# Alternative estimation framework: from single population to cause-of-death

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## Generalized Age Period Cohort framework

General framework for single population following [AMV<sup>+</sup>18] (StMoMo R package).

• the *random component*: The number of deaths  $D_{x,t}$  is assumed to follow a Poisson or a Binomial distribution:

$$D_{x,t} \sim \mathcal{P}(E_{x,t}^c m_{x,t}) \quad \text{or} \quad D_{x,t} \sim \mathcal{B}(E_{x,t}^0 q_{x,t}),$$

• the systematic component: the effects of age x, calendar year t and year-of-birth (cohort) c = t - x are captured through a *predictor*  $\eta_{x,t}$  defined as:

$$\eta_{x,t} = \alpha_x + \sum_i \beta_x^i \kappa_t^i + \beta_x^0 \gamma_{t-x}.$$

• The *link function g* associating the random and the systematic components:

$$g\left(\mathbb{E}\left(\frac{D_{x,t}}{E_{x,t}}\right)\right) = \eta_{x,t}.$$

• The set of parameter constraints: as most stochastic models are not identifiable, some constraints may be needed to ensure uniqueness of the parameters  $\alpha_x$ ,  $\kappa_t$ ,  $\gamma_{t-x}$ .

## Generalized Age Period Cohort framework - current limitations

Designed for a single-population

- Univariate Poisson or Binomial assumption;
- Multi-populations models are built based on a 2-steps procedure;
- Fail to handle multivariate distributions (e.g. for Causes-of-deaths modelling).

Relies on Maximum Likelihood Estimation (MLE)

- Involves Iterative Weighted Least Squares algorithm (no closed-form formulas);
- Can be computationally intensive.

## Leveraging categorical variables

- Assumptions: age and time are two categorical variables with dummy structure age ranges in [x<sub>1</sub>; x<sub>d1</sub>] and is encoded by (z<sub>x</sub><sup>(1)</sup>)<sub>x∈[x<sub>1</sub>:x<sub>d1</sub>]</sub>: z<sub>x<sub>j</sub><sup>(1)</sup></sub> = 1<sub>x=x<sub>j</sub></sub>; time ranges in [t<sub>1</sub>; t<sub>d2</sub>] and is encoded by (z<sub>x</sub><sup>(2)</sup>)<sub>r∈[t<sub>1</sub>:t<sub>d</sub>]</sub>: z<sub>t<sub>k</sub><sup>(2)</sup></sub> = 1<sub>t=t<sub>k</sub></sub>;
- Systematic component  $\eta_{x,t}$  can be rewritten as

$$\begin{split} \eta_{x,t} &= \theta_0 + \sum_{x' = x_1 \dots x_{d_1}} z_{x'}^{(1)} \theta_{x'} + \sum_{t' = t_1 \dots t_{d_2}} z_{t'}^{(2)} \theta_{t'} & \text{ intercept and single effect} \\ &+ \sum_{x' = x_1 \dots x_{d_1}} \sum_{t' = t_1 \dots t_{d_2}} z_{x'}^{(1)} z_{t'}^{(2)} \theta_{x',t'} & \text{ double effect.} \end{split}$$

$$\theta = (\theta_0, \theta_{x_1}, \dots, \theta_{x_{d_1}}, \theta_{t_1}, \dots, \theta_{t_{d_2}}, \theta_{1,1}, \dots, \theta_{x_{d_1}t_{d_2}}) \text{ parameters vector}$$

## Leveraging categorical variables

#### Theorem

Following results from [BDR20, BDR22], we derive the alternative estimators:

 $\tilde{\theta} = \left( Q^{\mathsf{T}} Q + R^{\mathsf{T}} R \right)^{-1} Q^{\mathsf{T}} \tilde{\eta},$ 

- ${\it Q}$  is the unique matrix such that  $\eta = {\it Q}\theta$  ;
- R a contrast matrix that ensures identifiability,  $R\theta = 0$  ;
- $\tilde{\eta}$  is an estimator of  $g(\mathbb{E}[D_{x,t}])$ .

#### Corollary

If  $D_{x,t} \sim \mathcal{B}(E_{x,t}^0, q_{x,t})$  with a logit link function, we have the single categorical estimator:

$$\forall x_j, t_k, \quad \tilde{\eta}_{x_j, t_k} = \text{logit}\left(\overline{q^{(x_j, t_k)}}\right), \quad \overline{q^{(x_j, t_k)}} = \frac{1}{m_{j,k}} \sum_x \sum_t q_{x,t} z_{x_j} z_{t_k}, \quad m_{j,k} = \sum_x \sum_t z_{x_j} z_{t_k}.$$

Example with x= 50..51 and t = 2023..2024 ( $d_1 = d_2 = 2$ ):

$$\eta = \begin{pmatrix} \eta_{50,2023} \\ \eta_{50,2024} \\ \eta_{51,2023} \\ \eta_{51,2024} \end{pmatrix}, \quad Q = \begin{pmatrix} 1 & 1 & 0 & 1 & 0 & 1 & 0 & 0 & 0 \\ 1 & 0 & 1 & 1 & 0 & 0 & 1 & 0 & 0 \\ 1 & 1 & 0 & 0 & 1 & 0 & 0 & 1 & 0 \\ 1 & 0 & 1 & 0 & 1 & 0 & 0 & 0 & 1 \end{pmatrix},$$

## Identifying classical models - APC (M3) model

We directly identify the coefficients of the M3 model, as  $\gamma_{t-x}$  play the same role as double effect coefficients  $\theta_{x,t}$ :

$$\eta_{x,t} = \alpha_x + \kappa_t + \gamma_{t-x} \quad (M3).$$

Example with  $d_1 = d_2 = 2$ :

$$Q_{M3} = \begin{array}{ccccc} x, t & \alpha_{x_1} & \alpha_{x_2} & \kappa_{t_1} & \kappa_{t_2} & \gamma_{c_{-1}} & \gamma_{c_0} & \gamma_{c_1} \\ x_1, t_1 \\ x_2, t_1 \\ x_1, t_2 \\ x_2, t_2 \end{array} \begin{pmatrix} 1 & 0 & 1 & 0 & 0 & 1 & 0 \\ 0 & 1 & 1 & 0 & 1 & 0 & 0 \\ 1 & 0 & 0 & 1 & 0 & 0 & 1 \\ 0 & 1 & 0 & 1 & 0 & 1 & 0 \end{pmatrix}$$
  

$$\begin{array}{c} constraints & \alpha_{x_1} & \alpha_{x_2} & \kappa_{t_1} & \kappa_{t_2} & \gamma_{c_{-1}} & \gamma_{c_0} & \gamma_{c_1} \\ \\ constraints & \alpha_{x_1} & \alpha_{x_2} & \kappa_{t_1} & \kappa_{t_2} & \gamma_{c_{-1}} & \gamma_{c_0} & \gamma_{c_1} \\ \\ \end{array}$$
  

$$\begin{array}{c} R_{M3} = \sum_{c} \gamma_{c} = 0 \\ \sum_{c} c \gamma_{c} = 0 \\ \sum_{c} c \gamma_{c} = 0 \end{array} \begin{pmatrix} 0 & 0 & 1 & 1 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 1 & 1 & 1 \\ 0 & 0 & 0 & 0 & c_{-1} & c_0 & c_1 \end{pmatrix}$$

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## Identifying classical actuarial mortality models

We identify as well the coefficients of the M5 model:

$$\eta_{x,t} = \kappa_t^{(1)} + \kappa_t^{(2)}(x - \bar{x})$$
 (M5).

Example with  $d_1 = d_2 = 2$ :

$$egin{aligned} & x,t & \kappa_{t_1}^{(1)} & \kappa_{t_2}^{(1)} & \kappa_{t_1}^{(2)} & \kappa_{t_2}^{(2)} \ & x_{1,t_1} & \left( egin{aligned} 1 & 0 & x_1 - ar{x} & 0 \ 1 & 0 & x_2 - ar{x} & 0 \ 1 & 0 & x_1 - ar{x} \ & x_2,t_1 & 0 & 1 & 0 & x_1 - ar{x} \ & 0 & 1 & 0 & x_2 - ar{x} \ \end{pmatrix}, & R_{M5} = 0. \end{aligned}$$

And similarly for

$$\eta_{x,t} = \kappa_t^{(1)} + \kappa_t^{(2)}(x - \bar{x}) + \gamma_{t-x}^{(3)}$$
(M6)

$$\eta_{x,t} = \kappa_t^{(1)} + \kappa_t^{(2)}(x - \bar{x}) + \kappa_t^{(3)}[(x - \bar{x})^2 - \hat{\sigma}_x^2] + \gamma_{t-x}^{(3)}$$
(M7).

#### Numerical examples

- Country: France
- Age band: 50-84
- Period: 2001-2020
- Sex: Females
- $\bullet$  Estimators: Maximum Likelihood Estimators (from StMoMo package) Vs Alternative Estimators  $\tilde{\theta}$
- Source: Human Mortality Database (INSEE)

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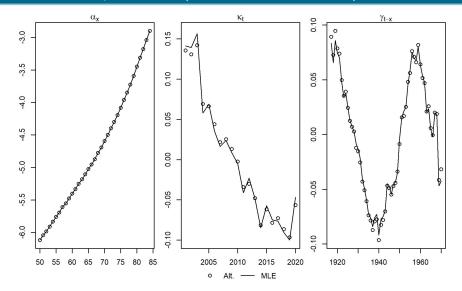
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## Numerical examples - Females, 50-84, 2001-2020

	Deviance	AIC	MSE	MAE	MAPE (%)
MLE.M3	8.41E+02	7.86E+03	6.85E-08	1.74E-04	1.75
Alt.M3	9.29E+02	7.94E+03	1.13E-07	1.99E-04	1.73
MLE.M5	5.06E+04	5.75E+04	6.20E-06	1.61E-03	1.42
Alt.M5	6.53E+04	7.22E+04	1.36E-05	1.93E-03	1.22
MLE.M6	1.84E+03	8.83E+03	1.17E-07	2.43E-04	2.70
Alt.M6	2.26E+03	9.24E+03	2.87E-07	3.11E-04	2.62
MLE.M7	7.37E+02	7.76E+03	6.30E-08	1.68E-04	1.63
Alt.M7	7.50E+02	7.77E+03	7.64E-08	1.76E-04	1.62

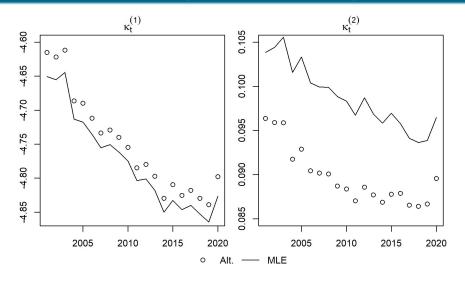
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## Numerical examples - M3 fit (Females, 50-84, 2001-2020)



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## Numerical examples - M5 fit (Females, 50-84, 2001-2020)







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## Possibility to build multi-populations models

We try here a multi-population model following ideas from [LL05]. Two populations indexed with  $g_1$  and  $g_2$ , with linear systematic component (here M3) and a common factor K(t) such that

$$\begin{cases} \eta_{\mathsf{x},t,\mathsf{g}_1} = \alpha_{\mathsf{x},\mathsf{g}_1} + \mathcal{K}(t) + \kappa_{t,\mathsf{g}_1} + \gamma_{t-\mathsf{x},\mathsf{g}_1} \\ \eta_{\mathsf{x},t,\mathsf{g}_2} = \alpha_{\mathsf{x},\mathsf{g}_2} + \mathcal{K}(t) + \kappa_{t,\mathsf{g}_2} + \gamma_{t-\mathsf{x},\mathsf{g}_2} \end{cases} \end{cases}$$

Problem equivalent to

- Adding a third categorical variable for country;
- Adding on observation of a country g which is the sum of the sub-populations g<sub>1</sub> and g<sub>2</sub>;
- Reference levels of  $\alpha_{x,g}$  and  $\gamma_{t-x,g}$  set to 0.

## Possibility to build multi-populations models

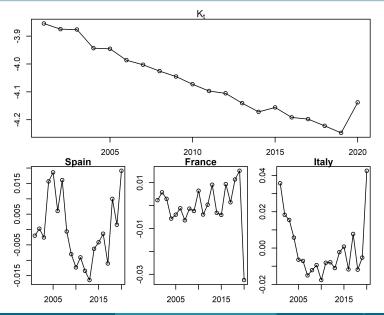
## Example with $d_1 = d_2 = 2$ :

$$Q = \begin{pmatrix} \theta_{g_1} & \theta_{g_2} & K_{t_1} & K_{t_2} \\ g_1 \begin{pmatrix} 0 & 1 & 0 \\ 0 & 1 & 0 \\ Q_{M3} & 0 & 0 & 1 \\ 0 & 0 & 1 \\ 0 & 0 & 1 \\ 0 & 0 & 1 & 0 \\ 0 & Q_{M3} & 0 & 1 \\ 0 & 0 & 1 & 0 \\ 0 & 0 & 1 & 0 \\ 0 & 0 & 1 & 0 \\ 0 & 0 & 1 & 0 \\ 0 & 0 & 0 & 1 \\ 0 & 0 & 0 & 1 \end{pmatrix}, \quad R = \begin{pmatrix} R_{M3} & 0 & 0 & 0 \\ 0 & R_{M3} & 0 & 0 \end{pmatrix}.$$

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## Possibility to build multi-populations models





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## Multivariate GAPC framework

Generalization of the GAPC framework based on multinomial distribution:

• the random component: The random vector of death counts  $D_{x,t}$  is assumed to follow a multinomial distribution

$$\left(D_{x,t}^{(1)},\ldots,D_{x,t}^{(d)}\right)\sim \mathcal{M}_d\left(E_{x,t}^0,\boldsymbol{q}_{x,t}=\left(q_{x,t}^{(1)},\ldots,q_{x,t}^{(d)}\right)\right).$$

• the systematic component: the effects of age x, calendar year t and year-of-birth (cohort) c = t - x are captured through a *predictor*  $\eta_{x,t}$  defined as

$$\boldsymbol{\eta}_{x,t} = (\eta_{x,t}^{(1)}, \dots, \eta_{x,t}^{(d)}), \quad \eta_{x,t}^{(j)} = \alpha_x^{(j)} + \sum_i \beta_x^{i,(j)} \kappa_t^{i,(j)} + \beta_x^{(0)} \gamma_{t-x}^{(j)}.$$

The formulation of the predictor may differ for each cause j.

• The *link function g* associating the random and the systematic components is the canonical link

$$g\left(\mathbb{E}\left(\frac{\boldsymbol{D}_{x,t}}{\boldsymbol{E}_{x,t}^{0}}\right)\right) = \boldsymbol{\eta}_{x,t}, \quad \text{with} \quad g: \boldsymbol{p} = (\boldsymbol{p}_{1}, \dots, \boldsymbol{p}_{d}) \rightarrow \left(\log\frac{\boldsymbol{p}_{1}}{\boldsymbol{p}_{d}}, \dots, \log\frac{\boldsymbol{p}_{d-1}}{\boldsymbol{p}_{d}}, 0\right).$$

• The *set of parameter constraints*: as most stochastic models are not identifiable, some constraints may be needed to ensure uniqueness of the estimates.

## Leveraging categorical variables for multinomial distribution

In multivariate context, we show that the alternative estimators still holds:

$$\tilde{\theta} = (Q^{\mathsf{T}}Q + R^{\mathsf{T}}R)^{-1} Q^{\mathsf{T}}\tilde{\eta}.$$

#### Theorem (from [BD24])

If  $\left(D_{x,t}^{(1)},\ldots,D_{x,t}^{(d)}\right) \sim \mathcal{M}_d\left(E_{x,t}^0, \boldsymbol{q}_{x,t} = \left(q_{x,t}^{(1)},\ldots,q_{x,t}^{(d)}\right)\right)$  with canonical link function, we have the single categorical estimator:

$$orall x_j, t_k, orall t \in [1; d], \quad ilde{\eta}_{x_j, t_k}^{(l)} = \log\left(rac{\overline{q_l^{x_j, t_k}}}{1 - \sum\limits_{l=1}^d \overline{q_l^{x_j, t_k}}}
ight), \quad \overline{q_l^{x_j, t_k}} = rac{1}{m_{j,k}} \sum_x \sum_t q_{x,t}^{(l)} z_{x_j} z_{t_k}.$$

## Numerical example

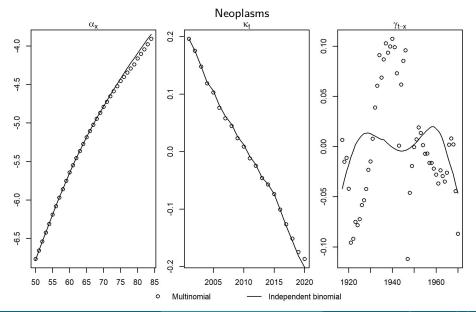
- Country: USA
- Age band: 50-84
- Period: 2001-2020
- Sex: Males
- Source: Human Mortality Database
- For each cause, M3 model with cause-specific parameters

$$\eta_{x,t}^{(j)} = \alpha_x^{(j)} + \kappa_t^{(j)} + \gamma_{t-x}^{(j)}.$$

 Estimators: comparison between univariate independent single population VS multinomial assumption.

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# Numerical Example - US Males 50-84 2001-2020

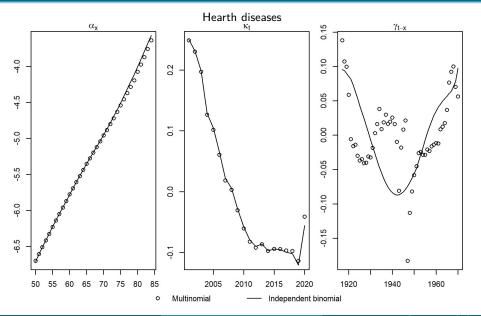




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## Numerical Example - US Males 50-84 2001-2020



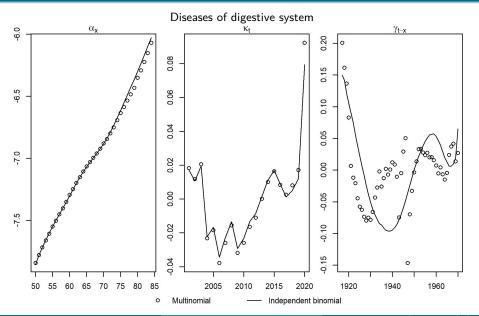
MGLM - CoD mortality



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# Numerical Example - US Males 50-84 2001-2020





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## Numerical Example - US Males 50-84 2001-2020

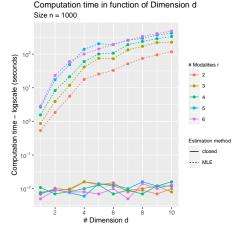
Parameters	Deviance	AIC	MSE	MAE	MAPE (%)
Independent univariate	11883	107342	2.55E-07	1.58E-04	7,03%
Multivariate	5149	97172	8.88E-09	2.56E-05	2.14%

Life Expectancy Comparison (all-causes)

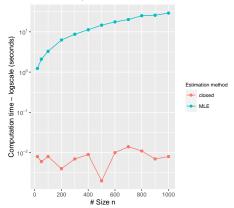
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## Computation time - closed form vs MLE for multinomial distribution



Computation time in function of Size n Dimension d = 5, Number of modalities r = 3



## Key takeaways

Objectives achieved:

- One-step framework for multivariate models (e.g. causes-of-death);
- Unified and flexible framework for mortality modelling;
- Faster computation time, especially in high dimensions.

Limits:

- Only linear models Lee-Carter model is out of scope due to the bilinear term  $\beta_{\rm X}\kappa_t$ ;
- Not MLE in general but not necessarily worse performance, depending on the metrics chosen.

Future work:

- Compare to other causes-of-deaths mortality models;
- Implement LeCam one-step procedure to improve performance of non-MLE estimators;
- Assess performance in forecasting;
- Parameters clustering with GLM-trees.

	Disease	ICD10
1	Certain infectious diseases	A00-B99
2	Neoplasms	C00-D48
3	Diseases of the blood and blood-forming organs	D50-D89
4	Endocrine, nutritional and metabolic diseases	E00-E88
5	Mental and behavioural disorders	F01-F99
6	Diseases of the nervous system and the sense organs	G00-G44, G47-H93
7	Heart diseases	100-151
8	Cerebrovascular diseases	G45, 160-169
9	Other and unspecified disorders of the circulatory system	170-199, K64
10	Acute respiratory diseases	J00-J22, U04, U07
11	Other respiratory diseases	J30-J98
12	Diseases of the digestive system	K00-K63, K65-K92
13	Diseases of the skin	L00-M99
14	Diseases of the genitourinary system	N00-O99
15	Certains conditions originating in the perinatal period	
16	External causes	V01-Y89
-		

- Villegas Andres, Pietro Millossovich, Kaishev Vladimir, et al., *Stmomo: Stochastic mortality modeling in r*, Journal of Statistical Software **84** (2018), no. 3, 1–38.
- Antoine Burg and Christophe Dutang, *Closed-form estimators for multivariate regressions models a single categorical variable approach*, submitted (2024).
- Alexandre Brouste, Christophe Dutang, and Tom Rohmer, *Closed-form maximum likelihood* estimator for generalized linear models in the case of categorical explanatory variables: application to insurance loss modeling, Computational Statistics **35** (2020), 689–724.
  - \_\_\_\_\_, A closed-form alternative estimator for glm with categorical explanatory variables, Communications in Statistics-Simulation and Computation (2022), 1–17.



Nan Li and Ronald Lee, *Coherent mortality forecasts for a group of populations: An extension of the lee-carter method*, Demography **42** (2005), 575–594.