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Collecting and Analysing Data to Understand Population Health of Ageing Populations: A Czech Perspective

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We do not have sufficient, good-quality data to study and adapt to population ageing

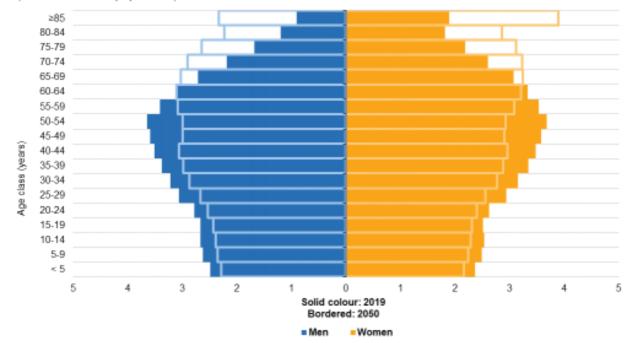


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Population Ageing: a Crisis or Opportunity?

Population pyramids, EU-27, 2019 and 2050

(% share of total population)



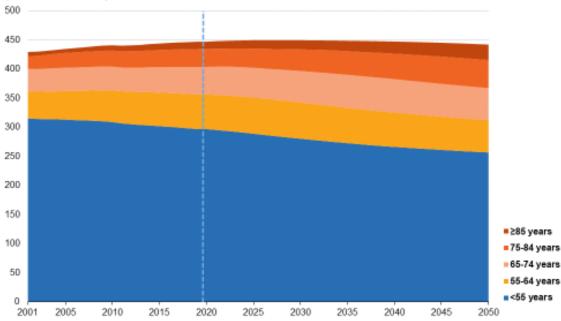
Note: all data as of 1 January. 2019: estimates and provisional. 2050: population according to the 2019 projections. baseline variant (EUROPOP2019).

Source: Eurostat (online data codes: demo plangroup and proj 19np)

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Population developments, by age class, EU-27, 2001-2050



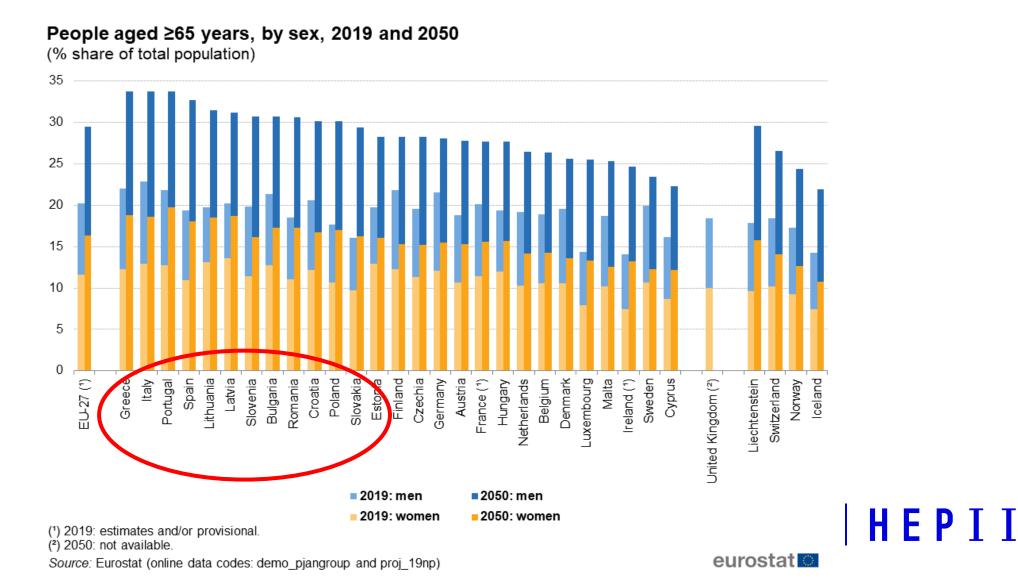


Note: all data as of 1 January. 2008, 2010-2012, 2014-2015 and 2017: breaks in series, 2019: provisional. 2020-2050: population according to the 2019 projections, baseline variant (EUROPOP2019). The vertical dotted line marks the divide between official historical data and EUROPOP2019 population projections. Source: Eurostat (online data codes: demo pjangroup and proj_19np)

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The share of people 65+ will grow from 20% to almost 30% in the next 20 years, a ~<u>50% increase!</u>



Data Collection of Interest (CZ case)

Administrative Health Data: Records from health services (e.g. hospital discharges, insurance claims), no real electronic health records outside of medication use registry.

Health Registries: Disease-specific registries (e.g. National Cancer Registry, other chronic disease registries).

Population Surveys: National and international surveys on health (EHIS, EU-SILC) and ageing (e.g. SHARE).

Long-Term Care (LTC) Data: Social sector statistics on home care, nursing homes, care allowance recipients.

Mortality & Morbidity Statistics: Vital statistics (life expectancy, causes of death) and illness incidence/prevalence from surveys and registries.

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Data Collection of Interest (CZ case)

Subjective Scoring

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Administrative/Claims & Registry Data

- National Health Information System (NHIS): Integrated system managed by ÚZIS (IHIS CZ).
 - Contains mandatory health claims covering the entire population, **limited use for research**
 - Example: the National Register of Hospitalised Patients captures every hospital stay nationwide
- Disease Registries
 - Long-established registries, e.g. the Czech National Cancer Registry (population-based since 1977)
 - Track incidence, prevalence, treatment and survival for major diseases (cancers, cardiovascular events, etc.)
 - Data from these registries inform epidemiologic research and policy
- **Vital Statistics:** Death Certificate Information System records all deaths and causes
 - Mortality data is of relatively high-quality and used for indicators like life expectancy and cause-specific death rates.
- Administrative Health Records: health insurance billing data (reimbursed services) and electronic health records, while collected, are very fragmented by provider.
 - They are not yet fully centralized for analysis, but contain rich information (diagnoses, prescriptions, visits) on patients' healthcare utilization.





Population Health Surveys & Studies

- European Health Interview Survey (EHIS)

- Czechia participates in EHIS to gather self-reported health status, chronic conditions, functional limitations, lifestyle, etc.,
- EHIS provides key indicators comparable across EU. However, EHIS excludes people in institutions.

- EU-SILC (EU Statistics on Income and Living Conditions)

- An annual household survey that includes some health questions (e.g. long-standing illnesses, self-care limitations).
- EU-SILC data are used to derive Healthy Life Years indicators. Interestingly, different survey contexts lead to different results in 2014, Czech Healthy Life Years at age 65 was 9 years for men via EU-SILC, but only ~5.7 years using EHIS data, due to higher reported disability in the dedicated health survey.
- Caveat: Like EHIS, it covers non-institutionalized population.

- SHARE (Survey of Health, Ageing and Retirement in Europe)

- Longitudinal panel of Europeans aged 50+. SHARE collects detailed data on health (objective and subjective), socioeconomic status, family support, and healthcare use.
- It enables analysis of ageing trajectories (e.g. onset of disability, retirement and health, intergenerational support). SHARE's longitudinal data provide insights into transitions (e.g. from independent living to needing care), but sample sizes per country are modest and the institutionalized elderly are largely missing.

- National Surveys & Studies

In the past, ÚZIS ran national Health Interview Surveys (HIS) and there are ad-hoc studies (e.g. the HAPIEE study in one region) that focused on older populations' health determinants. These contribute additional context but are not regular.

Surveys offer rich individual-level health details (including psychosocial factors) but may under-represent the oldest frail populations.



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Long-Term Care and Social Sector Data

Social Services Statistics

- The Ministry of Labour and Social Affairs collects data on long-term care providers and recipients which includes counts of seniors receiving in-home care services, numbers of nursing home residents, occupancy of senior homes, etc.
- However, data on *individual-level* LTC trajectories (who enters care, duration, outcomes) are limited.
- There is **no unified national client-level LTC registry** akin to health registries.

– Care Allowance and Informal Care:

- Czechia has a care allowance program for dependent seniors cared for at home. Administrative data exist on how many people receive these allowances (by dependency level), which can indirectly indicate people with care needs.
- The extensive role of family care is not well-captured in current official datasets.
- LTC and Health Data Linkage
 - The health and social care systems remain siloed. There is no routine linkage between hospital records and subsequent nursing home admission data. As a result, it's challenging to track the full care pathway of an older person (e.g. hospitalization followed by rehabilitation or LTC placement).
 - Data on quality of LTC (outcomes, staffing, etc.) are also **fragmentary**.
 - Some indicators (e.g. Healthy Life Years, disability-free life expectancy) attempt to quantify the balance between health and care needs, but their accuracy suffers from the data gaps noted.



Key Data Gaps & Limitations (I.)

- Regional Granularity:
 - Many health indicators for seniors are available only at national level.
 - Survey sample sizes and data aggregation mean we cannot reliably assess regional disparities in older population health. (E.g. life expectancy at 65 is published by region, but *healthy* life expectancy or disability prevalence by region is often unavailable or statistically unreliable).
 - Data gap: policy-makers lack region-specific ageing health profiles to target local needs.
- Institutionalized Population Missing:
 - Crucially, standard surveys (EHIS, EU-SILC) do not cover seniors in care institutions
 - This underestimates the prevalence of severe disability and chronic illness in the total elderly population. For instance, a frail 85-year-old in a nursing home is invisible in household surveys.
 - This bias can make Czech seniors appear "healthier" in data than they truly are, complicating planning for healthcare and LTC resources.
- Fragmented Data Silos:
 - Health, social care, and demographic data exist in separate silos with limited interoperability.
 - Although Czech citizens have a unique personal identifier ("rodné číslo"), data linkage across sectors is not routinely performed due to legal, privacy, and technical hurdles.
 - This prevents a holistic view e.g., linking hospital records with social care use or linking cause-of-death data with prior health status. Unlike Nordic countries where personal IDs enable seamless individual-level linkage across registries, Czech data remain fragmented.



Key Data Gaps & Limitations (II.)

- Inconsistencies & Quality Issues:
 - Different data sources yield diverging metrics (as seen with Healthy Life Years).
 - Some conditions among seniors may be underreported or misclassified in administrative data (e.g. dementia might be under-diagnosed on death certificates).
 - There is also a lack of routine quality assessments for some data collections fewer than onethird of health monitoring projects in EU countries applied standardized quality checks, and Czechia is no exception.
- Data Frequency and Timeliness:
 - Key surveys on older adults' health are infrequent (EHIS ~5-6 years, SHARE waves ~2 years).
 - Administrative data are continuous but published with delays.
 - Emerging issues (e.g. COVID-19's impact on elderly health) revealed the difficulty of quickly integrating data across systems (e.g. excess mortality in seniors was captured but linking that to comorbidities or care settings is challenging in real-time).
 - Overall, existing datasets, while rich in certain areas, have gaps in coverage, integration, and detail that limit their usefulness for comprehensive policy analysis.



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