A comparison of the prevalence of health problems among adults with and without intellectual disability: a total administrative population study

Martin McMahon^{1,2} & Chris Hatton¹

¹ Division of Health Research, Lancaster University, Lancaster ² Health and Community Services, Government of Jersey

Correspondence: Division of Health Research, Lancaster University, Lancaster <u>m.mcmahon2@lancaster.ac.uk</u>

Abstract

Introduction: There is considerable international research indicating health disparities between people with and without intellectual disabilities. It is important that comparative studies use representative population samples. This study compares a total administrative population of adults with intellectual disability to a random stratified general population sample in Jersey.

Methods: A total administrative population of 217 adults with intellectual disability and a random stratified sample of 2,350 adults without intellectual disability participated. A questionnaire using the International Classification of Diseases (ICD-10) Chapter Headings was administered to all participants to enable a like-for-like comparison across both populations.

Findings: Unadjusted comparisons identified that adults with intellectual disability have a greater prevalence of health problems. However, they were less likely to experience cancers and musculoskeletal diseases. The only significant impact of adjusting for between-group differences in age and gender was that a difference in genitourinary disorders became non-significant.

Conclusions: These findings are consistent with the hypothesis that adults with intellectual disabilities generally have greater prevalence rates of health problems than the general population.

Introduction

It is well documented that in high-income countries people with intellectual disability have poorer health than the general population (Emerson, Hatton, Baines & Robertson, 2016; Heslop & Glover, 2015), with people dying on average 20 years earlier than their non-disabled peers (O'Leary, Cooper, & Hughes-McCormack, 2018; Glover, Williams, Heslop, Oyinlola, & Grey, 2017; Heslop et al. 2014; Learning Disability Mortality Review, 2018; Troller, Srasuebkul, Xu, & Howlett, 2017; Lauer & McCallion, 2015; McCarron, Carroll, Kelly, & McCallion, 2015).

There is substantial variation in the prevalence rates of major health problems for people with intellectual disabilities reported across different studies and how they compare to people without intellectual disabilities. For example, studies that have investigated cancer (Bonell, 2010; Tyler & McGrother, 2009; Patja, Molsa Livanainen, 2001; Duff et al. 2001; Cooke 1997) diabetes (Mac Rae et al. 2015; de Winter et al. 2012; Tyler et al. 2010; McDermott, Platt, & Dasari, 2006) and mental health problems (Hughes-McCormack et al. 2017; Buckles, Luckasson, & Keefe, 2013) have reported varying prevalence rates in people with intellectual disabilities. A range of potential methodological reasons for this principally focus on the inconsistent definition of intellectual disabilities; the diverse diagnosis tools, and small sample sizes used in studies. Although there is a growing body of research that uses representative samples of people with and without intellectual disabilities (Balogh, Brownell, Ouellette-Kuntz, & Colantonio, 2010; Hosking et al., 2016; Hughes-McCormack et al. 2018; Morin, Mérineau-Côté, Ouellette-Kuntz, Tassé, & Kerr, 2012), this continues to be one of the most important methodological limitations in intellectual disability research more broadly (Emerson & Hatton, 2014; Hogg, & Tuffrey-Wijne, 2008; Hughes-McCormack et al., 2017).

Acknowledging such methodological limitations, the aim of this brief report was to build upon and integrate existing literature to estimate the current prevalence of health problems using ICD-10 classification headings in a total administrative population of adults with intellectual disabilities and a comparison random stratified general population sample in Jersey. The same variables were used to facilitate comparison across people with and without intellectual disabilities.

Method

Measures

A survey was developed based on ICD-10 (2015) English online version (https://icd.who.int/browse10/2015/en) chapter headings I to XV: viral or infective diseases; cancers, diseases of the blood; endocrine, nutritional or metabolic conditions; mental health illnesses or behavioural problems; neurological conditions; diseases of the eye; diseases of the ear; diseases of the circulatory system; diseases of the respiratory system; diseases of the digestive system; diseases of the skin; diseases of the musculoskeletal system; diseases of the genitourinary system; malformations or genetic problems; and injuries to your body as a result of trauma or poisoning. For the purpose of this paper, classification headings only were used to enable direct comparisons between groups in both populations. A dichotomous variable was created (yes/no) asking participants if they had diseases or disorders of the classification headings from these chapters. In each classification heading we provided examples of the most common diseases that were representative of that group. We included an open question for participants to record any other disease or disorders that they have not mentioned in the survey. For the intellectual disability sample, all electronic health and nursing notes held on Care Partner (an electronic health and social care database) by Jersey's Health and Community Services were reviewed. Demographic variables were collected on both surveys that mirrored the Jersey Opinions and Lifestyle Survey (States of Jersey, 2017). This data is reflective of the local population.

Ethics

Ethical approval was granted from Lancaster University and by the Government of Jersey, Health and Community Services Ethics Committee in January and March 2017. The consent process and accompanying documentation was designed using guidance from the Mental Capacity Act (2005) and the National Research Ethics Service (NRES) (<u>http://www.nres.nhs.uk/</u>). Full details of the consenting procedure for adults with an intellectual disability are outlined previous studies following the same methodology (x and y study - blinded for review).

Intellectual disability population

A total administrative sample of adults with intellectual disability known to services in Jersey were contacted to participate (i.e. people who were receiving, or had received, support from intellectual disability services in Jersey). 217 adults with intellectual disabilities participated (age range 18-85 [male n=122, female n=95]), a response rate of 76% (sampling frame n=285). Approximately 50% of participants were administratively defined by Jersey's Health and Community Services as having a mild intellectual disability (n =108), 25.8% (n = 56) as having a moderate intellectual disability, 15.7% as having a severe intellectual disability (n=34) and 8.8% (n=19) as having a profound intellectual disability.

All information was collected by face-to-face interviews with the participants themselves or through proxy respondents. In this regard, 132 (60.8%) adults were consented though proxy procedures and they answered on behalf of the person with an intellectual disability, whilst 85 (39.2%) participants consented and answered independently. All health records held on Care Partner were checked to corroborate findings. To receive a health and social service in Jersey individuals with an intellectual disability have a yearly assessment and they have a current care plan that includes a health assessment; therefore, this served as robust measure to identify the prevalence of disease in this population. However, in a pragmatic manner, where it was self-reported by the person or a proxy had a disease but there was no evidence to support this on Care Partner, their community nurse was requested to confirm. In this instance, if the finding was not corroborated it was excluded for our analysis.

General population

A random stratified sample approach was used to recruit general population adults. Jersey's 12 parishes were divided into strata. Each parish was weighted in terms of population considering the most recent population census and allowing for net inward migration (States of Jersey, 2011). Addresses were drawn at random from the list of residential, active addresses for each parish on the Jersey Land Property Index excluding any household which was sampled for one of the previous 2015, 2016, 2017 social surveys or the Disability Survey in 2015 - there were 28,000 households in the overall sampling frame. Eight thousand surveys were posted to cover the entire adult population at random. This was based on the initial estimation of having a +/-2 percentage point confidence interval and assuming a 30% response rate. The household member who next celebrated their birthday, and who was aged 18 years or over, was asked to complete the survey. A total of 2,415 (30.2% [age range 18 – 105, male n-941, female n-1,394]) surveys were returned with 65 of the these being unusable. In total, 2,350 general population responses were included in the analysis.

Analysis

Initially, descriptive statistics and the frequency of ICD-10 disease presentation in the two populations were examined. To investigate the scale of any differences in disease prevalence between the intellectual disability and general population, Odds Ratios with 95% confidence intervals were calculated. Secondly, binary logistic regression analysis was undertaken to estimate the strength of any differences in disease prevalence between the intellectual disability and general population groups (odds ratios), once gender (binary variable) and age (split at the median [over and under 57 years]) were taken into account. Thirdly, an interaction term was fitted to determine if the effects of age and or gender differed across the intellectual disability and general populations. Finally, we matched 206 participants according to age and gender to determine if there was a difference in the frequency of health problems in both populations. This matching procedure was undertaken in SPSS using the case matching procedure.

There were no missing data in the intellectual disability dataset and less than 3% (range 2%-2.7%) across the general population dataset. The pregnancy complications variable was excluded from analysis as no person with an intellectual disability was pregnant during the study. Data were analysed using SPSS 25 and graphs were produced in 'R'. Effect sizes for Odds Ratios for 2x2 comparisons are interpreted as; small (OR < =0.82 or > =1.22), medium (OR < =0.54 or > =1.86), large (OR < =0.33 or > =3.00) [Olivier & Bell, 2013].

Results

Bivariate comparisons of health problems

The first stage of analysis involved simple bivariate comparisons between participants with and without intellectual disability with regard to the ICD-10 Chapter Headings. Odds Ratios were calculated and associated 95% confidence intervals with significance levels.

********* Table 1 about here **********

In summary, our main results suggest participants with intellectual disability were more likely than the general population to have: viral or infective diseases; mental health illnesses and behavioural problems; neurological disorders; diseases of the genitourinary system and malformations or genetic problems. In contrast, participants with intellectual disability were statistically less likely than the general population to have cancers and diseases of the musculoskeletal system, representing a medium and small effect size respectively. It was not possible to distinguish between mental health and behavioural disorders due to the lack of comparative data. Nevertheless, 33.6% of the intellectual disability sample have had a mental health diagnosis at some stage in their life.

See the Figure 1 Forest Plot (malformations or genetic problems are excluded from the Forest Plot as the OR of 47.14 is extreme) for a representation of these differences.

+++ Insert Figure 1 Here +++

Binary logistic regression results

+++ Insert Table 2 Here +++

After adjusting for age, gender and presence of intellectual disability our principle results suggest that females are more likely to have cancers and circulatory disorders but less likely to have endocrine, nutritional and metabolic disorders mental lllness and behavioural disorders or neurological disorder. Females with an intellectual disability without were significantly more likely to have mental illness and behavioural disorders but less likely to have diseases of the ear than females without an intellectual disability. Furthermore, increasing age increased the chances of having cancer; endocrine and metabolic disorders; neurological disorders; disorders of the eye; disorders or he ear; disorders of the circulatory system; diseases of the digestive system and musculoskeletal disorders. In contrast, younger age increased the chances of having mental illnesses and behavioural disorders and disorders of the skin. Further statistically significant results are outlined in Table 2.

Matched sample comparison results

In the final stage of analysis, case control matching was used to compare the general and intellectual disability sample according to age and gender in an attempt to further minimise confounding and improve precision (Rothman, Greenland, &

Lash, 2008). A total of 206 individuals were matched on a like-for-like basis. It can be concluded that people with intellectual disabilities (n=206) had a greater number of health problems, median (IQR) 3 (2,6) than the general population (n=206), median (IQR) 1 (0,2) and the difference in these distributions is significant (U = 32836, p < .001) (Figure 2).

+++ Insert Figure 2 Here +++

We also compared age bands (less than 35 years, 35-55 years and over 55 years) across the two populations and used the cumulative number of ICD-10 conditions as the dependent outcome variable. Across all three age band categories, people with intellectual disabilities had a greater prevalence of ICD-10 conditions and these were statistically significant: less than 35 years (U = 3048, p < .001); 35-55 years (U = 5182, p < .001); over 55 years (U = 3027, p < .001) (Figure 3).

+++ Insert Figure 3 Here +++

Discussion

Consistent with the results of previous epidemiological research our results indicate that in unadjusted comparisons, adults with intellectual disabilities have considerably greater prevalence rates of viral or infective diseases; diseases of the blood; endocrine, nutritional and metabolic conditions; mental health illnesses and behavioural disorders; neurological disorders; diseases of the eye; diseases of the respiratory system; diseases of the digestive system; diseases of the skin; diseases of the genitourinary system and malformations or generic problems (Heslop et al. 2014; Bonell, 2010; Robertson et al. 2015; Straetmans *et al.* 2007; Hughes-McCormack et al. 2017; Timmeren et al. 2017; Henderson et al. 2009; Janicki &

Dalton, 1998). Nevertheless, adults with intellectual disability were less likely to have cancers and diseases of the musculoskeletal system. No difference was observed between prevalence rates for diseases of the ear, diseases of the circulatory system or injuries to the body as a result of trauma or poisoning.

These results are consistent with previous research and are reflective of the health inequalities that adults with intellectual disabilities experience (Emerson & Baines, 2011; Emerson & Hatton, 2014; Krahn & Fox, 2014). Only diseases of the genitourinary system became non-significant after accounting for age and sex. Further adjusted comparisons identified a different topography of prevalence with regard to gender with cancers and circulatory disorders being more prevalent in females. In contrast, endocrine, nutritional and metabolic disorders, mental illness and behavioural disorders and neurological disorders were more prevalent in males. Our analysis only found two significant associations in the interaction component insofar as females with an intellectual disability were more likely to have mental illnesses and behavioural disorders but less likely to have diseases of the ear than their non-disabled peers. The age adjustment finding is not unique and suggests that older age increased the chances of having certain diseases. Notwithstanding, the age interaction effect between the general population and intellectual disability population identified that increasing age in the intellectual disability population increases the incidence of disorders of the eye, whereas reduced age in the general population identifies a lower prevalence of viral and infective diseases, mental illnesses and behavioural disorders and disorders of the skin. The matched sample analysis also highlights that people with an intellectual disability experience greater levels of ill health at a younger age and this trajectory continues throughout their life.

These results consolidate and extend existing knowledge about the health inequalities faced by people with intellectual disability in a number of ways. First, the use of a total administrative population in the intellectual disability sample is a strength of this study. Having access to participants' health records ensure accuracy of health data. Similarly, the random stratified sample that covered the whole residential address population of Jersey ensured a representative general population comparison sample of considerable numbers, although we were unable to check health data on the health system database due to large numbers of respondents and lack of consent.

Second, this study supports other evidence that cancer is less prevalent in the intellectual disability population (Cooke, 1997, Bonell, 2010) whilst mental health and behavioural disorders are more prevalent (Cooper et al. 2007; Hughes-McCormack et al. 2017; Bowring et al. 2017). This analysis did not distinguish between mental health and behaviours that challenge to ensure like-for-like comparison with the general population. The 33.6% prevalence rate for mental health disorders reported in this study is higher than two of most influential papers in this area that cite a 22.4% (Cooper et al. 2007) and 23.4% (Hughes-McCormack et al. 2017) prevalence rate respectively. This may be due to this study's total administrative population approach insofar as those known to services may have more health-related problems. In addition to the increased prevalence rate of the other conditions, these findings are not new and support the consistently highlighted poorer health of this population (Hoskings et al. 2016; Heslop et al. 2014) that are aligned to well-known determinants of health and wellbeing (Emerson & Hatton, 2014). In addition to this, the trajectory of ill-health and disease in the intellectual disability population needs to be considered from an age perspective. There is clear evidence in this study that people at a younger age experience a greater number of health problems. Medical advancements have meant that sustaining life in infancy has become more achievable and children who were born extremely premature or with complex needs are now living into adulthood where once they would have died. The consequence of such treatment can have a marked impact of these persons' health meaning they experience many morbidities earlier which continue throughout life. This potentially polarises the finding that younger age in the general population may not be a protective factor for people with an intellectual disability. Future research should use population level longitudinal evidence from universally standardised health coding systems to identify the burden of ill-health in both children and adults with an intellectual disability.

Four principle limitations need to be kept in mind when considering these results. Firstly, the ICD-10 classification structure used in this study does not specify what specific disease the person has as it groups disorders under an anatomical and physiological systems approach. Although examples of specific illnesses were used to assist the general population to correctly identify and match their disease to the correct heading, we acknowledge there is the potential for error as we could not cross-check results as it was an anonymous postal questionnaire. Second, although the use of a random stratified sampling approach ensured that the sampling frame is highly representative of the general population, there was only a 30% response rate. Third, this study has used two different methods to recruit participants. Although we acknowledge that this is a significant limitation in itself, we are also of the firm belief that general population cohort surveys are wholly exclusive for individuals with intellectual disabilities with greater needs. Therefore, in making reasonable adjustments to include as many people as possible with intellectual disabilities, we have produced this limitation. Fourth, this study has included adults known to services and there may be a 'hidden majority' such as adults with mild intellectual disability who do not access intellectual disability services (Emerson & Hatton, 2014).

Although these four limitations introduce a source of methodological bias into the findings, there is a substantial evidence base that substantiates the prevalence of the reported disease in this study as it is broadly similar to other Jersey estimates over the last ten years (States of Jersey, 2012; 2014; 2016). Additionally, there was no evidence of any nonresponse variable correlation (Johnson & Wislar, 2012), and missing values were less than 2.7% across all variables. This goes a significant way to mitigate against the first and second limitations. Concerning the third and fourth limitation, the evidence-base in intellectual disability research continues to be challenged over how should individuals with an intellectual disability be included in general population cohort surveys (Hughes-McCormack et al. 2017; Emerson et al. 2014). Overcoming such challenges is inevitably going to create issues where sampling procedures are disconnected to a certain degree. Therefore, the use of a total population sample is considered an appropriate response to include people with intellectual disabilities in comparative research who have significant needs while ensuring the general population is equally representative. Our findings are suggestive of it appropriateness as is substantiates and integrates the existing literature.

References

Bonell, S. (2010), "Neoplasms", in O'Hara, J., McCarthy, J.E. and Bouras, N. (Eds), Intellectual Disability and III Health: A Review of the Evidence, Cambridge University Press, Cambridge, pp. 127-36.

Bowring, D. L., Totsika, V., Hastings, R. P., Toogood, S., & Griffith, G. M. (2017). Challenging behaviours in adults with an intellectual disability: A total population study and exploration of risk indices. *British Journal of Clinical Psychology*, *56*(1), 16-32.

Buckles, J., Luckasson, R., & Keefe, E. (2013). A systematic review of the prevalence of psychiatric disorders in adults with intellectual disability, 2003–2010. *Journal of Mental Health Research in Intellectual Disabilities*, *6*(3), 181-207.

Cooke, L. B. (1997). Cancer and learning disability. *Journal of Intellectual Disability Research*, *41*(4), 312-316.

Cooper, S. A., Smiley, E., Morrison, J., Williamson, A., & Allan, L. (2007). Mental illhealth in adults with intellectual disabilities: prevalence and associated factors. *The British Journal of Psychiatry*, *190*(1), 27-35.

De Winter, C. F., Bastiaanse, L. P., Hilgenkamp, T. I. M., Evenhuis, H. M., & Echteld, M. A. (2012). Cardiovascular risk factors (diabetes, hypertension, hypercholesterolemia and metabolic syndrome) in older people with intellectual disability: results of the HA-ID study. *Research in developmental disabilities*, *33*(6), 1722-1731.

Emerson, E., & Baines, S. (2011). Health inequalities and people with learning disabilities in the UK. *Tizard Learning Disability Review*, *16*(1), 42-48.

Emerson, E., Hatton, C., Robertson, J., & Baines, S. (2014). Perceptions of neighbourhood quality, social and civic participation and the self rated health of British adults with intellectual disability: cross sectional study. *BMC Public Health*, *14*(1), 1252.

Emerson, E., & Hatton, C. (2014). *Health Inequalities and People with Intellectual Disabilities*. Cambridge: Cambridge University Press.

Emerson, E., Hatton, C., Baines, S., & Robertson, J. (2016). The physical health of British adults with intellectual disability: cross sectional study. *International journal for equity in health*, *15*(1), 11.

Glover, G., Williams, R., Heslop, P., Oyinlola, J., & Grey, J. (2017). Mortality in people with intellectual disabilities in England. *Journal of Intellectual Disability Research*, *61*(1), 62-74.

Henderson, C. M., Rosasco, M., Robinson, L. M., Meccarello, J., Janicki, M. P., Turk, M. A. & Davidson, P. W. (2009). Functional impairment severity is associated with health status among older persons with intellectual disability and cerebral palsy. Journal of Intellectual Disability Research, 53(11), 887-897.

Heslop, P., & Glover, G. (2015). Mortality of people with intellectual disabilities in England: a comparison of data from existing sources. *Journal of applied research in intellectual disabilities*, *28*(5), 414-422.

Heslop, P., Blair, P. S., Fleming, P., Hoghton, M., Marriott, A., & Russ, L. (2014). The Confidential Inquiry into premature deaths of people with intellectual disabilities in the UK: a population-based study. *The Lancet*, *383*(9920), 889-895.

Hogg, J., & Tuffrey-Wijne, I. (2008). Cancer and intellectual disability: a review of some key contextual issues. *Journal of Applied Research in Intellectual Disabilities*, *21*(6), 509-518.

Hosking, F. J., Carey, I. M., Shah, S. M., Harris, T., DeWilde, S., Beighton, C., & Cook, D. G. (2016). Mortality among adults with intellectual disability in England:

comparisons with the general population. *American Journal of Public Health*, *106*(8), 1483-1490.

Hosmer D. W., Jr., Lemeshow S. & Sturdivant R. X. (2013) Model-Building Strategies and Methods for LogisticRegression, in Applied Logistic Regression. In: AppliedLogistic Regression, Third edn, pp.89–151. John Wiley & Sons, New Jersey.

Hughes-McCormack, L. A., Rydzewska, E., Henderson, A., MacIntyre, C., Rintoul, J., & Cooper, S. A. (2017). Prevalence of mental health conditions and relationship with general health in a whole-country population of people with intellectual disabilities compared with the general population. *BJPsych open*, *3*(5), 243-248.

Janicki, M. P., & Dalton, A. J. (1998). Sensory impairments among older adults with intellectual disability. *Journal of Intellectual and Developmental Disability*, *23*(1), 3-11.

Krahn, G. L., & Fox, M. H. (2014). Health disparities of adults with intellectual disabilities: what do we know? What do we do?. *Journal of Applied Research in Intellectual Disabilities*, 27(5), 431-446.

Lauer, E., & McCallion, P. (2015). Mortality of people with intellectual and developmental disabilities from select US state disability service systems and medical claims data. *Journal of Applied Research in Intellectual Disabilities*, *28*(5), 394-405.

Learning Disability Mortality Review. (2018) The Learning Disabilities Mortality Review (LeDeR) Programme: Annual Report. Retrieved from http://www.bristol.ac.uk/university/media/press/2018/leder-annual-report-final.pdf

MacRae, S., Brown, M., Karatzias, T., Taggart, L., Truesdale-Kennedy, M., Walley, R., ... & Davies, M. (2015). Diabetes in people with intellectual disabilities: a systematic review of the literature. *Research in developmental disabilities*, *47*, 352-374.

McCarron, M., Carroll, R., Kelly, C., & McCallion, P. (2015). Mortality rates in the general Irish population compared to those with an intellectual disability from 2003 to 2012. *Journal of Applied Research in Intellectual Disabilities*, *28*(5), 406-413.

McDermott, S., Moran, R., Platt, T., & Dasari, S. (2006). Variation in health conditions among groups of adults with disabilities in primary care. *Journal of Community Health*, *31*(3), 147-159.

O' Leary, L., Cooper, S. A., & Hughes-McCormack, L. (2018). Early death and causes of death of people with intellectual disabilities: A systematic review. *Journal of Applied Research in Intellectual Disabilities*, *31*(3), 325-342.

Olivier, J., & Bell, M. L. (2013). Effect sizes for 2× 2 contingency tables. *PLoS One*, *8*(3), e58777.

Patja, K., Mölsä, P., & livanainen, M. (2001). Cause-specific mortality of people with intellectual disability in a population-based, 35-year follow-up study. *Journal of Intellectual Disability Research*, *45*(1), 30-40.

Robertson, J., Hatton, C., Baines, S., & Emerson, E. (2015). Systematic reviews of the health or health care of people with intellectual disabilities: a systematic review to identify gaps in the evidence base. *Journal of Applied Research in Intellectual Disabilities*, *28*(6), 455-523.

Rothman, K. J., Greenland, S., & Lash, T. L. (2008). *Modern Epidemiology* (Vol. 3). Philadelphia: Wolters Kluwer Health/Lippincott Williams & Wilkins.

States of Jersey (2011) Population and census statistics. Retrieved from :https://www.gov.je/Government/JerseyInFigures/Population/Pages/index.aspx

States of Jersey (2012) Health Profile for Jersey 2010. Retrieved from : https://www.gov.je/News/2012/Pages/HealthProfile2010Report.aspx

States of Jersey (2014) Health Profile for Jersey 2014. Retrieved from:

https://www.gov.je/Government/Pages/StatesReports.aspx?ReportID=1055

States of Jersey (2016) Health Profile for Jersey 2016. Retrieved from: https://www.gov.je/Government/Pages/StatesReports.aspx?ReportID=2464

States of Jersey (2017) Jersey Opinions and Lifestyle Survey. Retrieved from https://www.gov.je/Government/Pages/StatesReports.aspx?ReportID=3908

Straetmans, J. M., van Schrojenstein Lantman-de, H. M., Schellevis, F. G., & Dinant, G. J. (2007). Health problems of people with intellectual disabilities: the impact for general practice. *Br J Gen Pract*, *57*(534), 64-66.

The Mental Capacity Act (2005) available from https://www.legislation.gov.uk/ukpga/2005/9/contents (Retrieved 28/4/19)

Timmeren, E. A., van der Schans, C. P., van der Putten, A. A. J., Krijnen, W. P., Steenbergen, H. A., van Schrojenstein Lantman-de Valk, H. M. J. & Waninge, A. (2017). Physical health issues in adults with severe or profound intellectual and motor disabilities: A systematic review of cross-sectional studies. Journal of Intellectual Disability Research, 61(1), 30-49.

Trollor, J., Srasuebkul, P., Xu, H., & Howlett, S. (2017). Cause of death and potentially avoidable deaths in Australian adults with intellectual disability using retrospective linked data. *BMJ open*, *7*(2), e013489.

Tyrer, F., & McGrother, C. (2009). Cause-specific mortality and death certificate reporting in adults with moderate to profound intellectual disability. *Journal of Intellectual Disability Research*, *53*(11), 898-904.

Table 1: Prevalence of Diseases in the Intellectual Disability and GeneralPopulation with Associated ORs and 95% CI and P-Values

Variable		Intellectual	General	Odds	95% CI	P Value
Variable		Disability	Population	Ratio		
Participants		n = 217	n = 2,350			
Viral or infective diseases	Yes	n = 17 (7.8%)	n = 57 (2.5%)			
	No	n = 200 (92.2%)	n = 2284 (97.5%)	3.3	1.90-5.81	p < 0.001
	Missing data	n = 0 (0%)	n = 66 (2.8%)			
Cancers	Yes	n = 5 (2.3%)	n = 130 (5.7%)			
	No	n = 212 (97.7%)	n = 2164 (94.3%)	0.39	0.16-0.97	p = 0.036
	Missing data	n = 0 (0%)	n = 56 (2.4%)			
Diseases of the blood	Yes	n = 16 (7.4%)	n = 70 (3.1%)			
	No	n = 201 (92.6%)	n = 2217 (96.9%)	2.52	1.44-4.42	p < 0.001
	Missing data	n = 0 (0%)	n = 63 (2.7%)			
Endocrine, nutritional or metabolic	Yes	n = 67 (30.9%)	n = 456 (19.9%)			
conditions	No	n = 150 (69.1%)	n = 1837 (80.1%)	1.80	1.33-2.44	p < 0.001
	Missing data	n = 0 (0%)	n = 57 (2.4%)			
Mental health illnesses or behavioural	Yes	n = 114 (52.5%)	n = 343 (15%)			
problems	No	n = 103 (47.5%)	n = 1950 (85%)	6.29	4.70-8.41	p < 0.001
	Missing data	n = 0 (0%)	n = 56 (2.4%)			
Neurological conditions	Yes	n = 65 (30%)	n = 108 (4.7%)			
	No	n = 152 (70%)	n = 2185 (95.3%)	8.65	6.10-12.26	p < 0.001
	Missing data	n = 0 (0%)	n = 57 (2.4%)			
Diseases of the eye	Yes	n = 41 (18.9%)	n = 201 (8.8%)			
	No	n = 176 (81.1%)	n = 2093 (91.2%)	2.43	1.67-3.51	p < 0.001
	Missing data	n = 0 (0%)	n = 56 (2.4%)			
Diseases of the ear	Yes	n = 42 (19.4%)	n = 383 (16.6%)			
	No	n = 175 (80.6%)	n = 1919 (83.4%)	1.20	0.84-1.71	p = 0.307
	Missing data	n = 0 (0%)	n = 48 (2%)			
Diseases of the circulatory system	Yes	n = 49 (22.6%)	n = 514 (22.4%)			
	No	n = 168 (77.4%)	n = 1784 (77.6%)	1.01	0.73-1.41	p = 0.943
	Missing data	n = 0 (0%)	n = 52 (2.2%)			

Diseases of the respiratory system	Yes	n = 42 (19.4%)	n = 308 (13.4%)			
	No	n = 175 (80.6%)	n = 1989 (86.6%)	1.55	1.08-2.21	p = 0.016
	Missing data	n = 0 (0%)	n = 53 (2.3%)			
Diseases of the digestive system	Yes	n = 75 (34.6%)	n = 350 (15.2%)			
	No	n = 175 (65.4%)	n = 1949 (84.8%)	2.94	2.17-3.98	p < 0.001
	Missing data	n = 0 (0%)	n = 51 (2.2%)			
Diseases of the skin	Yes	n = 67 (30.9%)	n = 332 (14.5%)			
	No	n = 150 (69.1%)	n = 1957 (85.5%)	2.63	1.93-3.59	p < 0.001
	Missing data	n = 0 (0%)	n = 61 (2.6%)			
Diseases of the musculoskeletal	Yes	n = 76 (35%)	n = 1014 (44%)			
system	No	n = 141 (65%)	n = 1288 (56%)	0.69	0.51-0.91	p = 0.010
	Missing data	n = 0 (0%)	n = 48 (2%)	2		
Diseases of the genitourinary system	Yes	n = 65 (30%)	n = 190 (8.3%)			
	No	n = 152 (70%)	n = 2101(91.7%)	4.73	3.41-6.55	P < .001
	Missing data	n = 0 (0%)	n = 59 (2.5%)			
Malformations or genetic problems	Yes	n = 64 (29.5%)	n = 20 (0.9%)			
	No	n = 153 (70.5%)	n = 2267 (99.1%)	47.41	27.96-	p < 0.001
	Missing data	n = 0 (0%)	n = 63 (2.7%)		00.40	
Injuries to your body as a result of	Yes	N = 24 (11.1%)	n = 215 (9.4%)			
trauma or poisoning	No	n = 193 (88.9%)	n = 2074 (90.6%)	1.20	0.77-1.88	p = 0.561
	Missing data	n = 0 (0%)	n = 61 (2.6%)			
c, C)	r			<u> </u>	
R						

Figure 1: Forest Plot of ICD Chapter Headings and Associated Odds Ratios (with 95% CI)



Malformations or Generic Problems OR is 47.14 (95% CI 27.96-80.40) and had been omitted from this Forest Plot as it distorts interpretation

Table 2: Logistic Regression Model with Statistically Significant Results

		Nagelkerke	β	S.E.	Wald's X ²	Sig.	OR	95% CI for	Odds Ratio
		R ²			(df 1)			Lower	Upper
					C				
Viral & Infective	General/ Intellectual Disability	0.035	-1.185	.593	4.00	*	0.30	0.10	0.97
Diseases	Population*Age								
_	Gender		0.391	.183	4.572	*	1.48	1.03	2.11
Cancers		0.069							
	Age		-1.276	.213	35.804	***	0.28	0.18	0.42
Diseases of the Blood	General/ Intellectual Disability	0.027	-1.226	.461	7.062	**	0.29	0.12	0.72
Fude arine Nutritienel 9	Gender		328	.111	8.725	**	0.72	0.58	0.89
Endocrine Nutritional & Metabolic Disorders	Age	0.064	937	.111	71.463	***	0.39	0.32	0.49
	General/ Intellectual Disability		840	.257	10.688	**	0.43	0.26	0.71
	Gender		-451	.127	12.571	***	0.64	0.50	0.82
	Age		.611	.122	24.953	***	1.84	1.45	2.34
Mental Illness &	General/ Intellectual Disability	0.122	-1.853	.226	66.976	***	0.16	0.10	0.24
Behavioural Disorders	General/ Intellectual Disability		.785	.306	6.572	*	2.20	1.20	3.99
	Population*Gender								
	General/ Intellectual Disability		-1.318	.361	13.339	***	0.27	0.13	0.54
	Population*Age								
	Gender		491	.214	5.240	*	0.61	0.40	0.93
Neurological	Age	0.137	589	.204	8.353	**	0.56	0.37	0.83
	General/ Intellectual Disability		-2.592	.313	68.744	***	0.08	0.04	0.14
	Age		-1.351	.174	60.282	***	0.26	0.18	0.36
Еуе	General/ Intellectual Disability	0.077	-1.564	.310	25.547	***	0.21	0.11	0.38

	General/ Intellectual Disability		1.072	.434	6.093	*	2.92	1.24	6.85
	Population*Age								
	Age		-1.056	.122	65.477	***	0.35	0.27	0.44
Ear	General/ Intellectual Disability	0.061	930	.373	6.174	*	0.40	0.19	0.82
	Population*Gender								
	Gender		.233	.107	4.757	*	1.26	1.02	1.56
Circulatory Disorders	Age	0.145	-1.626	.118	190.147	***	0.20	0.16	0.25
Respiratory Disorders	Age	0.013	329	.124	7.059	**	0.72	0.57	0.92
Digestive Disorders	Age		665	.121	30.251	***	0.51	0.41	0.65
	General/ Intellectual Disability	0.057	996	.250	15.981	***	0.37	0.23	0.60
	Age		0.300	.120	6.208	*	1.35	1.07	1.71
Skin	General/ Intellectual Disability	0.031	537	.250	4.606	*	0.59	0.36	0.93
	General/ Intellectual Disability Population*Age		849	.361	5.541	*	0.43	0.21	0.87
Musculoskeletal	Age	0.067	927	.087	113.840	***	0.40	0.33	0.47
Disorders									
Malformations & Genetic	General/ Intellectual Disability	0.379	3.647	.465	61.614	***	0.03	0.01	0.07
Problems									

*p < 0.05, **p < 0.01, ***p < 0.001

Note: Any variables with a p-value > 0.05 are excluded. Each of the final models was assessed against the Hosmer–Lemeshow goodness of fit test statistic (Hosmer,Lemeshow, & Sturdivant, 2013). For each model, apart from diseases of the genitourinary system, a p-value above .10 was observed along with a small test statistic identifying that the models provided a good fit to the data.

Injuries to your body as a result of trauma or poisoning excluded as they were not statistically significant

P.Cox

Figure 2: A Matched Comparison Sample (n-206) Identifying the Cumulative Number of ICD-10 Conditions





