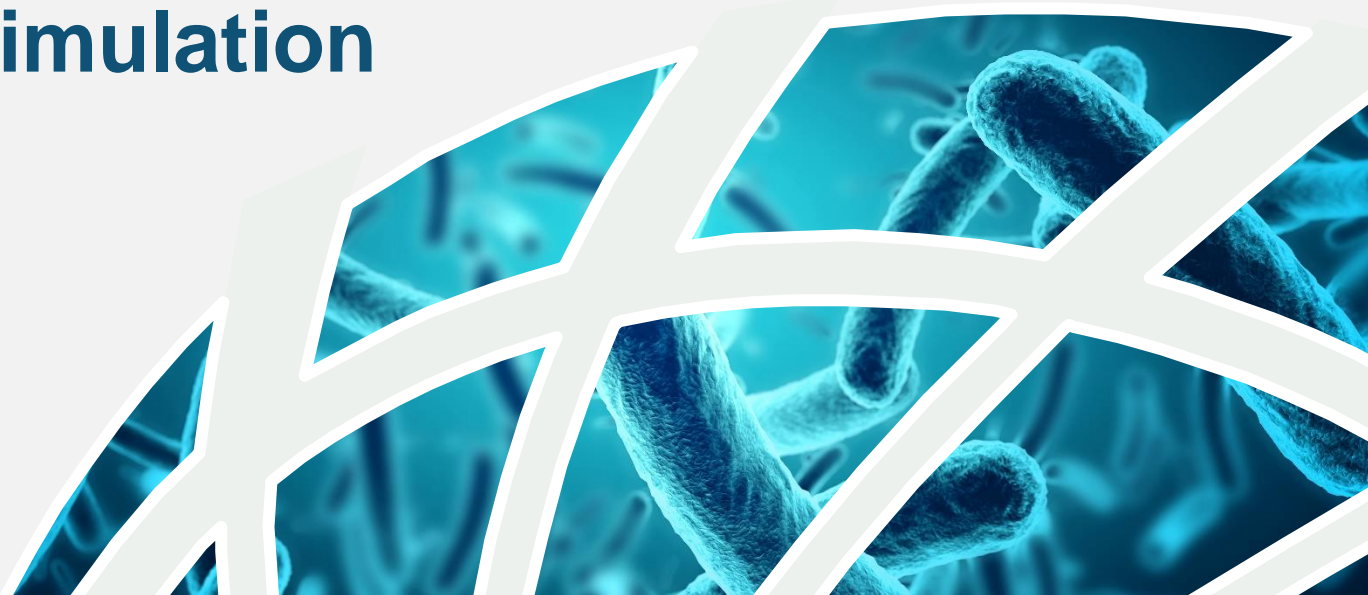


Water quality modelling - Part 2

Pathogen simulation

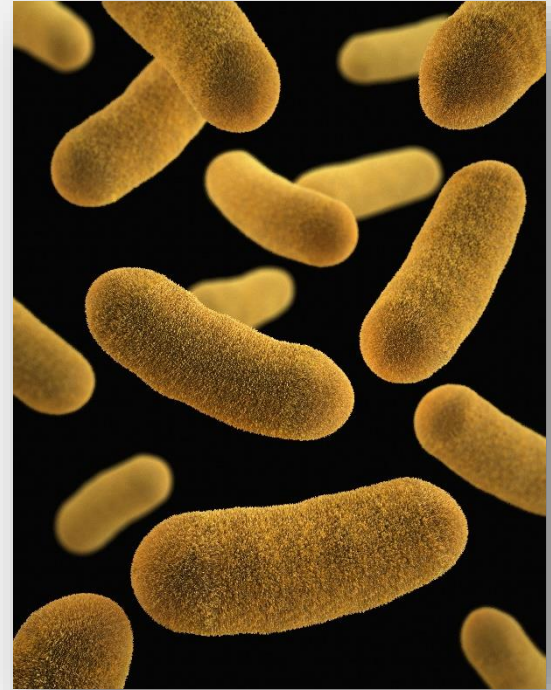
Dr Michael Barry
Emma McCall



Overview

Today we'll cover

- What are pathogens?
- Why are pathogens of interest?
- How do we model pathogens?
- Example model



What are pathogens?

Pathogens are

- Microbes (not all microbes are pathogens)
 - Viruses, bacteria, single and multi-cell eukaryotes
 - Beneficial and harmful
- Everywhere on Earth!
 - Estimated to be 10^{31} organisms
 - Ten billion times the number of stars in the universe
 - Each human carries 30 trillion



What are pathogens?

Pathogens are

- Some familiar examples
 - Measles, smallpox, zika, Ebola, SARS, COVID
- Some common aquatic environment examples
 - Cryptosporidium or “crypto” (not the currency!)
 - Escherichia coli or “E. coli”
 - Norovirus
 - Campylobacter



Why are pathogens of interest?

Pathogens are of interest because they

- Can cause human disease
- Can be fatal – up to 20% of global deaths in 2019
- Human and economic cost
 - US\$1bn annually

Our concern today

- Environmental waterborne pathogens



Why are pathogens of interest?

Common environmental sources

- Human waste
 - Sewers, combined sewers (CSOs) and onsite treatment systems
- Mammalian waste (agricultural and urban)

Common environmental receivers

- Freshwater creeks, streams, rivers and lakes
- Reservoirs (including water supply), natural and urban lakes
- Coastal oceans



Why are pathogens of interest?

Environmental pathways to humans

- Ingestion of contaminated drinking water
 - Water supply contamination
- “Primary contact” with contaminated water
 - e.g. swimming, bathing, diving, water skiing
- “Secondary contact” with contaminated water
 - e.g. boating, fishing, canoeing
- Consumption of contaminated food

Québec

Home < Health < Health issues < A to Z < Health problems associated with bathing waters

Health problems associated with bathing waters

GOV.UK

Home > Health and social care > Public health > Health protection > Chemical and environmental hazards > Swim healthy

Environment Agency

Public Health England

Guidance

Swim healthy

Updated 24 June 2019

COVID-19 IDSA Academy MyIDSA Journals News Science Speaks Multimedia

Guidelines Public Health Clinical Practice Research Professional

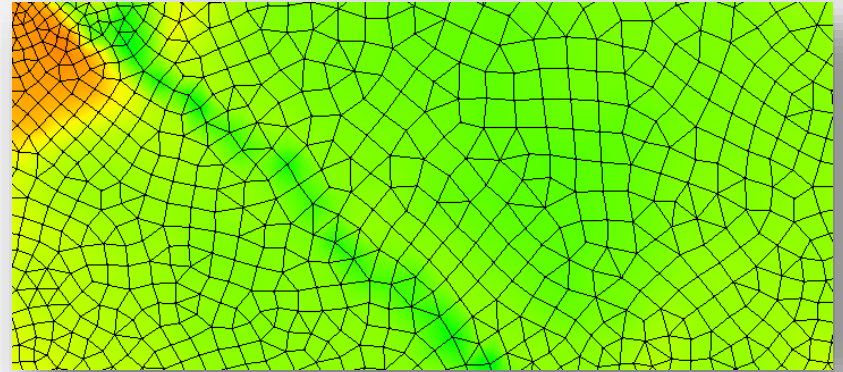
News Releases and Statements > CDC Alerts > Outbreak of E. coli Infections Linked to Romaine Lettuce

Outbreak of E. coli Infections Linked to Romaine Lettuce

Why are pathogens of interest?

Aquatic environmental

- Look to numerical modelling
 - Risk elimination / minimisation
 - Public health
- QMRAs
 - Quantitative microbial risk assessment



World Health Organization

Home Health Topics Countries Newsroom Emergencies Data

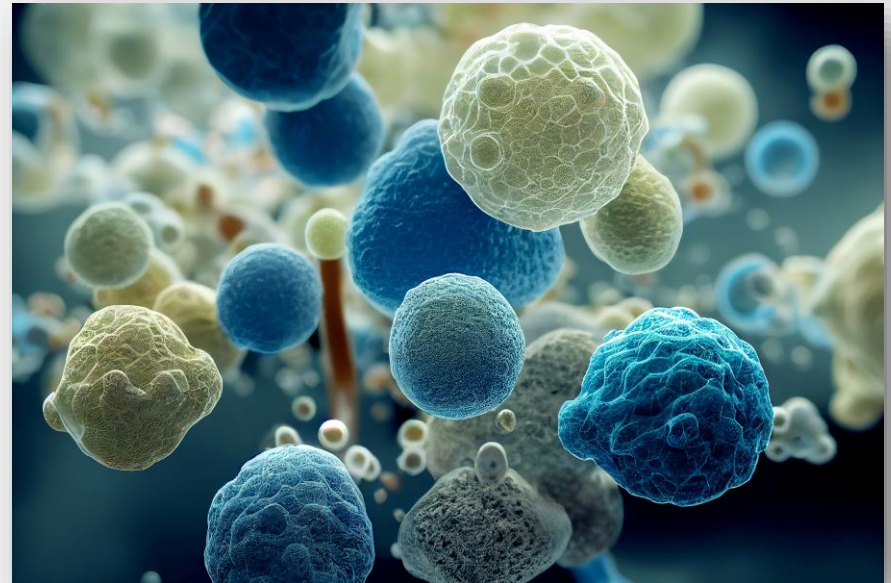
Home / Publications / Overview / Quantitative microbial risk assessment: Application for water safety management

Quantitative microbial risk assessment: Application for water safety management

How do we model pathogens?

On waterborne release, pathogens transform via

- Natural mortality
- Light inactivation
- Settling
- Growth
- Predation
- Attachment / detachment to sediment

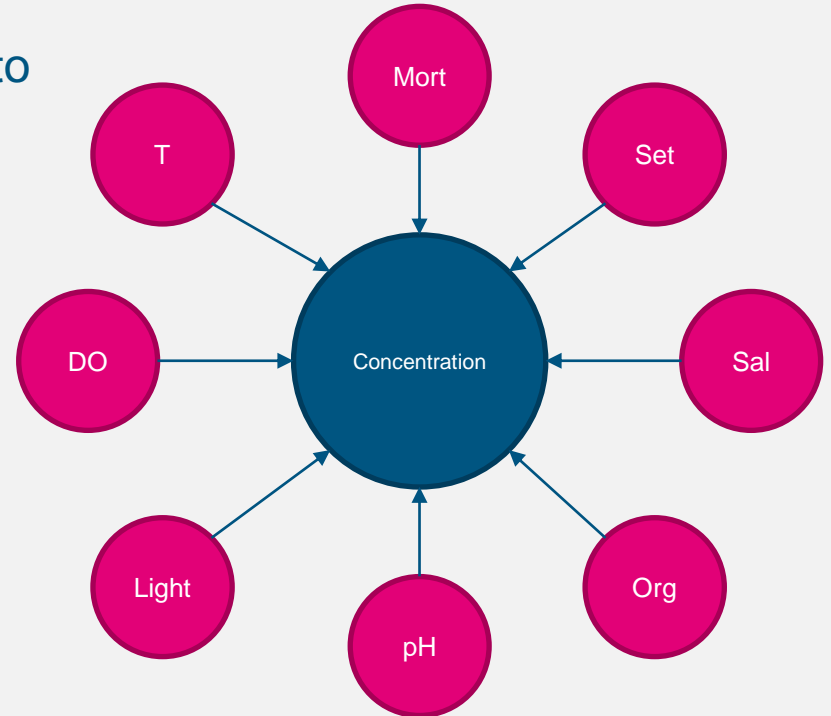


How do we model pathogens?

On waterborne release, pathogens respond to

- Temperature
- Light (visible, UV-A, UV-B, infrared)
- Salinity
- Oxygen
- pH
- Organic material

Not pretending this is straightforward!



How do we model pathogens?

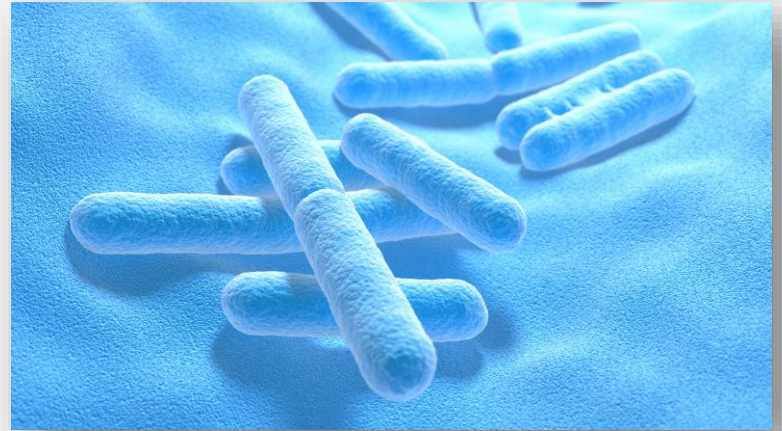
Inputs - concentrations

- Initial conditions
- Boundary inflows
 - Diffuse sources (e.g. catchment inflows)
 - Point sources (e.g. sewage outfalls)

Inputs - parameters

- Many!

```
! Pathogens
pathogen model == attached, crypto
> alive min max == 0.0, 1e7
> mortality == 0.048, 0.0, 6.1, 1.0, 1.14
> visible inactivation == 0.0, 0.0667, 0.1
> uva inactivation == 0.01, 0.00667, 0.1
> uvb inactivation == 0.02, 0.00667, 0.1
> settling == -0.01, -0.2
> target attached fraction == 0.5
end pathogen model
```



How do we model pathogens?

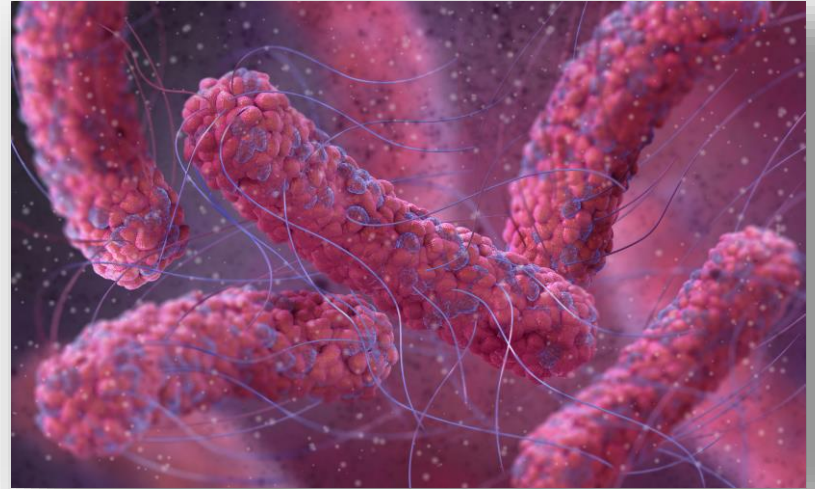
Outputs - concentrations

- Transformation under environmental modification
- Concentrations are the result of multiple processes

Outputs – diagnostics (fluxes)

- Which processes dominate? Why?
- Ultimately seek understanding, not just concentrations

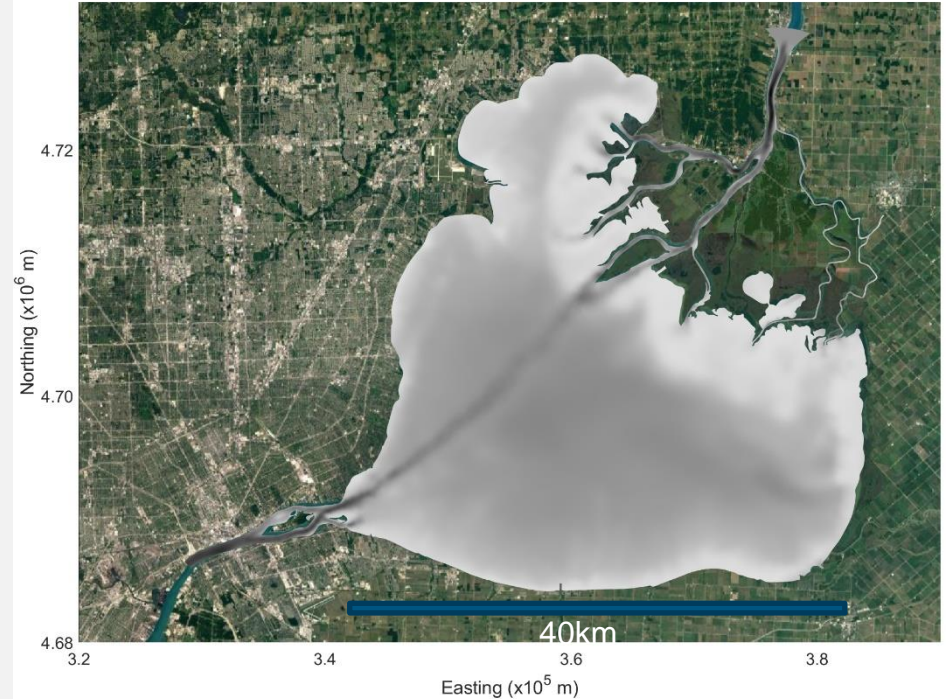
Lets look at an example



Lake St Clair

Large North American lake

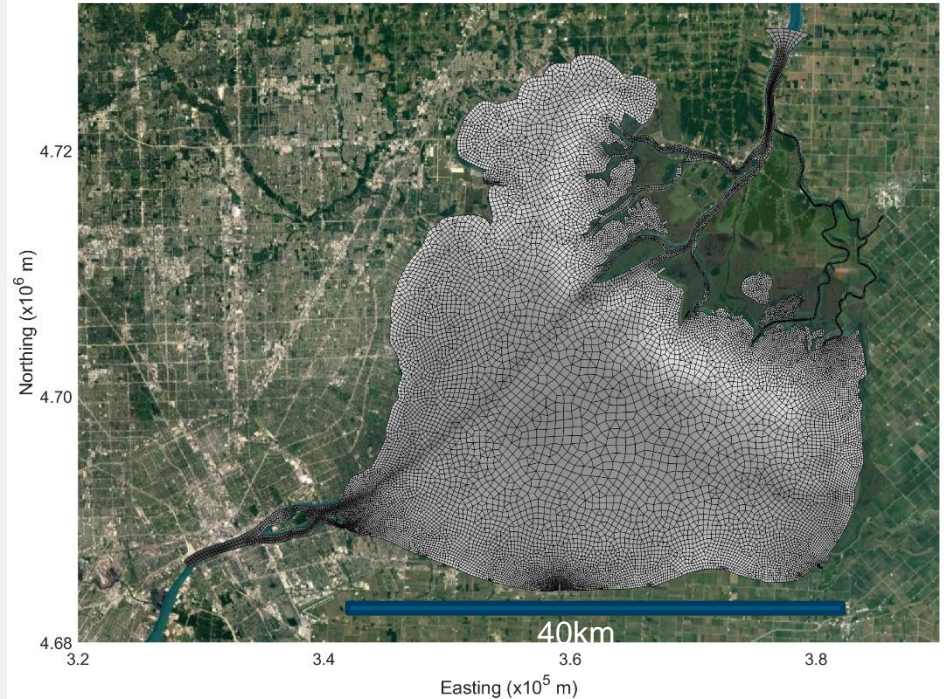
- ~40 kilometres at widest
- ~10 metres at deepest
- ~1,200 km² area
- Bathymetry is grey
 - Darker is deeper
 - Lighter is shallower



Lake St Clair

Model

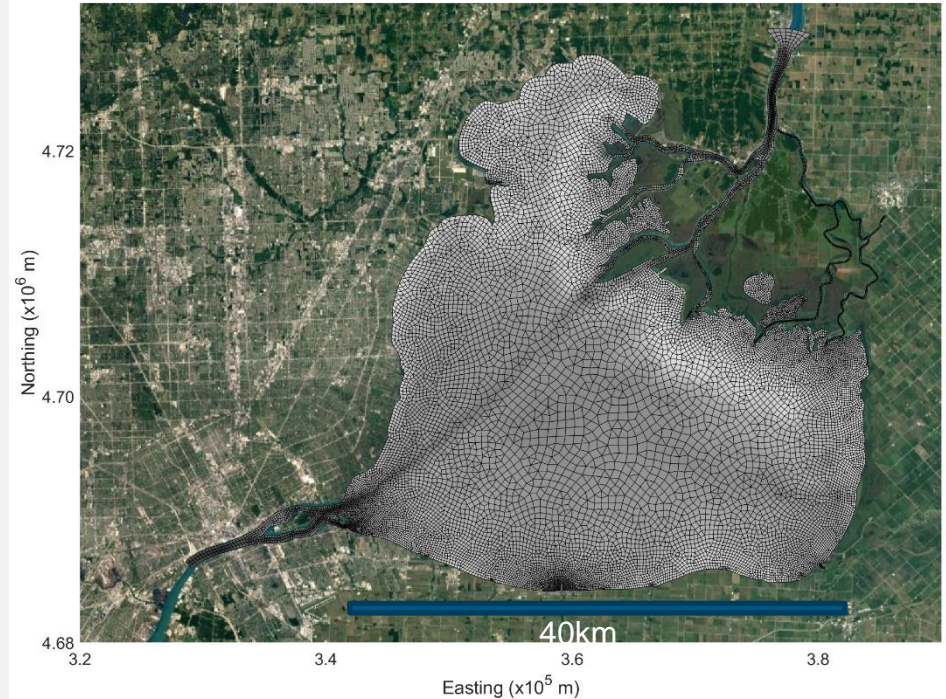
- TUFLOW FV
 - Hydrodynamics
 - Advection dispersion
 - 3D
 - Water quality



Lake St Clair

Model

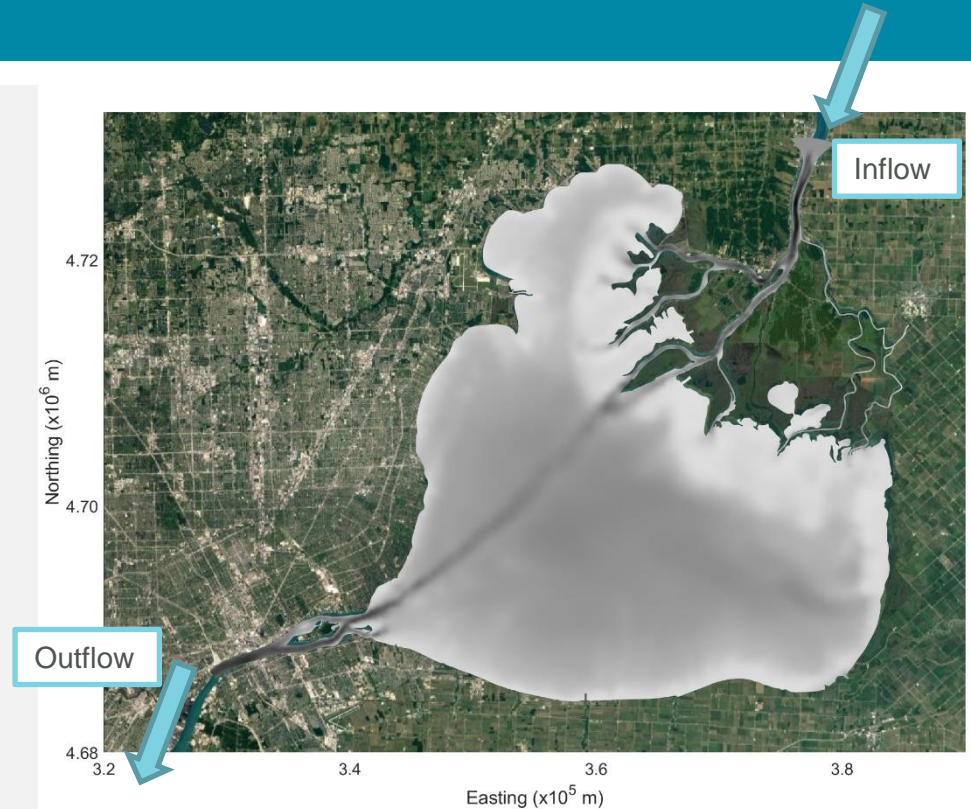
- ~21,000 2D cells
- Developed by University of Windsor, Ontario
- Used with the kind permission of Dr Mohammad Madani



Lake St Clair

Environmental setting

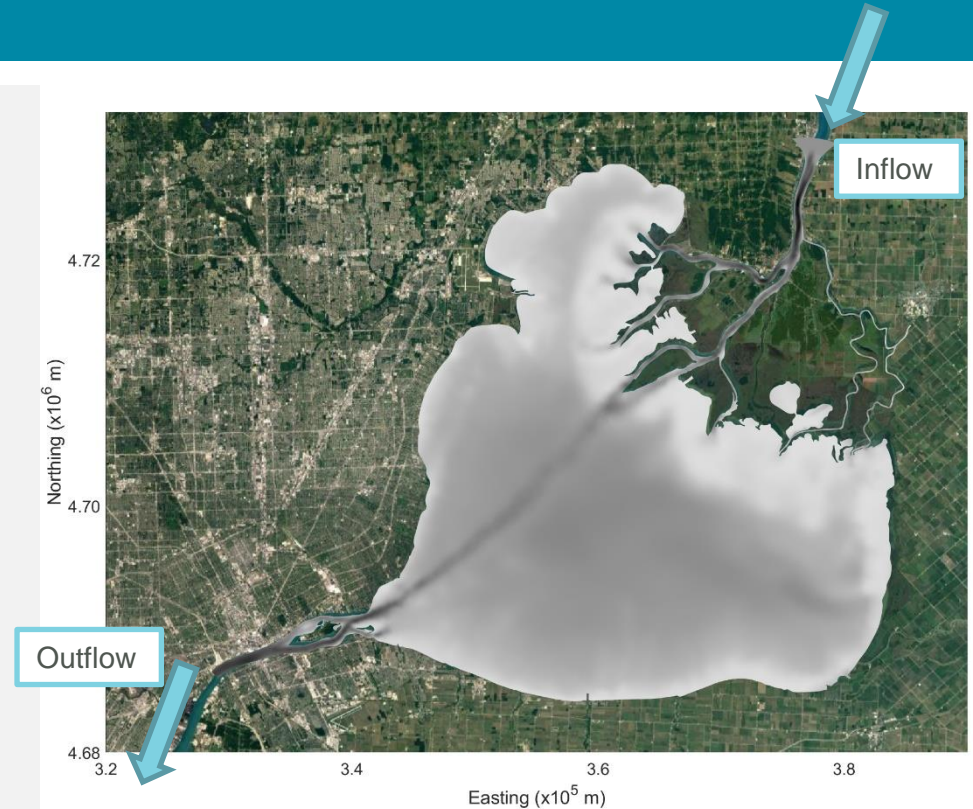
- Receives pathogens from
 - Upstream inflows
 - Fringing catchments
- Sees some recreational use
 - Boating
 - Fishing
 - Water sports



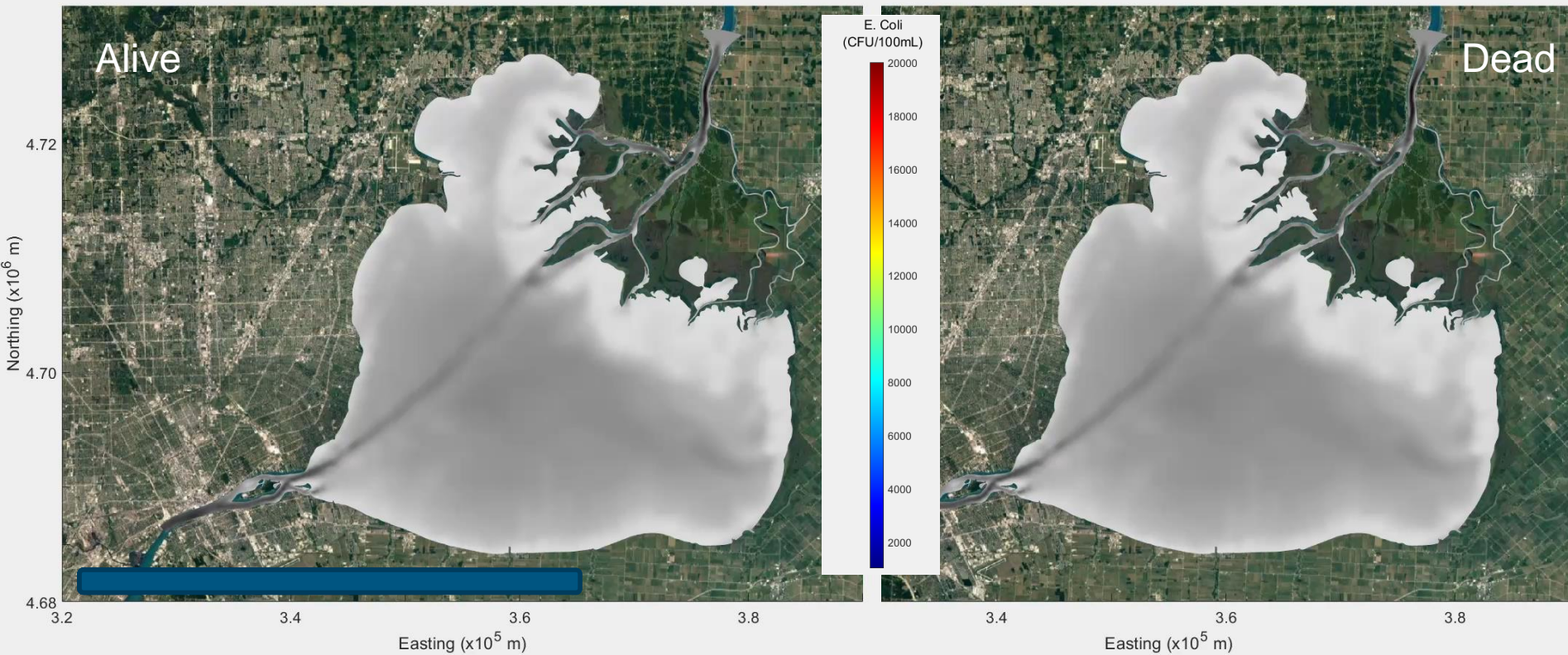
Lake St Clair

Simulation

- ~1 month
- Examine pathogen concentrations
 - Surface alive (active)
 - Surface dead (inactive)
- Entirely hypothetical
 - Not a real case - example only
- Included light and natural mortality



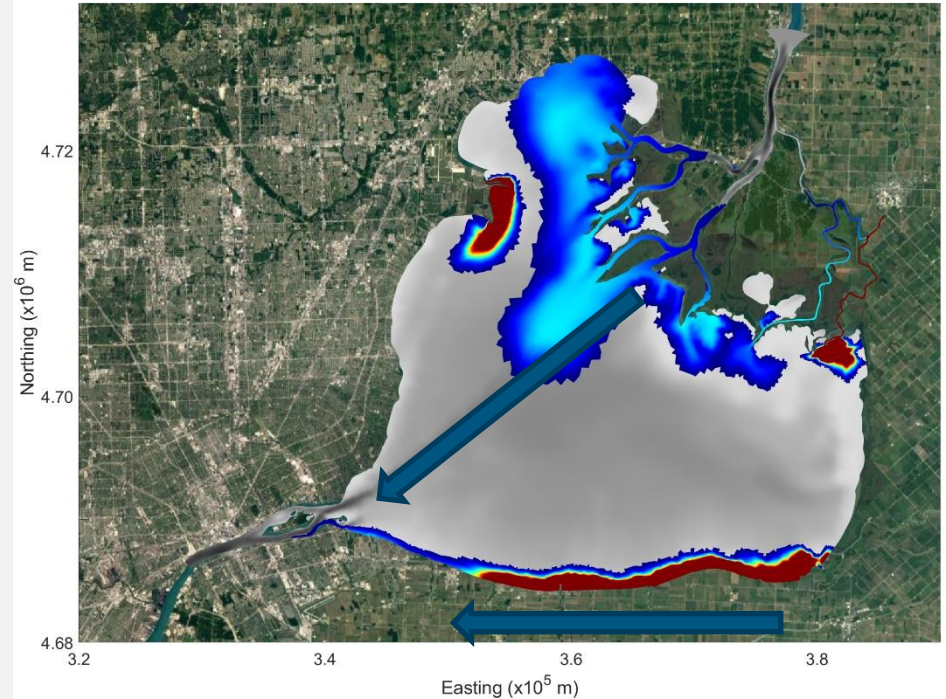
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Qualitatively

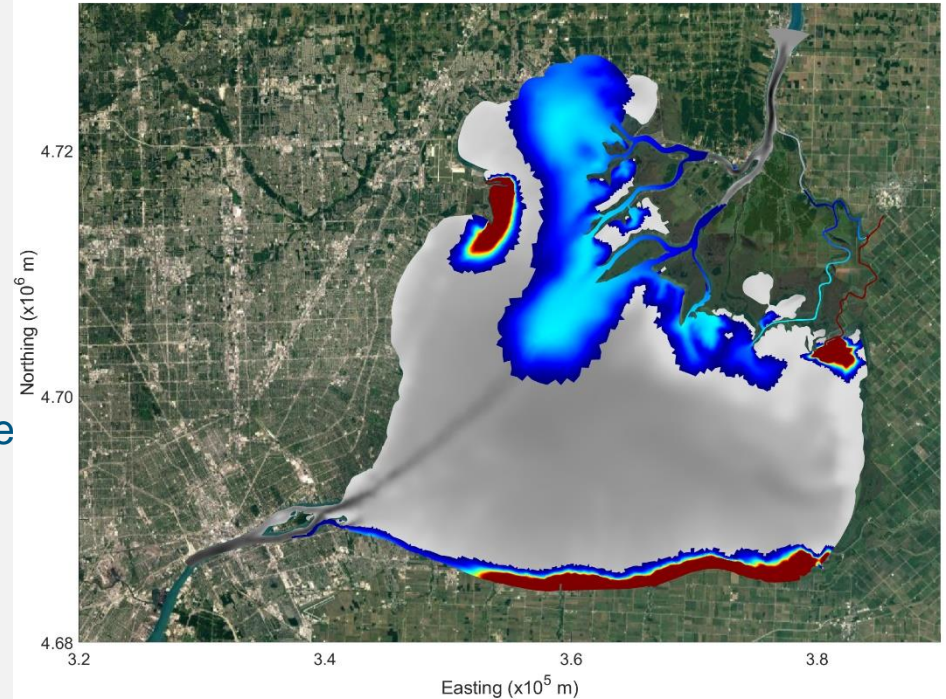
- Alive pathogens do not persist
- Dead pathogens accumulate
- Long shore east to west transport
- But what process dominates
 - Natural mortality?
 - Light deactivation?
 - ...?



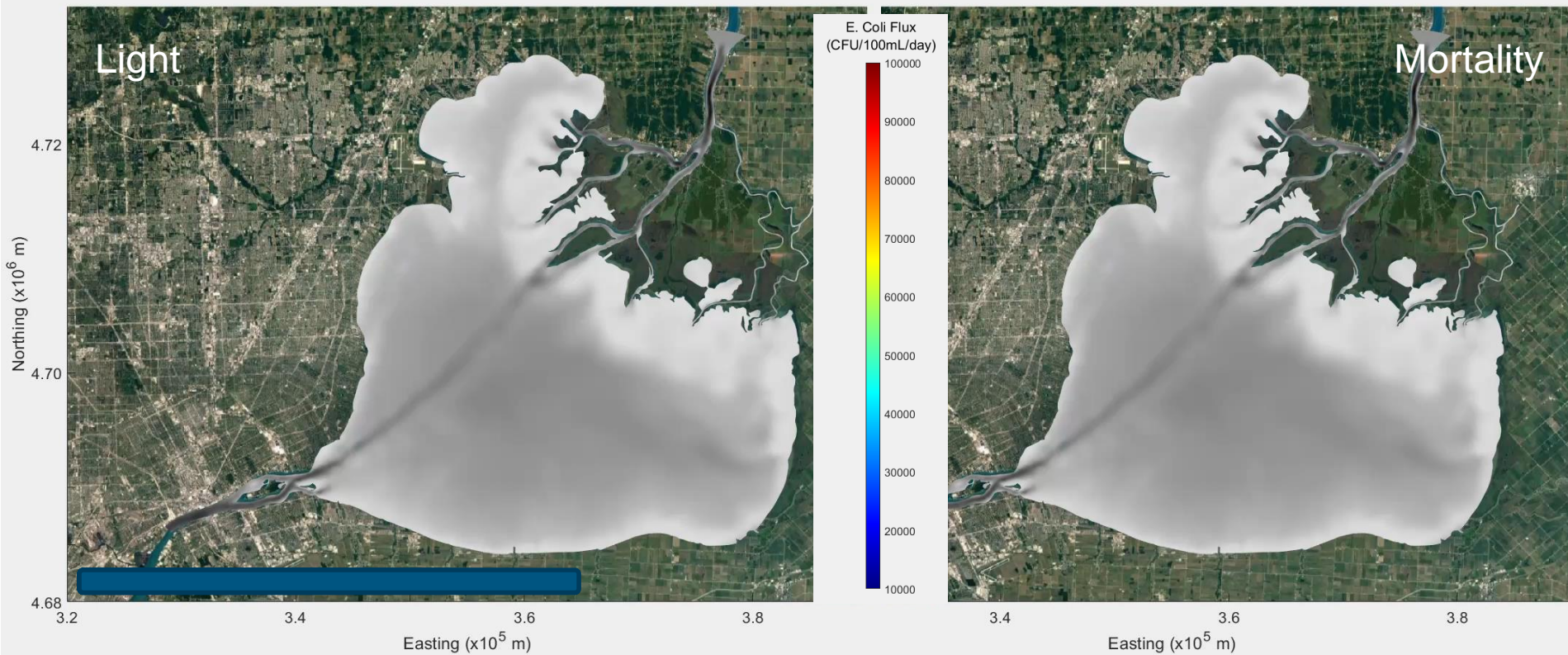
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Diagnostics (fluxes)

- Same ~1 month
- Examine fluxes
 - Light deactivation and natural mortality
- Surface contours of fluxes
CFU / 100 mL /day on same colour scale
 - A positive flux is from alive to dead
- Not presenting concentrations



Lake St Clair



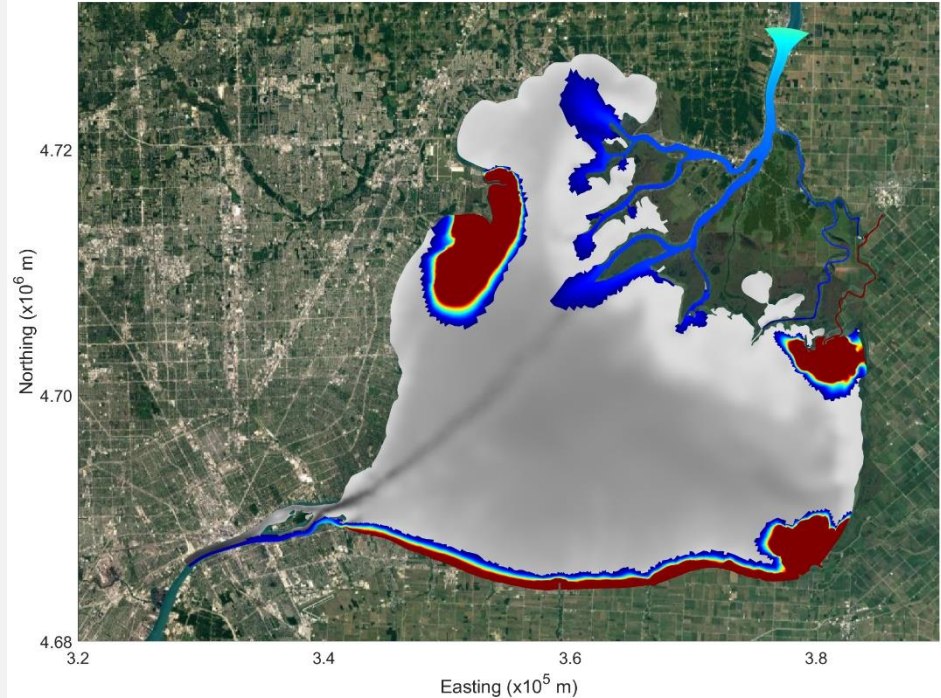
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Qualitatively

- Light inactivation fluxes dominate
 - Obviously diurnal (night and day)
- Natural mortality less so

Quantitatively

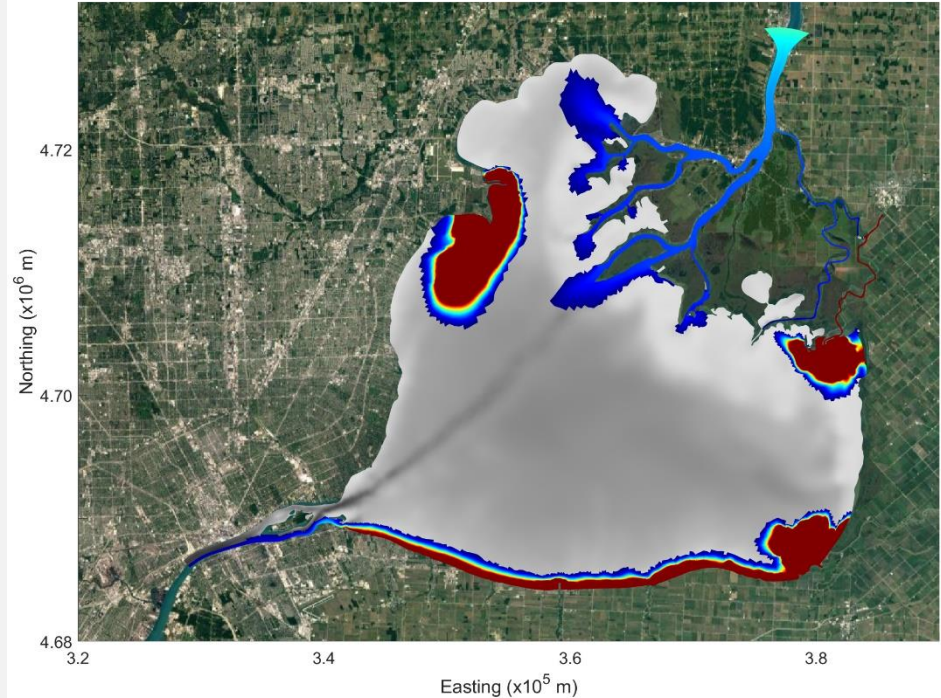
- Light inactivation orders of magnitude greater than natural mortality
- Dominant flux pathway
 - Check and support parameterisation



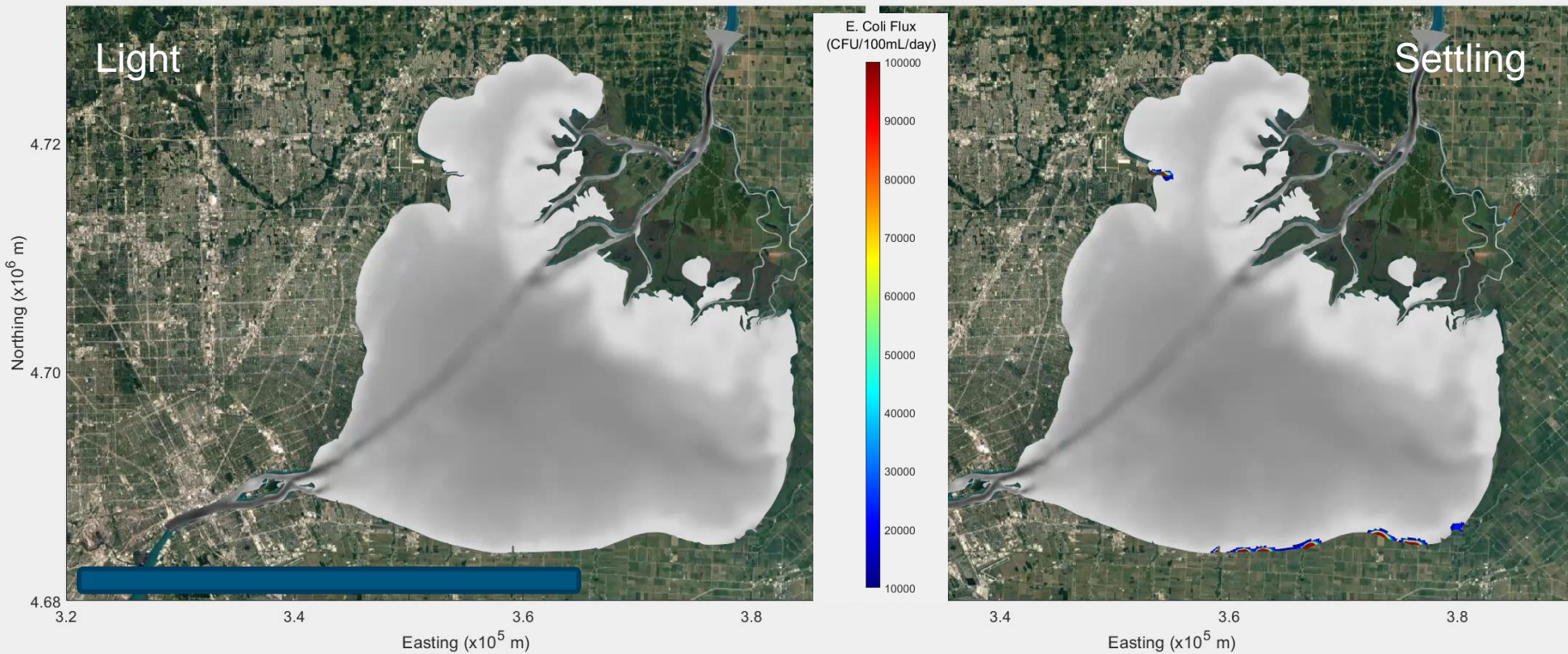
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What about settling?

- Set typical settling rate
 - Alive and dead pathogens
- No resuspension
- Compare with light inactivation
- Positive flux is from water column to bed



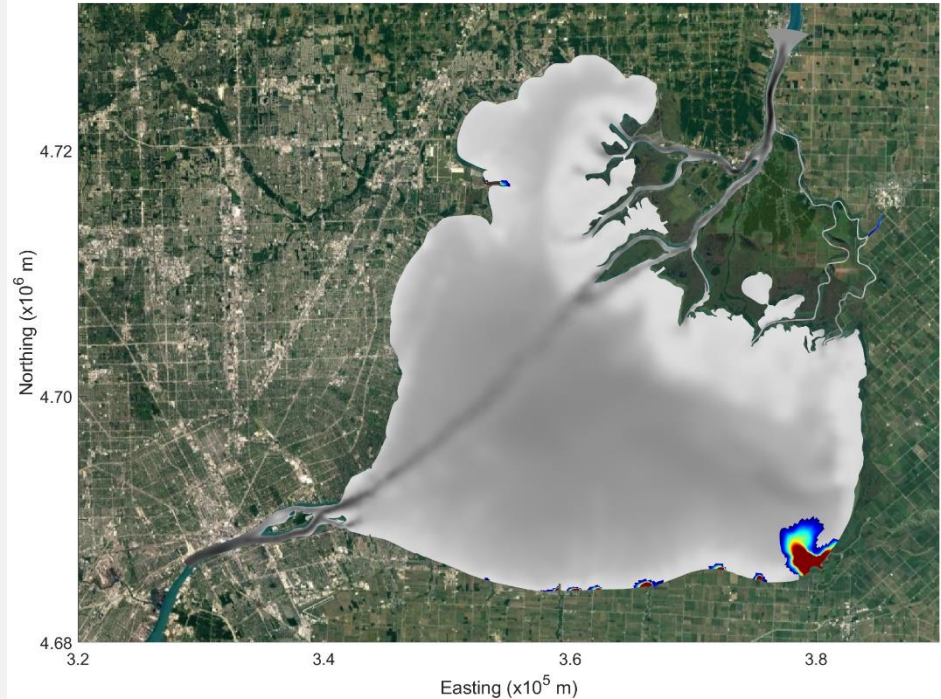
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What about settling?

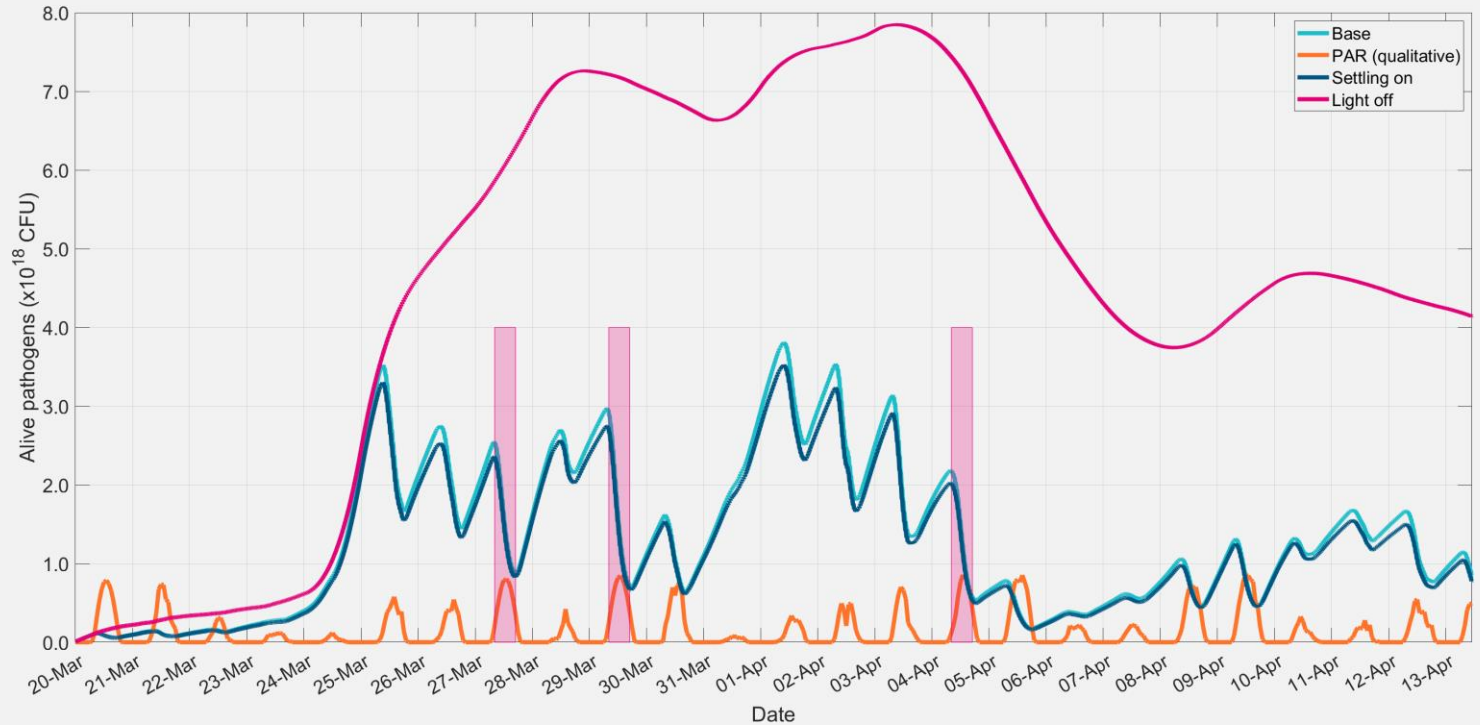
- Is not comparable in flux to light inactivation
- But more quantitatively
 - Use additional TUFLOW FV outputs
 - Total masses summed across the entire model domain in time



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Quantitatively

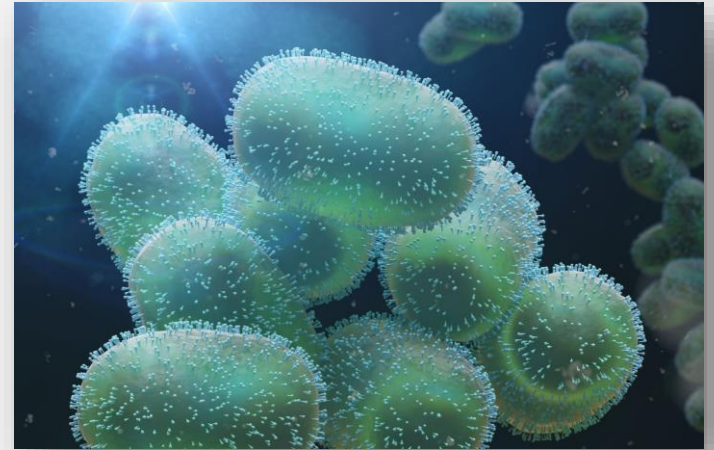
- Different scale



Lake St Clair

What have we learned?

- Light inactivation is clearly the dominant process in this system
- To promote understanding, we have used
 - TUFLOW FV water quality module diagnostics / fluxes
 - TUFLOW FV mass reporting
 - Multiple lines of evidence beyond trying to interpret concentrations
- Is a general technique for expediting water quality model calibration
 - **Do not guess: understand!**



Summary

Modelling environmental pathogens

- No simple task!
- Usually report concentrations (benchmarks)
- But: explore fluxes / diagnostics
 - Seek understanding
 - Why? To support What?
- Applies equally to large and small systems
 - Urban wetlands



Questions?

