# Supporting Information

### 2 SI 1: Implementation of the Crofton Method

The algorithm for fitting the Crofton Method (Crofton 1971) proceeds as follows. 3 First, obtain a dataset with n hosts where each host has some parasite intensity 0 4 5 to  $p_{max}$ . Starting with the full dataset, guess a vector of pre-mortality parameters  $(N_{p1}, \mu_{p1}, k_{p1})$  where  $N_{p1}$  is the total number of hosts before mortality,  $\mu_{p1}$ 6 is the parasite intensity before mortality, and  $k_{p1}$  is the parasite aggregation 7 before mortality. Given these parameters, use a negative binomial distribution 8 to calculate the predicted number of hosts with  $0, 1, 2, \dots p_{max}$  parasites. Compare 9 the expected number of hosts with  $0, 1, 2, \dots p_{max}$  parasites to the observed 10 number hosts with  $0, 1, 2, \dots p_{max}$  parasites and calculate the  $\chi^2$ -squared statistic 11 associated with the observed and predicted vectors. In reality, one often has to bin 12 the parasite intensity data because all parasite intensities are not represented in 13 the dataset. Continue to guess  $(N_{p1}, \mu_{p1}, k_{p1})$  vectors until a set of parameters is 14 found that minimizes the  $\chi^2$ - squared statistic. 15 Second, choose a truncation value  $(t_2)$  such that  $t_2 < p_{max}$ . Truncate the 16 data such that  $data_{truncated} \ll t_2$  and repeat the above iterative procedure to 17 calculate another set of parameters  $(N_{p2},\mu_{p2},k_{p2})$  that minimizes the  $\chi^2$ -squared 18 statistic on the truncated data. Choose a new truncated value  $t_3 < t_2$  and repeat 19 20 the first two steps. Continue to truncate the dataset until it only contains hosts with 0, 1, and 2 parasites (or 3 bins). Because the method attempts to estimate 21 three parameters, at least 3 classes are needed for all 3 parameters to be identifiable 22 (Royce and Rossignol 1990). 23 Once the iterative procedure has been completed, parasite-induced host 24 mortality is traditionally identified by plotting the different truncation values  $t_i$ 25 against the different values of  $N_{pi}$  and looking for a distinct "kink" in the resulting 26

plot. Once the "kink" as occurred, the values of  $N_{pi}$  will typically remain close to constant as  $t_i$  is decreased further. The "true" pre-mortality parameters  $N_{pt}$ ,  $\mu_{pt}$ , and  $k_{pt}$  are taken to be at the point where the "kink" occurs.

We provide an implementation and unit tests of the Crofton Method in Supplementary Information 4. Figure 10 visually shows that our implementation of the Crofton Method agrees with results previously published by Crofton (1971).

### SI 2: Implementation of the Adjei Method

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The Adjei Method for estimating PIHM has two steps (Adjei et al. 1986). The first 34 step is to estimate the parameters of the pre-mortality host-parasite distribution 35 using the Crofton Method (see SI 1). The three parameters estimated are the 36 total number of hosts before mortality  $N_p$ , the mean number of parasites per host 37 before mortality  $\mu_p$ , and the aggregation of parasites before mortality given by the 38 parameter  $k_p$  from a negative binomial distribution. When  $k_p$  is small, parasites 39 are highly aggregated among hosts and when  $k_p$  is large parasites are more evenly 40 41 distributed across hosts (Wilson et al. 2002). The implementation of the Crofton Method has been discussed at length elsewhere (e.g. Royce and Rossignol 1990; 42 Lester 1984, and in SI 1) and we provide a tested implementation of the method 43 in SI 4. 44 The second step of the Adjei Method is to make the assumption that 45 46 infection, host mortality, and sampling occur in that order and are temporally separate (Adjei et al. 1986). Next, Adjei et al. assume that the host survival 47 function follows the logistic form 48

$$h(\text{survival}; x, a, b) = \frac{\exp(a - b\log(x))}{1 + \exp(a - b\log(x))}$$
(1)

where x is the parasite intensity in a given host and a and b are the two parameters

of the logistic function. Generally, a larger a allows for hosts to tolerate larger parasite intensities before experiencing parasite-induced mortality and a larger b leads to a more rapid decline in the probability of host survival as parasite intensity increases. The value  $\exp(a/b)$  is referred to as the  $LD_{50}$ . Individuals with loads higher than this will have a greater than 50% chance of death.

By taking the first and second derivatives of equation 1, one can easily find that the maximum rate of decline in host survival probability with increasing parasite intensity occurs at the  $LD_{50}$  and has a value of b/4. This is in many ways analogous to the parasite pathogenicity parameter  $\alpha$  given in classic macroparasite models, which specifies the slope of the linear relationship between between host death rate and parasite intensity (Anderson and May 1978; Isham 1995). The parameter a is easily interpreted by holding b constant and looking at how a one unit change in a affects the log parasite intensity at which some percentage p of hosts experience mortality. Letting  $a_1$  and  $a_2$  be two different values of a and  $a_3$  and  $a_4$  be two different parasite intensities, a bit of rearranging of equation 1 gives.

$$\log \frac{p}{1-p} = a_1 - b \log x_1$$

$$-\log \frac{p}{1-p} = a_2 - b \log x_2$$

$$0 = a_1 - a_2 - b \log x_1 + b \log x_2$$

$$a_2 - a_1 = b(\log x_2 - \log x_1)$$

If  $a_2 - a_1 = 1$ , than the change in log parasite intensity at which p percentage of hosts survive is 1/b.

To estimate the parameters in equation 1, the Adjei Method first calculates the expected number of hosts with a given parasite load x by using the equation  $g(x; \mu_p, k_p) * N_p$ , where  $g(x; \mu_p, k_p)$  is the negative binomial pre-mortality distribution. Second, the observed and predicted number of hosts with x parasites are paired as a single data point and the method then assumes that this data point follows a binomial distribution with the total number of "trials" equal to the predicted number of hosts and the total number of "successes" equal to the observed number of hosts. In some cases, the observed number of hosts is greater than the expected number of hosts and the Adjei Method alters the data so that the observed is equal to the predicted (Adjei et al. 1986). After this questionable manipulation, the (observed, predicted) pairs are fit to a standard Generalized Linear Model (McCullagh and Nelder 1989) with a binomial response variable and a logistic link function given by equation 1. This model provides estimates for parameters a, b and  $LD_{50}$ .

While not included in the original implementation of the Adjei Method, a  $\chi^2$  test with a degrees of freedom of 1 can be used to assess whether a GLM model that includes parasite intensity as a predictor of host survival probability is a "better" model than a GLM without this predictor. This allows the Adjei Method to determine whether PIHM is a significant factor in a host-parasite system.

The Adjei Method's most glaring deficiency is the need to alter the observed data in order to fit the model into the binomial GLM framework. A second more subtle problem with the Adjei Method is the potential need to bin data in order to predict greater than one host in a given parasite intensity class. For example, if the total number of hosts pre-mortality was 50, the mean number of parasites per host pre-mortality was 100 and the aggregation parameter was 1, applying the equation  $g(x; \mu_p = 100, k_p = 1) * 50$  would result in less than 1 individual in all parasite intensities x. In other words, the Adjei Method cannot be applied to samples with either very high mean parasite loads, small sample sizes, or both without some sort of binning of the data. While this is not a flaw per se, it does add a certain level of subjectivity (i.e. which bins should you use?) to a method that already has serious potential issues. In this analysis, we always assume the

- Adjei Method is not binning the data, though we provide code for applying the
- 99 binning method in SI 4.

### 100 SI 3: Additional Figures

101 See Figures 1 - 9.

## SI 4: Code and unit tests for estimating parasite-induced host

#### 103 mortality

- 104 Python code, unit tests, and a help file for the Crofton Method, the Adjei Method
- and the Likelihood Method can be found at https://github.com/mqwilber/
- 106 parasite\_mortality

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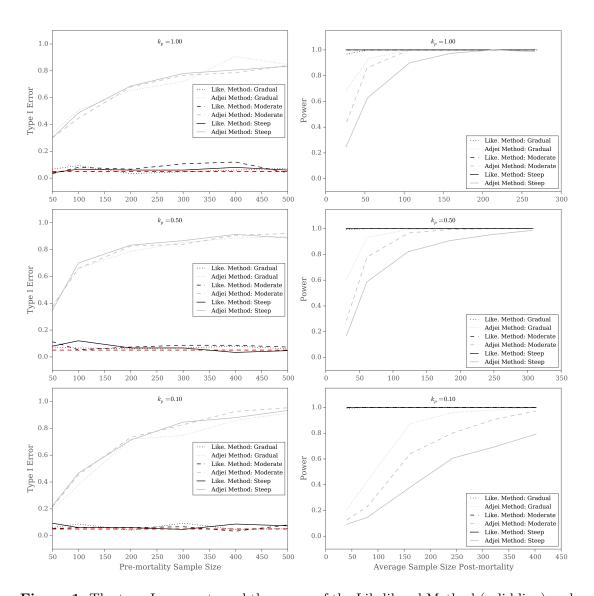


Figure 1: The type I error rate and the power of the Likelihood Method (solid line) and the Adjei Method (dashed lines) when  $\mu_p = 10$  for various shapes of the host survival function and levels of aggregation  $k_p$ . The first column gives the type I error rate of each method for falsely detecting PIHM when none is present. The red line gives the the pre-set type I error rate of  $\alpha = 0.05$ . The second column gives the power of a given method to detect PIHM when it is actually occurring.

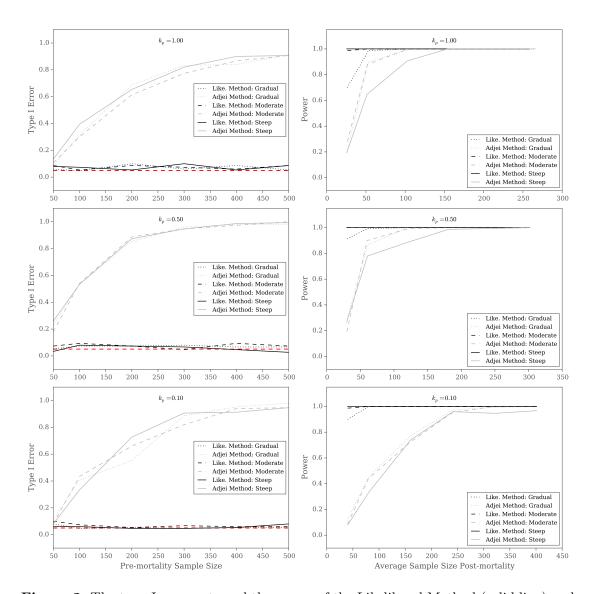


Figure 2: The type I error rate and the power of the Likelihood Method (solid line) and the Adjei Method (dashed lines) when  $\mu_p = 50$  for various shapes of the host survival function and levels of aggregation  $k_p$ . The first column gives the type I error rate of each method for falsely detecting PIHM when none is present. The red line gives the the pre-set type I error rate of  $\alpha = 0.05$ . The second column gives the power of a given method to detect PIHM when it is actually occurring.

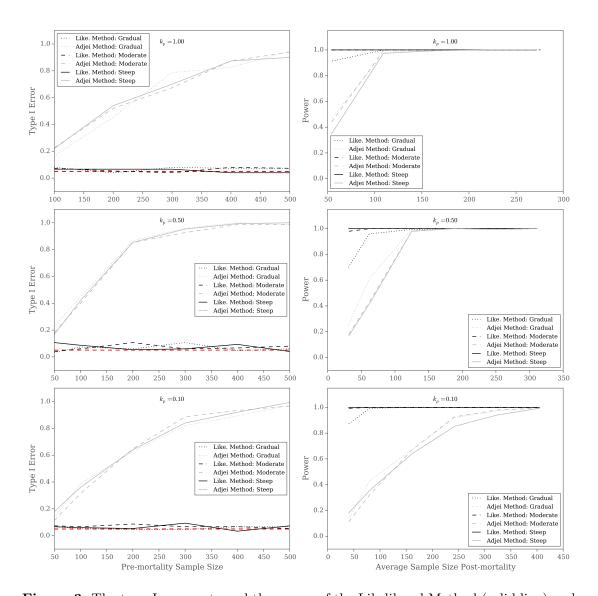


Figure 3: The type I error rate and the power of the Likelihood Method (solid line) and the Adjei Method (dashed lines) when  $\mu_p = 100$  for various shapes of the host survival function and levels of aggregation  $k_p$ . The first column gives the type I error rate of each method for falsely detecting PIHM when none is present. The red line gives the the pre-set type I error rate of  $\alpha = 0.05$ . The second column gives the power of a given method to detect PIHM when it is actually occurring.

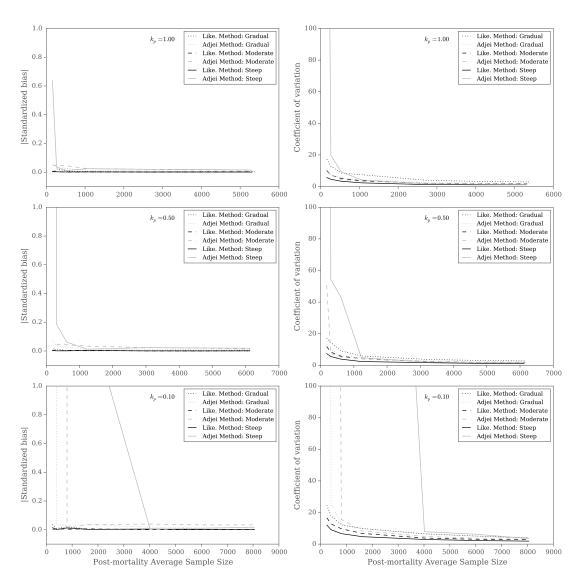


Figure 4: The bias and the precision of the Likelihood Method (solid line) and the Adjei Method (dashed lines) when  $\mu_p=10$  for various shapes of the host survival function and levels of aggregation  $k_p$  when estimating  $LD_{50}$ . The first column gives the bias of each method's  $LD_{50}$  estimate over 150 simulations. The second column gives the precision of each method's  $LD_{50}$  estimate over 150 simulations.

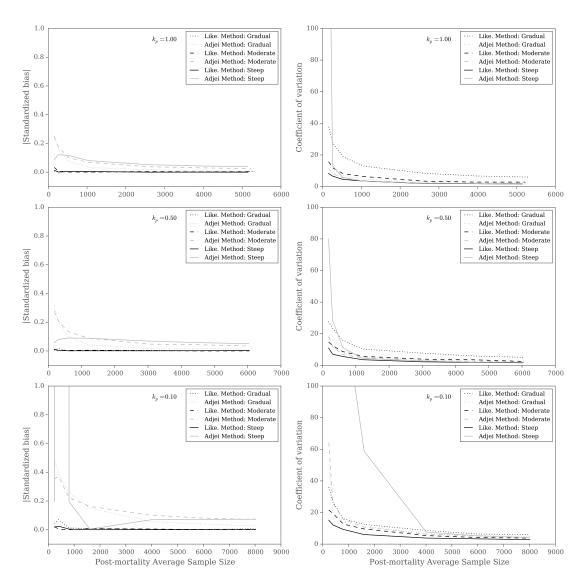


Figure 5: The bias and the precision of the Likelihood Method (solid line) and the Adjei Method (dashed lines) when  $\mu_p=50$  for various shapes of the host survival function and levels of aggregation  $k_p$  when estimating  $LD_{50}$ . The first column gives the bias of each method's  $LD_{50}$  estimate over 150 simulations. The second column gives the precision of each method's  $LD_{50}$  estimate over 150 simulations.

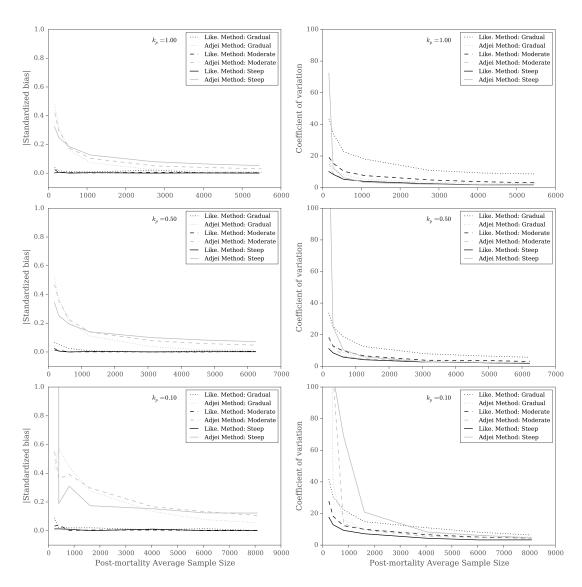


Figure 6: The bias and the precision of the Likelihood Method (solid line) and the Adjei Method (dashed lines) when  $\mu_p = 100$  for various shapes of the host survival function and levels of aggregation  $k_p$  when estimating  $LD_{50}$ . The first column gives the bias of each method's  $LD_{50}$  estimate over 150 simulations. The second column gives the precision of each method's  $LD_{50}$  estimate over 150 simulations.

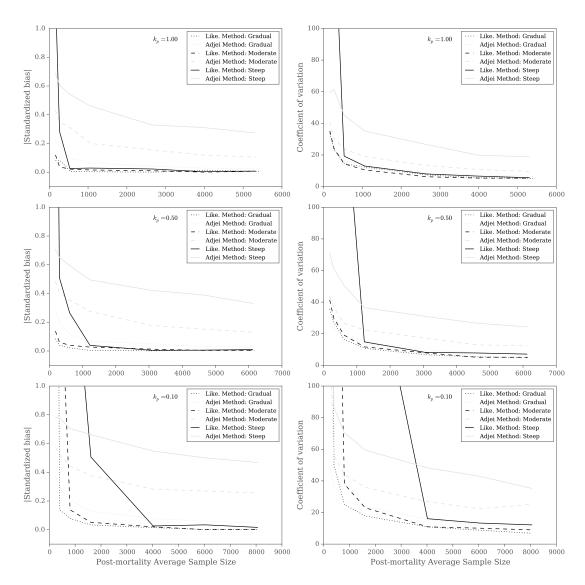


Figure 7: The bias and the precision of the Likelihood Method (solid line) and the Adjei Method (dashed lines) when  $\mu_p=10$  for various shapes of the host survival function and levels of aggregation  $k_p$  when estimating the a parameter of the host survival function. The first column gives the bias of each method's a estimate over 150 simulations. The second column gives the precision of each method's a estimate over 150 simulations.

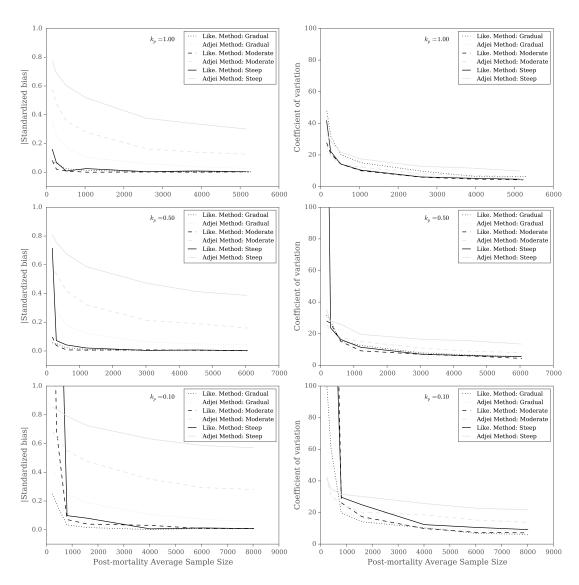


Figure 8: The bias and the precision of the Likelihood Method (solid line) and the Adjei Method (dashed lines) when  $\mu_p=50$  for various shapes of the host survival function and levels of aggregation  $k_p$  when estimating the a parameter of the host survival function. The first column gives the bias of each method's a estimate over 150 simulations. The second column gives the precision of each method's a estimate over 150 simulations.

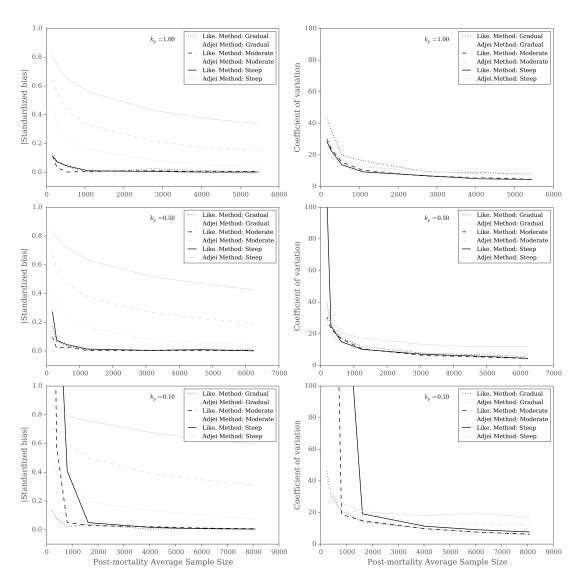


Figure 9: The bias and the precision of the Likelihood Method (solid line) and the Adjei Method (dashed lines) when  $\mu_p=100$  for various shapes of the host survival function and levels of aggregation  $k_p$  when estimating the a parameter of the host survival function. The first column gives the bias of each method's a estimate over 150 simulations. The second column gives the precision of each method's a estimate over 150 simulations.

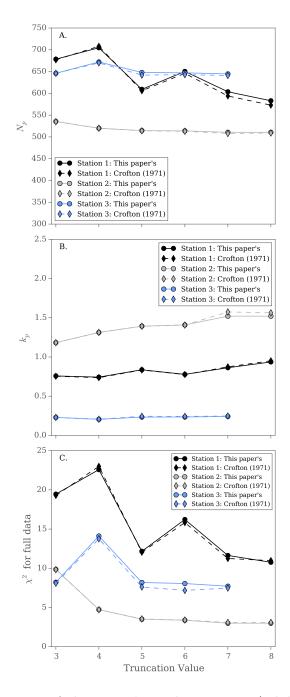


Figure 10: A comparison of this paper's implementation (solid line, circles) of the Crofton Method with the results given in Crofton (1971) (dashed line, diamonds). Figure A compares the predicted number of hosts in a population pre-mortality  $(N_p)$ . Figure B compares the predicted parasite aggregation pre-mortality  $(k_p)$ . Figure C compares the  $\chi^2$  statistic for each implementation. Three of the 6 stations fit by Crofton are shown here and all show that our implementation gives very similar results to those given by Crofton.