

**S2 File. Derivation of the basic reproduction number ( $R_0$ ) using the next-generation matrix method.**

The expressions for the basic reproduction number ( $R_0$ ) were derived using the next-generation matrix (NGM) method originally defined by van den Driessche & Watmough [1]. It is straightforward to verify that models 1 and 2 satisfy the conditions (A1)—(A5) in [1] needed to apply this method. Indeed, (A1) the directed transfer of individuals between compartments can be expressed as the difference between positive-valued functions  $F_i$  and  $V_i$ ,  $F_i - V_i$ . (A2) If a compartment is empty, then there can be no transfer out of the compartment. (A3) The incidence of infection for uninfected compartments is zero. (A4) There is no immigration of infected individuals into the system. (A5) In the absence of new infections, the disease-free equilibrium (DFE) is stable.

To construct the next-generation matrix,  $(FV^{-1})$ , we defined the matrices  $F$  and  $V$  as:

$$F = \left[ \frac{\partial F_i(x)}{\partial x_j} \right] x = x_0 \text{ and } V = \left[ \frac{\partial V_i(x)}{\partial x_j} \right] x = x_0,$$

where the  $(i,j)$  entry of matrix  $F$  was the rate at which infected individuals in compartment  $j$  produce new infections in compartment  $i$  and the  $(i,j)$  entry of  $V$  was the net rate of change of animals in compartment  $i$  by any other means. We considered that  $E$ ,  $I_A$  and  $I_C$  are the infected states but that new infections occurred only in the  $E$  compartment. The rates were evaluated at the disease-free equilibrium  $x = x_0$ .

Setting  $\vec{I} = (E, I_A, I_C)^T$  for model 1, it follows that

$$\frac{d\vec{I}}{dt} = J\vec{I} = (F - V)\vec{I},$$

where  $J$  denotes the Jacobian matrix evaluated at the DFE and  $F$  and  $V$  matrices are:

$$F = \begin{bmatrix} 0 & \beta_A S_0 & \beta_C S_0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{bmatrix}$$

$$V = \begin{bmatrix} d + h + \alpha & 0 & 0 \\ -\alpha p_1 & d + h + \mu_A + \delta & 0 \\ -\alpha(1 - p_1) & -\delta(1 - p_2) & d + h + \mu_C + \gamma \end{bmatrix}$$

Following [1,2], the basic reproduction number  $R_0$  is defined as the spectral radius (dominant eigenvalue) of matrix  $FV^{-1}$ . That is,

$$R_0 = \rho(FV^{-1}) = R_{0A} + R_{0C}, \quad (\text{A1})$$

where

$$R_{0A} = \frac{\alpha p_1 \beta_A S_0}{(d + h + \alpha)(d + h + \mu_A + \delta)}$$

$$R_{0C} = \left[ \frac{\alpha(1 - p_1)}{(d + h + \alpha)} + \frac{\alpha p_1 \delta(1 - p_2)}{(d + h + \alpha)(d + h + \mu_A + \delta)} \right] \frac{\beta_C S_0}{(d + h + \mu_C + \gamma)}$$

The expression for  $R_{0A}$  can be interpreted as follows: a fraction  $\alpha p_1/(d + h + \alpha)$  of exposed hosts  $E$  progress to state  $I_A$  and spend an average of  $1/(d + h + \mu_A + \delta)$  days in state  $I_A$  over the expected duration of infection. Multiplying by  $\beta_A S_0$  gives the expected number of secondary infections resulting from interactions between susceptible and acutely-infected hosts.

Similarly, the expression for  $R_{0C}$  can be interpreted as follows: a fraction  $\alpha p_1/(d + h + \alpha)$  of exposed hosts  $E$  progress to state  $I_A$ , and of these acutely-infected hosts, a fraction  $\delta(1 - p_2)/(d + h + \mu_A + \delta)$  progress to state  $I_C$  spending an average of  $1/(d + h + \mu_C + \gamma)$  days in state  $I_C$  over the expected duration of infection. Alternatively, a fraction  $\alpha(1 - p_1)/(d + h + \alpha)$  of exposed hosts progress directly to state  $I_C$  and spend an average of  $1/(d + h + \mu_C + \gamma)$  days in state  $I_C$ . Multiplying by  $\beta_C S_0$  gives the expected number of secondary infections resulting from interactions between susceptible and chronically-infected hosts.

For model 2, we considered that  $E_j, I_{Aj}, I_{Cj}, E_a, I_{Aa}$ , and  $I_{Ca}$  are the infected states but that new infections occur only when a susceptible bird  $S_j$  or  $S_a$  became exposed  $E_j$  or  $E_a$ . Thus,  $F$  and  $V$  matrices are:

$$F = \begin{bmatrix} 0 & \beta_{Aj} S_{j0} & \beta_{Cj} S_{j0} & 0 & \beta_{Aa} S_{j0} & \beta_{Ca} S_{j0} \\ 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & \beta_{Aj} S_{a0} & \beta_{Cj} S_{a0} & 0 & \beta_{Aa} S_{a0} & \beta_{Ca} S_{a0} \\ 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 0 & 0 \end{bmatrix},$$

$$V = \begin{bmatrix} c_1 & 0 & 0 & 0 & 0 & 0 \\ -\alpha p_1 & c_2 & 0 & 0 & 0 & 0 \\ -\alpha(1-p_1) & -\delta(1-p_1) & c_3 & 0 & 0 & 0 \\ -\Omega & 0 & 0 & c_4 & 0 & 0 \\ 0 & -\Omega & 0 & -\alpha p_1 & c_5 & 0 \\ 0 & 0 & -\Omega & -\delta(1-p_1) & -\delta(1-p_2) & c_6 \end{bmatrix},$$

where

$$\begin{aligned} c_1 &= d_j + h_j + \Omega + \alpha, \\ c_2 &= d_j + h_j + \Omega + \mu_{Aj} + \delta, \\ c_3 &= d_j + h_j + \Omega + \mu_{Cj} + \gamma, \\ c_4 &= d_a + h_a + \alpha, \\ c_5 &= d_a + h_a + \mu_{Aa} + \delta, \\ c_6 &= d_a + h_a + \mu_{Ca} + \gamma. \end{aligned}$$

Thus,  $R_0$  is defined as:

$$R_0 = \rho(FV^{-1}) = \frac{1}{2} [R_{01} + R_{02} + \sqrt{(R_{01} - R_{02})^2 + 4R_{03}R_{04}}], \quad (A2)$$

where

$$\begin{aligned} R_{01} &= \frac{\alpha p_1 \beta_{Aj} S_{j0}}{c_1 c_2} + \frac{\alpha(1-p_1) \beta_C S_{j0}}{c_1 c_3} + \frac{\alpha p_1 \delta(1-p_2) \beta_C S_{j0}}{c_1 c_2 c_3} + \frac{\Omega \alpha p_1 \beta_{Aa} S_{j0}}{c_1 c_4 c_5} + \frac{\Omega \alpha p_1 \beta_{Aa} S_{j0}}{c_1 c_2 c_5} + \\ &\quad \frac{\Omega \alpha(1-p_1) \beta_C S_{j0}}{c_1 c_4 c_6} + \frac{\Omega \alpha(1-p_1) \beta_C S_{j0}}{c_1 c_3 c_6} + \frac{\Omega \alpha p_1 \delta(1-p_2) \beta_C S_{j0}}{c_1 c_4 c_5 c_6} + \frac{\Omega \alpha p_1 \delta(1-p_2) \beta_C S_{j0}}{c_1 c_2 c_5 c_6} + \frac{\Omega \alpha p_1 \delta(1-p_2) \beta_C S_{j0}}{c_1 c_2 c_3 c_6}, \\ R_{02} &= \frac{\alpha p_1 \beta_{Aj} S_{a0}}{c_1 c_2} + \frac{\alpha(1-p_1) \beta_C S_{a0}}{c_1 c_3} + \frac{\alpha p_1 \delta(1-p_2) \beta_C S_{a0}}{c_1 c_2 c_3} + \frac{\Omega \alpha p_1 \beta_{Aa} S_{a0}}{c_1 c_4 c_5} + \frac{\Omega \alpha p_1 \beta_{Aa} S_{a0}}{c_1 c_2 c_5} + \\ &\quad \frac{\Omega \alpha(1-p_1) \beta_C S_{a0}}{c_1 c_4 c_6} + \frac{\Omega \alpha(1-p_1) \beta_C S_{a0}}{c_1 c_3 c_6} + \frac{\Omega \alpha p_1 \delta(1-p_2) \beta_C S_{a0}}{c_1 c_4 c_5 c_6} + \frac{\Omega \alpha p_1 \delta(1-p_2) \beta_C S_{a0}}{c_1 c_2 c_5 c_6} + \frac{\Omega \alpha p_1 \delta(1-p_2) \beta_C S_{a0}}{c_1 c_2 c_3 c_6}, \\ R_{03} &= \frac{\alpha p_1 \beta_{Aa} S_{j0}}{c_4 c_5} + \frac{\alpha(1-p_1) \beta_C S_{j0}}{c_4 c_6} + \frac{\alpha p_1 \delta(1-p_2) \beta_C S_{j0}}{c_4 c_5 c_6}, \\ R_{04} &= \frac{\alpha p_1 \beta_{Aa} S_{a0}}{c_4 c_5} + \frac{\alpha(1-p_1) \beta_C S_{a0}}{c_4 c_6} + \frac{\alpha p_1 \delta(1-p_2) \beta_C S_{a0}}{c_4 c_5 c_6}. \end{aligned}$$

The terms  $R_{01}$  and  $R_{02}$  represent the average number of secondary juvenile or adult infections, respectively, produced by one exposed juvenile  $E_j$  during its entire infectious period. The terms  $R_{03}$  and  $R_{04}$  represent the average number of secondary juvenile or adult infections produced by one exposed adult  $E_a$  during its entire infectious period, respectively.

## References

1. van den Driessche P, Watmough J (2002) Reproduction numbers and sub-threshold endemic equilibria for compartmental models of disease transmission. *Mathematical Biosciences* 180: 29-48.
2. Diekmann O, Heesterbeek JAP, Metz JAJ (1990) On the definition and the computation of the basic reproduction ratio ( $R_0$ ) in models for infectious diseases in heterogeneous populations. *Journal of Mathematical Biology* 28: 365-382.