#### Lecture slides

Philip Murray

## Lecture 1

## A general model

Consider the first order difference equation

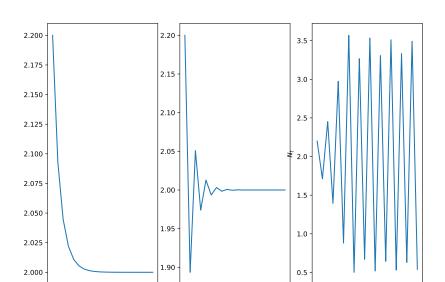
$$N_{t+1} = N_t f(N_t) = H(N_t),$$
 (1)

where  $f(N_t)$  is a function that defines the per capita growth rate. The function  $H(N_t)$  describes the total (net) growth rate.

# The Malthusian model

#### The Ricker model

$$N_{t+1} = N_t e^{r(1-\frac{N_t}{K})}.$$



# Steady state calculation

Compute the steady states of the system of ODEs

$$\begin{aligned} \frac{du}{dt} &= 1 - u, \\ \frac{dv}{dt} &= 1 - uv - v. \end{aligned}$$

Suppose  $(u^*, v^*)$  is a steady state. Hence

$$0 = 1 - u^*$$

and

$$0 = 1 - u^*v^* - v^*$$

The steady state is (1, 1/2).

## Linear stability calculation

Deduce, by considering the form for the eigenvalues that, for example, the conditions  $\det(A)>0$ ,  $\operatorname{tr}(A)>0$  with  $\operatorname{tr}(A)^2<4\det(A)$  imply that the steady state is an unstable spiral.

The eigenvalues are given by

$$\lambda = \frac{\operatorname{tr} A \pm \sqrt{\operatorname{tr} A^2 - 4 \det A}}{2}$$

## Jacobian calculation

Compute the Jacobian matrix for the system of ODEs

$$\begin{aligned} \frac{du}{dt} &= 1 - u, \\ \frac{dv}{dt} &= 1 - uv - v. \end{aligned}$$

Evaluate the Jacobian matrix at the steady state and hence determine its linear stability.

The Jacobian is given by

$$A = \left( \begin{array}{cc} -1 & 0 \\ -v & -u - 1 \end{array} \right).$$

At (1,1/2)

$$A = \left( \begin{array}{cc} -1 & 0 \\ -\frac{1}{2} & -2 \end{array} \right).$$

In this case

$$tr(A) = -3$$

and

#### Nullclines calculation

Sketch the nullclines of the system of ODEs

$$\begin{split} \frac{du}{dt} &= 1 - u, \\ \frac{dv}{dt} &= 1 - uv - v. \end{split}$$

The u nullcline is u = 1. The v nullcline is v = 1/(1+u).

#### Biochemical kinetics I

Suppose A and B react to produce C. Hence

$$A + B \xrightarrow{k} C$$
.

The law of mass action states that the rate of the reaction is

Using the reaction rates, we write down ordinary differential equations that describe how concentrations of a given molecule will change in time. Hence

$$\frac{d[C]}{dt} = k[A][B].$$

#### Biochemical kinetics

Consider the reversible reaction

$$A + B \stackrel{k_+}{\rightleftharpoons} C.$$

Define dependent variables, identify reaction rates and derive ordinary differential equations that describe how concentrations evolved in time.

The dependent variables are:

.

Applying the law of mass action yields the reaction rates:

$$k_+[A][B]$$
 and  $k_-[C]$ 

.

Ctd.

The ODEs are

$$\begin{split} \frac{d[A]}{dt} &= -k_{+}[A][B] + k_{-}[C], \\ \frac{d[B]}{dt} &= -k_{+}[A][B] + k_{-}[C], \\ \frac{d[C]}{dt} &= k_{+}[A][B] - k_{-}[C]. \end{split}$$

For a given set of initial conditions,

$$[A](t=0) = [A]_0, \quad [B](t=0) = [B]_0, \quad [C](t=0) = [C]_0,$$

the ODEs can be solved and hence the concentrations of the different molecules described as time evolves.