Association of disability with mortality in the Spanish adult non-institutionalised population

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ABSTRACT

Background There are scant studies focused on measuring the association between disability and all-cause mortality based on large representative national samples of the community-dwelling adult population; moreover, the number of such studies which also include cause-specific mortality is yet lower.

Methods Longitudinal cohort study that used baseline data from 162,381 adults who participated in a countrywide disability survey (2008). A nationally representative sample was selected and interviewed in their homes. We present data on people aged ≥18 years. Disability was considered as any substantial limitation found on a list of 44 life activities that have lasted or are expected to last more than 1 year and originate from an impairment. Cause-specific mortality data were obtained from the Spanish Statistical Office. Subjects contributed follow-up time from baseline interview until death or the censoring date (31 December 2017). We computed standardised rate ratios (SRRs), with age, sex, living with a partner and education level distribution of the total group as standard population.

Results Adults with disability (11%) had an adjusted mortality rate more than twice as high as adults without disability (SRR 2.37, 95% CI 2.24 to 2.50). The increased mortality risk remained over the 10-year follow-up period. Mortality due to diseases of the nervous system (SRR 4.86, 95% CI 3.93 to 6.01), diseases of the musculoskeletal system (SRR 3.45, 95% CI 2.18 to 5.47), infectious diseases (SRR 3.38, 95% CI 2.27 to 5.01) and diabetes mellitus (SRR 3.56, 95% CI 2.71 to 4.68) was particularly high in those with disability.

Conclusions All-cause mortality rates are markedly higher among adults with disability. Preventive measures and health promotion initiatives are needed to reduce mortality risk in this population. Special attention should be paid to disabled people with certain specific diseases.

INTRODUCTION

Disability is an umbrella term, covering impairments, activity limitations and participation restrictions. Impairments are problems in body function or structure; an activity limitation is a difficulty in executing a task or action; whereas participation restriction refers to problems in involvement in life situations.1 There are many types of disabilities, such as those that affect vision, hearing, movement, thinking, remembering, learning, communicating, mental health and social relationships.2 Disability, in either of its different types, could appear in relation to very different conditions, injuries or diseases.2

The prevalence of disability is largely heterogeneous because of real differences in the prevalence of its determinants but also due to different definitions and sources. It can range between 10% and 15% of the population.3 The association between disability and health is also complex. Disability can be the consequence of disease or health conditions but also their cause, since it has been reported that 87% of people with disability have at least one secondary health condition.4 So, different disabilities have been identified as determinants for multiple conditions that are potentially related to an increased risk of death, such as depression,5 decubitus ulcers,6 a reduced immune function,7 pulmonary infections,8 obesity9 and weight problems in general,4 bowel and bladder problems,4 asthma,4 cardiovascular problems,3 anxiety10 and more propensity to injuries and falls,4 among others. Disabilities can also cause difficulty getting out into the community,1 sleep problems4 and muscle spasms1 that could be related to poorer mental health including higher risk of suicide.7 Moreover, despite advances in personalised care, health professionals are not always properly prepared to diagnose and treat the secondary conditions derived from disability,1 which can lead to a worse prognosis. Thus, it is reasonable to expect that disability may be associated with an increased risk of death.

Finally, as disability rates are strongly increasing in Europe,1 it becomes important to provide epidemiological information about potential consequences of this trend, such as mortality, all-cause and cause-specific. Many studies show associations between disability and mortality in older populations, but there are few studies that primarily focus on measuring this association in representative samples of adult populations.

In 2008, the Spanish Statistical Office (INE) carried out a large nationwide survey on disability.9 This survey formed the baseline cohort for a subsequent follow-up study on mortality.

The main objective of this research was to measure the association of disability with all-cause and cause-specific mortality in a representative sample of the adult population in Spain.

METHODS

Study population

Baseline data came from the 2008 Spanish Survey on Disabilities, Personal Autonomy and Dependency (Encuesta Sobre Discapacidades, Autonomía Personal y Situaciones de Dependencia (EDAD2008)).9

The study population covered by EDAD2008 consisted of a two-step, stratified, random sample of the community-dwelling population, representative of each province: first, a sample of census tracts was drawn, and a sample of family dwellings was then randomly selected within each tract. Thereafter, all households within the dwelling were group screened, and one member of each household was interviewed as the main informant. Subjects identified with a possible disability were then interviewed in depth. The interviews were conducted between November 2007 and February 2008. A total of 258,187 people living in 91,290 households were thus screened for disability. Data on mortality could be obtained for 207,529 subjects (80.4%). Of these, 162,381 people aged 18 years or over formed the study sample. Missing data on mortality were due to difficulties in identifying subjects.

Study variables
In the EDAD2008 survey, disability was considered as an important limitation to carrying out an activity, which had lasted or was expected to last more than 1 year and whose origin was an impairment. A person was considered to have a disability even if it had been overcome with the use of external technical aids, or with the help and/or supervision of another person. For screening purposes, a list of 44 questions about possible disabilities—read verbatim—was presented to the main informant of each household (see online supplemental material). The list addressed the following eight domains: vision; audition; communication; learning and application of knowledge and performance of tasks; mobility; self-care; domestic life; interaction and interpersonal relationships. People who answered affirmatively to at least one of these questions formed the disability group. To assess disability severity, we used two methods. First, a reported score based on the International Classification of Functioning Disability and Health (ICF) Checklist. Twenty-nine EDAD2008 items were backcoded to specific domains of the ICF Checklist and by rating the recorded difficulty performing specific tasks. A global ICF score was derived, although lacking data for domains d1, ‘Learning and applying knowledge’, d8, ‘Major Life Areas’ and d9, ‘Community, Social and Civic Life’. Disability severity was categorised into mild, moderate, severe and complete disability according to ICF categories. The categories of severe and complete disability were collapsed together; consequently, a four-category variable was analysed (no disability; mild; moderate; severe/complete). A detailed explanation of these procedures can be found elsewhere. As a second step, the number of disabilities identified for each person was grouped together to produce four categories: 0, 1–3, 4–9 and 10–42.

Age, marital status and educational level attained were obtained by interview. Mortality data were provided by the INE. Underlying causes of death were coded using the International Statistical Classification of Diseases and Related Health Problems, 10th Revision reduced list. These include 102 different causes of death grouped into 17 types. Subjects contributed follow-up time from their baseline interview until death or the censoring date, 31 December 2017.

Analysis
Overall baseline proportions, means and crude rates were weighted with sampling weights. These were computed as the inverse of each participant’s baseline selection probability, corrected for non-response.

To estimate the adjusted association of disability with mortality—all-cause and cause-specific—we computed standardised rate ratios (SRRs) via inverse probability weighting, taking the baseline distribution of age (18–39, 40–64, 65–74, ≥75 years), sex (women or men), living with a partner (yes or no) and educational level (less than primary, primary, secondary, preuniversity, university) for the total population as the standard. Weights for subjects were computed as the inverse of the conditional probability of being in their own exposure category, computed through sampling-weighted logistic regression models with disability as dependent variable, and the independent variables above as covariates. In the case of the four-category disability variables (ICF severity and number of disabilities), these probabilities were estimated through multinomial logistic regression models. Weights were then multiplied by the unconditional probability of being in their own exposure category. These stabilised weights were finally multiplied by the sampling weights. We computed weighted Poisson regression models for estimating standardised rates and rate ratios with these final weights, with disability as the independent variable and the logarithm of each observed person-year as the offset variable. To evaluate time-dependent effects, we computed 10 SRRs with increasing follow-up time, from 1 to 10 years.

Population attributable fraction for disability was calculated with the formula \(\hat{a} = \frac{E_i - E_n}{E} \), where \(E_i\) is the fraction with disability among deaths and \(E\) is the standardised mortality ratio. SMR was computed via inverse probability weighting using the baseline distribution of the disability group as standard. In this case, the weight for the \(i\) subject was estimated by fitting a sampling-weighted logistic regression model with disability status (E) as dependent variable and age, sex, living with a partner and educational level as independent variables (Z), and computed as:

\[ w_i = \frac{P(E=1|Z=z_i)}{P(E=0|Z=z_i)} \]

Thus, weights were 1 for disability subjects, and the conditional disability odds for non-disability. Final weights were the product of sampling weights and standardisation weights and applied in a Poisson model to obtain the SMR. CIs for the population attributable fraction were calculated with the formula proposed by Greenland.

Due to the complex sampling design, we used appropriate methods to account for the effect of stratification and clustering on SEs. Analyses were run with Stata V.15.

RESULTS
Baseline characteristics of the study population are shown in table 1. The prevalence (95% CI) of disability was 11.0% (10.7% to 11.2%), and was higher in older people, women, participants not living with a partner, and those with lower educational levels. There were 17,132 deaths in 1,541,910 person-years follow-up. The median follow-up was 10.0 years. The sampling weighted mortality rates (95% CI) per 1000 person-years, were 10.0 (10.1 to 10.6), for the overall population, 30.8 (49.3 to 52.3) for the disability group and 6.4 (6.2 to 6.6) for the non-disability group, for a crude rate ratio of 7.93 (7.63 to 8.24).

Table 2 shows the standardised mortality rates for the whole group and for subgroups of sex and age. The standardised rates per 1000 person-years were 20.03 for the disability group and 8.47 for non-disability group, for an SRR of 2.37. The standardised rate difference (SRD) between the mortality of the disability group and the non-disability group was 11.56 per 1000 person-years. There was little difference in the relative estimates.
The association between disability and more specific causes of death can be found in online supplemental table 1. Some of these estimates are unstable due to small number of cases (for a simple measure of precision for rate ratio estimates, we provide the ratio of the upper to lower confidence limits; as a somewhat arbitrary guide, estimates with ratios below 2 are deemed very precise and above 10 are very unstable. Diseases showing reasonable stable estimates (UL/LL less than 5) and strong associations (SRR of 2.5 or more) deserve particular attention.

**DISCUSSION**

In this large follow-up study, we found mortality to be more than twice higher in persons with disability, with a population attributable fraction of 21%. This could be translated to a figure of 67 309 annual deaths attributable to disability in the Spanish adult population (415 757 total deaths in people aged ≥18 years, estimated for 2019). We also found that the risk remains high over time. Interestingly, we found a clear increase in risk for almost all causes of death.

A limited number of works have primarily measured mortality associated with disability in representative samples. In general, the estimates presented by these studies are similar to those of the present article. Majer et al found 65% higher mortality in the Dutch population over 55 years of age (average adjusted HR=1.65). Forman-Hoffman et al report a 50% higher risk of mortality in people with disabilities, in a representative study of the American population aged 18 or over. Wu et al report an adjusted HR of 2.23 in a follow-up study of a representative sample of American older people. Finally, Park et al found 57% higher adjusted mortality in the disabled group in a large all-age Korean population.

The study of mortality associated with disability is complex due to the nature of the determinants of disability and its consequences. These determinants are both numerous and varied by sex, with SRR of 2.37 for women and 2.38 for men. There was a clear gradient by age with younger people showing higher SRRs and older groups with notably higher SRDs.

The SMR (95% CI) was 1.92 (1.83 to 1.99) and the fraction of persons with disability among the deceased was 0.44, for a population attributable fraction (95% CI) of 0.21 (0.20 to 0.22).

Online supplemental figure 1 and 2 show the estimates for increasing follow-up years. The SRRs were very high in the first years of follow-up but remained high and over 2 throughout the 10 years. SRDs also declined with follow-up time.

Tables 3 and 4 show the association of disability severity with mortality, measured both by the ICF score and the number of disabilities suffered by each subject. In both cases, a strong gradient was evident.

Table 5 shows the cause-specific mortality associated with disability. Disability was clearly associated with mortality in all groups. The SRR was particularly high in the following groups: diseases of the nervous system (4.86), diseases of the musculoskeletal system (3.45), infectious diseases (3.38), diseases of the genitourinary system (3.10), and endocrine diseases (3.08); and was lower for neoplasms (1.74). In terms of absolute impact, as measured by the SRD, diseases of the circulatory system and neoplasms showed higher figures, with 3.24 and 2.13 deaths per 1000 person years, respectively.

The association between disability and more specific causes of death can be found in online supplemental table 1. Some of these estimates are unstable due to small number of cases (for a simple measure of precision for rate ratio estimates, we provide the ratio of the upper to lower confidence limits; as a somewhat arbitrary guide, estimates with ratios below 2 are deemed very precise and above 10 are very unstable. Diseases showing reasonable stable estimates (UL/LL less than 5) and strong associations (SRR of 2.5 or more) deserve particular attention.
and the causal paths equally intricate. Some diseases lead to visual and auditory impairments. The interplay between activity ICF score was mainly based on ICF activities and did not include a role for activity limitations. The 10-disability among older people, that is, dementia, stroke and depression, among others. The SRR being relatively invariant across sexes and higher at ages mortality is mediated by disability itself and a smaller part leads to death by paths that do not include disability.

The analysis appears to reveal some systematic patterns. With the SRR being relatively invariant across ages and higher at ages <65 years, it would appear that disability-related causes of death in the adult population were different than the most notable causes of death among older people, that is, dementia, stroke and depression, among others. The 10-fold SRR increase we found for severity when the latter was measured with our ICF score might indicate a role for activity limitations as risk-of-death determinants; our ICF score was mainly based on ICF activities and did not include visual and auditory impairments. The interplay between activity limitations and medical care has yet to be mapped as a crucial part of coordinating health and social services, which is frequently mentioned as a prominent but relatively little-known field of primary care. Understanding the relevance of specific activity limitations for determining risk of death when interacting with environmental (living alone, institutionalisation, homebound, etc) or personal (literacy) factors may constitute grounds for increased surveillance of non-assisted disability persons, as has been proposed for primary care. In addition, large-scale epidemiological research could identify limitations in mobility, domestic activities or self-care associated with risk of death, as well as their potential underlying pathophysiological mechanisms.

This study has strengths that deserve to be mentioned. First, the very large sample size notably limits random errors. Second, interviewers specifically trained for the survey obtained disability assessments in the subjects' homes and we obtained mortality data from the INE. All of this greatly limits the possibility of bias. Finally, the representativeness of the sample improves the external validity of the study. Our study has several limitations. First, there was no information on mortality for 20% of the sample. However, we believe that the expected biasing impact on the estimates will be small, mainly because these missing data were due to problems in the identification of subjects, and thus we can reasonably assume that they are randomly distributed. Second, we were unable to make the adjustment more comprehensive, and our estimates may consequently suffer from residual confounding. However, we do not expect this problem to be quantitatively important, when considered in comparison with the estimates found in similar studies with adjustment for health variables. In addition, we found that the mortality rate increased in the disability group for almost every cause of death. Third, individuals' disability status was self-reported, and this can be a source of both over-reporting and under-reporting. However, it is expected that this potential misclassification would have diluted the strength of the estimations. Finally, we could not include the institutionalised population. Neglecting the institutionalised severely disabled, 1/5 of the overall group may have led to an underestimation of SRD and the attributable fraction.

### Table 4: Association of number of disabilities with mortality

<table>
<thead>
<tr>
<th>No of disabilities</th>
<th>SR*</th>
<th>SRD</th>
<th>SRR</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>8.49 (8.24–8.74)</td>
<td>0 (ref)</td>
<td>1 (ref)</td>
</tr>
<tr>
<td>1–3</td>
<td>12.53 (12.43–14.63)</td>
<td>5.04 (3.94–6.14)</td>
<td>1.59 (1.47–1.73)</td>
</tr>
<tr>
<td>4–9</td>
<td>19.20 (17.48–20.92)</td>
<td>10.71 (8.98–12.43)</td>
<td>2.26 (2.06–2.48)</td>
</tr>
<tr>
<td>10–42</td>
<td>42.54 (38.37–46.72)</td>
<td>34.06 (29.88–38.23)</td>
<td>5.01 (4.53–5.54)</td>
</tr>
</tbody>
</table>

*SR per 1000 person-years, with the baseline total population distribution of age, sex, living with a partner and education as the standard.

SR, standardised rate; SRD, standardised rate difference; SRR, standardised rate ratio.

### Table 5: Association of disability with mortality by group of causes

<table>
<thead>
<tr>
<th>Group of causes</th>
<th>SR (95% CI)*</th>
<th>SRR (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Certain infectious and parasitic diseases</td>
<td>0.28 (0.14 to 0.42)</td>
<td>3.38 (2.27 to 5.01)</td>
</tr>
<tr>
<td>Neoplasms</td>
<td>2.13 (1.65 to 2.61)</td>
<td>1.74 (1.57 to 1.92)</td>
</tr>
<tr>
<td>Diseases of the blood and blood-forming organs and certain disorders involving the immune mechanism</td>
<td>0.02 (−0.00 to 0.05)</td>
<td>1.75 (0.94 to 3.26)</td>
</tr>
<tr>
<td>Endocrine, nutritional and metabolic diseases</td>
<td>0.45 (0.33 to 0.57)</td>
<td>3.08 (2.43 to 3.90)</td>
</tr>
<tr>
<td>Mental and behavioural disorders</td>
<td>0.40 (0.28 to 0.53)</td>
<td>2.43 (1.91 to 3.07)</td>
</tr>
<tr>
<td>Diseases of the nervous system</td>
<td>1.46 (1.13 to 1.79)</td>
<td>4.86 (3.93 to 6.01)</td>
</tr>
<tr>
<td>Diseases of the circulatory system</td>
<td>3.24 (2.78 to 3.71)</td>
<td>2.32 (2.12 to 2.55)</td>
</tr>
<tr>
<td>Diseases of the respiratory system</td>
<td>1.42 (1.17 to 1.66)</td>
<td>2.69 (2.35 to 3.07)</td>
</tr>
<tr>
<td>Diseases of the digestive system</td>
<td>0.56 (0.38 to 0.73)</td>
<td>2.25 (1.84 to 2.75)</td>
</tr>
<tr>
<td>Diseases of the skin and subcutaneous tissue</td>
<td>0.03 (0.00 to 0.05)</td>
<td>2.38 (1.22 to 4.64)</td>
</tr>
<tr>
<td>Diseases of the musculoskeletal system and connective tissue</td>
<td>0.20 (0.09 to 0.31)</td>
<td>3.45 (2.18 to 5.47)</td>
</tr>
<tr>
<td>Diseases of the genitourinary system</td>
<td>0.51 (0.37 to 0.65)</td>
<td>3.10 (2.44 to 3.94)</td>
</tr>
<tr>
<td>Congenital malformations, deformations and chromosomal abnormalities</td>
<td>0.05 (0.01 to 0.10)</td>
<td>28.23 (5.86 to 135.96)</td>
</tr>
<tr>
<td>Symptoms, signs and abnormal clinical and laboratory findings, not elsewhere classified</td>
<td>0.32 (0.19 to 0.46)</td>
<td>2.73 (2.01 to 3.71)</td>
</tr>
<tr>
<td>External causes</td>
<td>0.47 (0.26 to 0.69)</td>
<td>2.68 (1.98 to 3.65)</td>
</tr>
</tbody>
</table>

*SR per 1000 person-years, with the baseline total population distribution of age, sex, living with a partner and education as the standard.

SR, standardised rate; SRD, standardised rate difference; SRR, standardised rate ratio.

### What is already known on this subject

- There are many studies that present data on mortality associated with disability, but few have focused on measuring this association in large representative samples of populations. These few studies have found mortality rates between 50% and 100% higher in disability populations as compared with non-disability populations.

### What this study adds

- In a large, long-term follow-up study, we provide information on the risk of all-cause and cause-specific mortality associated with disability, including analyses of associations with disability severity. We report that clearly higher mortality was associated with disability, both in absolute and relative measures. Thus, preventive and health promotion measures are imperative to reduce the mortality in these populations.
diseases and diabetes mellitus. In absolute terms, the number of deaths attributable to disability may also be considered very high. Thus, it would be advisable to implement adequate preventive measures and health promotion initiatives, to help both improve survival and offer better quality of life to people with disability.

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