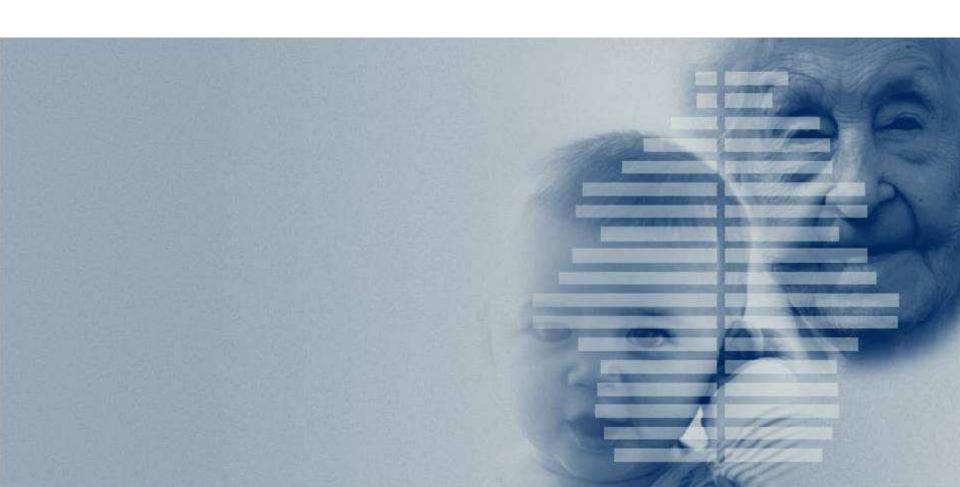


MAX-PLANCK-INSTITUT FÜR DEMOGRAFISCHE FORSCHUNG

MAX PLANCK INSTITUTE FOR DEMOGRAPHIC RESEARCH





RESEARCH

MAX-PLANCK-INSTITUT FÜR DEMOGRAFISCHE FORSCHUNG

Healthy Life Expectancy,
Mortality,
and Age Prevalence of Morbidity
Alyson van Raalte, Tim Riffe



Expected life years with disability (DLY)

- Most often measured by Sullivan method
- DLY is sum product of lifetable survival L_x and morbidity age-specific prevalence π_x
- DLY can change because of changes in L_x or π_x



Expected life years with disability (DLY)

- Most often measured by Sullivan method
- DLY is sum product of lifetable survival L_x and morbidity age-specific prevalence π_x
- DLY can change because of changes in L_x or π_x



Expected life years with disability (DLY)

- Most often measured by Sullivan method
- DLY is sum product of lifetable survival L_x and morbidity age-specific prevalence π_x
- DLY can change because of changes in L_x or π_x

- Disability prevalence at each age
- Stock variable: slow to react to abrupt health innovations since it depends on past cohort experiences with sickness (Barendregt et al. 1994)
- Prevalence can vary by age, time-to-death, lifespan, or combinations of these things.
- This complicates comparisons of period DLY (or HLE) across populations with different mortality.
- Since π_x changes across mortality regimes, attributing between-population differences in DLY to mortality and morbidity is problematic.

- Disability prevalence at each age
- Stock variable: slow to react to abrupt health innovations since it depends on past cohort experiences with sickness (Barendregt et al. 1994)
- Prevalence can vary by age, time-to-death, lifespan, or combinations of these things.
- This complicates comparisons of period DLY (or HLE) across populations with different mortality.
- Since π_x changes across mortality regimes, attributing between-population differences in DLY to mortality and morbidity is problematic.

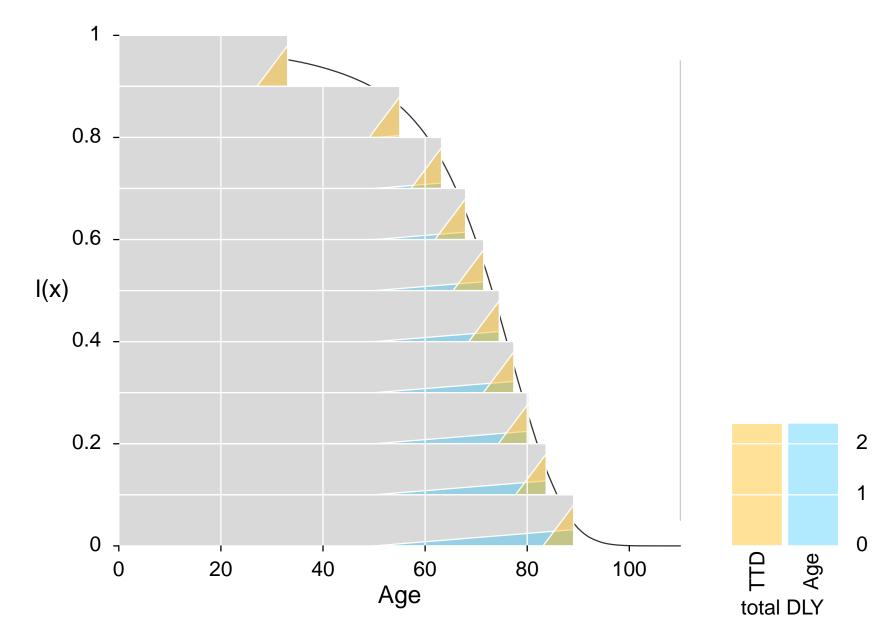
- Disability prevalence at each age
- Stock variable: slow to react to abrupt health innovations since it depends on past cohort experiences with sickness (Barendregt et al. 1994)
- Prevalence can vary by age, time-to-death, lifespan, or combinations of these things.
- This complicates comparisons of period DLY (or HLE) across populations with different mortality.
- Since π_x changes across mortality regimes, attributing between-population differences in DLY to mortality and morbidity is problematic.

- Disability prevalence at each age
- Stock variable: slow to react to abrupt health innovations since it depends on past cohort experiences with sickness (Barendregt et al. 1994)
- Prevalence can vary by age, time-to-death, lifespan, or combinations of these things.
- This complicates comparisons of period DLY (or HLE) across populations with different mortality.
- Since π_x changes across mortality regimes, attributing between-population differences in DLY to mortality and morbidity is problematic.

- Disability prevalence at each age
- Stock variable: slow to react to abrupt health innovations since it depends on past cohort experiences with sickness (Barendregt et al. 1994)
- Prevalence can vary by age, time-to-death, lifespan, or combinations of these things.
- This complicates comparisons of period DLY (or HLE) across populations with different mortality.
- Since π_x changes across mortality regimes, attributing between-population differences in DLY to mortality and morbidity is problematic.

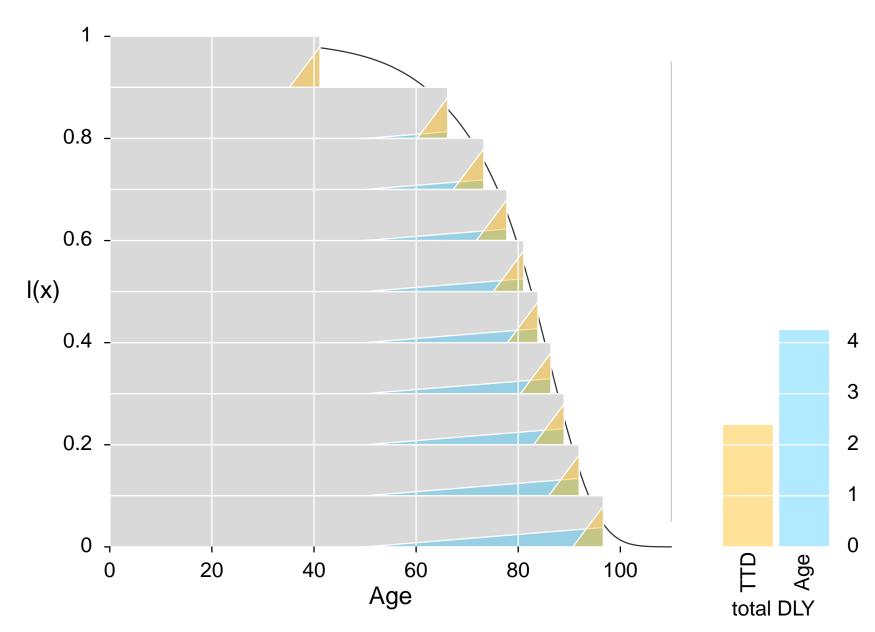


A simple illustration



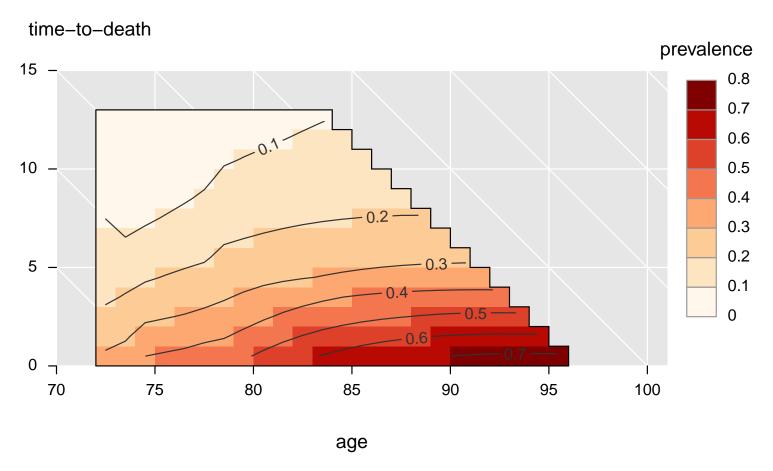


A simple illustration





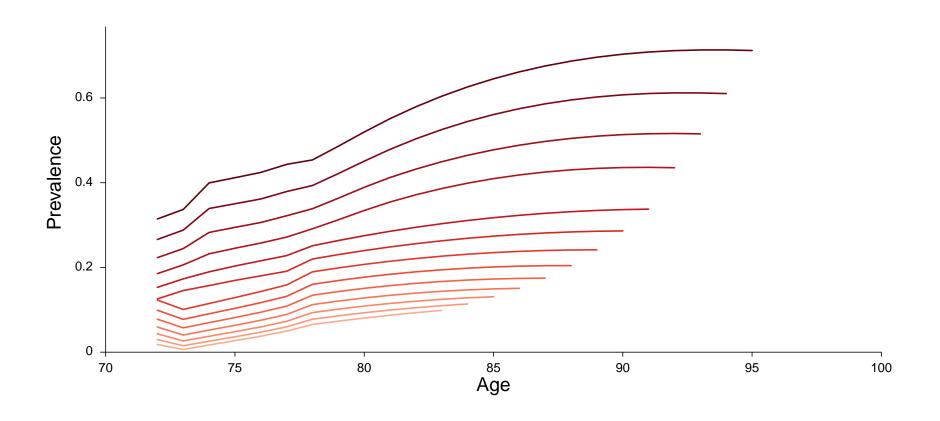
Disablity broken down by age and time to death



Proportion of USA males from the 1915-1919 cohort with at least 1 of 5 IADLs

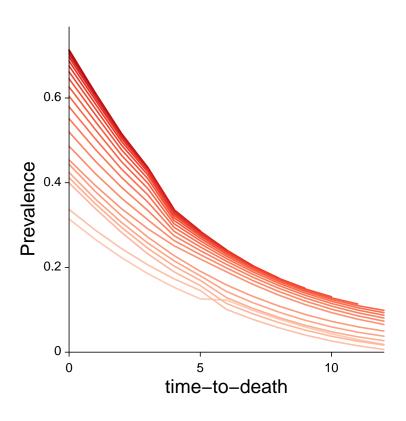


Disablity broken down by age and time to death



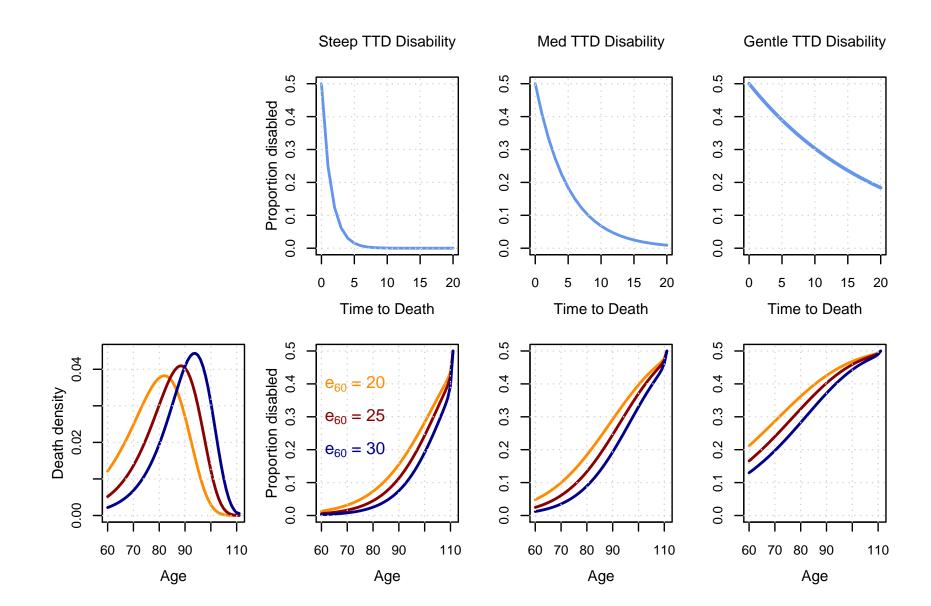


Disablity broken down by age and time to death





Proportion disabled by TTD and mortality level





- Are differences in DLY from mortality or morbidity?
- Decomposition methods isolate the effects of changes in L_x and changes in π_x
- These are considered as *mortality* and *morbidity* effects (Nusselder and Looman 2004, Andreev et al. 2002)
- Interpretation problem: mortality can change π_{\times} all by itself if disability is patterned by time-to-death



- Are differences in DLY from mortality or morbidity?
- Decomposition methods isolate the effects of changes in \mathcal{L}_{x} and changes in π_{x}
- These are considered as *mortality* and *morbidity* effects (Nusselder and Looman 2004, Andreev et al. 2002)
- Interpretation problem: mortality can change π_{\times} all by itself if disability is patterned by time-to-death



- Are differences in DLY from mortality or morbidity?
- Decomposition methods isolate the effects of changes in L_x and changes in π_x
- These are considered as mortality and morbidity effects (Nusselder and Looman 2004, Andreev et al. 2002)
- Interpretation problem: mortality can change π_{\times} all by itself if disability is patterned by time-to-death



- Are differences in DLY from mortality or morbidity?
- Decomposition methods isolate the effects of changes in L_x and changes in π_x
- These are considered as *mortality* and *morbidity* effects (Nusselder and Looman 2004, Andreev et al. 2002)
- Interpretation problem: mortality can change π_{\times} all by itself if disability is patterned by time-to-death



Estimating the upper magnitude of bias of morbidity differences from mortality decline

- Estimated average TTD profile for different disability types, based on USA HRS data, quinquennial cohorts 1905-1930
- Calculated apparent period age prevalence of morbidity for HMD countries had they experienced the US TTD morbidity
- Assumed all populations were stationary
- Decomposed differences between all population pairs in 1980, 1990, 2000 into apparent mortality and morbidity components
- Same for within-population changes over 10-year periods, 1950-2010



Estimating the upper magnitude of bias of morbidity differences from mortality decline

- Estimated average TTD profile for different disability types, based on USA HRS data, quinquennial cohorts 1905-1930
- Calculated apparent period age prevalence of morbidity for HMD countries had they experienced the US TTD morbidity
- Assumed all populations were stationary
- Decomposed differences between all population pairs in 1980, 1990, 2000 into apparent mortality and morbidity components
- Same for within-population changes over 10-year periods, 1950-2010



Estimating the upper magnitude of bias of morbidity differences from mortality decline

- Estimated average TTD profile for different disability types, based on USA HRS data, quinquennial cohorts 1905-1930
- Calculated apparent period age prevalence of morbidity for HMD countries had they experienced the US TTD morbidity
- Assumed all populations were stationary
- Decomposed differences between all population pairs in 1980, 1990, 2000 into apparent mortality and morbidity components
- Same for within-population changes over 10-year periods, 1950-2010



Estimating the upper magnitude of bias of morbidity differences from mortality decline

- Estimated average TTD profile for different disability types, based on USA HRS data, quinquennial cohorts 1905-1930
- Calculated apparent period age prevalence of morbidity for HMD countries had they experienced the US TTD morbidity
- Assumed all populations were stationary
- Decomposed differences between all population pairs in 1980, 1990, 2000 into apparent mortality and morbidity components
- Same for within-population changes over 10-year periods, 1950-2010

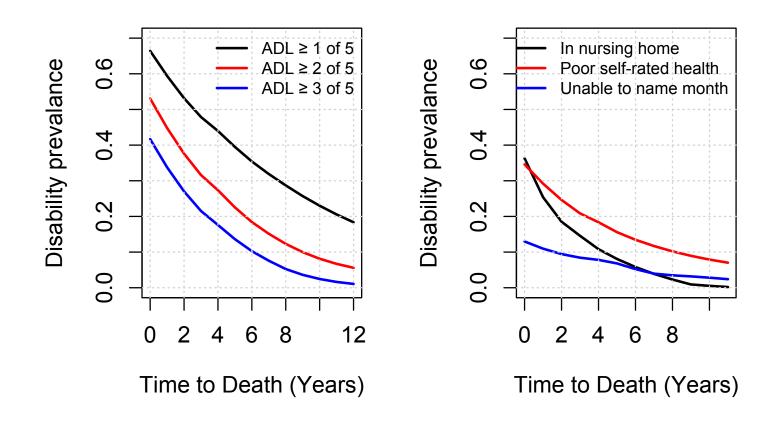


Estimating the upper magnitude of bias of morbidity differences from mortality decline

- Estimated average TTD profile for different disability types, based on USA HRS data, quinquennial cohorts 1905-1930
- Calculated apparent period age prevalence of morbidity for HMD countries had they experienced the US TTD morbidity
- Assumed all populations were stationary
- Decomposed differences between all population pairs in 1980, 1990, 2000 into apparent mortality and morbidity components
- Same for within-population changes over 10-year periods, 1950-2010

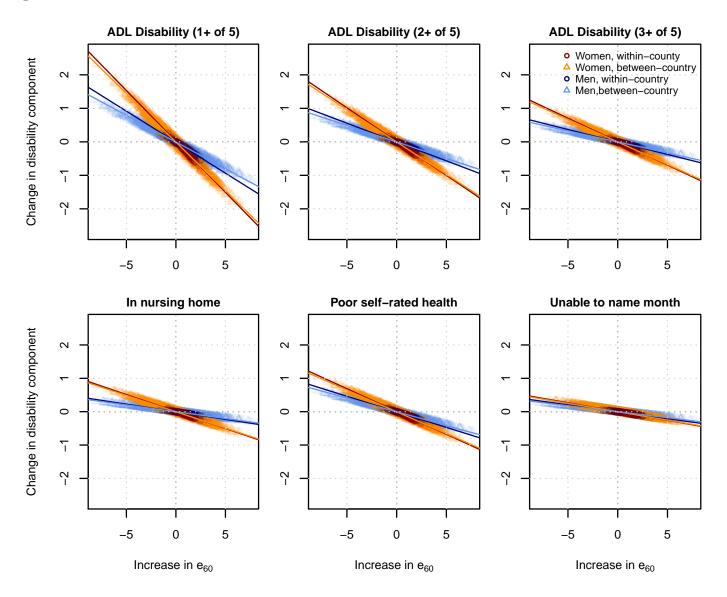


TTD disability prevalence for different disability types





Decomposition: Change in disability component





- True value of the change in disability component is zero by design
- Deviation is result of differences in mortality
- If *e*₆₀ increases by 5 years, up to 1 year of decrease in DLY attributed to disability component could be from decrease in mortality (Female ADL 3 or more)
- Departure from upper bound depends on patterns of π_x , how well US pattern applies, departure from stationarity.
- Different slopes partly from differences in final π_{\times} between disability types and the sexes



- True value of the change in disability component is zero by design
- Deviation is result of differences in mortality
- If *e*₆₀ increases by 5 years, up to 1 year of decrease in DLY attributed to disability component could be from decrease in mortality (Female ADL 3 or more)
- Departure from upper bound depends on patterns of π_x , how well US pattern applies, departure from stationarity.
- Different slopes partly from differences in final π_x between disability types and the sexes



- True value of the change in disability component is zero by design
- Deviation is result of differences in mortality
- If e₆₀ increases by 5 years, up to 1 year of decrease in DLY attributed to disability component could be from decrease in mortality (Female ADL 3 or more)
- Departure from upper bound depends on patterns of π_{\times} , how well US pattern applies, departure from stationarity.
- Different slopes partly from differences in final π_x between disability types and the sexes



- True value of the change in disability component is zero by design
- Deviation is result of differences in mortality
- If *e*₆₀ increases by 5 years, up to 1 year of decrease in DLY attributed to disability component could be from decrease in mortality (Female ADL 3 or more)
- Departure from upper bound depends on patterns of π_{\times} , how well US pattern applies, departure from stationarity.
- Different slopes partly from differences in final π_x between disability types and the sexes



- True value of the change in disability component is zero by design
- Deviation is result of differences in mortality
- If *e*₆₀ increases by 5 years, up to 1 year of decrease in DLY attributed to disability component could be from decrease in mortality (Female ADL 3 or more)
- Departure from upper bound depends on patterns of π_{\times} , how well US pattern applies, departure from stationarity.
- Different slopes partly from differences in final π_x between disability types and the sexes

Considerations

- Considering morbidity prevalence as a function of time to death does not imply that morbidity incidence is a time to death
- Modeling prevalence as TTD requires no specification of process
- In reality morbidity varies over both chronological age and time-to-death

Considerations

- Considering morbidity prevalence as a function of time to death does not imply that morbidity incidence is a time to death
- Modeling prevalence as TTD requires no specification of process
- In reality morbidity varies over both chronological age and time-to-death

Considerations

- Considering morbidity prevalence as a function of time to death does not imply that morbidity incidence is a time to death
- Modeling prevalence as TTD requires no specification of process
- In reality morbidity varies over both chronological age and time-to-death

- HLE or DLY provide an important snapshot of expected life years lived in good or poor health
- Difficulty in interpreting period differences in these quantities between populations
- Chronological age pattern of disability can change solely as a function of mortality change even when the underlying morbidity function is held constant
- Could partly explain why mortality levels and disability prevalence are related (Van Oyen et al. 2013, Luy and Minagawa 2014)

- HLE or DLY provide an important snapshot of expected life years lived in good or poor health
- Difficulty in interpreting period differences in these quantities between populations
- Chronological age pattern of disability can change solely as a function of mortality change even when the underlying morbidity function is held constant
- Could partly explain why mortality levels and disability prevalence are related (Van Oyen et al. 2013, Luy and Minagawa 2014)

- HLE or DLY provide an important snapshot of expected life years lived in good or poor health
- Difficulty in interpreting period differences in these quantities between populations
- Chronological age pattern of disability can change solely as a function of mortality change even when the underlying morbidity function is held constant
- Could partly explain why mortality levels and disability prevalence are related (Van Oyen et al. 2013, Luy and Minagawa 2014)

- HLE or DLY provide an important snapshot of expected life years lived in good or poor health
- Difficulty in interpreting period differences in these quantities between populations
- Chronological age pattern of disability can change solely as a function of mortality change even when the underlying morbidity function is held constant
- Could partly explain why mortality levels and disability prevalence are related (Van Oyen et al. 2013, Luy and Minagawa 2014)



Thanks!

