





Figures and figure supplements

Nationally-representative serostudy of dengue in Bangladesh allows generalizable disease burden estimates

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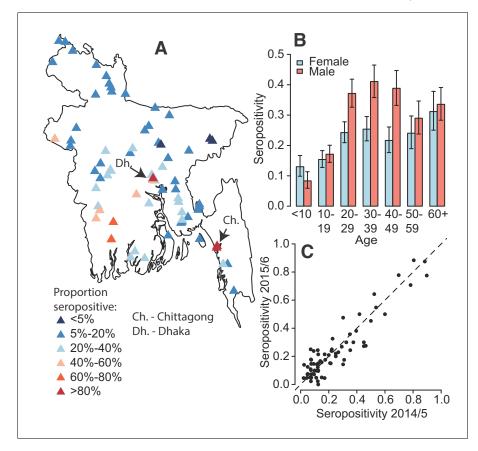


Figure 1. Dengue seropositivity in the sampled communities. (A) Locations of sampled communities and the estimated seroprevalence by community. (B) Proportion seropositive by age and sex with 95% confidence intervals. (C) Seropositivity in Y1 compared to seropositivity in Y2 for each community.

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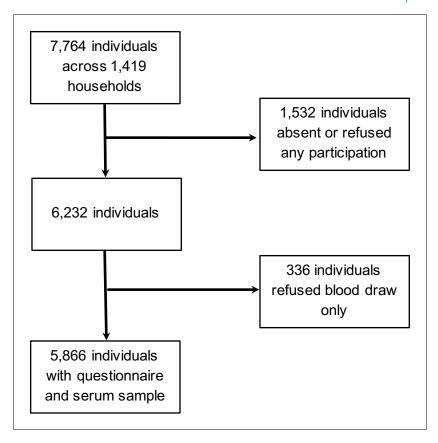


Figure 1—figure supplement 1. Participants included in study.



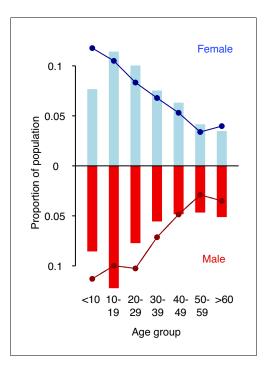


Figure 1—figure supplement 2. Distribution of age groups in years and sex (solid bars) in the study population compared to the 2011 census (lines). DOI: https://doi.org/10.7554/eLife.42869.005



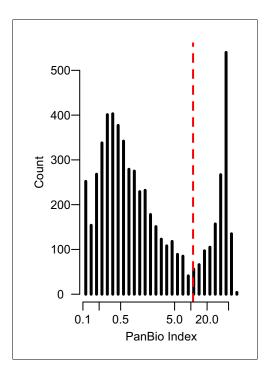


Figure 1—figure supplement 3. Histogram of PanBio Units derived from the optical densities of the PanBio assay with the manufacturer recommended cutpoint (PanBio Unit of 11) in dashed red. Those to the left of the dashed line are considered seronegative and those to the right of the line are considered seropositive.

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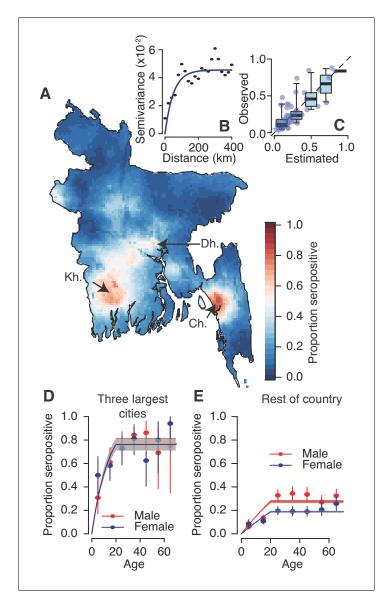


Figure 2. Modelled dengue seropositivity across Bangladesh and across age groups. (A) Spatial predictions of seropositivity for the whole country. Kh. = Khulna, Dh. = Dhaka, Ch. = Chittagong (B) Semivariogram showing spatial dependence between the proportion seropositive between communities as a function of distance between them. (C) Observed versus predicted levels of seropositivity by community from leave one out cross validation. (D) Observed (points) and fitted seropositivity by age for the sampled communities within the three largest cities (Khulna, Chittagong and Dhaka) for both males and females. (E) Observed and fitted seropositivity for the remaining communities by sex.



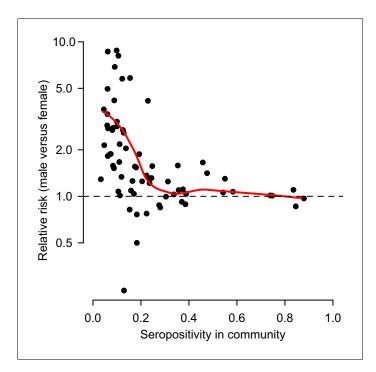


Figure 2—figure supplement 1. Relative risk of being seropositive for males versus females as a function of the overall proportion seropositive in the community. The red line represents a LOESS curve fit through the 70 points. DOI: https://doi.org/10.7554/eLife.42869.010



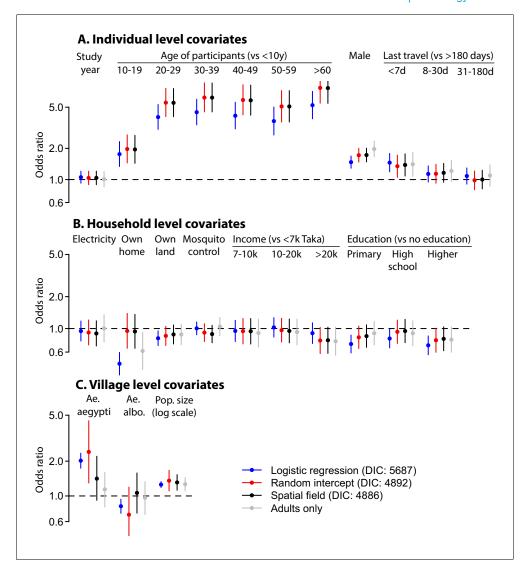


Figure 2—figure supplement 2. Differences in coefficient estimates in multivariable models run using logistic regression with no random intercept or spatial covariance (blue), with random intercepts at the household and community level (red) and with random intercepts at the household and community level and a Matern spatial covariance matrix (black, base model). In addition the results of the spatial field model but restricted to individuals over 20y is shown (grey). The bars represent 95% confidence intervals each time.



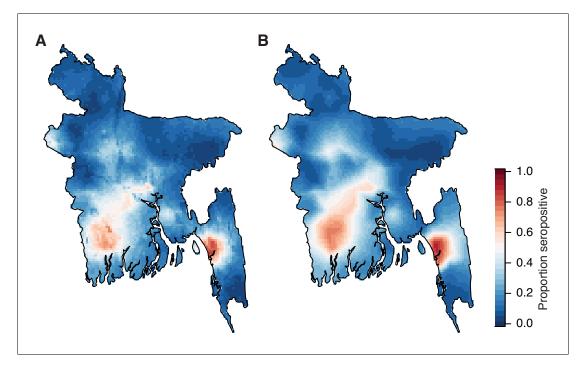


Figure 2—figure supplement 3. Comparison of risk maps using different prediction methods. (A) Spatial prediction map using age, sex and population size and a Matern covariance structure. (B). Spatial prediction using a Matern spatial covariance only. DOI: https://doi.org/10.7554/eLife.42869.012



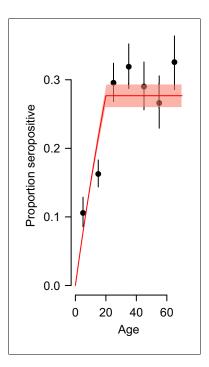


Figure 2—figure supplement 4. Observed proportion seropositive by age group with 95% confidence intervals (black) and the fit using the force of infection estimated by the catalytic model (red). We used the probability of being seropositive as a function of age from the whole country to estimate the proportion of the susceptible population (individuals that have never been exposed to dengue) that get infected each year. DOI: https://doi.org/10.7554/eLife.42869.013



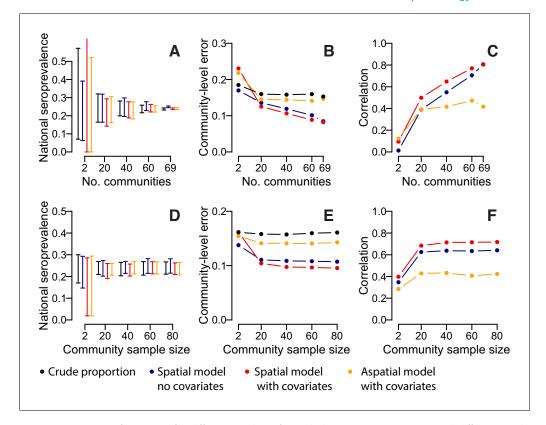


Figure 3. Accuracy of estimates for different number of sampled communities (top row) and different numbers of sampled individuals per community (bottom row) using different estimation methods. (A) 95% range of estimates of overall seroprevalence from 100 repeated iterations when data from a random subset of communities is used. (B) Mean absolute error among heldout communities over repeated iterations. (C) Mean correlation between predicted and observed seroprevalence among heldout communities. (D) 95% range of estimates of overall seroprevalence from 100 repeated iterations when data from a random number of individuals from 50 communities is used. (E) Mean absolute error among 20 randomly selected heldout communities over repeated iterations. (F) Mean correlation between predicted and observed seroprevalence among 20 randomly selected heldout communities. The different estimation methods are overall proportion seropositive (black), spatial correlation model using Matern covariance structure and no covariates (blue), spatial correlation model using Matern covariance structure and population size as covariates, logistic regression using age, sex and population size as covariates with no spatial component (orange).