

6A – Aging Measurement and Mortality Modeling 2

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Short- and Long-Term Dynamics of Cause-specific Mortality Rates using the Cointegration Analysis

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Figure: Total age-standardized mortality rates, US males, WHO

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 \rightarrow International Classification of Diseases

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Figure: Cause-specific age-standardized mortality rates, US males, WHO

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Figure: Cause-specific age-standardized mortality rates, US males, WHO

 \rightarrow Build a model which preserves the information on different causes of death



Figure: Cause-specific age-standardized mortality rates, US males, WHO

 \rightarrow Build a model which preserves the information on different causes of death \rightarrow and takes into account their dependency.

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Datasets



(World Health Organization database)

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Datasets



(World Health Organization database)

 \rightarrow Identify patterns/trends and compare them across datasets

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$$\mathbf{y}_{t} = \begin{pmatrix} log(m_{t}^{IP}) \\ log(m_{t}^{Cancer}) \\ log(m_{t}^{Circulatory}) \\ log(m_{t}^{Respiratory}) \\ log(m_{t}^{External}) \end{pmatrix}$$

▶ Data dimenstion: n = 5, t = 1, ..., 55-65 years

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- Cointegration relation = linear combination of non-stationary variables that is stationary
- Information on the long-term equilibrium between the causes

Error Correction Model representation for the cause-specific mortality rates vector \mathbf{y}_t :

$$\Delta \mathbf{y}_{t} = \mathbf{c} + \mathbf{dt} + \alpha \beta' \mathbf{y}_{t-1} + \sum_{n=1}^{p} \mathbf{\Gamma}_{i} \Delta \mathbf{y}_{t-i} + \epsilon_{t}$$

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Equation valid only if αβ'y_{t-1} is stationary

- β is a matrix of rank r where r is the number of cointegration relations
- $\blacktriangleright \alpha$ is a loading matrix

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• Equation valid only if $\alpha \beta' \mathbf{y}_{t-1}$ is stationary

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• Long-term dynamics via the cointegrated term $\alpha\beta'\mathbf{y}_{t-1}$

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Short-term dynamics via the matrices Γ_i

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Algorithm

Following procedure was applied to every dataset :

- Test formally for unit roots
- Test for the number of lags in a vector ECM
- Test for the number of cointegration relations r and the form of the deterministic elements (Johansen, 1994)
- Calculate the matrices α, β and Γ_i
- Check the residuals for autocorrelation and normality

Example of a vector ECM - US males

The following model was chosen as the best describing the data:

 $\Delta \mathbf{y}_t = \mathbf{c} + \mathbf{dt} + \mathbf{\Gamma}_1 \Delta \mathbf{y}_{t-1} + \alpha \beta' \mathbf{y}_{t-1} + \epsilon_t =$

$$= \begin{bmatrix} -2.772 \\ -0.150 \\ -0.243 \\ -2.057 \\ 0.156 \end{bmatrix} + \begin{bmatrix} 0.010 \\ 0.000 \\ 0.005 \\ 0.000 \end{bmatrix} \mathbf{t} + \begin{bmatrix} -0.121 & -0.707 & -0.188 & 0.174 & 0.323 \\ -0.004 & 0.015 & -0.166 & -0.008 & 0.133 \\ -0.043 & -0.121 & 0.034 & -0.089 & 0.196 \\ -0.135 & -0.294 & -0.085 & -0.381 & 1.119 \\ 0.042 & -0.323 & 0.198 & -0.146 & 0.227 \end{bmatrix} \Delta \mathbf{y}_{t-1} + \mathbf{e}_t$$

Significant coefficients (at 5% significance level) are in bold.

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Impulse-response analysis (1)

- What is the response of every cause-specific mortality rate to a shock from a particular cause (e.g. IP)?
- \rightarrow Vector of starting values: $\mathbf{y}_0 = (\text{sdt.dev}(\text{IP}), 0, 0, 0, 0)$



ightarrow All causes show weak reaction to the shock from IP mortality rate

Impulse-response analysis (2)

Alternative way: how sensitive is a particular cause-specific mortality rate (e.g. IP) to a random shock from other causes (one at a time)?



 $\rightarrow\,$ IP mortality rate shows important reaction to the shocks from every other cause

Short- vs. long-term dynamics

What drives this behavior of the IP mortality rate?

- Not the short term: the corresponding coefficients in the F₁ matrix are not significant
- Long-term: the cointegration relation enters the equation for the IP mortality rate with a significant coefficient α₁

Overview : impulse-response analysis

- Same procedure applied to the rest of the datasets
- High-level summary of responses of the mortality rate Y to the shock given to the rate X:

$X\setminusY$	IP	Cancer	Circulatory	Respiratory	External
IP		Low	Low	Low	Low
Cancer	High		Med	High	High
Circulatory	High	Low		High	Med
Respiratory	Med	Low	Low		Low
External	High	Low	Low	High	

Overview: the short term

 $\pmb{\Gamma}_1$ coefficients which are significantly different from zero, significance level of 5%:

Dataset	ΔIP_t	$\Delta Canc_t$	$\Delta Circ_t$	$\Delta Resp_t$	$\Delta E x t_t$
US/M	-	Circ,Ext	-	Resp,Ext	Resp
JP/M	-	Canc	Resp	Circ,Resp	-
FR/M	-	-	Resp	Circ,Resp	Circ,Resp
E&W/M	Ext	-	-	IP,Circ, Resp	Ext
AU/M	IP	Canc, Ext	Circ	-	Ext
US/F	Canc,Ext	Ext	Canc	Resp	Canc,Resp
JP/F	IP,Resp	-	Resp	-	Circ,Resp,Ext
FR/F	Canc,Circ	-	-	Canc,Circ,Resp	IP
E&W/F	-	-	Resp	Resp	Resp
AU/F	IP,Resp	Canc	Circ	Circ,Resp	-

Overview: the long term

Equations to which the long-term component (i.e., cointegration relation) enters with a coefficient α_i significantly different from zero, significance level of 5%:

Country		Males	Females	
US	α_i	$\Delta IP_t, \Delta Resp_t$	$\Delta IP_t, \Delta Canc_t, \Delta Circ_t, \Delta Resp_t$	
JP	α_{1i}	$\Delta IP_t, \Delta Canc_t$	$\Delta Circ_t, \Delta Resp_t$	
	α_{2i}	$\Delta IP_t, \Delta Canc_t, \Delta Resp_t$	ΔIP_t , $\Delta Canc_t$, $\Delta Circ_t$, $\Delta Resp_t$	
FR	α_i	$\Delta IP_t, \Delta Canc_t, \Delta Resp_t$	-	
	α_i	-	ΔIP_t , $\Delta Canc_t$, $\Delta Resp_t$	
E&W	α_i	$\Delta Canc_t, \Delta Resp_t, \Delta Ext_t$	$\Delta IP_t, \Delta Circ_t, \Delta Ext_t$	
AU	α_i	-	ΔIP_t , $\Delta Circ_t$, ΔExt_t	
	α_i	ΔIP_t , $\Delta Circ_t$, $\Delta Resp_t$	-	

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In the short term

- Development of Circulatory, Respiratory, and External mortality rates depends on other cause-specific mortality rates;
- ▶ IP and Cancer mortality rates seem to be less impacted by other causes.

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In the long term

- IP and Respiratory mortality rates are the most impacted by the cointegration relation;
- Cancer and Circulatory mortality rates are less impacted;
- External causes seem to be totally independent from it.

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Objective

Set more informed assumptions on the future development of mortality.

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Objective

- Set more informed assumptions on the future development of mortality. Next steps
 - Study common stochastic trends shared by the cause-specific mortality rates.
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Thank you for your attention!

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2020 Living to 100 Symposium Rotation of the Age Pattern of Mortality Improvements in EU Member States Péter Vékás, Ph.D. (Corvinus University of Budapest)

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Introduction

- If human lifetimes are three years longer than expected (in line with underestimations in the past), costs of aging will increase by 50% of GDP in advanced economies and 25% of GDP in emerging economies (IMF, 2012).
- Mortality forecasting is invaluable for actuaries.
- Model assumptions may make a huge difference in the long run!



Contents

- 1. Time-invariance assumption of the Lee–Carter (1992) model
- 2. Rotation of the age pattern of mortality improvements (Li–Lee–Gerland, 2013)
- 3. Proposed methodology to assess rotation (Vékás, 2019)
- 4. Results on EU data



Lee-Carter (1992) model

- "Gold standard" of mortality forecasting.
- Log-mortality rates modeled as $\ln m_{xt} = a_x + b_x k_t + \varepsilon_{xt}$.
- As k_t declines over time, the coefficients b_x regulate the rates of improvement by age.
- •Age-specific improvement rates b_x assumed to be independent of time (not b_{xt})!



Rotation

- Rotation of the age pattern of mortality improvements (Li-Lee-Gerland, 2013): mortality improvements tend to slow down in younger ages and speed up in older ages.
- Possible reasons:
 - little room left for spectacular advances in preventing child mortality,
 - improved, costly medical technology to cope with serious illness and extend life.



Illustration





Rotation in theory and practice

- Plenty of sporadic evidence for rotation in the literature, mostly based on *ad hoc* methods.
- •Li–Lee–Gerland (2013) have created the LC model including rotation (LCR variant).
- Applied by Vékás (2018) on Hungarian data to see the long-term impact of rotation.



Rotation of b_x in LCR model (Hungary, 2018-2100)





Projection with and without rotation (Hungary, 2018-2100)









To rotate or not to rotate?

- Forecasts ignoring rotation systematically underestimate longevity risk!
- Errors may be tremendous in the long run.
- It is crucial to assess whether there is rotation.
- •Vékás (2019) proposes a methodology to measure and statistically test rotation and applies it on historical data of 28 EU countries.



Mortality improvement rates and acceleration rates

- Mortality improvement rates (x: age, t: period, c: country, g: gender) $r_{xt}^{cg} = -\log\left(\frac{m_{x,t+1}^{cg}}{m_{xt}^{cg}}\right)$
- •Long-term acceleration = slope of linear trend of mortality improvement rates over time:

$$\beta_x^{cg} = \frac{\sum_{t=1}^{12} (r_{xt}^{cg} - \bar{r}_x^{cg})(t - \bar{t})}{\sum_{t=1}^{12} (t - \bar{t})^2}$$



Strength of rotation

 Measured by Spearman's ρ between acceleration and age, weighted by population sizes of age groups:

$$\rho^{cg} = \frac{\sum_{i=1}^{22} P_{x_i}^{cg} (\operatorname{rank}(\beta_{x_i}^{cg}) - \mu^{cg}) (i - v^{cg})}{\sqrt{\sum_{i=1}^{22} P_{x_i}^{cg} (\operatorname{rank}(\beta_{x_i}^{cg}) - \mu^{cg})^2} \sqrt{\sum_{i=1}^{22} P_{x_i}^{cg} (i - v^{cg})^2}} ((c, g) \in \{c_1, c_2, \dots, c_{28}\} \times \{M, W\})$$
$$\mu^{cg} = \frac{\sum_{i=1}^{22} P_{x_i}^{cg} \operatorname{rank}(\beta_{x_i}^{cg})}{\sum_{i=1}^{22} P_{x_i}^{cg}} \quad \text{and} \quad v^{cg} = \frac{\sum_{i=1}^{22} P_{x_i}^{cg} i}{\sum_{i=1}^{22} P_{x_i}^{cg}}$$



Data

- •UN World Population Prospects 2017
- Mortality rates, life expectancies at birth and population counts
- •22 age groups, both genders
- •13 periods (1950–1955 up to 2010–2015)
- •28 member states of the EU



Strength of rotation by country and gender





Strongest rotation: Cyprus (earliest vs. latest periods)



Potential predictors of rotation

- Gender
- Former political bloc (East or West)
- e_0 (Li-Lee-Gerland, 2013: rotation is more prevalent in low-mortality countries)
- Possibly also e_{60} , or improvement of e_0 between 1950 and 2015



Impact of gender and political bloc

- Significantly more rotation in women's data!
- More rotation in the former Eastern bloc, but difference is not significant.

Gender (region)	$\rho^b(g)$	<i>p</i> -value	
Men (EU)	0.246		
Women (EU)	0.498		
Gender difference (EU)	-0.252	0.001	**
Men (West)	0.199	n	
Women (West)	0.466		
Gender difference (West)	-0.267	0.016	*
Men (East)	0.43		
Women (East)	0.619		
Gender difference (East)	-0.189	0.011	*
Regional difference (Men)	-0.231	0.202	
Regional difference (Women)	-0.153	0.396	



Rotation only correlated with e_0 **in the East**





Conclusions

- •New, simple, data-driven methodology to assess rotation and its relationships with other variables.
- •Rotation is far from universal: only in some member states.
- More rotation in women's data.



Conclusions

- •Somewhat more in rotation in former Eastern bloc.
- Only related to e_0 in the East.
- Methodology may be used to decide whether to use LC model or LCR variant.



Thank you very much for your attention!



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