

Risk-assessment of *Aedes*-borne diseases and need for modelling-tools for decision making

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Climate-Sensitive Vector Dynamics Modelling Workshop
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Risk assessment questions



probability of transmission (in the EU)

Risk = X

impact (severity and amount of disease)

Modified from <https://www.ecdc.europa.eu/sites/default/files/documents/operational-tool-rapid-risk-assessment-methodology-ecdc-2019.pdf>

- What is the current/future probability of an outbreak happening in a geographic area?
- What is the expected size of an outbreak, if it occurs? (we do not yet look at impact in terms of e.g. DHF)
- How could the public health response (vector control/case finding) mitigate the probability and impact?

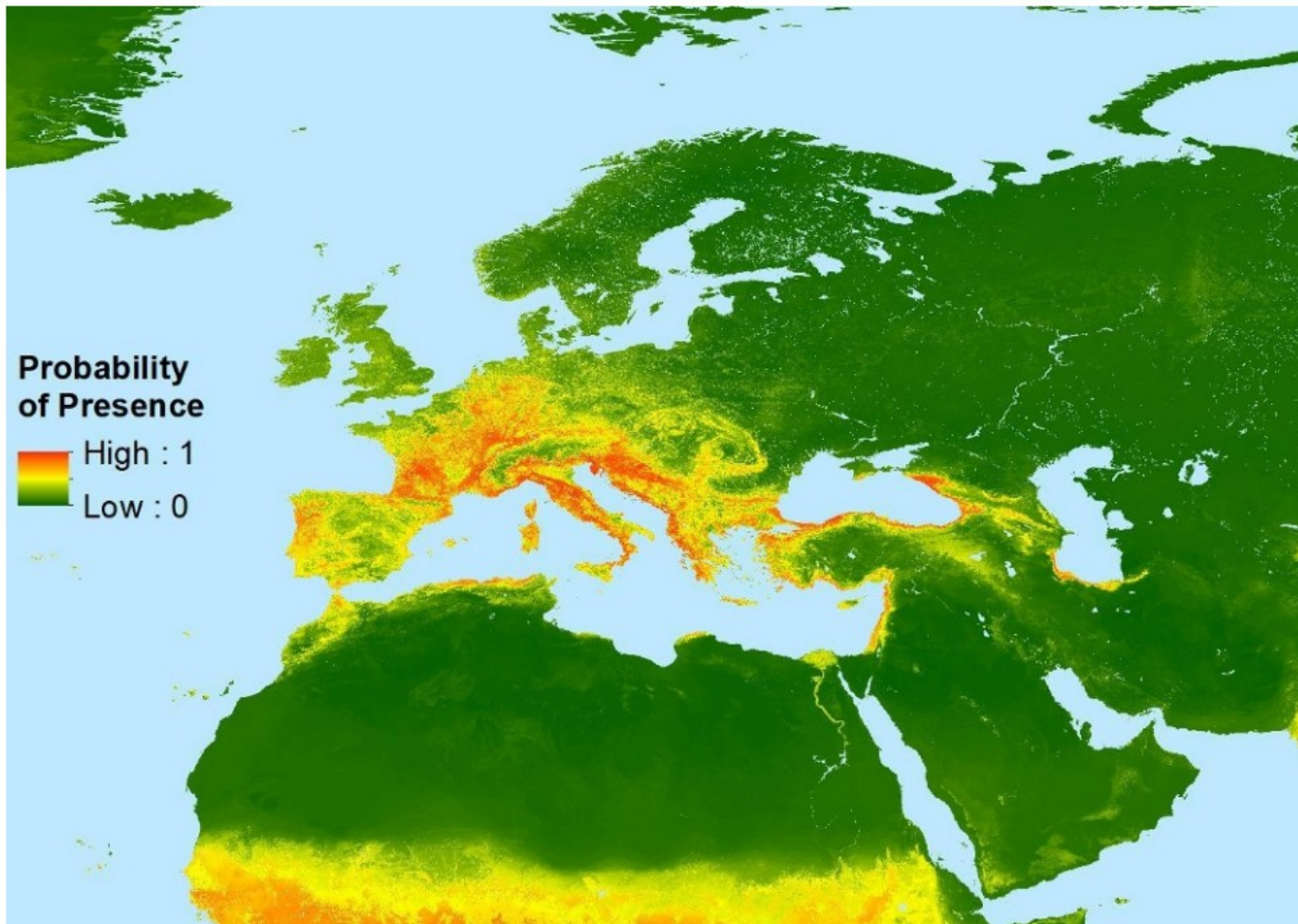
Statistical and mathematical modelling



Where vectors are not present:

- Are the environmental and climatic conditions appropriate for the establishment of competent vectors in a particular area?
- Model ecological niche suitability of the area

Wint et al., 2020 doi: 10.2903/sp.efsa.2020.EN-1800



Masked predictions for *Ae. albopictus*.

Wint et al., 2020 doi:10.2903/sp.efsa.2020.EN-1847

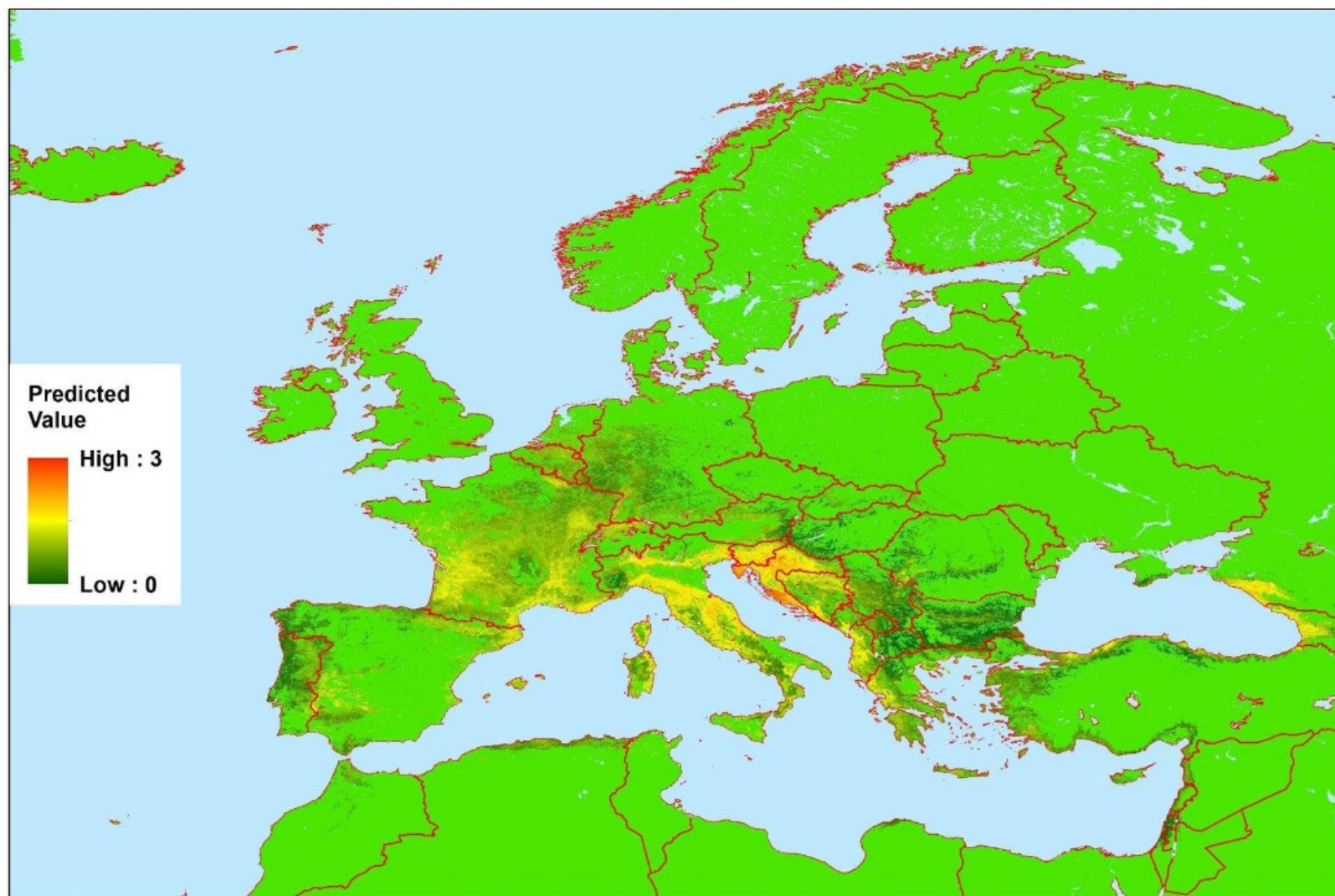


Figure 3: Predicted abundance (log egg equivalent), masked, 1 km aggregation for *Aedes albopictus*.

Statistical and mathematical modelling



Where vectors are not present:

- Are the environmental and climatic conditions appropriate for the establishment of competent vectors in a particular area?
- Model ecological niche suitability of the area

Where vectors are present:

- What is the receptivity of the vectors in an area and at a timepoint to transmit the disease in case of importation?
- Model the vectorial capacity (number of expected onward transmissions if an infectious individual is exposed to the vector population for one day) of the vectors in an area at a given time.
- What is the vulnerability (rate of importation of viraemic individuals)?
- Statistical prediction models of imported cases in space and time dependent on travel data and prevalence/incidence in source countries.
- What is the receptivity in terms of capacity of the health system to reduce exposure of mosquitoes to imported cases
- Model controlled reproduction number (R_c) based on duration of infection (and case-detection-action delays)

Statistical and mathematical modelling (continued)



Where local transmission is likely, and depending on the past and future weather conditions:

- What is the expected size of an outbreak, if it occurs?
- Model R_c based on health system response (including preventive vector control)
- What are the most effective and cost-effective (accepted) public-health strategies for mitigating the probability and impact of local transmission?
- Scenario-modelling combined with intervention cost-effectiveness data

Risk modelling at ECDC



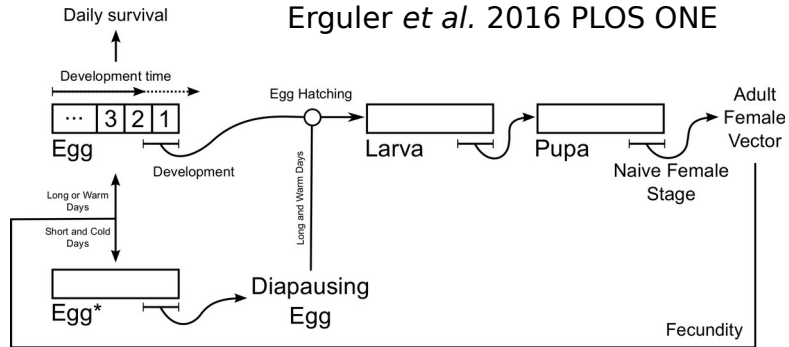
ArboRisk – a tool in development to provide ‘near-real time’ maps of risk at NUTS3 spatial resolution and weekly temporal resolution in terms of:

- Outbreak risk (autochthonous transmission happening)
- Outbreak impact (the size of a possible outbreak)

Using:

- Virus importation probability, based on:
 - Historically imported cases
 - Human population density
 - Distance to airports
 - IATA passenger data
- Vector dynamic model, based on:
 - Vector presence
 - Rainfall
 - Temperature
 - Photoperiod
 - Human population density
- Dynamic transmission model, based on:
 - Temperature
 - Vector model
 - Virus importation
 - Autochthonous transmission data for scaling of results

Aedes albopictus-borne chikungunya transmission model

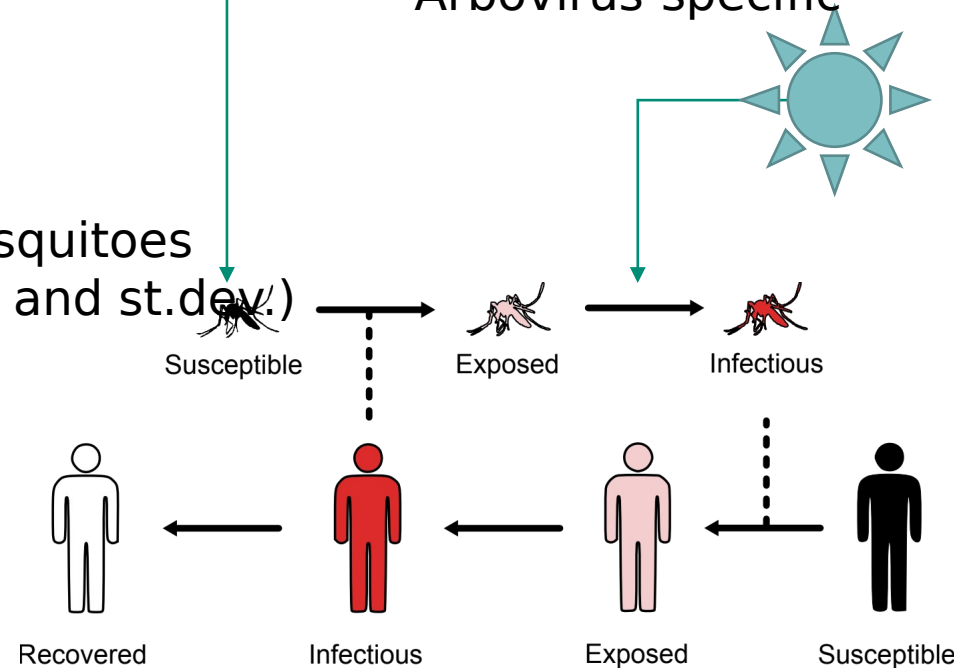


Extrinsic incubation period (EIP)

- Temperature-driven
- Arbovirus-specific

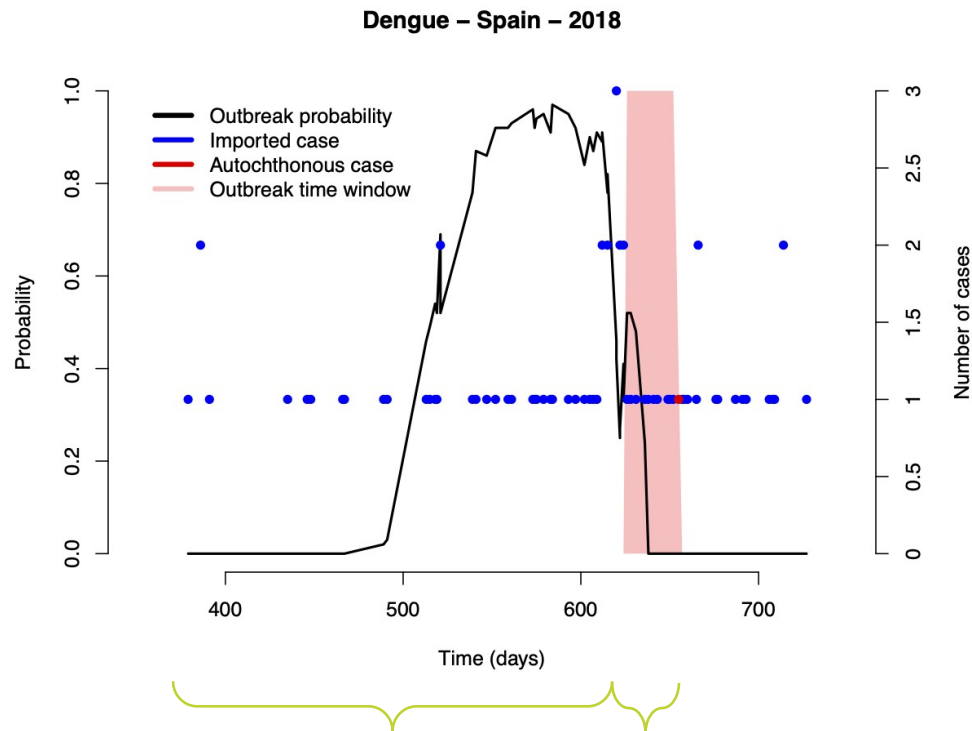
Transmission model input

(Relative) number of adult female mosquitoes
Adult female mosquito lifetime (mean and st.dev.)
Fecundity
Mean air temperature



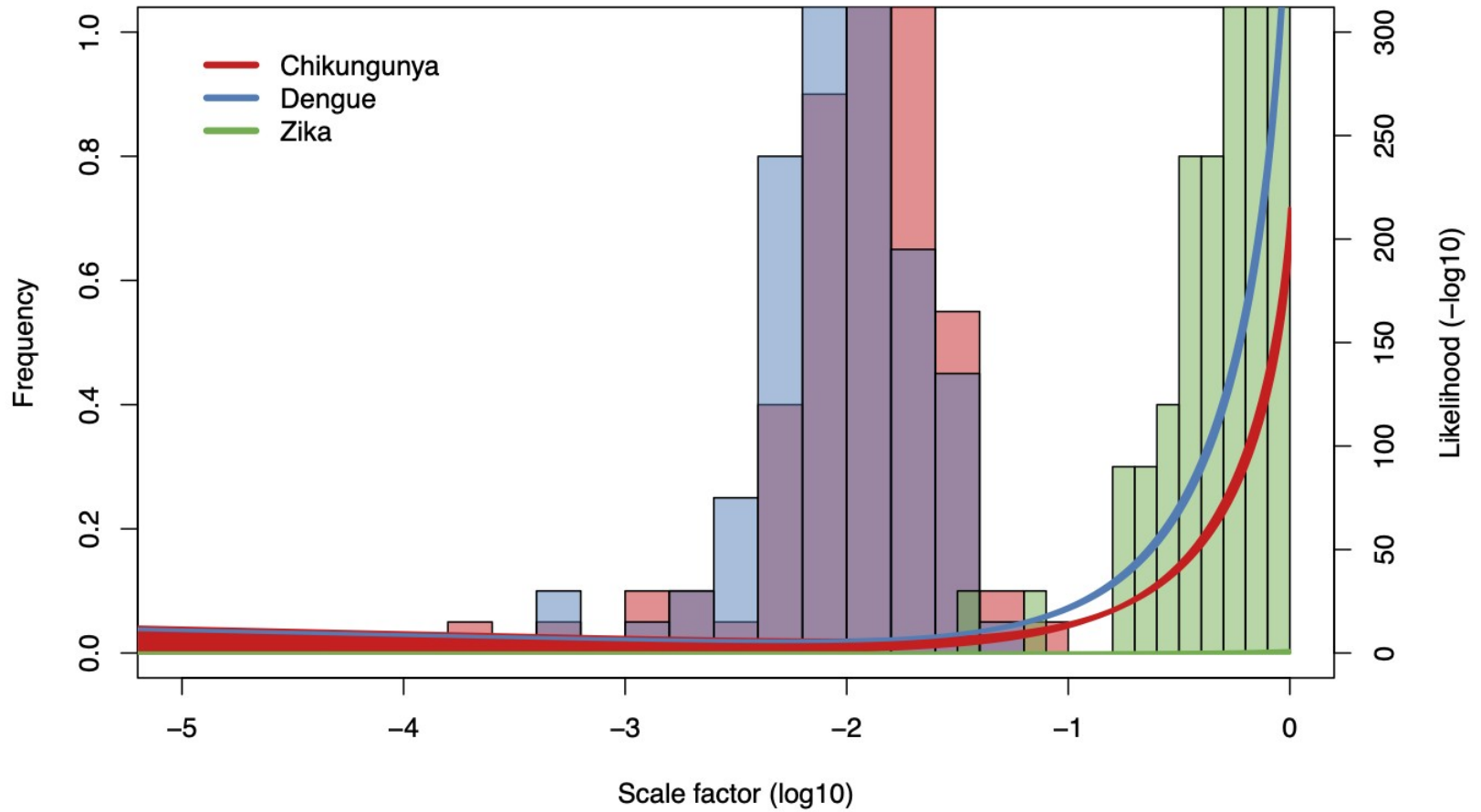
Modification from Erguler *et al.* 2017 PLOS ONE

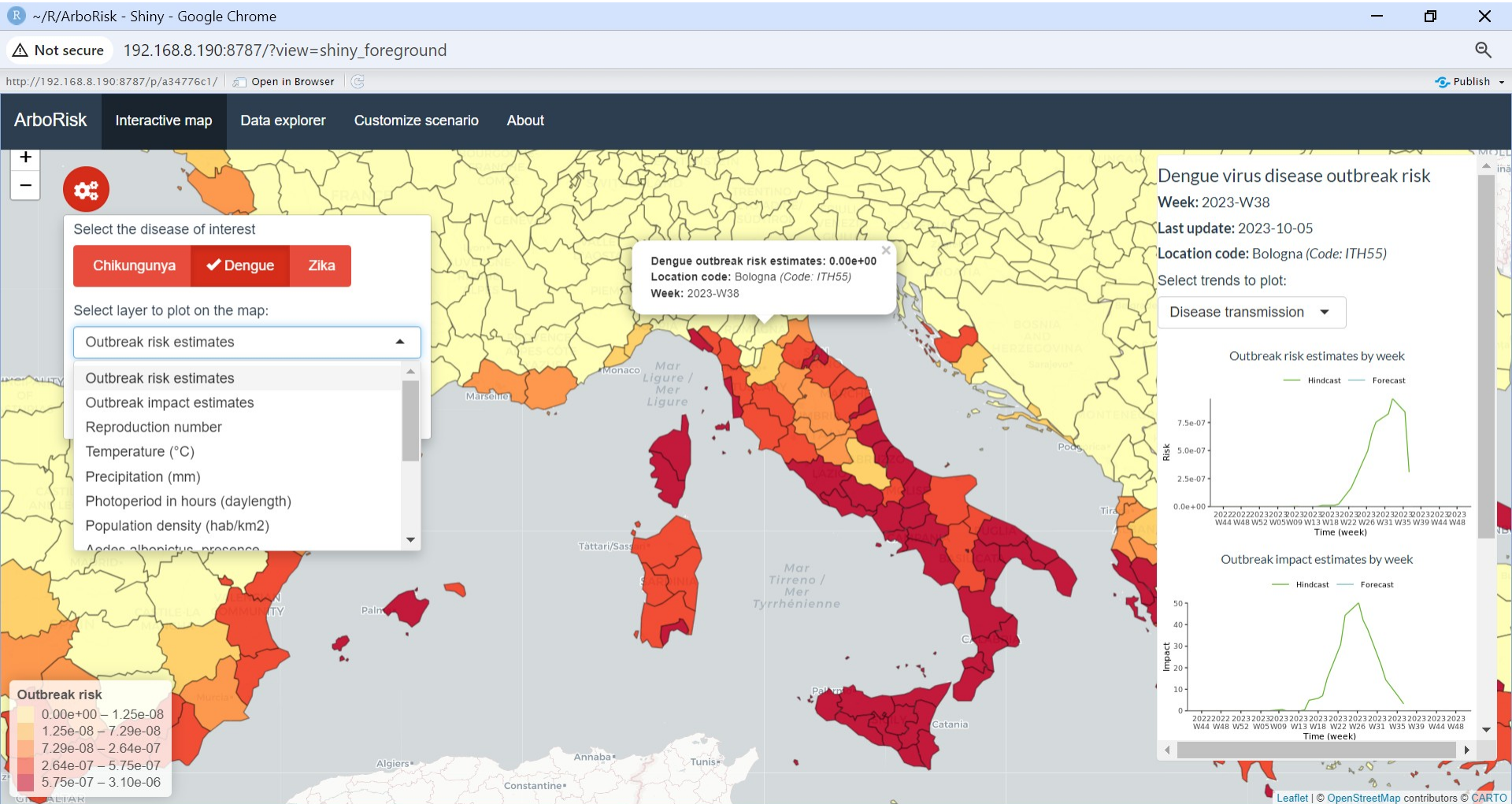
Validating and scaling



Probability of none to cause an outbreak Probability of at least one to cause an outbreak

Validating and scaling

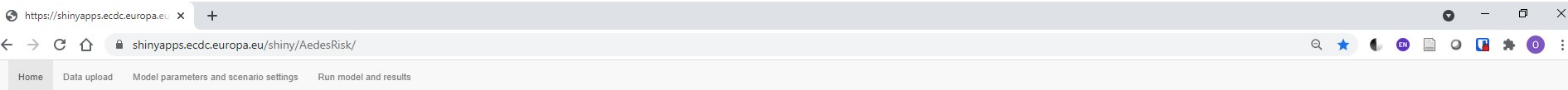




Scenario modelling at ECDC



<https://shinyapps.ecdc.europa.eu/shiny/AedesRisk/>



AedesRisk Version 1.0

A tool to support the decision making process for surveillance and vector control of dengue, chikungunya and Zika virus disease in Europe

Tool source directory: /home/shiny/ShinyApps/AedesRisk

in collaboration with

for

Description

This tool comprises of a compartmental stochastic model at a population-scale, which is based on three different components: entomological dynamics, transmission dynamics in the human population, and (optionally) spatial dynamics. Mosquitoes and humans reside in patches where their respective densities are assumed to be homogeneous. Humans may move between patches, propagating viruses spatially if infected or infectious. The mosquito dynamics is rainfall and temperature driven, and the transmission is temperature driven. The user may experiment with vector control affecting immature or mature mosquito stages, and vary virus importation in humans.

Manual

[Download the manual](#)

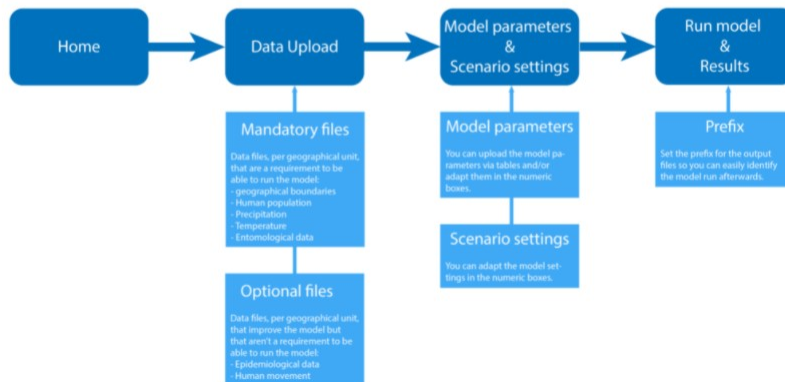
Model code (R)

[Download the model code](#)

For experienced R users who want to run the model on their own computers, the R code of this shiny app is made available for download.

Workflow

For a more detailed description please refer to the tool's manual.

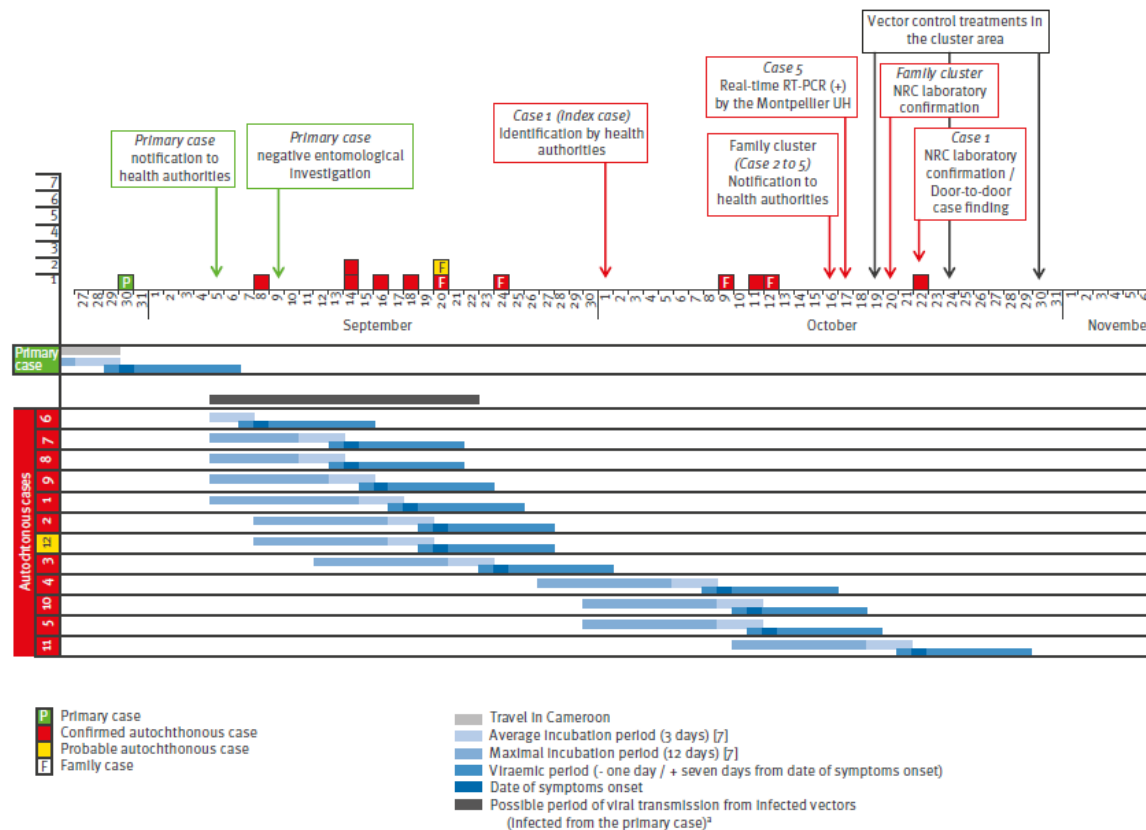


Chikungunya outbreak in Montpellier, France, September to October 2014

E Delisle (delisle.elsa@gmail.com)¹, C Rousseau¹, B Broche², I Leparc-Goffart³, G L'Ambert⁴, A Cochet¹, C Prat³, V Foulongne⁵, J B Ferré⁴, O Catelinois¹, O Flusin³, E Tchernonog⁵, I E Moussion², A Wiegandt², A Septfonds⁶, A Mendy², M B Moyano², L Laporte², J Maurel², F Jourdain⁷, J Reynes⁵, M C Paty⁷, F Golliot¹

FIGURE

Timeline of symptoms onset for imported and autochthonous cases of chikungunya and epidemiological features, Montpellier, France, September–October 2014 (n = 13)

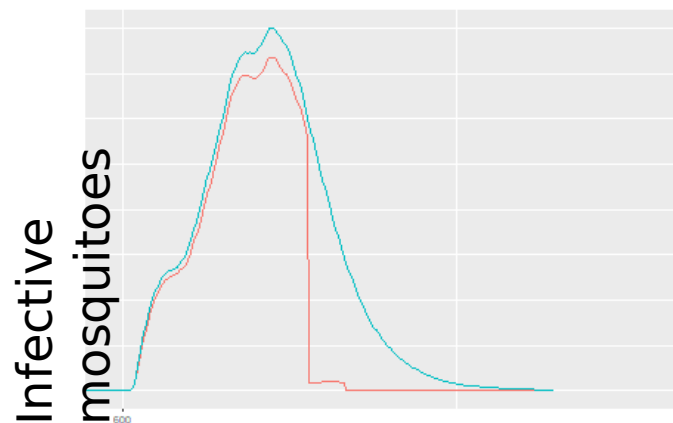


NRC: National Reference Center; RT-PCR: reverse transcriptase-PCR; UH: University Hospital.

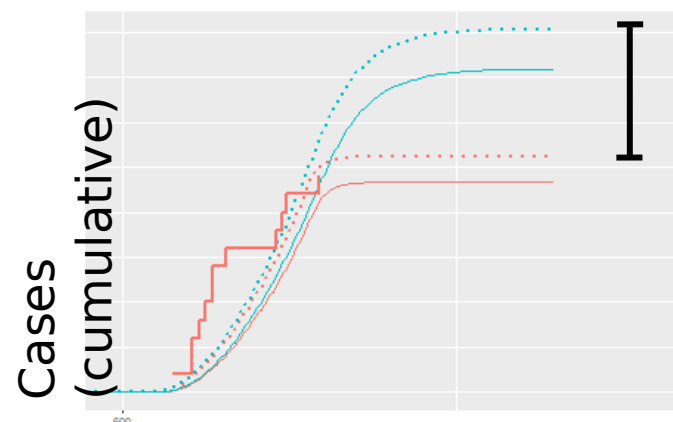
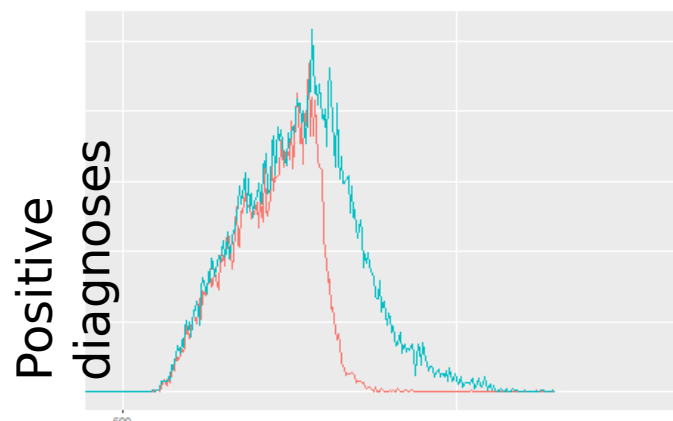
Cases numbered by order of identification.

Source : French Institute for Public Health Surveillance (Institut de veille sanitaire), 2014.

^a Possible period of viral transmission from infected vector (infected from the primary case):- mosquitoes biting the primary case between the first day and the last day of his viraemic period, extrinsic incubation period: seven days [8], mosquito lifespan: 10 days [9].



Red: intervention as done
Blue: non-intervention
counterfactual



cases prevented

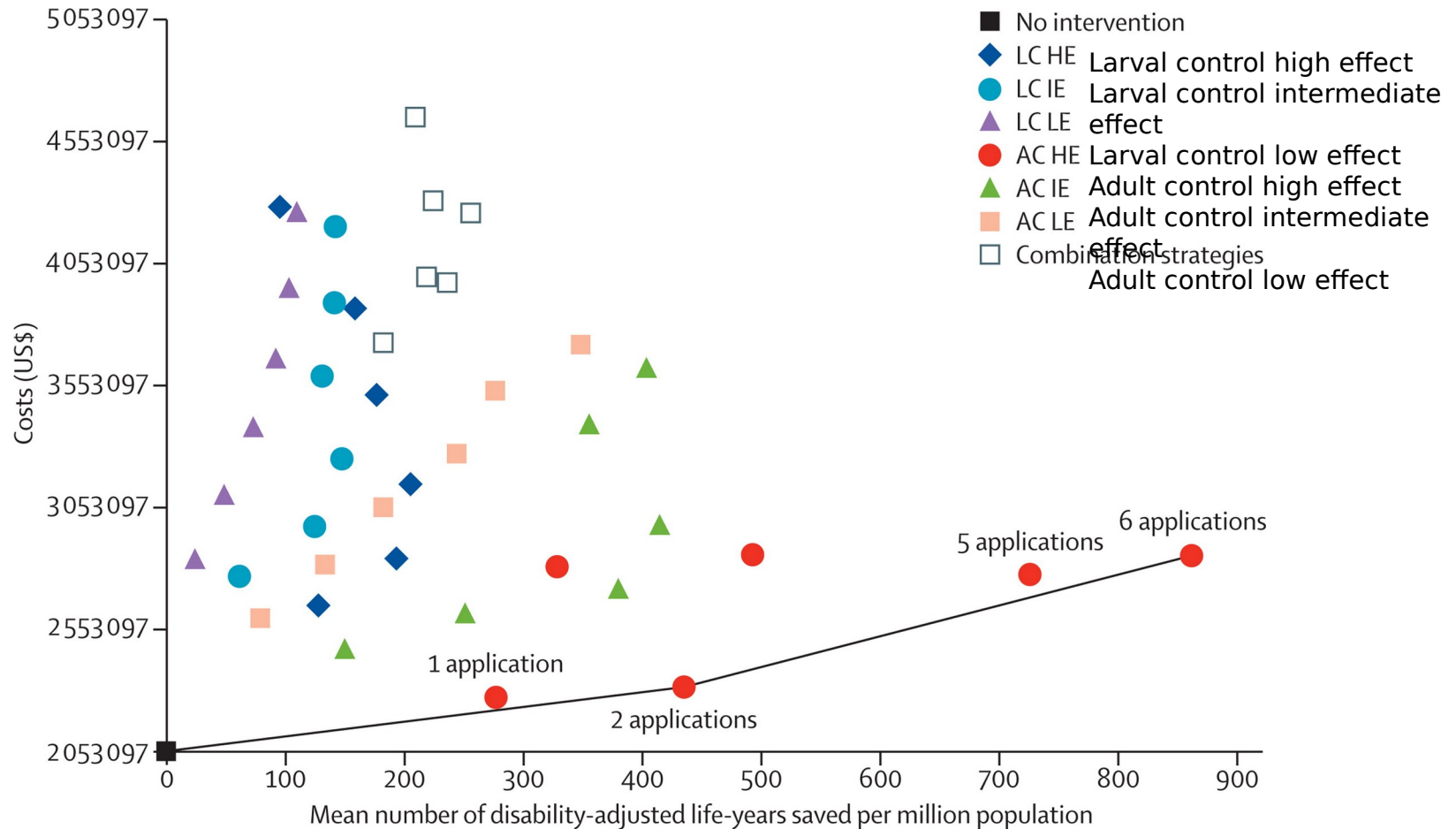
Thick red line: observed cases
Dotted lines: infections in humans
Solid lines: positive diagnoses

Cost-effectiveness

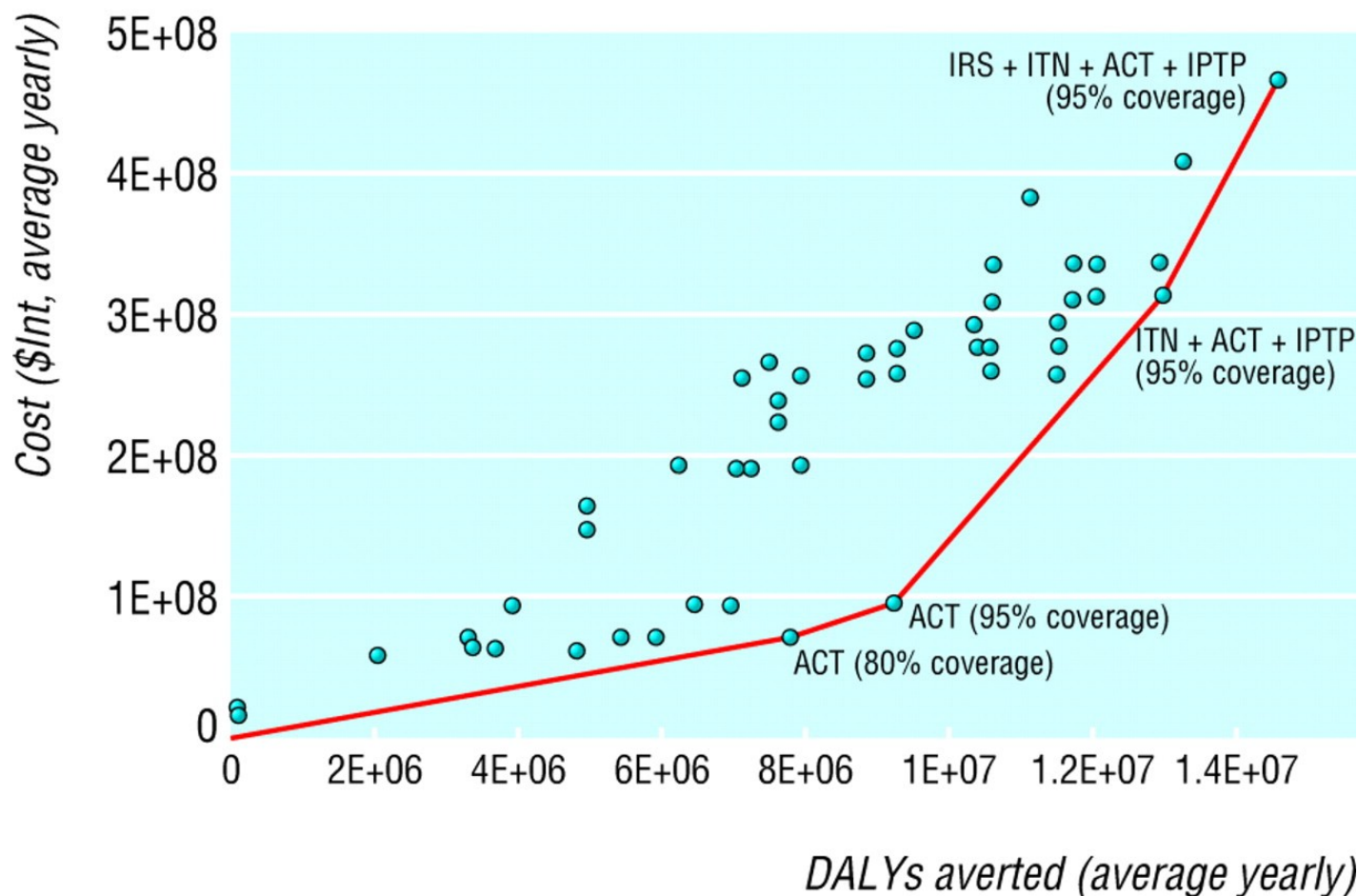


- What are the most effective and cost-effective (accepted) public-health strategies for mitigating the probability and impact of local transmission?
- Scenario-modelling combined with intervention cost-effectiveness data

Dengue vector control strategies in an urban setting: an economic modelling assessment



Cost-effectiveness plane showing 60 interventions against malaria analysed (20 interventions individually and combined at three assumed levels of coverage) and the expansion path, Afr-D



ACT = Artemisinin combination therapy

IRS = Indoor residual spraying

ITN = Insecticide treated nets

IPTP = Intermittent preventive treatment during pregnancy

<https://www.euro.who.int/en/publications/abstracts/manual-on-prevention-of-establishment-and-control-of-mosquitoes-of-public-health-importance-in-the-who-european-region-with-special-reference-to-invasive-mosquitoes-2018>

**Manual on prevention of establishment
and control of mosquitoes
of public health importance
in the WHO European Region
(with special reference to invasive mosquitoes)**

Willem Takken
Henk van den Berg



2019

“Recent systematic reviews of the **effectiveness** of vector control methods against *Ae. aegypti* and *Ae. albopictus* in the context of dengue control have concluded that there is a **paucity of reliable evidence**: there are few rigorous studies available on the impact of vector control on the vector population or on dengue incidence, and there is a need for standardized and comparative studies (Erlanger et al., 2008; Bowman et al., 2016). As a result, we do not have a clear understanding of which of the currently available interventions actually work, nor of the conditions under which they work.”

“Phase III studies should be designed around ***epidemiological endpoints*** to demonstrate the public health value of the intervention. ***Entomological outcomes cannot be used on their own for this purpose***, although they can be combined with epidemiological outcomes to evaluate a claimed entomological effect” (World Health Organization, [2017](#))

For Aedes-borne diseases in Europe, this is prohibitive: in order for epidemiological endpoints to provide significant results, this would require impossibly large and expensive studies, because of the low number of locally transmitted reported cases.

EXTERNAL SCIENTIFIC REPORT

VectorNet European Network
for Medical and
Veterinary Entomology

APPROVED: 23 June 2021

doi:10.2903/sp.efsa.2021.EN-6954

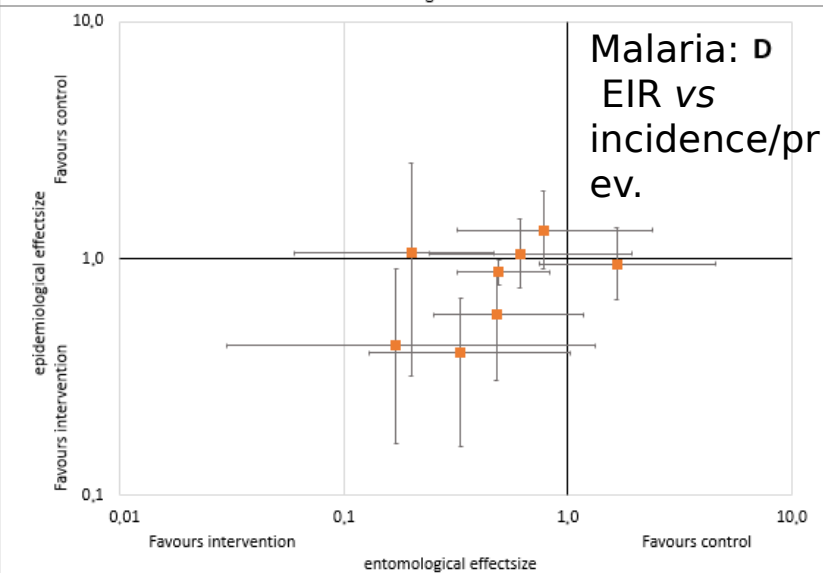
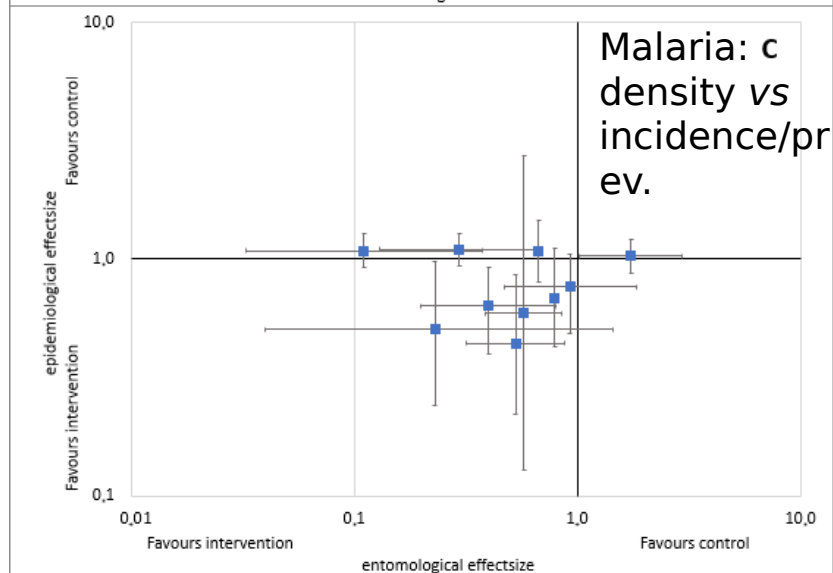
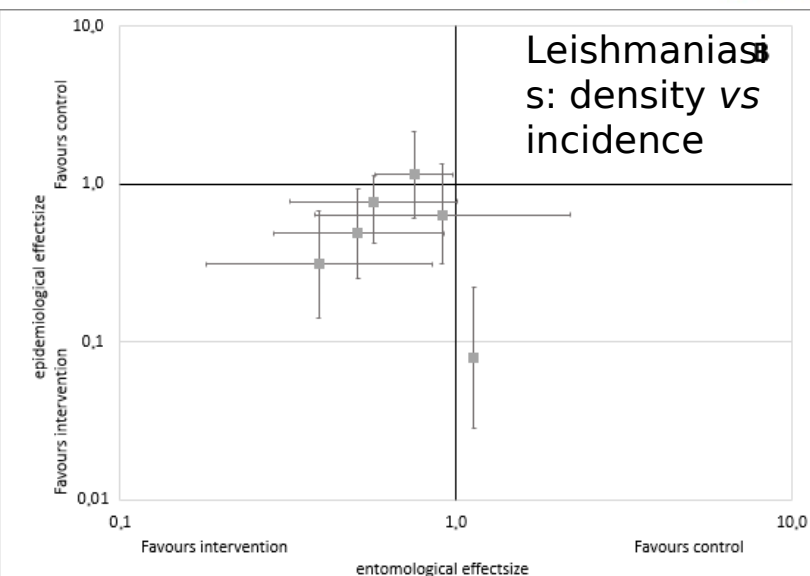
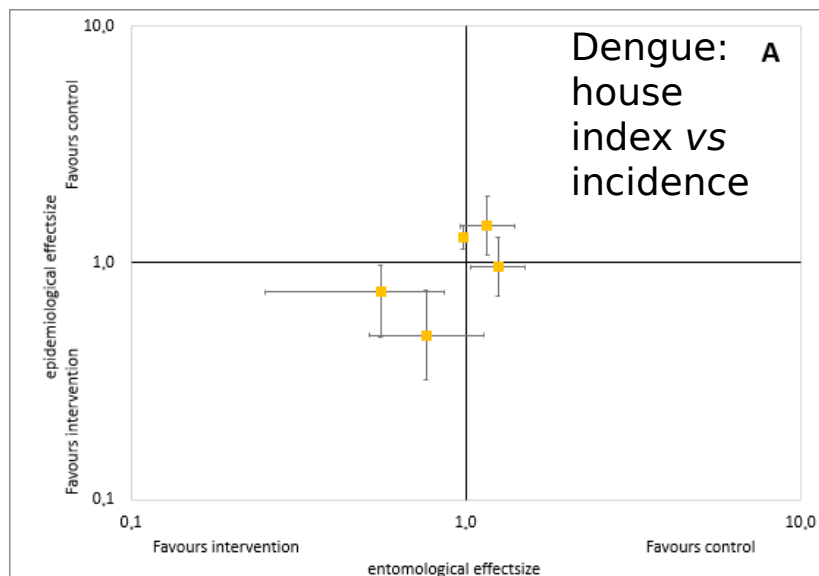
A systematic review to understand the value of entomological endpoints for assessing the efficacy of vector control interventions

Nick Van Hul¹, Marieta Braks², Wim Van Bortel¹

1. Institute of Tropical Medicine, Antwerp, Belgium

2. National Institute for Public Health and the Environment, Bilthoven, the Netherlands

Effectiveness: Entomological vs epidemiological endpoints



Cost-effectiveness analysis

- Is complex (and therefore error-prone), often requires modelling, and many assumptions.
- Needs to be specifically tailored to the setting and options under consideration
- Cost-effectiveness is not the only criterion for selection. Examples of other criteria:
 - Equity and other ethical considerations
 - Environmental considerations
 - Feasibility

Effectiveness of larviciding (diflubenzuron) for *Ae. albopictus* in Chioggia, Italy

An environment where catch basins are an important breeding site for *Ae. albopictus*



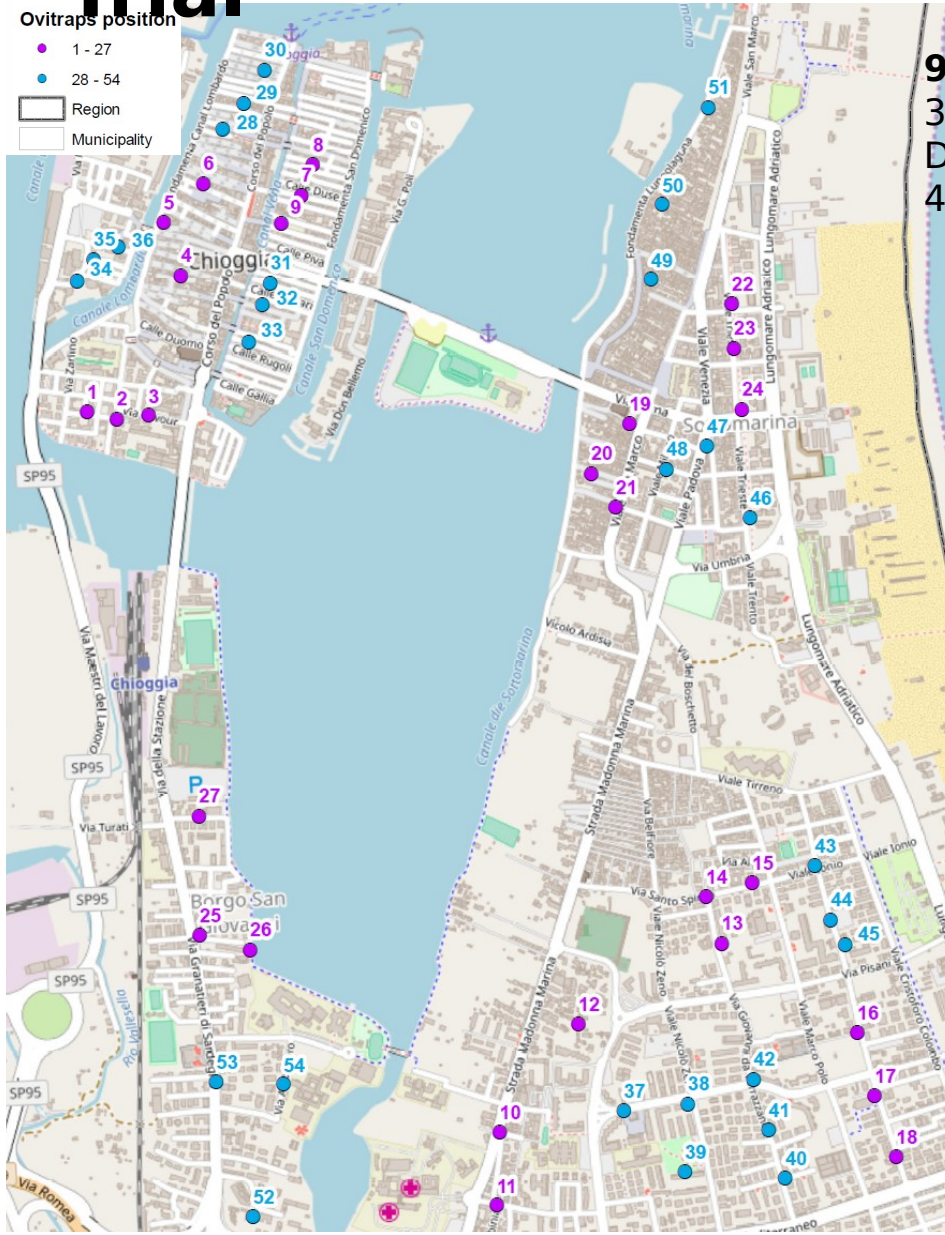
Chioggia: Entomological Pair-wise Cluster Randomised Controlled Trial

Ovitrap position

- 1 - 27
- 28 - 54

Region

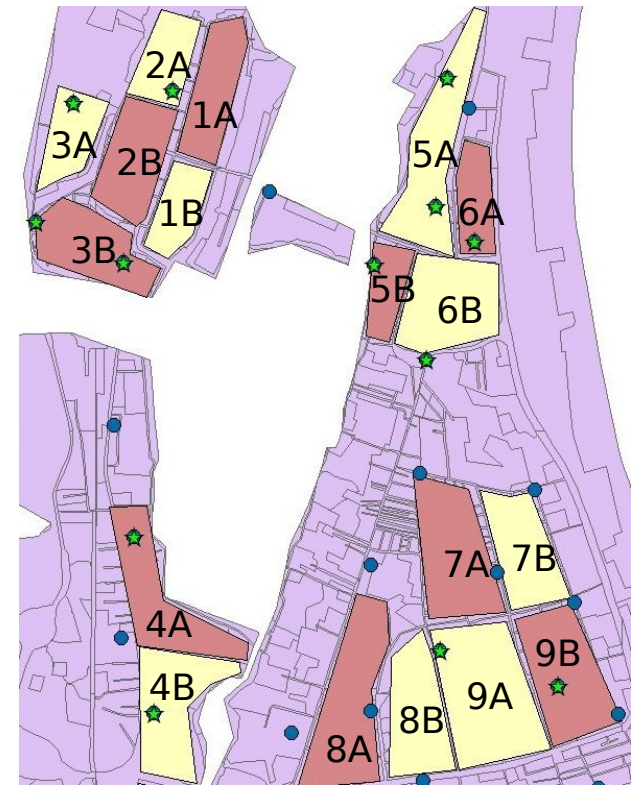
Municipality



9 clusters: 9 treated areas, 9 untreated (control) area
3 ovitraps in each

Duration: 6 weeks (07/08 to 16/10).

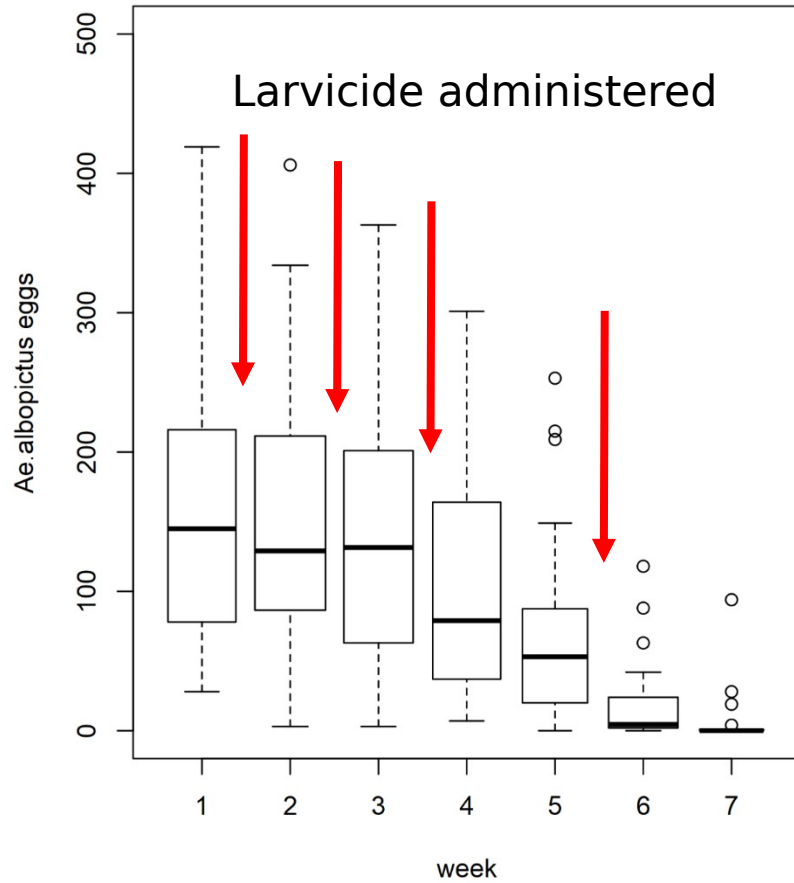
4 Treatments: 08/08, 28/08, 13/09, 15/10



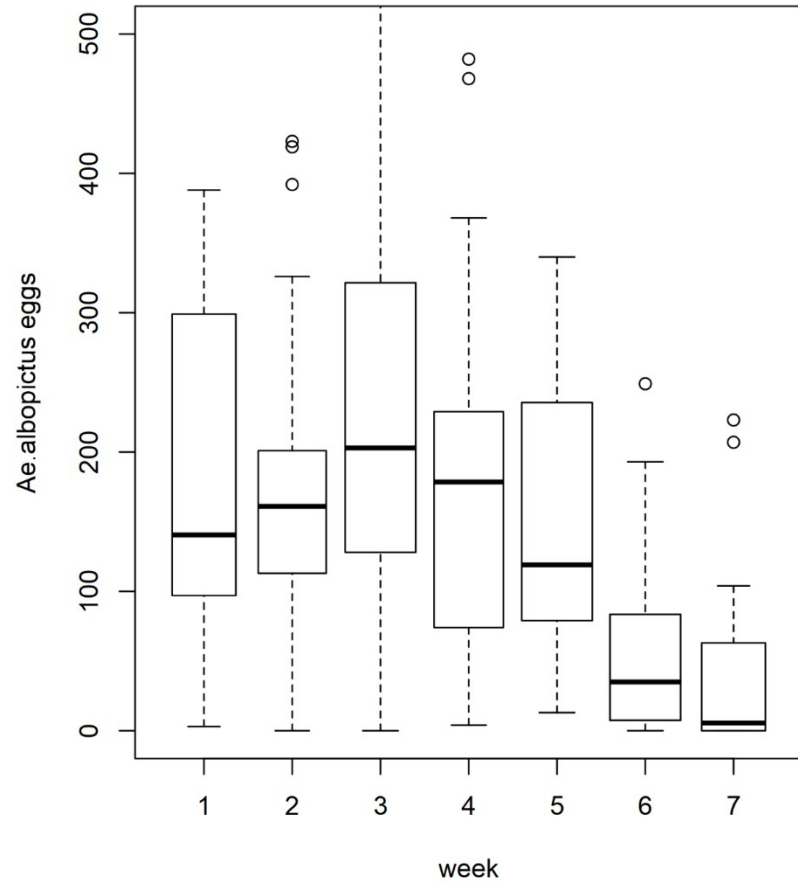
Light brown: treated / Yellow: untreated

Ovitrap temporal results

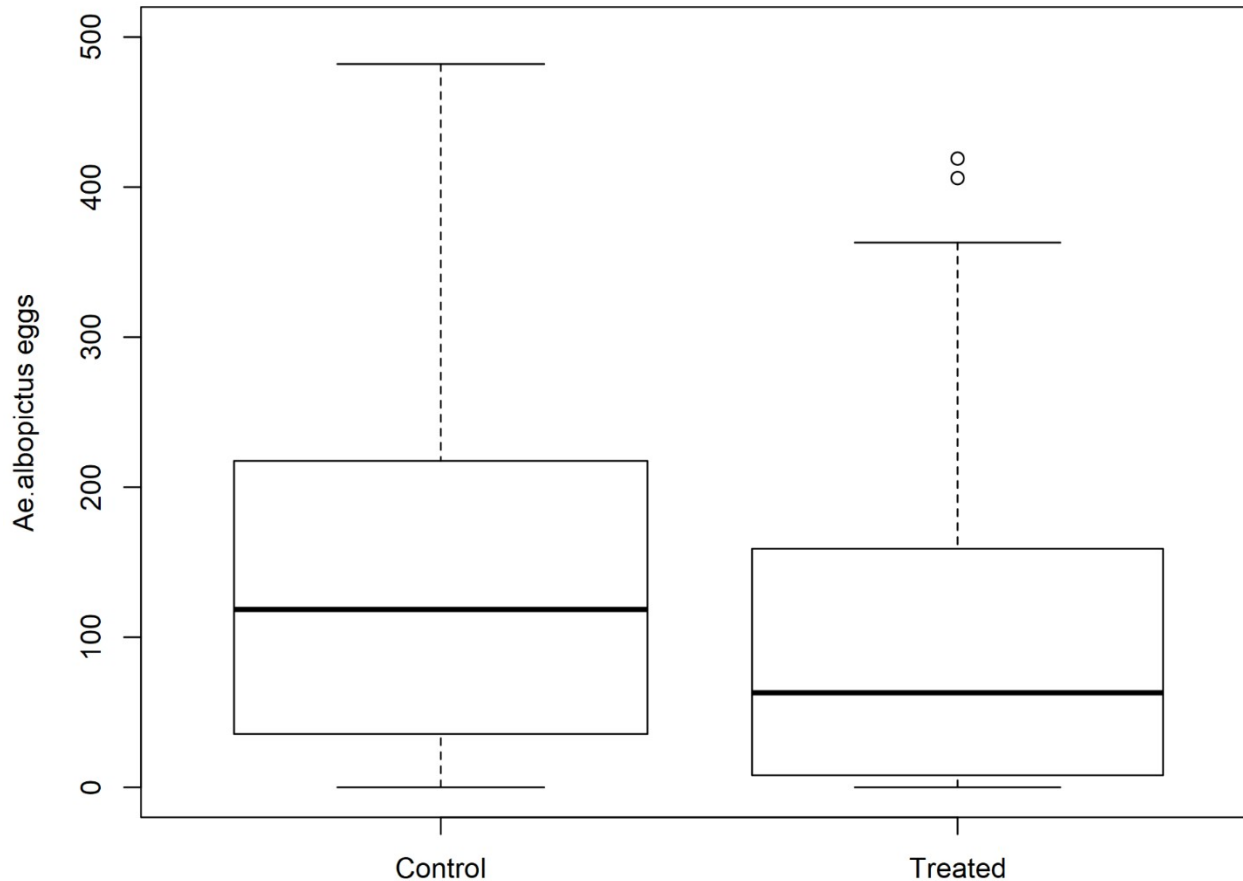
Treated



Control



Ovitrap regression results

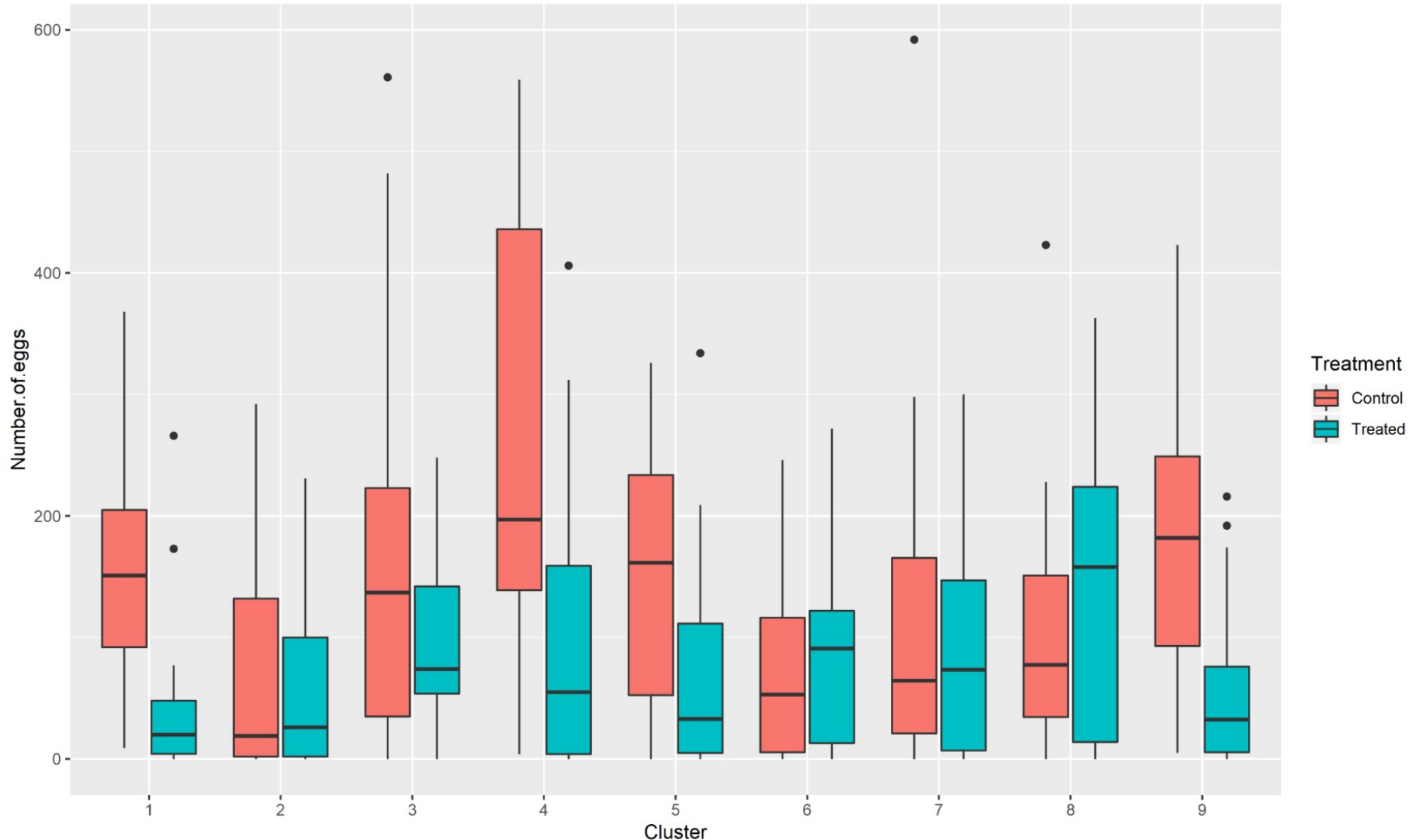


Negative binomial zero-inflated generalized linear mixed model with Cluster/Trap and Week as random effects

Incidence Rate Ratio = 0.46

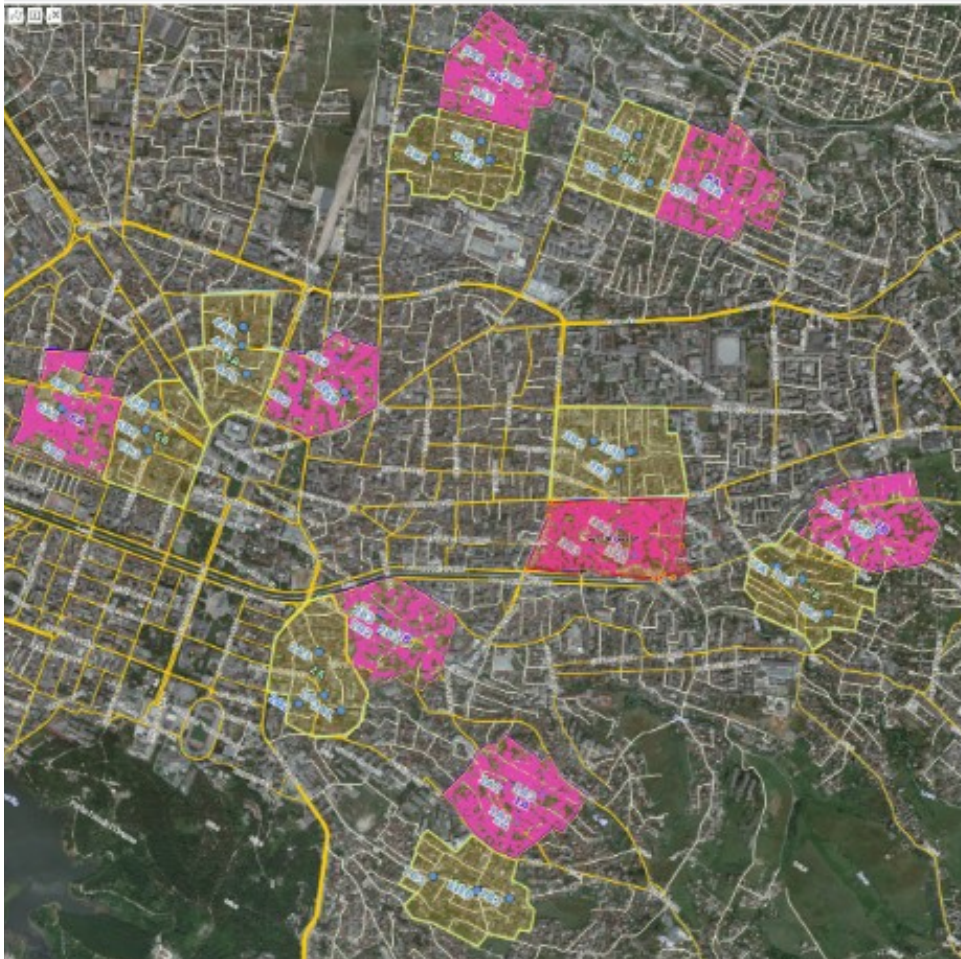
p=0.00003

Differences in effect among cluster-pairs after first treatment: large variation



Entomological Pair-wise Cluster Randomised Controlled Trial in Tirana, Albania

Design



Eight cluster pairs

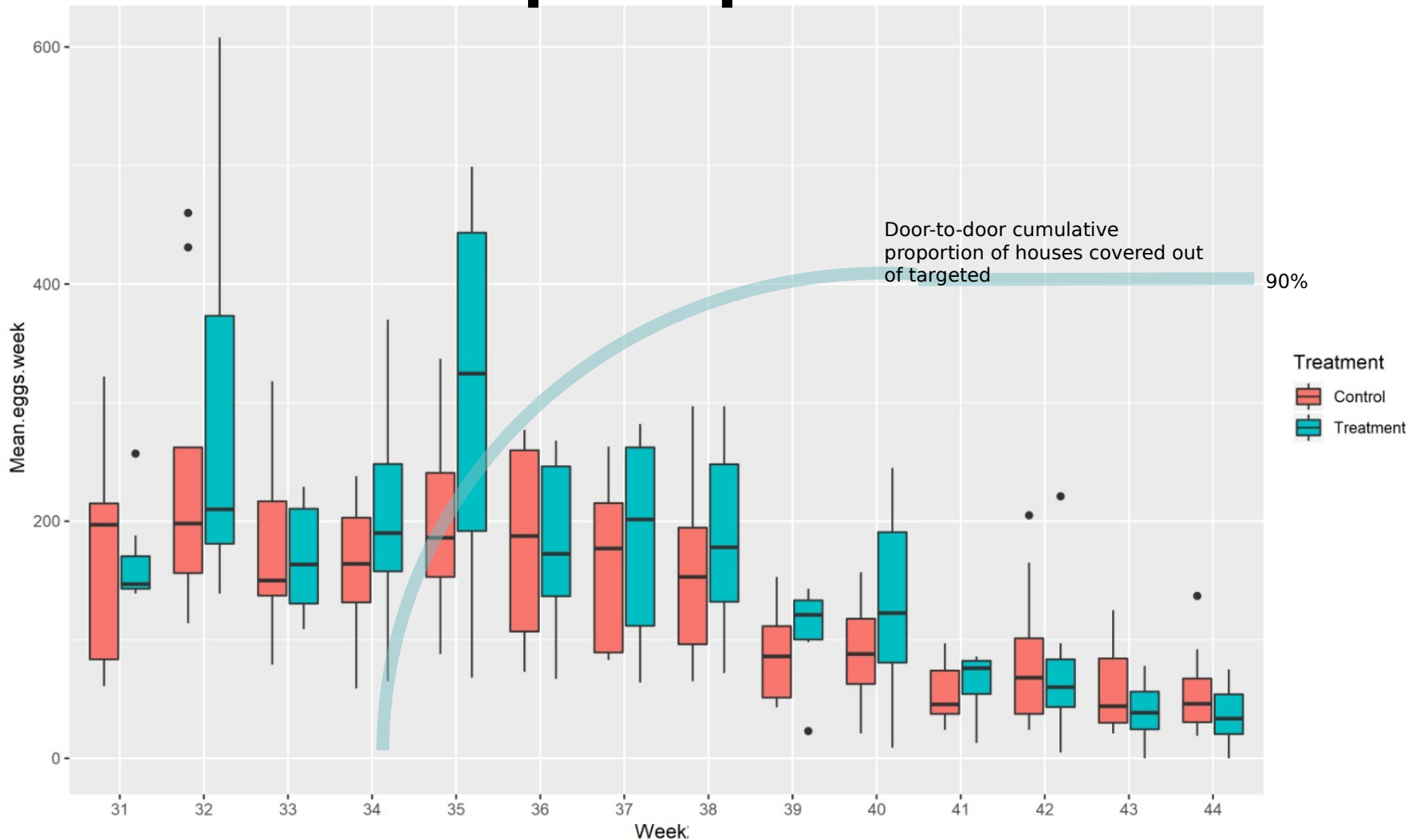
- 8 door-to-door/IVM areas
- 8 untreated (control) areas

Monitoring:

3 ovitraps and 1 BG-Sentinel trap in each cluster

Duration: 13 weeks monitoring

Ovitrap temporal results



Negative binomial Zero Inflated GLMM with random effects for Cluster/trap and Week

1 week lag: IRR=0.75, 95%CI 0.56–0.99

2 week lag: IRR=0.73, 95%CI 0.54–0.98

Lag between the interventions and the ovitrap sampling, related to the biological larval development time. IRR= incidence rate ratio

Conclusions

- Modelling plays an important role in risk assessment and planning of mitigation strategies
- There is insufficient evidence for **existing** entomological interventions
- Field studies with entomological outcomes specific to European setting could still be very useful as data input for models to aid strategic planning of vector control

Thank you for your attention

and:

Colleagues from ECDC, EFSA, VectorNet, ArboRisk and AedesRisk projects