis, homoclinic loop, numerical artefact?)
 - How do host-only oscillations become stabilized by parasitoid invasion, at smaller values of a ? (transition from periodic or chaotic attractor on the boundary to a stable fixed point in the interior?)
- Type III seems to differ from type II when combined with Beverton-Holt
- Arnold tongues: structure in the periodicities?