Simple Epidemic Models with Segmentation Can Be Better than Complex Ones

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Overview

- 1. Introduction
- 2. Backgrounds
- 3. Method
- 4. Experiments
- 5. Conclusion



Problem Definition

- **Understanding and predicting epidemic spreads** are important for prediction and effective decision making.
 - How many people will be infected within a week?
 - How will lockdowns affect the spread?
- To answer these questions, we require a method that is **simple & expressive** method to model and predict the spread of infectious diseases.

Is a Single Model Enough?

- Epidemic models describe dynamics of epidemic spreads.
- However, describing **long-term dynamics of epidemics** is challenging.
 - Unpredictability & abruptness of real-world events
 - E.g., lockdowns or the capability to perform tests

Experiments

Conclusion

Is a Single Model Enough? (cont.)

- Can a single model describe the long-term epidemic event sequence?
 - The data is extremely **complex**.
 - The model can **overfit** the data.

Our Idea

Properly divide an epidemic event sequence into **multiple segments** and fit a

simple epidemic model to each segment.

Ordinary Differential Equations (ODEs)

- Many epidemic models are based on **ordinary differential equations** (**ODEs**).
- Some of the earliest epidemic models are SIS, **SIR**, and SEIR.
 - They are based on human knowledge.
- Other data-driven models: LLD and NLLD
 - They model time-series data without relying on human knowledge.

Conclusion

SIR: Susceptible-Infectious-Recovered Model

- A closed population *P* is divided into three states:
 - S (susceptible), I (infectious), and R (recovered)
- At timestamp t, the number of individuals of each state is S(t), I(t), and R(t).



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LLD: Linear Latent Dynamics Model [Matsubara and Sakurai, KDD 2016]

- v(t): *d*-dimensional **observable** event sequence
 - In our case, we use 2-dimensional data v(t) = [I(t), R(t)].
- w(t): k-dimensional **latent** event sequence

$$\frac{\mathrm{d}w(t)}{\mathrm{d}t} = \mathbf{p} + \mathbf{Q}w(t)$$
$$v(t) = \mathbf{u} + \mathbf{V}w(t)$$

- **p** and **Q** describe dynamics between latent factors
 - **p** ($\in \mathbb{R}^k$): linear dynamics
 - **Q** ($\in \mathbb{R}^{k \times k}$): exponential dynamics
- $\mathbf{u} \in \mathbb{R}^d$ and $\mathbf{V} \in \mathbb{R}^{k \times d}$ project latent factors to the observed events

NLLD: Non-Linear Latent Dynamics Model [Matsubara and Sakurai, KDD 2016]

- v(t): *d*-dimensional **observable** event sequence
 - In our case, we use 2-dimensional data v(t) = [I(t), R(t)].
- w(t): k-dimensional **latent** event sequence

$$\frac{\mathrm{d}w(t)}{\mathrm{d}t} = \mathbf{p} + \mathbf{Q}w(t) + \mathbf{A}w(t)^2$$
$$v(t) = \mathbf{u} + \mathbf{V}w(t)$$

- **p**, **Q**, and **A** describe dynamics between latent factors
 - **p** ($\in \mathbb{R}^k$): linear dynamics
 - **Q** ($\in \mathbb{R}^{k \times k}$): exponential dynamics
 - **A** ($\in \mathbb{R}^k$): **non-linear** dynamics
- $\mathbf{u} \in \mathbb{R}^d$) and $\mathbf{V} \in \mathbb{R}^{k \times d}$) project latent factors to the observed events

Our Method

Our Idea

Properly divide an epidemic event sequence into **multiple segments** and fit a

simple epidemic model to each segment.

Q1. How to divide the epidemic event sequence?

Q2. How to automatically find the best segmentation?

Q3. How to control the trade-offs between the model complexity and fitness?



Description Length

• Given a sequence X and a model M, the description length (in bits) of X is:

 $Cost(X) \coloneqq Cost(M) + Cost(X|M)$

- Model cost *Cost(M)*: the number of bits required to describe the model *M*
- Data cost Cost(X|M): the number of bits to encode X given M

Model Cost

<u>SIR Model</u>	LLD Model	NLLD Model
$2 \cdot C_F$ bits	$(k^2 + (2+d) \cdot k + d) \cdot C_F$ bits	$(k^2 + (3+d) \cdot k + d) \cdot C_F$ bits
• β : C_F bits	• $w_0 \in \mathbb{R}^k$: $k \cdot C_F$ bits	• $w_0 \in \mathbb{R}^k : k \cdot C_F$ bits
• γ : C_F bits	• $p \in \mathbb{R}^k$: $k \cdot C_F$ bits	• $p \in \mathbb{R}^k$: $k \cdot C_F$ bits
	• $\boldsymbol{Q} \in \mathbb{R}^{k imes k}$: $\boldsymbol{k}^2 \cdot \boldsymbol{C}_F$ bits	• $\boldsymbol{Q} \in \mathbb{R}^{k imes k}$: $\boldsymbol{k}^2 \cdot \boldsymbol{C}_F$ bits
	• $\boldsymbol{u} \in \mathbb{R}^d$: $\boldsymbol{d} \cdot \boldsymbol{C}_F$ bits	• $\pmb{A} \in \mathbb{R}^k$: $\pmb{k} \cdot \pmb{C}_F$ bits
	• $\boldsymbol{V} \in \mathbb{R}^{k imes d}$: $\boldsymbol{k} \cdot \boldsymbol{d} \cdot \boldsymbol{C}_F$ bits	• $\boldsymbol{u} \in \mathbb{R}^d$: $\boldsymbol{d} \cdot \boldsymbol{C}_F$ bits
		• $V \in \mathbb{R}^{k \times d}$: $k \cdot d \cdot C_F$ bits

 C_F is the number of bits to encode a real number.



Data Cost

- **Data cost** is the number of bits to encode *X* given *M*.
 - *X*: observed event sequence
 - *V*: estimated event sequence by the model *M*
- The number of bits required is the negative log-likelihood under a Gaussian distribution $\mathcal{N}(0, \sigma^2)$:

$$Cost(X|M) = -\log P(X - V) = -\log \prod_{t=1}^{n} \prod_{i=1}^{d} \frac{1}{\sqrt{2\pi\sigma^2}} e^{-\frac{(x_i(t) - v_i(t))^2}{2\sigma^2}}$$

Conclusion

Trade-off: Model Complexity vs. Fitness



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Trade-off: Model Complexity vs. Fitness (cont.)

• We divide the sequence into **multiple segments** and fit a **simple model** to each segment.



Q. How to **automatically** divide the sequence that leads to better trade-offs

between model complexity and fitting error?



Minimum Description Length (MDL) Principle

• An epidemic event sequence $X (= X_{1:n})$ of length n is divided into r segments:

$$X_{S_1:e_1} \oplus \cdots \oplus X_{S_r:e_r}$$

• Apply model f_i to each segment $X_{s_i:e_i}$.

<i>s</i> ₁		e_1	$s_2 e_2$	$s_3 e_3$	$S_4 e_4$
	$X_{s_1:e_1}$		$X_{s_2:e_2}$	$X_{s_3:e_3}$	$X_{s_4:e_4}$
	1		1	1	1
	f_1		f_2	f_3	f_4



Minimum Description Length (MDL) Principle (cont.)

• The description length in bits of $X_{s_1:e_1} \oplus \cdots \oplus X_{s_r:e_r}$ is:

$$Cost(X_{s_1:e_1} \oplus \cdots \oplus X_{s_r:e_r}) = (r-1) \cdot \log_2(n) + \sum_{i=1}^r \left(Cost(f_i) + Cost(X_{s_i:e_i}:f_i) \right)$$

$$(1)$$

1 Bits required to encode r - 1 splitting points.

2 The description length of each segment.



Segmentation Search

- Given an event sequence $X_{1:n}$, there are 2^n ways of segmentation.
- We propose a greedy segmentation scheme.



Find a splitting point where the description length is minimized.

Splitting the segment is no more beneficial. Stop splitting.

Experimental Settings

• We consider **70 countries** with the most confirmed cases of **COVID-19** as of the end of March 2021.

Argentina, Armenia, Austria, Azerbaijan, Bangladesh, Belarus, Belgium, Bolivia, Brazil, Bulgaria, Canada, Chile, Colombia, Costa Rica, Croatia, Czech, Denmark, Dominican Republic, Ecuador, Egypt, France, Georgia, Germany, Greece, Guatemala, Honduras, Hungary, India, Indonesia, Iran, Iraq, Ireland, Israel, Italy, Japan, Jordan, Kazakhstan, Kuwait, Lebanon, Lithuania, Malaysia, Mexico, Moldova, Morocco, Nepal, Netherlands, Pakistan, Panama, Paraguay, Peru, Philippines, Poland, Portugal, Qatar, Romania, Russia, Saudi Arabia, Serbia, Slovakia, Slovenia, South Africa, Spain, Sweden, Switzerland, Tunisia, Turkey, United Arab Emirates, United Kingdom, Ukraine, United States

Public datasets: <u>https://www.kaggle.com/sudalairajkumar/novel-corona-virus-2019-dataset</u>

Q1. Effectiveness of Segmentation

Simple epidemic models with segmentation provide more concise and accurate

description of the spread of COVID-19 than complex models without segmentation.



NLLD (India)



Fitting with Segmentation (Proposed)

Fitting without Segmentation

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Q1. Effectiveness of Segmentation (cont.)

Simple epidemic models with segmentation provide more concise and accurate

description of the spread of COVID-19 than complex models without segmentation.





Q2. Effectiveness of Our Segmentation Scheme

Our segmentation scheme yields better segmentation scheme than in the

incremental method [Matsubara and Sakurai, KDD 2016].



O Greedy Segmentation (Proposed)

Incremental Segmentation

NLLD (India)

Q3. Accuracy of Forecasting

Segmentation is helpful to accurate prediction of the spread of COVID-19.



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and fit a **simple model** to each segment.

Our methodology is:

- ✓ Automatic: All parameters are tuned automatically based on MDL principle.
- ✓ Model-agnostic: Any ODE-based epidemic models can be used.
- ✓ Effective: It fits and predicts well in COVID-19 datasets.

Code & datasets: <u>https://github.com/geonlee0325/covid_segmentation</u>