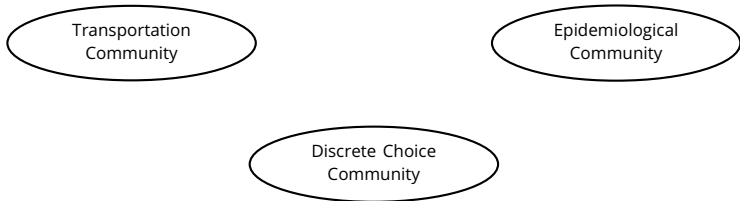


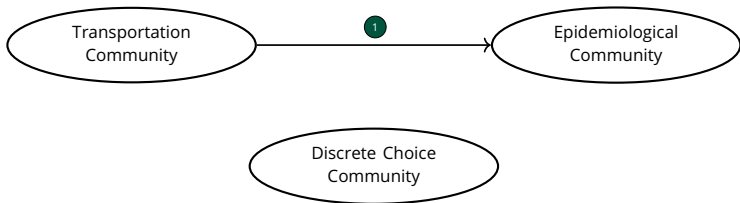


## How do we link these communities?



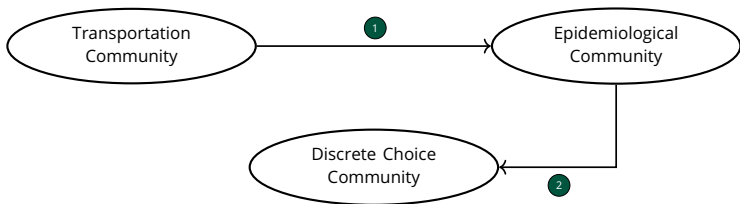
## How do we link these communities?

- 1 Activity-travel behavior impacts the spread of a disease.



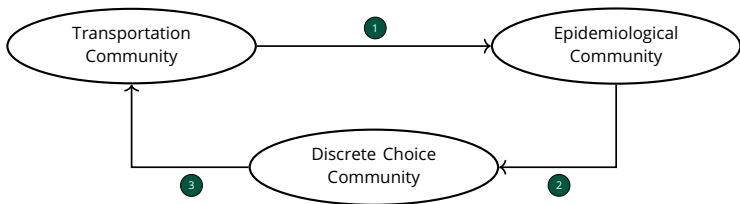
## How do we link these communities?

- 1 Activity-travel behavior impacts the spread of a disease.
- 2 Being infected is not a choice, testing is.

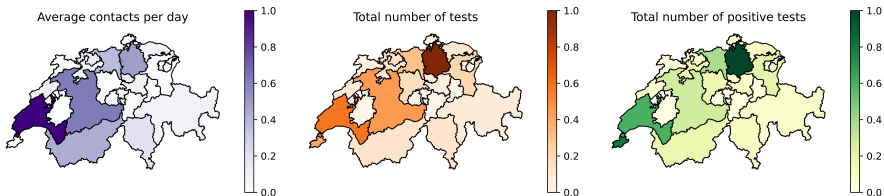


## How do we link these communities?

- 1 Activity-travel behavior impacts the spread of a disease.
- 2 Being infected is not a choice, testing is.
- 3 Testing choices change the activity-travel behavior of the individuals, and therefore the spread of a disease.



# Research Question



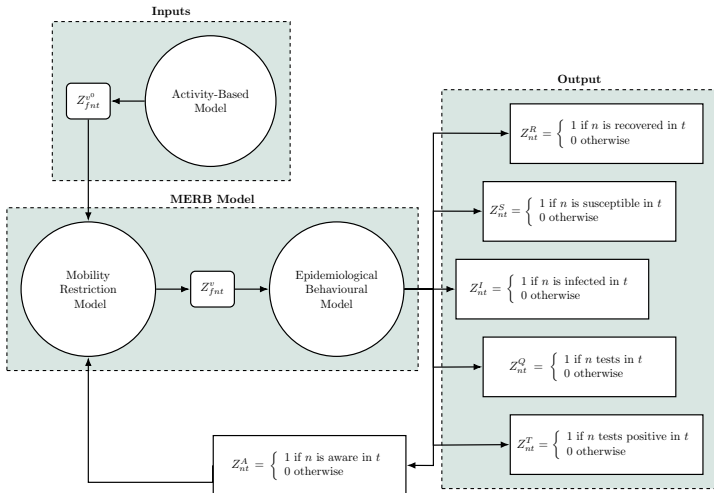
**“How do individual testing choices influence activity-travel behavior, and therefore, the dynamics of disease spread?”**

# Research Gaps

- Traditional models often overlook the **heterogeneity of individual actions** and their impact on the spread of disease (Hackl and Dubernet 2019; Eubank et al. 2004; Perez and Dragicevic 2009).
- Existing research does not fully integrate **socioeconomic and health factors** influencing mobility and epidemiological outcomes.
- There is a **lack of emphasis on individual choices**, particularly regarding testing and the subsequent behavioral adjustments (Cui, Ni, and Shen 2021; Brotherhood et al. 2020).
- Need for models that can **dynamically capture individual decisions** and their effects on the pandemic's trajectory (Tuomisto et al. 2020).

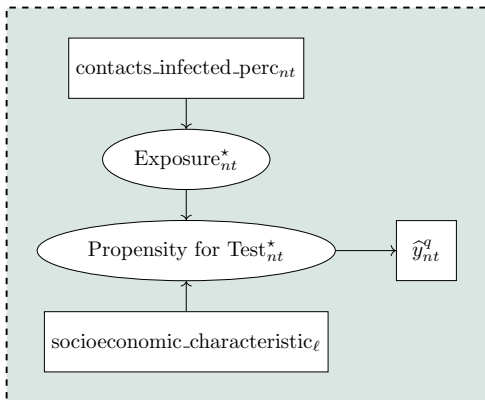
**Main Problem:** Lack of disaggregated data available.

# Overall framework





# Epidemiological-Behavioural Model



Structural model

- $L$  socioeconomic characteristics: age, employment, income, level of education, etc.
- $\hat{y}_{nt}^q$  binary observed data, 1 if the individual  $n$  tested at timestep  $t$ .
- $\text{contacts\_infected\_perc}_{nt}$ : the proportion of infected individuals that individual  $n$  encounters in a facility  $f$  at timestep  $t$ , where

$$x_{nt}^v = \sum_f \sum_{m \neq n} Z_{fnt}^v Z_{fnt}^v Z_{fnt}^i$$

# Epidemiological-Behavioural Model: Structural equations

## Exposure State

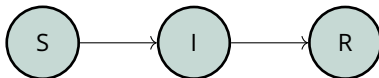
$$E_{nt}^* = \beta_V x_{nt}^V + \varepsilon_{E^*}$$

## Propensity to Test

$$Q_{nt}^* = \beta_{Q^*}^0 + \sum_{\ell=1}^{L-1} \beta_{\ell} x_n^{\ell} + \eta_{E^*} E_{nt}^* + \varepsilon_{Q^*}$$

- $x_n^{\ell}$ :  $L$  socioeconomic characteristics of individual  $n$ .
- $x_{nt}^V$ : is the number of contacts\_infected\_perc $_{nt}$ .
- $\beta_{Q^*}^0$ ,  $\beta_V$ ,  $\beta_{\ell}$ ,  $\eta_{E^*}$ ,  $\varepsilon_{E^*}$ , and  $\varepsilon_{Q^*}$  are parameters to be calibrated.

# Dynamics of the MERB model: Health State Transition



A **hidden Markov chain** is employed to model the **transitions** between **health states**. The **state transition matrix**  $\mathbb{B}$  is:

$$\begin{bmatrix} P(Z_{n(t+1)}^s = 1 | Z_{nt}^s = 1) & P(Z_{n(t+1)}^i = 1 | Z_{nt}^s = 1) & 0 \\ 0 & P(Z_{n(t+1)}^i = 1 | Z_{nt}^i = 1) & P(Z_{n(t+1)}^r = 1 | Z_{nt}^i = 1) \\ P(Z_{n(t+1)}^s = 1 | Z_{nt}^r = 1) & 0 & P(Z_{n(t+1)}^r = 1 | Z_{nt}^r = 1) \end{bmatrix}$$

# Dynamics of the MERB model: Health State Transition

Is the individual **becoming infected or remaining susceptible**?

$$P(Z_{n(t+1)}^i = 1 | Z_{nt}^s = 1) = E_{nt}^* \quad (1)$$

$$P(Z_{n(t+1)}^s = 1 | Z_{nt}^s = 1) = 1 - P(Z_{n(t+1)}^i = 1 | Z_{nt}^s = 1) \quad (2)$$

$$Z_{n(t+1)}^i = \begin{cases} 1, & \text{if } \pi < P(Z_{n(t+1)}^i = 1 | Z_{nt}^s = 1) \\ 0, & \text{otherwise.} \end{cases} \quad (3)$$

$\pi$ : random uniform value

# Dynamics of the MERB model: Health State Transition

Is the individual **becoming recovered or remaining infected?**

$$P(Z_{n(t+1)}^r = 1 | Z_{nt}^i = 1) = \Phi_{\gamma_n}(t - t_n^i)$$

$\Phi_{\gamma_n}$ : cumulative distribution function of the log-normal distribution of  $\gamma_n$  (recovery rate (Kerr et al. 2020)).

$t - t_n^i$ : number of timesteps that satisfy  $Z_n^i = 1$ .

For each infected individual  $n$  at time  $t$ , we decide if they will be recovered by time  $t$  given that they were infected at time  $t_n^i$ , by computing:

$$Z_{n(t+1)}^r = \begin{cases} 1, & \text{if } \pi < P(Z_{n(t+1)}^r = 1 | Z_{nt}^i = 1) \\ 0, & \text{otherwise.} \end{cases} \quad (4)$$

# Dynamics of the MERB model: Health State Transition

Finally, we assume that an individual who recovers remains recovered until the end of the simulation.

$$P(Z_{n(t+1)}^r = 1 | Z_{nt}^r = 1) = 0,$$

$$P(Z_{n(t+1)}^s = 1 | Z_{nt}^r = 1) = 0.$$

# Dynamics of the MERB model: Testing process

## What about the testing process?

We model the probability of an individual to test through a logit:

$$P(Z_{n(t+1)}^q = 1) = \frac{1}{1 + e^{-\mu Q_{nt}^*}}, \quad (5)$$

and each individual  $n$  at time  $t$ , decides to get tested by generating  $\pi$ , and following:

$$Z_{n(t+1)}^q = \begin{cases} 1, & \text{if } \pi < P(Z_{nt}^q) \\ 0, & \text{otherwise.} \end{cases} \quad (6)$$

# Dynamics of the MERB model: Mobility Restriction Model - What about awareness?

## Assumption:

Only people that are infected and test positive will quarantine.

$$Z_{nt}^a = Z_{nt}^i Z_{nt}^+ \quad (7)$$

Now, we apply the mobility restriction model, and the new dynamics can be written as:

$$Z_{fn(t+1)}^v = \begin{cases} Z_{fnt}^v & \text{if individual's } n \text{ outcome is 0, and} \\ 0 & \text{otherwise.} \end{cases} \quad (8)$$



# Results: Case Study - Population of Vaud

## Study Focus

- Our study examines the population of **Vaud in Switzerland (823'456 individuals)**.
- A **synthetic population** is generated to simulate **socio-economic characteristics** and **daily schedules** for each individual (Horl and Balac 2021).

## Data requirements

- **Open-source data** including **number of tested individuals** ( $\hat{y}_{gw}^q$ ) and **positive tests** ( $\hat{y}_{gw}^+$ ) individuals ( CloudPlatform 2021).
- Data from **Federal Office of Public Health** including the **tested positive** individuals ( $\hat{y}_{hjk\ell}^+$ ) with their **age, gender and municipality** information ( Riou et al. 2021).

# Results: Case Study - Population of Vaud

## Data and Model Development

Due to the unavailability of disaggregated data:

- The exposure level of an individual ( $E_{nt}^*$ ) is calculated as:

$$E_{nt}^* = \beta^v x_{nt}^v, \quad (9)$$

- The propensity to test of an individual ( $Q_{nt}^*$ ) is calculated as:

$$Q_{nt}^* = \beta_{Q^*}^0 + \beta_{age} x_n^{age} + \beta_{employment} x_n^{employment} + \eta_{E^*} E_{nt}^*, \quad (10)$$

where:

- $x_n^{age}$  represents the age, and  $x_n^{employment}$  represents the employment status of the individual.
- $\beta_{age}$ ,  $\beta_{employment}$ ,  $\beta^v$ ,  $\beta_{Q^*}^0$  and  $\eta$  are parameters to be estimated.

## Measurement equations

Since available **data** tends to be **aggregated**,

we link the tests performed for each individual and timestep, with the observed number of tests by defining:

$$P_{gw}^q = \sum_{n \in g} \sum_{t=1}^{7T} P(Z_{nt}^q = 1), \quad (11)$$

$$P_{hjk\ell}^+ = \sum_{i \in h,j,k\ell} \sum_{t=1}^T P(Z_{nt}^+ = 1). \quad (12)$$

**Beware:** The lack of panel data presents a challenge because the modeling of testing choices is influenced by preceding ones.

# Epidemiological-Behavioural Model: Measurement equations

## Data & Likelihood functions

We use the **negative binomial distribution** for modeling the count data.

$\mathcal{L}_1(\theta)$  - testing

$$\begin{aligned}
 &= \sum_{i \in g} \sum_{w \in W} \left( \log \Gamma(\hat{y}_{gw}^q + r_1) - \log \Gamma(r_1) \right. \\
 &\quad \left. - \log \Gamma(\hat{y}_{gw}^q + 1) + r_1 \cdot \log \left( \frac{r_1}{r_1 + P_{gw}^q} \right) \right. \\
 &\quad \left. + \hat{y}_{gw}^q \cdot \log \frac{P_{gw}^q}{r_1 + P_{gw}^q} \right),
 \end{aligned}$$

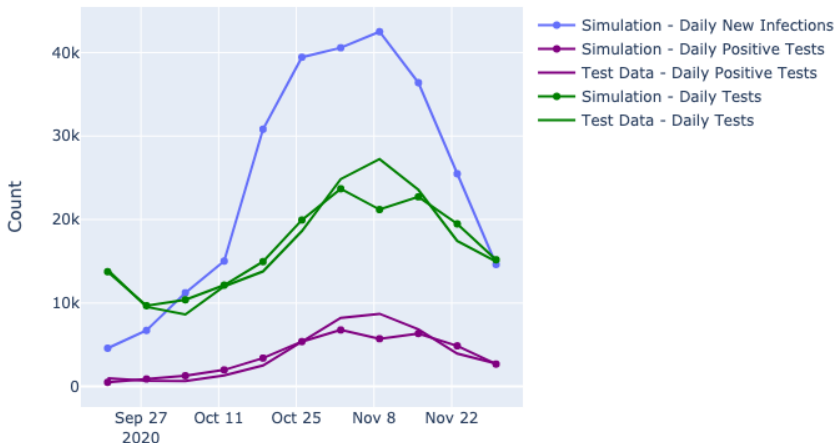
$\mathcal{L}_2(\theta)$  - testing positive

$$\begin{aligned}
 &= \sum_{i \in h, j, k, \ell} \sum_{\ell \in L} \left( \log \Gamma(\hat{y}_{hjk\ell}^+ + r_2) - \log \Gamma(r_2) \right. \\
 &\quad \left. - \log \Gamma(\hat{y}_{hjk\ell}^+ + 1) + r_2 \cdot \log \left( \frac{r_2}{r_2 + P_{hjk\ell}^+} \right) \right. \\
 &\quad \left. + \hat{y}_{hjk\ell}^+ \cdot \log \frac{P_{hjk\ell}^+}{r_2 + P_{hjk\ell}^+} \right).
 \end{aligned}$$

where  $r_1$  and  $r_2$  are the parameters of the negative binomial,  $\hat{y}_{gw}^+$  are the positive tests per age group,  $g$ , and week  $w$ , and  $\hat{y}_{hjk\ell}^+$  are the positive tests per age,  $h$ , gender,  $j$ , and municipality  $k$ , per day  $\ell$ .

# Calibrated model: aggregated validation

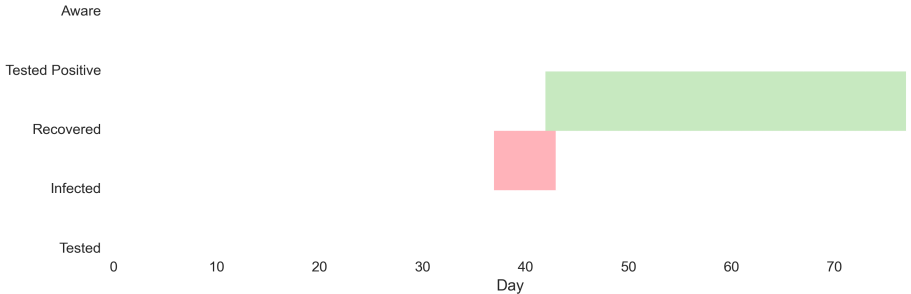
## Epidemic Incidence and COVID19 Testing Trajectory



# Results across geographical area

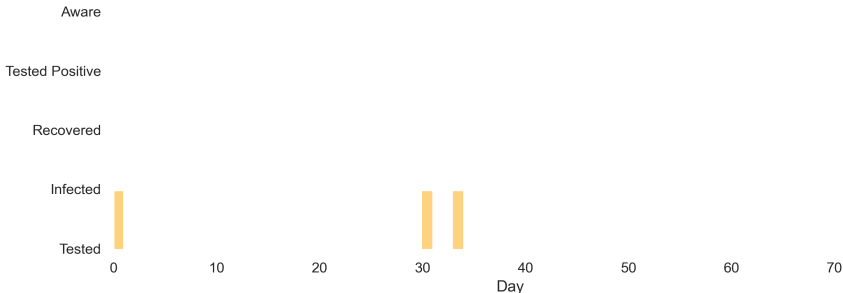
# Results across individuals: insights on behavior

## The 'unaware' behavior:



# Results across individuals: insights on behavior

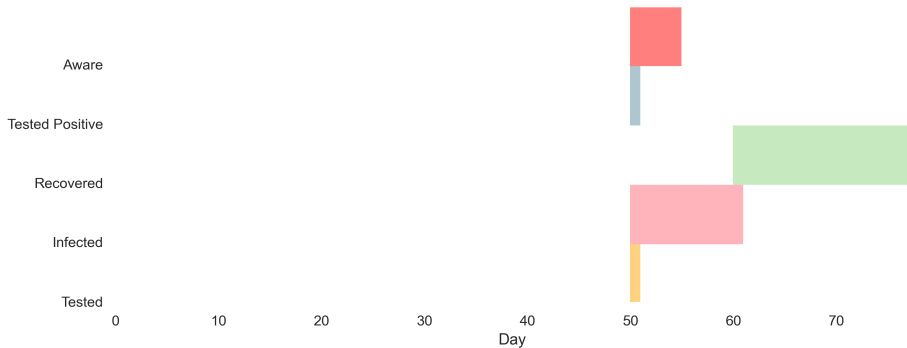
## The 'cautious' behavior:



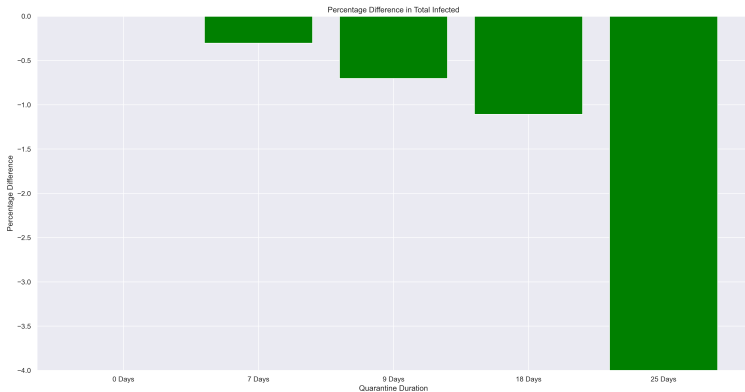


# Results across individuals: insights on behavior

## The 'full package' behavior:



# Results by applying different confinements



# Conclusion and Future Work

## Conclusions:

- 1 Tool that effectively models the individual dynamic choice of testing, and the impact on activity-travel behaviour.
- 2 We are able to predict the actual infected individuals (in our case study are around 3 times higher than the tracked data).
- 3 A computationally efficient tool (1m 11s for 800k individuals and 90 days with a timestep of 30m).
- 4 Lack of disaggregated data always makes it hard to calibrate the models.
- 5 Bridging epidemiology, transportation, and discrete choice communities for a interdisciplinary model that can better explain how and why a spreading occurs.

## Future work:

- 1 Include health characteristics and calibrate the model.
- 2 Run this model together with the policy optimization framework from Cortes Balcells 2021.
- 3 Run the model for different cantons and see how the testing behavior impacts the spreading.



Thank you for your attention

## Testing process

The outcome of the test is computed by:

$$\begin{aligned}
 P(Z_{nt}^+ = 1 | Z_{nt}^i = 1) &= P(Z_{nt}^+ = 1 | Z_{nt}^q = 1 \text{ and } Z_{nt}^i = 1) P(Z_{nt}^q = 1 \text{ and } Z_{nt}^i = 1) = \\
 P(Z_{nt}^+ = 1 | Z_{nt}^q = 1 \text{ and } Z_{nt}^i = 1) P(Z_{nt}^i = 1 | Z_{nt}^q = 1) P(Z_{nt}^q = 1) &= \\
 P(Z_{nt}^+ = 1 | Z_{nt}^q = 1 \text{ and } Z_{nt}^i = 1) P(Z_{nt}^i = 1) P(Z_{nt}^q = 1), &
 \end{aligned}$$

$$\begin{aligned}
 P(Z_{nt}^+ = 1 | Z_{nt}^s = 1) &= P(Z_{nt}^+ = 1 | Z_{nt}^q = 1 \text{ and } Z_{nt}^s = 1) P(Z_{nt}^q = 1 \text{ and } Z_{nt}^s = 1) = \\
 P(Z_{nt}^+ = 1 | Z_{nt}^q = 1 \text{ and } Z_{nt}^s = 1) P(Z_{nt}^s = 1 | Z_{nt}^q = 1) P(Z_{nt}^q = 1) &= \\
 P(Z_{nt}^+ = 1 | Z_{nt}^q = 1 \text{ and } Z_{nt}^s = 1) P(Z_{nt}^s = 1) P(Z_{nt}^q = 1), &
 \end{aligned}$$

$$\begin{aligned}
 P(Z_{nt}^+ = 1 | Z_{nt}^r = 1) &= P(Z_{nt}^+ = 1 | Z_{nt}^q = 1 \text{ and } Z_{nt}^r = 1) P(Z_{nt}^q = 1 \text{ and } Z_{nt}^r = 1) = \\
 P(Z_{nt}^+ = 1 | Z_{nt}^q = 1 \text{ and } Z_{nt}^r = 1) P(Z_{nt}^r = 1 | Z_{nt}^q = 1) P(Z_{nt}^q = 1) &= \\
 P(Z_{nt}^+ = 1 | Z_{nt}^q = 1 \text{ and } Z_{nt}^r = 1) P(Z_{nt}^r = 1) P(Z_{nt}^q = 1). &
 \end{aligned}$$

$P(Z_{nt}^+ = 1 | Z_{nt}^q = 1 \text{ and } Z_{nt}^i = 1)$ ,  $P(Z_{nt}^+ = 1 | Z_{nt}^q = 1 \text{ and } Z_{nt}^s = 1)$ ,  $P(Z_{nt}^+ = 1 | Z_{nt}^q = 1 \text{ and } Z_{nt}^r = 1)$

are taken from Ai et al. 2020. 