

# Quantitative Models to Assess the Operational Impact of Biosurveillance and Early Warning of Infectious Disease

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## Summary

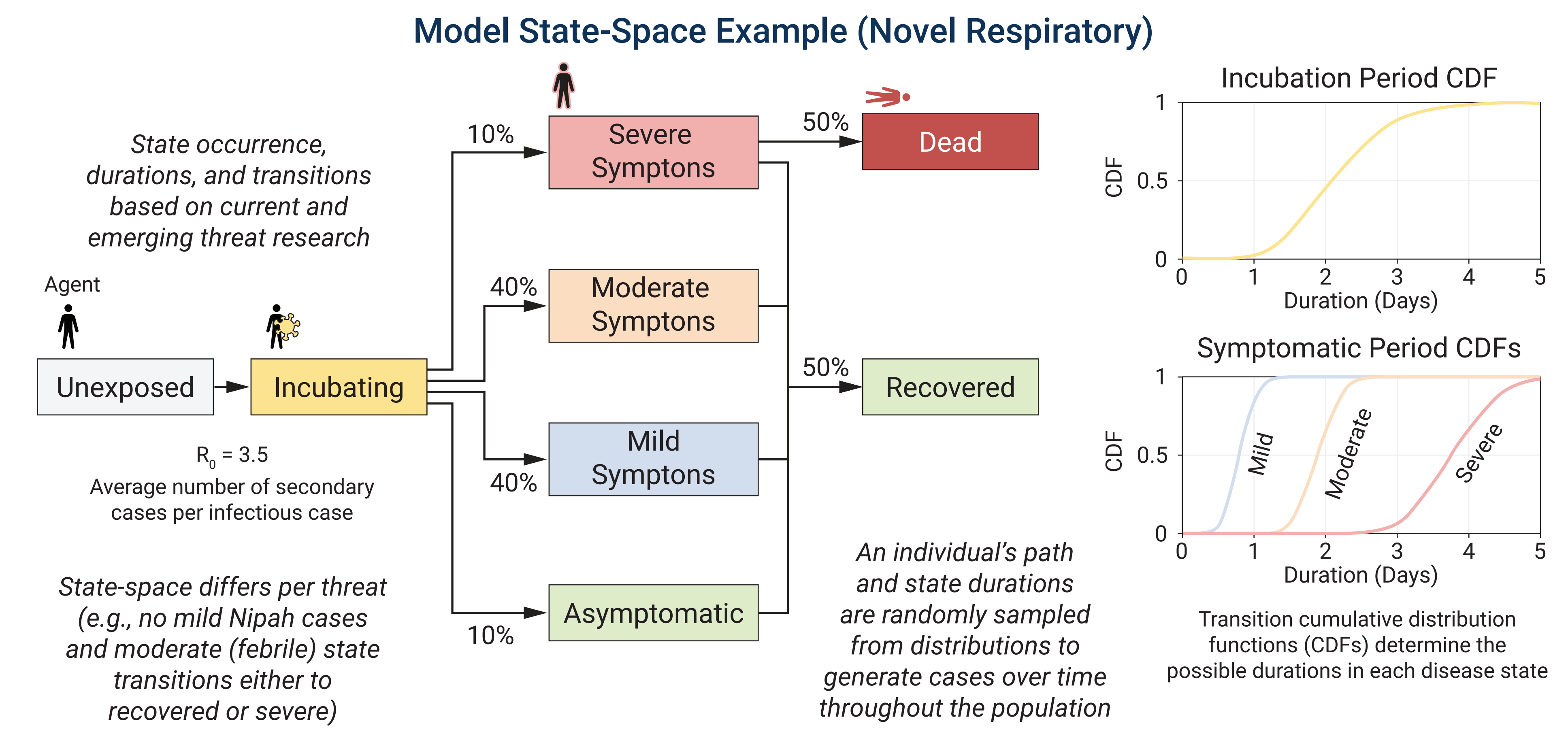
- Biosurveillance systems need to respond to a range of scenarios
- A stochastic disease model has been developed that is adaptable for different scenarios and enables quantification of optimal information timing and response utility
- These outputs support optimization of biosurveillance architecture development and prioritization of R&D

## Motivation

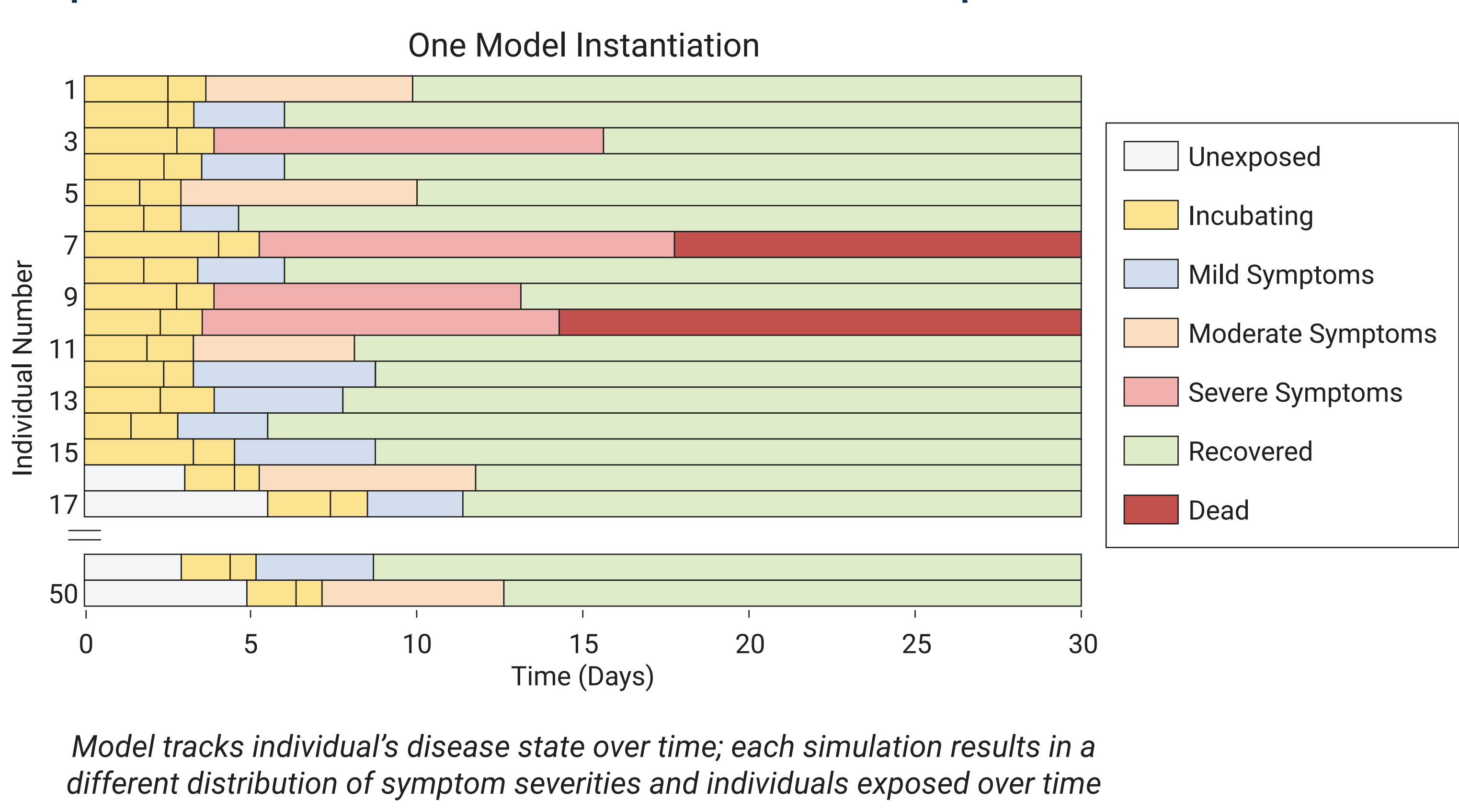
Force readiness depends on our ability to detect, understand, and respond to a variety of biological threats before they compromise mission success. It is a priority of the DoW to develop biosurveillance capabilities to protect force health and mission effectiveness.

- To support this priority:
- A stochastic model has been developed to simulate disease progression and to quantify operational impact and response effectiveness as a function of response initiation time
  - Outputs from the modeling framework support the development and refinement of biosurveillance architectures and the prioritization of capability development to maximize force readiness.

## Disease Model

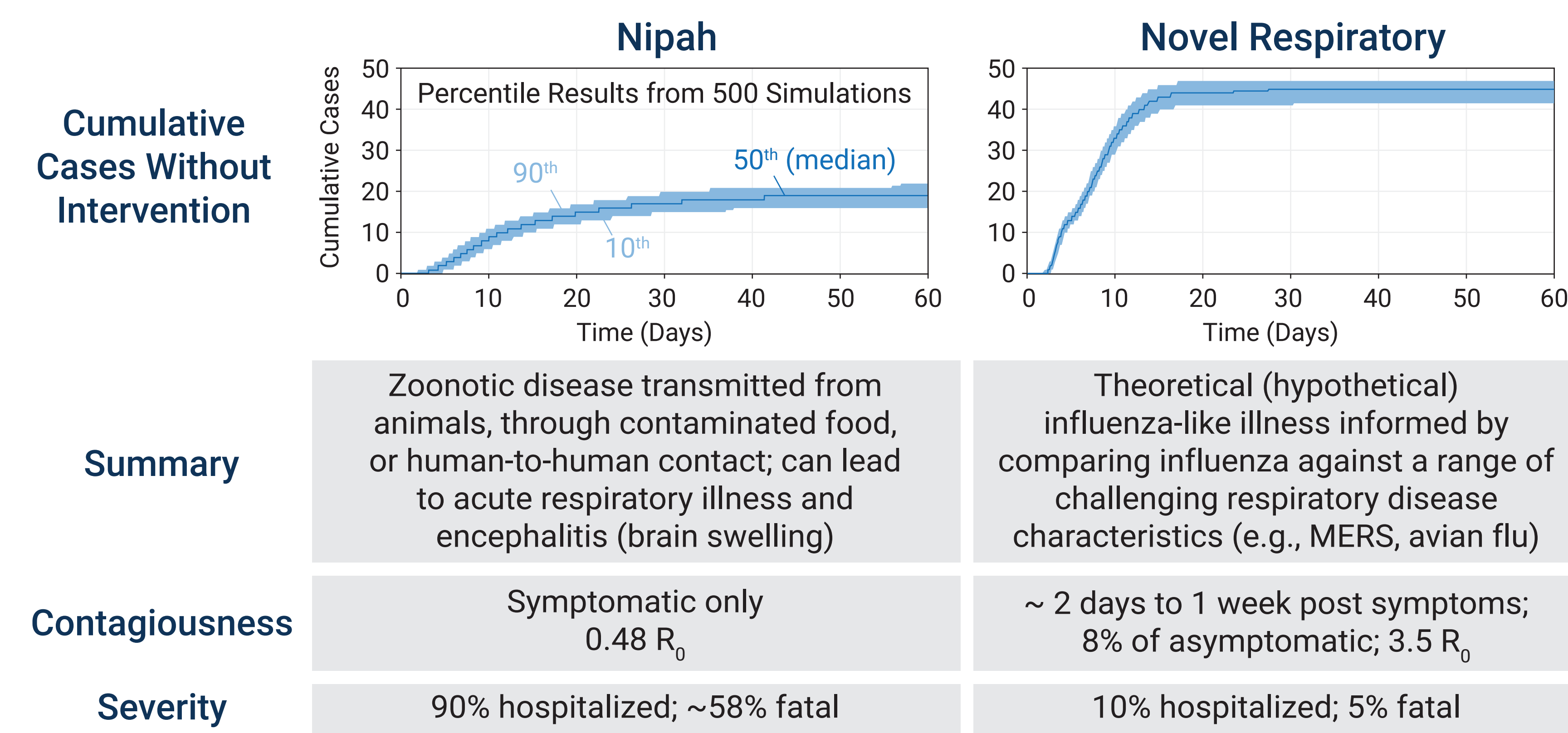


### Operational Scenario: 50 Soldiers with 15 Initial Exposures



## Threats Modeled

Mission impact, response options, and response timelines differ by disease, highlighting the need to assess biosurveillance across the full range of disease threats



## Detect and Mitigate

Shared biosurveillance information will lag initial detections which depend on specific threat progression

Response options differ depending on threat and current state of countermeasures

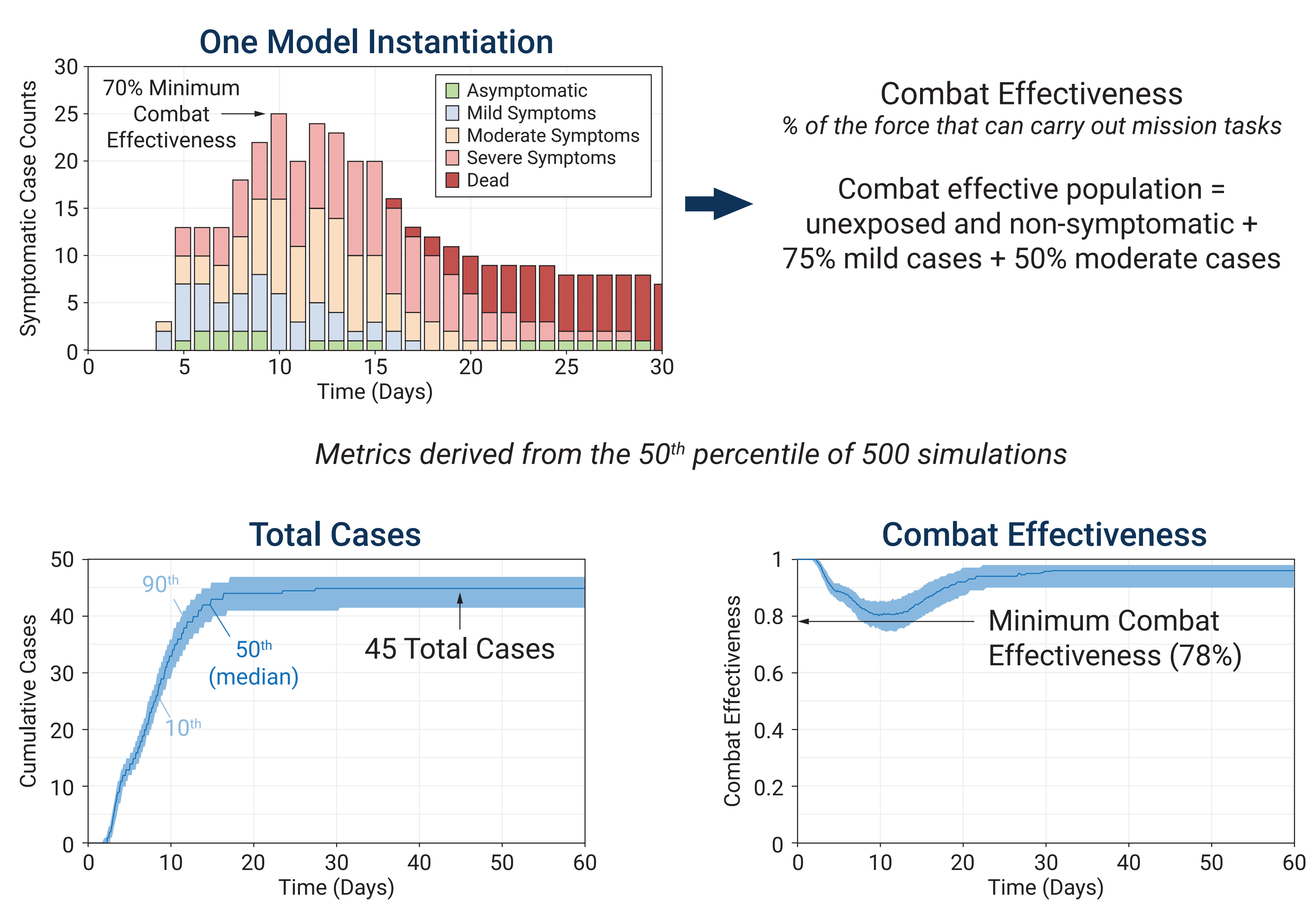
| Earliest Detection    |                                                                                                                                                                         |  |
|-----------------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------|--|
| Data/Info Type        | Detection Timing Assumptions                                                                                                                                            |  |
| Environmental Sensors | Potentially available at exposure (depending on pathogen)                                                                                                               |  |
| Syndromic             | Reporting occurs upon symptom onset                                                                                                                                     |  |
| Clinical Diagnosis    | <ul style="list-style-type: none"><li>• First severe case (field forward) + time to access diagnostic technologies</li><li>• First fatality + reporting delay</li></ul> |  |
| Sequencing            | First severe case + time to sequence and analyze results                                                                                                                |  |

*Estimate timelines for different information types to become available based on scenario context, resource availability, and specific threat progression.*

| Response Options                      |                                                               |                                                   |                                                  |
|---------------------------------------|---------------------------------------------------------------|---------------------------------------------------|--------------------------------------------------|
| Option                                | Estimated Effectiveness                                       | Assumptions                                       |                                                  |
|                                       |                                                               | Nipah                                             | Novel Respiratory                                |
| Non-Pharmacological (NP) Intervention | 80% reduced infectious contact (e.g., N95 masking)            | Applied 100% compliant population; proper use     | Applied 90% compliant population; proper use     |
| Cohorting/Isolation                   | 100% reduced infectious contact                               | Applied by 100% compliant, symptomatic population | Applied by 90% compliant, symptomatic population |
| Antiviral Treatment                   | 36% reduction in mortality and 1-day reduced symptom duration | Not applicable (No treatment exists)              | Administered to symptomatic                      |

## Operational Impact Metrics

### Metrics Example (Novel Respiratory Without Intervention)



## Estimated Impact of Information Timing

Modeling results enable quantitative assessment of biological event impacts, response utility, and optimal information timing to maximize force readiness.

