

# Aladynoulli: A Dynamic Model for Disease Progression

## Understanding Cardiovascular Risk Through Temporal Pattern Analysis

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# Outline

- 1 Model Overview
- 2 Mathematical Framework
- 3 Clinical Example
- 4 Model Benefits
- 5 Future Directions

# The Big Picture

## Key Ideas

- Individual risk profiles evolve over time
- Diseases share underlying risk factors
- Genetic factors influence progression
- Time-varying disease relationships

## Clinical Relevance

- Personalized risk trajectories
- Natural disease progression
- Multiple disease interactions
- Treatment planning implications

# Individual Risk Profiles ( $\lambda$ )

## Individual-Specific Topic Scores

For each individual  $i$ , topic  $k$ , and time  $t$ :

$$\lambda_{ik}(t) \sim \mathcal{GP}(\Gamma_k^T g_i, \Sigma_k)$$

### Components:

- $g_i$ : Genetic covariates
- $\Gamma_k$ : Genetic effects
- $\Sigma_k$ : Temporal covariance

### Clinical Meaning:

- Personal trajectories
- Genetic influence
- Smooth evolution

# Topic Proportions ( $\theta$ )

## From Scores to Proportions

Via softmax transformation:

$$\theta_{ik}(t) = \frac{\exp(\lambda_{ik}(t))}{\sum_{j=1}^K \exp(\lambda_{ij}(t))}$$

### Properties:

- $\theta_{ik}(t) \in (0, 1)$
- $\sum_k \theta_{ik}(t) = 1$
- Smooth changes

### Interpretation:

- Relative risk weights
- Competing factors
- Dynamic profiles

# Disease Topic Loadings ( $\phi$ )

## Disease-Topic Relationships

For each disease  $d$  and topic  $k$ :

$$\phi_{kd}(t) \sim \mathcal{GP}(\mu_d, \Omega_k)$$

### Components:

- $\mu_d$ : Base disease risk
- $\Omega_k$ : Topic covariance

### Clinical Meaning:

- Topic-disease links
- Disease patterns
- Time variation

# Disease Probabilities ( $\pi$ )

## Individual Disease Risk

Probability for individual  $i$ , disease  $d$ , at time  $t$ :

$$\pi_{id}(t) = \sum_{k=1}^K \theta_{ik}(t) \cdot \text{sigmoid}(\phi_{kd}(t))$$

### Components:

- Personal risk profile
- Topic contributions
- Temporal dynamics

### Clinical Use:

- Risk prediction
- Trajectory planning
- Intervention timing

# Cardiovascular Risk Profiles

## Three Key Risk Domains

### 1 Metabolic Risk

- Strong diabetes link
- Gradual vascular impact

### 2 Vascular Risk

- Primary CAD driver
- Early HTN effects

### 3 Inflammatory Risk

- Accelerated progression
- Late complications



# Disease Progression Patterns

## Temporal Sequence

### ① Early Stage

- Hypertension
- Metabolic changes

### ② Mid Stage

- Type 2 Diabetes
- Early CAD signs

### ③ Late Stage

- Clinical CAD
- Heart Failure

# Key Advantages

## Clinical Strengths:

- Personalization
- Natural progression
- Multiple diseases
- Interpretable

## Statistical Power:

- Temporal smoothing
- Uncertainty measures
- Principled inference

## Practical Impact

- Better risk stratification
- Informed treatment planning
- Early intervention opportunities

# Next Steps

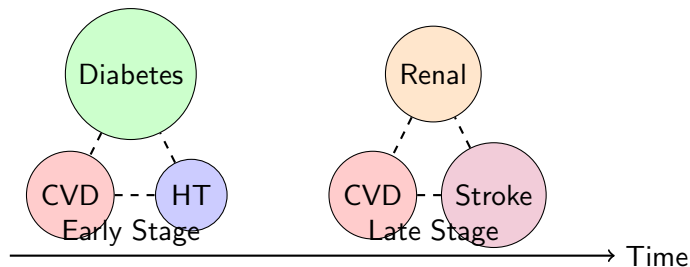
## Clinical Applications

- Risk stratification tools
- Treatment optimization
- Prevention strategies

## Model Extensions

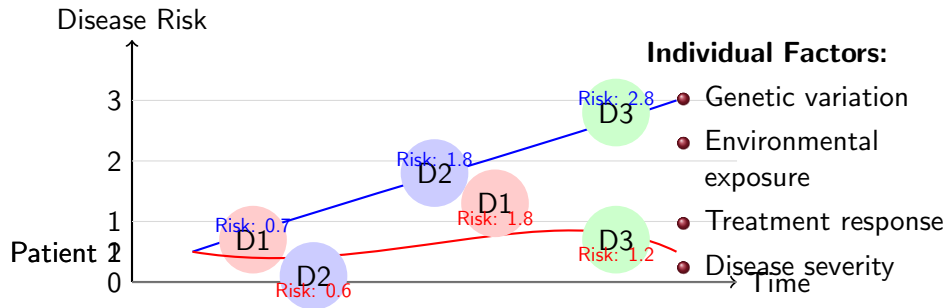
- Treatment effects
- Environmental factors
- Additional outcomes
- Real-world validation

## Problem 1: Topics Need to Evolve



Standard topic models assume fixed disease relationships within topics, but disease associations change over disease progression

## Problem 2: Individuals Need to Evolve



Topic models struggle with individual-specific temporal evolution and varying disease sequences