

1 **S1 Text. SI Methods & Results**

2

3 **Drivers for Rift Valley fever emergence in Mayotte:**

4 **a Bayesian modelling approach**

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11 **SI Methods**

12 **Natural history of disease and demographics.** The age-stratified SEIR model
13 diagram is presented in S2 Fig.

14 **Parameters: survival rates using demographic data.** In the absence of disease, the
15 demographic model is:

16 For the zero-to-one year old animals (i.e. age group $a=1$):

17
$$N_{1,t+1} = \alpha(1-\delta)N_{1,t} + (1-\alpha)\sum_{a=1}^9 N_{a,t} + (1-\alpha_{10})N_{10,t} \quad (S1)$$

18 For the one to nine years old animals (i.e. age groups $a \in [2-9]$):

19
$$N_{a,t+1} = \alpha(1-\delta)N_{a,t} + \alpha\delta N_{a-1,t} \quad (S2)$$

20 For animals of 9 years-old and older (i.e. age group $a=10$):

21
$$N_{10,t+1} = \alpha_{10}N_{10,t} + \alpha\delta N_{9,t} \quad (S3)$$

22

23

24 Age-groups presented in the above equations and S3 Fig were defined as follow:

25 Age-group 1 (a=1): Animals \leq 12 months

26 Age-group 2 (a=2): $>$ 12 months-old to \leq 24 months-old

27 Age-group 3 (a=3): $>$ 24 months-old to \leq 36 months-old

28 Age-group 4 (a=4): $>$ 36 months-old to \leq 48 months-old

29 Age-group 5 (a=5): $>$ 48months-old to \leq 60 months-old

30 Age-group 6 (a=6): $>$ 60 months-old to \leq 72 months-old

31 Age-group 7 (a=7): $>$ 72 months-old to \leq 84 months-old

32 Age-group 8 (a=8): $>$ 84 months-old to \leq 96 months-old

33 Age-group 9 (a=9): $>$ 96 months-old to \leq 108 months-old

34 Age-group 10 (a=10): $>$ 108 months-old

35

36 Where δ is the weekly ageing factor (S1 Table), and α and α_{10} are the survival rates
37 for age groups 1 to 9 and age-group 10, respectively. N_a is the number of animals in
38 each age group ($a \in [1-10]$) (S3 Fig - with the number of small ruminants and cattle in
39 each age group being summed up), and the population size N is equal to 30,000 and
40 assumed constant:

41
$$N = \sum_{a=1}^{10} N_a \quad (S4)$$

42 At the stable equilibrium, the number \bar{N}_a of animals in each age group a can be
43 expressed as a function of the population size N , δ , α and α_{10} :

44
$$\bar{N}_1 = \frac{N}{\theta} \quad \text{with } \theta = 1 + \sum_{x=1}^8 \left[\frac{\delta\alpha}{1-(1-\delta)\alpha} \right]^x + \left[\frac{\delta\alpha}{1-\alpha_{10}} \right] \left[\frac{\delta\alpha}{1-(1-\delta)\alpha} \right]^8 \quad (S5)$$

45
$$\bar{N}_{10} = \left[\frac{\delta\alpha}{1-\alpha_{10}} \right] \left[\frac{\delta\alpha}{1-(1-\delta)\alpha} \right]^8 \frac{N}{\theta} \quad (\text{S6})$$

46 And for $a \in [2-9]$:

47
$$\bar{N}_a = \left[\frac{\delta\alpha}{1-(1-\delta)\alpha} \right]^{a-1} \frac{N}{\theta} \quad (\text{S7})$$

48

49 N and δ are fixed, $N=30,000$ and $\delta=0.021$, and the parameters α and α_{10} are estimated

50 by maximising a Poisson likelihood function: the observed number of animals n_a in

51 each age-group a follows a Poisson distribution of mean \bar{N}_a , $n_a \sim \text{Poisson}(\bar{N}_a)$, and

52 the likelihood function is expressed as:

53
$$L = \prod_a \frac{\bar{N}_a^{n_a} e^{-\bar{N}_a}}{n_a!} \quad (\text{S8})$$

54 The estimated values of α and α_{10} are shown in S1 Table. The reported n_a and

55 simulated number of animals at equilibrium \bar{N}_a of animals in each age category are

56 both shown in S3 Fig.

57

58 **Parameters.** S2 Table shows the results if the IgM testing on imported animals

59 caught in 2008.

60

61 **Model fitting and parameter estimation.** Parameter estimation was done by fitting

62 the age-specific simulated proportion of immune animals $p_{a,i}$, for each

63 epidemiological year i , to RVF serological data (Oct 2004-Jun 2016), as presented in

64 Metras et al. 2016 [25], such as (black dots on Fig 3A-H and S6A-H Fig):

65
$$p_{a,i} = R_{a,i}/N_a, a \text{ in } [1, 10] \quad (\text{S9})$$

66

67 Note however that for Oct 2004-Jun 2008, serological data were only available for the
 68 whole population so we aggregated model outputs by month (blue dots on Fig 2, S4
 69 Fig, and S5 Fig):

$$70 \quad p_i = \sum_{a=1}^{10} p_{a,i} \quad (\text{S10})$$

71 We sampled from the posterior distribution of all six parameters
 72 $\theta = \{imm_t0, a, b, t_{imp}, P, p_{seized}\}$ using a Monte Carlo Markov Chain Metropolis-Hastings
 73 (MCMC-MH) algorithm [49]. For all parameters, we assumed uniform priors (Table
 74 1). The number of IgG positive animals $x_{a,i}$ among the $n_{a,i}$ tested animals in the age-
 75 group a during period i followed a binomial distribution;

$$76 \quad x_{a,i} \sim Bin(n_{a,i}, p_{a,i}) \quad (\text{S11})$$

77 The log-likelihood of the data was therefore given by:

$$78 \quad l(data|\theta) = \sum_i \sum_a l_{a,i}(x_{a,i}, n_{a,i} | \theta) \quad (\text{S12})$$

$$79 \quad l_{a,i}(x_{a,i}, n_{a,i} | \theta) = x_{a,i} \log p_{a,i} + (n_{a,i} - x_{a,i}) \log(1 - p_{a,i}) \quad (\text{S13})$$

80 We ran two independent MCMC-MH chains of 100,000 iterations, and visually
 81 checked that both chains converged to the same stationary distribution. We discarded
 82 the first 5,000 iterations of the burn-in periods and thinned at a ratio of 1:20 to
 83 eliminate auto-correlation (S10A-F Fig), finally we combined both chains. The
 84 median, 95% Credible Interval (CrI) and interquartile range (IQR) of the posterior
 85 distributions were computed (Table 3). The best model among the two had the lowest
 86 deviance information criterion value (DIC) [50]:

$$87 \quad DIC = D(\theta_m) + 2p_D \quad (\text{S14})$$

$$88 \quad D(\theta) = -2l(data|\theta) + k \quad (\text{S15})$$

$$89 \quad p_D = \frac{1}{2} \text{var}(D(\theta)) \quad (\text{S16})$$

90 Where $D(\theta)$ is the deviance of the model for a parameter set θ , $l(data|\theta)$ is the
91 likelihood of the data given θ , and k is a constant that cancel out when comparing
92 models; θ_m is the posterior mean of θ and p_D is the effective number of model
93 parameters, computed as half the variance of the deviance under the posterior.

94

95 **SI Results**

96 S4 Fig shows the fit of both linear and exponential models with no import at all over
97 the study period (models 2a and 2b), and of model 3 with NDVI not tied to
98 transmission. S7 Fig presents the resulting relationships between NDVI and R_s for
99 both the exponential and linear models. Both fits yielded very similar results (S5 Fig
100 and S6A-H Fig compared with Fig 2 and Fig 3A-H). Finally, the Forecasts 2-5 and
101 Forecasts 7-10 of the different scenarios of infectious animal imports (1, 10, 20, 30
102 infectious animals introduced in October 2016 and April 2017) are presented in S8A-
103 D Fig and S9A-D Fig. S10A-F Fig presents the autocorrelation plots for the 6
104 parameters of a MCMC chain for Model 1b.