

Original citation:

GBD 2015 Disease and Injury Incidence and Prevalence Collaborators (Including: Uthman, Olalekan A.). (2016) Global, regional, national incidence, prevalence and years lived with disability for 310 acute and chronic diseases and injuries 1990-2015: a systematic analysis for the Global Burden of Disease Study 2015. Lancet, 388 (10053). pp. 1545-1602.

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Global, regional, and national incidence, prevalence, and years lived with disability for 310 diseases and injuries, 1990–2015: a systematic analysis for the Global Burden of Disease Study 2015



GBD 2015 Disease and Injury Incidence and Prevalence Collaborators*

Background Non-fatal outcomes of disease and injury increasingly detract from the ability of the world's population to live in full health, a trend largely attributable to an epidemiological transition in many countries from causes affecting children, to non-communicable diseases (NCDs) more common in adults. For the Global Burden of Diseases, Injuries, and Risk Factors Study 2015 (GBD 2015), we estimated the incidence, prevalence, and years lived with disability for diseases and injuries at the global, regional, and national scale over the period of 1990 to 2015.

Methods We estimated incidence and prevalence by age, sex, cause, year, and geography with a wide range of updated and standardised analytical procedures. Improvements from GBD 2013 included the addition of new data sources, updates to literature reviews for 85 causes, and the identification and inclusion of additional studies published up to November, 2015, to expand the database used for estimation of non-fatal outcomes to 60 900 unique data sources. Prevalence and incidence by cause and sequelae were determined with DisMod-MR 2.1, an improved version of the DisMod-MR Bayesian meta-regression tool first developed for GBD 2010 and GBD 2013. For some causes, we used alternative modelling strategies where the complexity of the disease was not suited to DisMod-MR 2.1 or where incidence and prevalence needed to be determined from other data. For GBD 2015 we created a summary indicator that combines measures of income per capita, educational attainment, and fertility (the Socio-demographic Index [SDI]) and used it to compare observed patterns of health loss to the expected pattern for countries or locations with similar SDI scores.

Findings We generated 9.3 billion estimates from the various combinations of prevalence, incidence, and YLDs for causes, sequelae, and impairments by age, sex, geography, and year. In 2015, two causes had acute incidences in excess of 1 billion: upper respiratory infections (17 · 2 billion, 95% uncertainty interval [UI] 15 · 4-19 · 2 billion) and diarrhoeal diseases (2.39 billion, 2.30-2.50 billion). Eight causes of chronic disease and injury each affected more than 10% of the world's population in 2015; permanent caries, tension-type headache, iron-deficiency anaemia, age-related and other hearing loss, migraine, genital herpes, refraction and accommodation disorders, and ascariasis. The impairment that affected the greatest number of people in 2015 was anaemia, with 2.36 billion (2.35-2.37 billion) individuals affected. The second and third leading impairments by number of individuals affected were hearing loss and vision loss, respectively. Between 2005 and 2015, there was little change in the leading causes of years lived with disability (YLDs) on a global basis. NCDs accounted for 18 of the leading 20 causes of age-standardised YLDs on a global scale. Where rates were decreasing, the rate of decrease for YLDs was slower than that of years of life lost (YLLs) for nearly every cause included in our analysis. For low SDI geographies, Group 1 causes typically accounted for 20-30% of total disability, largely attributable to nutritional deficiencies, malaria, neglected tropical diseases, HIV/AIDS, and tuberculosis. Lower back and neck pain was the leading global cause of disability in 2015 in most countries. The leading cause was sense organ disorders in 22 countries in Asia and Africa and one in central Latin America; diabetes in four countries in Oceania; HIV/AIDS in three southern sub-Saharan African countries; collective violence and legal intervention in two north African and Middle Eastern countries; iron-deficiency anaemia in Somalia and Venezuela; depression in Uganda; onchoceriasis in Liberia; and other neglected tropical diseases in the Democratic Republic of the Congo.

Interpretation Ageing of the world's population is increasing the number of people living with sequelae of diseases and injuries. Shifts in the epidemiological profile driven by socioeconomic change also contribute to the continued increase in years lived with disability (YLDs) as well as the rate of increase in YLDs. Despite limitations imposed by gaps in data availability and the variable quality of the data available, the standardised and comprehensive approach of the GBD study provides opportunities to examine broad trends, compare those trends between countries or subnational geographies, benchmark against locations at similar stages of development, and gauge the strength or weakness of the estimates available.

Funding Bill & Melinda Gates Foundation.

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Lancet 2016; 388: 1545-602

This online publication has been corrected. The corrected version first appeared at the lancet.com on January 5, 2017

See Editorial page 1447

See Comment pages 1448 and 1450

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Introduction

Although substantial progress has been made toward reducing mortality and extending life expectancy throughout the world over the past few decades, the epidemiological transition is manifest in the growing importance of non-fatal diseases, outcomes, and injuries which pose, partly as a consequence of decreasing death rates, a rising challenge to the ability of the world's population to live in full health. Complementing information on deaths by age, sex, cause, geography, and time with equally detailed information on disease incidence, prevalence, and severity is key to a balanced debate in health policy. For this reason, the Global Burden of Disease (GBD) Study uses the disabilityadjusted life-year (DALY), combining years of life lost (YLLs) due to mortality and years lived with disability (YLDs) in a single metric. One DALY can be thought of as one lost year of healthy life. The sum of DALYs in a population can be thought of as the gap between the population's present health status and an ideal situation where the entire population lives to an advanced age, free of disease. Assessments of how different diseases lead to multimorbidity and reductions in functional health status are important for both health system planning1 and a broader range of social policy issues such as the appropriate age for retirement in some countries.2,3 Many challenges in making standardised estimates of non-fatal health outcomes are similar to those affecting mortality estimates (including variations in case definitions, data collection methods, variable quality of data collection, conflicting data, and missing data) but are compounded by more sparse and varied data sources, the need to characterise each disease by its disabling sequelae or consequence(s), and the need to quantify the severity of these consequences. The standardised approach of the annual GBD updates addresses these measurement problems to enhance comparability between causes by geography and over time.

See Online for appendix

The estimates from GBD 2013 drew attention to large increases in the number of YLDs over the previous decade, whereas rates of YLDs for most causes remained stable or showed only small decreases.4 The GBD 2013 assessment largely attributed increases in the number of YLDs to musculoskeletal disorders, mental and substance use disorders, neurological disorders, and chronic respiratory diseases, as well as population growth and ageing. GBD 2013 also brought attention to increased differences in trends between mortality and morbidity for many causes. YLDs as a proportion of DALYs increased globally, a manifestation of the continuing epidemiological transition in low-income and middleincome countries. Decreases in mortality from diseases such as pneumonia, diarrhoea, maternal and neonatal disorders, and an absence of progress in reducing YLD rates continued to drive a transition toward a greater global number of YLDs.

Along with broad recognition that data from some regions were sparse and that more and higher quality data in general would probably improve estimation, useful debates on the GBD results have been published. These debates have focused on the analysis or presentation of individual diseases, such as changes over time in GBD estimates of dementia,5 the accuracy of HIV incidence estimates, 67 the absence of sepsis as a disease, 89 the quality of some cancer registry data,10 and the absence of mental disorders as sequelae of neglected tropical diseases.11 The GBD empirical approach to measuring the public's view of health state severity has generated substantial interest with questions about the relative importance of different dimensions of health, 12,13 the quantification of health loss, 14,15 and discussions of the transferability of judgments about relative health to conventional notions of disability and dependence.⁵ In each cycle of the GBD, we seek to improve the estimates, reflecting published and unpublished critique through the acquisition of new data, expansion of the network of collaborators, changes in how data are corrected for bias, advances in modelling techniques, and the targeted expansion of the GBD cause list.

The primary objective of this component of the GBD was to use all available data of sufficient quality to generate reliable and valid assessments of disease and injury sequelae incidence, prevalence, and YLDs for all 310 causes in the GBD cause hierarchy for 591 locations in the GBD study during 1990–2015. We describe the change over time and between populations in relation to where countries fall on the development continuum. Continuing efforts to improve data and code transparency are an important part of the GBD cycle. These results thus supersede any previous publications about the GBD on disease incidence, prevalence, and YLDs.

Methods

Overall approach

We estimated incidence and prevalence by age, sex, cause, year, and geography using a wide range of updated and standardised analytical procedures. The overall logic of our analytical approach is shown for the entire non-fatal estimation process in figure 1. The appendix provides a single source for detail of inputs, analytical processes, and outputs and methods specific to each cause. This study complies with the Guidelines for Accurate and Transparent Health Estimates Reporting (GATHER) recommendations (methods appendix pp 1, 608–10).¹⁷

Geographies in GBD 2015

The geographies included in GBD 2015 have been arranged into a set of hierarchical categories composed of seven super-regions and a further nested set of 21 regions containing 195 countries and territories. Eight additional subnational assessments were done for Brazil, China, India, Japan, Kenya, Saudi Arabia, South Africa, Sweden, and the USA (methods appendix pp 611–24). For this study we present data at the national and territory level.

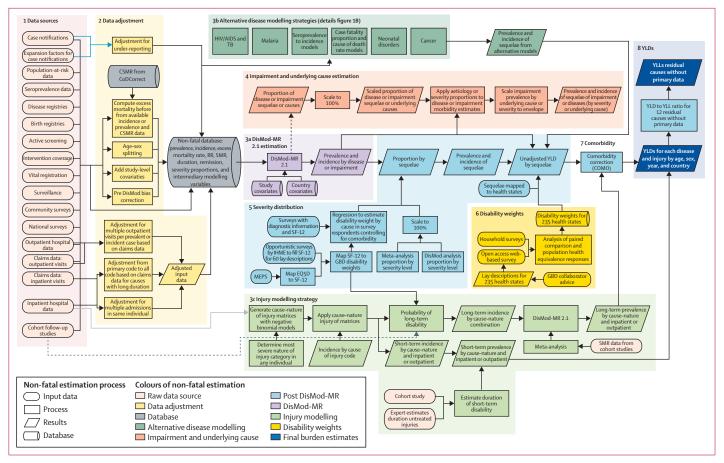


Figure 1: Analytical flow chart for the estimation of cause-specific YLDs by location, age, sex, and year for GBD 2015

Ovals represent data inputs, square boxes represent analytical steps, cylinders represent databases, and parallelograms represent intermediate and final results. The flow chart is colour-coded by major estimation component: raw data sources, in pink; data adjustments, in yellow; DisMod-MR 2.1 estimation, in purple; alternative modelling strategies, in light green; injury modelling strategy, in dark green; estimation of impairments and underlying causes, in brown; post-DisMod-MR and comorbidity correction, in blue; disability weights, in orange; and cause of death and demographic inputs, in grey. GBD=Global Burden of Disease. TB=tuberculosis. SF-12=Short Form 12 questions. MEPS=Medical Expenditure Panel Surveys. CSMR=cause-specific mortality rate. SMR=standardised mortality ratio. YLDs=years lived with disability. YLLs=years of life lost. IHME=Institute for Health Metrics and Evaluation.

List of causes and sequelae

The GBD cause and sequelae list is organised hierarchically (methods appendix 625-53). At Level 1 there are three cause groups: communicable, maternal, neonatal, and nutritional diseases (Group 1 diseases); non-communicable diseases; and injuries. These Level 1 aggregates are subdivided at Level 2 of the hierarchy into 21 cause groupings. The disaggregation into Levels 3 and 4 contains the finest level of detail for causes captured in GBD 2015. Sequelae of diseases and injuries are organised at Levels 5 and 6 of the hierarchy. The finest detail for all sequelae estimated in GBD is at Level 6 and is aggregated into summary sequelae categories (Level 5) for causes with large numbers of sequelae. Sequelae in GBD are mutually exclusive and collectively exhaustive, and thus our YLD estimates at each level of the hierarchy sum to the total of the level above. Prevalence aggregations are estimated at the level of individuals who might have more than one sequela or disease and therefore are not additive.

The cause and sequelae list was expanded based upon feedback after the release of GBD 2013 and input from GBD 2015 collaborators. Nine causes for which non-fatal outcomes are estimated were added: Ebola virus disease, motor-neuron disease, environmental heat and cold exposure, four subtypes of leukaemia, and two subtypes of non-melanoma skin cancer (methods appendix pp 625-53). The incorporation of these changes expanded the cause list from the 301 causes with non-fatal estimates examined in GBD 2013, to 310 causes with non-fatal estimates and from 2337 to 2619 unique sequelae at Level 6 of the hierarchy. At the newly created Level 5 of the hierarchy there were 154 summary sequela categories. The methods appendix (pp 654-61) provides a list of International Classification of Diseases version 9 (ICD-9) and version 10 (ICD-10) codes used in the extraction of hospital and claims data, mapped to GBD 2015 non-fatal causes, impairments, and nature of injury categories.

For data visualisations at vizhub see http://vizhub.

healthdata.org/gbd-compare

Period of analysis

A complete set of age-specific, sex-specific, cause-specific, and geography-specific incidence and prevalence numbers and rates were computed for the years 1990, 1995, 2000, 2005, 2010, and 2015. In this study we focus on trends for main and national results over the past decade, from 2005 to 2015, together with more detailed results for 2015. Online data visualisations at vizhub provide access to results for all GBD metrics.

Non-fatal modelling strategies vary substantially between causes. Figure 1 outlines the general process of non-fatal outcome estimation from data inputs to finalisation of YLD burden results; step 3b of that process identifies alternative modelling approaches used for specific causes (methods appendix pp 603, 604). The starting point for non-fatal estimation is the compilation of data sources identified through systematic analysis and extractions based on predetermined inclusion and exclusion criteria (methods appendix p 603). As part of the inclusion criteria, we defined disease-specific or injury-specific reference case definitions and study methods, as well as alternative allowable case definitions and study methods which were adjusted for if we detected a systematic bias. We used 15 types of primary data sources representing disease prevalence, incidence, mortality risk, duration, remission, or severity in the estimation process (oval shapes in figure 1).

Data sources

For this iteration of the study, we updated data searches through systematic data and literature reviews for 85 causes published up to Oct 31, 2015. For other causes, input from GBD collaborators resulted in the identification and inclusion of a small number of additional studies published after January, 2013. Data were systematically screened from household surveys archived in the Global Health Data Exchange, sources suggested to us by in-country experts, and surveys identified in major multinational survey data catalogues and Ministry of Health and Central Statistical Office websites. Case notifications reported to WHO were updated up to and including 2015. Citations for all data sources used for nonfatal estimation in GBD 2015 are provided in searchable form through a new web tool. A description of the search terms used for cause-specific systematic reviews, inclusion and exclusion criteria, and the preferred and alternative case definitions and study methods are detailed by cause in the methods appendix (pp 26-601).

Hospital inpatient data were extracted from 284 country-year and 976 subnational-year combinations from 27 countries in North America, Latin America, Europe, and New Zealand. Outpatient encounter data were available from the USA, Norway, Sweden, and Canada for 48 country-years. For GBD 2015, we also accessed aggregate data derived from claims information in a database of US private and public insurance schemes for the years 2000, 2010, and 2012. From the linked

claims data, we generated several correction factors to account for bias in health service encounter data from elsewhere, which were largely available to us aggregated by ICD code and by primary diagnosis only. First, for chronic disorders, we estimated the ratio between prevalence from primary diagnoses and prevalence from all diagnoses associated with a claim. Second, we used the claims data to generate the average number of outpatient visits per disorder. Similarly, we generated per person discharge rates from hospital inpatient data in the USA and New Zealand, the only sources with unique patient identifiers available for GBD 2015.

In GBD 2013, we calculated a geographical and temporal data representativeness index (DRI) of non-fatal data sources for each cause or impairment. The DRI represents the fraction of countries for which any incidence, prevalence, remission, or mortality risk data were available for a cause. This metric quantifies data availability, not data quality. The overall DRI and period-specific DRI measures for each cause and impairment are presented in the methods appendix (pp 662-68). DRI ranged from 90% for nine causes, including tuberculosis and measles, to less than 5% for acute hepatitis C and the category of other exposures to mechanical forces. Required case reporting resulted in high DRI values for notifiable infectious diseases; the network of population-based registries for cancers resulted in a DRI of above 50%. DRI values ranged from 6.1% in North Korea to 91.3% in the USA. Many high-income countries, as well as Brazil, India, and China, had DRI values above 63%; data availability was low in several countries, including Equatorial Guinea, Djibouti, and South Sudan.

Non-fatal disease models

In addition to the corrections applied to claims and hospital data, a number of other adjustments were applied including age-sex splitting, bias correction, adjustments for under-reporting of notification data, and computing expected values of excess mortality. In GBD 2013, we estimated expected values of excess mortality from prevalence or incidence and cause-specific mortality rate data for a few causes only, including tuberculosis and chronic obstructive pulmonary disease. In order to achieve greater consistency between our cause of death and non-fatal data, we adopted this strategy systematically for GBD 2015. We matched every prevalence data point (or incidence datapoint for short duration disorders) with the cause-specific mortality rate value corresponding to the age range, sex, year, and location of the datapoint. The ratio of cause-specific mortality rate to prevalence is conceptually equivalent to an excess mortality rate.

To estimate non-fatal health outcomes in previous iterations of GBD, most diseases and impairments were modelled in DisMod-MR, a Bayesian meta-regression tool originally developed for GBD 2010 (step 3a in figure 1).¹⁸ DisMod-MR was designed to address statistical challenges in estimation of non-fatal health outcomes, and for

For Global Health Data Exchange

see http://ghdx.healthdata.org

For **data in GBD 2015** see http:// ghdx.healthdata.org/globalburden-disease-study-2015 synthesis of often sparse and heterogeneous epidemiological data. For GBD 2015, the computational engine of DisMod-MR 2.1 remained unchanged, but we substantially rewrote the code that organises the flow of data and settings at each level of the analytical cascade. The sequence of estimation occurs at five levels: global, superregion, region, country, and where applicable, subnational locations (appendix pp 611–24). At each level of the cascade, the DisMod-MR 2.1 computational engine enforces consistency between all disease parameters. For GBD 2015, we generated fits for the years 1990, 1995, 2000, 2005, 2010, and 2015. We log-linearly interpolated estimates for the intervening years in each 5-year period. Greater detail on DisMod-MR 2.1 is available at Global Health Data Exchange and the methods appendix (pp 7–11).

In previous iterations of GBD, custom models were created for a short list of causes for which the compartment model underpinning DisMod (susceptible, diseased, and dead) was insufficient to capture the complexity of the disease or for which incidence and prevalence needed to be derived from other data. Step 3b of figure 1 describes the development of custom models with greater detail shown in the methods appendix figure 1B (p 604, and for associated write-ups pp 26–601) for HIV/AIDS, tuberculosis, malaria, cancer, neonatal disorders, infectious diseases for which we derived incidence from seroprevalence data, and infectious diseases for which we derived incidence from cause of death rates and pooled estimates of the case fatality proportion.

In GBD 2013, we estimated the country-age-sex-year prevalence of nine impairments (step 4 of figure 1). Impairments in GBD are disorders or specific domains of functional health loss that are spread across many GBD causes as sequelae and for which there are better data to estimate the occurrence of the overall impairment than for each sequela based on the underlying cause. Overall impairment prevalence was estimated with DisMod-MR 2.1 except for anaemia, for which spatiotemporal Gaussian Process regression methods were applied. We constrained cause-specific estimates of impairments, such as in the 19 causes of blindness, to sum to the total prevalence estimated for that impairment. Anaemia, epilepsy, hearing loss, heart failure, and intellectual disability were estimated at different levels of severity.

Severity distributions

In step 5, sequelae were further defined in terms of severity for 194 causes at Level 4 of the hierarchy (figure 1A). We generally followed the same approach for estimating the distribution of severity as in GBD 2013. For Ebola virus disease, we created a health state for the infectious disease episode with duration derived from average hospital admission times, and a health state for ongoing postinfection malaise and joint problems based on four follow-up studies^{19–22} from which we derived an average duration. The health states for the subtypes of leukaemia and non-melanoma skin cancer were the

same as the general cancer health states. For motorneuron disease we accessed the Pooled Resource Open-Access ALS Clinical Trials (PROACT) database containing detailed information on symptoms and impairments for more than 8500 patients who took part in the trials.²³

Disability weights

We used the same disability weights as in GBD 2013 (see methods appendix pp 669–94 for a complete listing of the lay descriptions and values for the 235 health states used in GBD 2015).

Comorbidity

In step 7, we estimated the co-occurrence of different diseases by simulating 40 000 individuals in each geography-age-sex-year combination as exposed to the independent probability of having any of the sequelae included in GBD 2015 based on disease prevalence. We tested the contribution of dependent and independent comorbidity in the US Medical Expenditure Panel Surveys (MEPS) data, and found that independent comorbidity was the dominant factor even though there are well known examples of dependent comorbidity. Age was the main predictor of comorbidity such that age-specific microsimulations accommodated most of the required comorbidity correction. Taking dependent comorbidity into account changed the overall YLDs estimated in the MEPS data by only 2.5% (and ranging from 0.6% to 3.4% depending on age) in comparison to assuming independent comorbidity (methods appendix pp 18-20).24

YLD computation

We report 95% uncertainty intervals (95% UI) for each quantity in this analysis using 1000 samples from the posterior distribution of prevalence and 1000 samples of the disability weight to generate 1000 samples of the YLD distribution. The 95% UI is reported as the 25th and 975th values of the distribution. We report significant changes in disease estimates between countries or over time if the change was noted in more than 950 of the 1000 samples computed for each result. For GBD 2015, we computed age-standardised prevalence YLD rates from the updated world population age standard developed for GBD 2013.25 Less common diseases and their sequelae were included in 35 residual categories (methods appendix pp 695–97). For 22 of these residual categories, estimates were made from epidemiological data for incidence or prevalence. For 13 residual categories, we estimated YLDs by multiplying the residual YLL estimates by the ratio of YLDs to YLLs from the estimates for explicitly modelled Level 3 causes in the same disease category.

Socio-demographic Index

In GBD 2013, a sociodemographic status variable was computed based on a principal components analysis of income per capita, educational attainment, average age of the population, and the total fertility

For **DisMod-MR 2.1 engine and the code** see http://ghdx. healthdata.org/global-burdendisease-study-2015

1550

	Incidence (thousands)		Percentage change (%)
	2005	2015	_
Upper respiratory infections	15 624 257	17 230 659	10·3
	(13 851 237-17 411 199)	(15 351 516-19 172 439)	(9·0 to 11·6)*
Diarrhoeal diseases	2 2 3 5 7 3 9	2392517	7·0
	(2 1 3 9 0 5 0 - 2 3 4 8 4 0 7)	(2301101—2503094)	(6·0 to 8·0)*
Permanent caries	426 963	487 629	14·2
	(371 216-484 332)	(423 507-552 895)	(13·1 to 15·4)*
Otitis media	429 820	471 027	9.6
	(353 915-525 059)	(386 606–577 286)	(7.9 to 11.2)*
Lower respiratory infections	273 131	291759	6.8
	(257 286–288 078)	(276 244–307 004)	(5.6 to 8.1)*
Malaria	339 275	286 859	-15·4
	(261 417-447 605)	(219 712-377 332)	(-23·0 to -8·1)*
Gastritis and duodenitis	185 250	213729	15·4
	(167 471–204 974)	(192486–236339)	(10·8 to 17·2%)*
Pyoderma	178 382	207 452	16·3
	(172 199–184 147)	(200 498–213 936)	(15·6% to 17·0)*
Gonococcal infection	138 220	172 676	24·9
	(107 106–184 385)	(129 731-235 737)	(17·9 to 31·0)*
Interstitial nephritis and urinary tract infections	127 380	152 295	19·6
	(124 308 – 130 683)	(148 748–156 177)	(19·2 to 19·9)*
Varicella and herpes zoster	128 678	142 413	10·7
	(124 298–133 090)	(137 804–147 181)	(10·0 to 11·4)*
Trichomoniasis	121 948	140781	15·4
	(104 825–141 284)	(121207-163163)	(14·5 to 16·5)*
Acute hepatitis A	109 609	114212	4·2
	(101 813–117 648)	(103349-124776)	(-7·2 to 17·0)
Hepatitis B	98 277	111 212	13·2
	(86 507-112 527)	(97 410–126 251)	(-4·3 to 35·2)
Gallbladder and biliary diseases	88 215	104322	18·3
	(79 276–96 495)	(93074-114430)	(16·0 to 20·7)*
Peptic ulcer disease	83 388	87 410	4·8
	(77 760 – 89 435)	(80 343–94 506)	(2·6 to 7·0)*
Dengue	32749	79 609	143·1
	(18879–68335)	(53 784–169 704)	(-0·3 to 564·7)
Other sense organ diseases	60 659	69 945	15·3
	(58 721-62 579)	(67 856–72 080)	(14·6 to 16·0)*
Chlamydial infection	56 976	61 173	7·4
	(45 489–70 839)	(48 871-76 698)	(5·5 to 9·4)*
Maternal abortion, miscarriage, and ectopic pregnancy	53 942 (43 630–67 168)	53 958 (43 224-67 417)	0·0 (-3·8 to 4·0)
Syphilis	43 515	45 413	4·4
	(37 479-51 054)	(37 787-54 921)	(-0·3 to 8·1)
Deciduous caries	41353	43 688	5.6
	(28723-58131)	(29 630-62 543)	(2.1 to 8.1)*
Genital herpes	29 112	39 791	36·7
	(25 131-33 432)	(35 569-44 572)	(32·4 to 41·6)*
Urolithiasis	17 875	22 080	23.5
	(16 320-19 728)	(20 183-24 295)	(21.2 to 25.6)*
Cellulitis	17312	21 211	22·5
	(15 988-18739)	(19 582–22 985)	(21·0 to 24·1)*
Maternal hypertensive disorders	20 416	20731	1·5
	(17 593-23 417)	(17355-24379)	(−3·0 to 6·3)
Acute hepatitis E	18 869	19 525	3·5
	(17 340-20 580)	(18 011-21 273)	(2·2% to 4·7)*
Whooping cough	22 457	16 298	-27·4
	(17 322–28 268)	(12 599-20 445)	(-29·5 to -25·2)*
	,		continues on next pag

rate.²⁶ For GBD 2015, we excluded mean age of the population because it is directly affected by death rates. To improve interpretability for GBD 2015, we computed a Socio-demographic Index (SDI) similar to the computation of the human development index.²⁷ In the SDI, each component was weighted equally and rescaled from zero (for the lowest value observed during 1980–2015) to one (for the highest value observed) for income per capita and average years of schooling, and the reverse for the total fertility rate. The final SDI score was computed as the geometric mean of each of the components. SDI ranged from 0.060 in Mozambique in 1987 to 0.978 in District of Columbia, USA, in 2015.

Role of the funding source

The funder of the study had no role in study design, data collection, data analysis, data interpretation, or writing of the report. The corresponding author had full access to all the data in the study and had final responsibility for the decision to submit for publication.

Results

Global incidence and prevalence

We generated over 9·3 billion outcomes of incidence, prevalence, and YLDs for 310 diseases, injuries, and aggregate categories; 2619 unique and aggregate sequelae; nine impairments; 63 age–sex groups; 591 geographies; and 26 individual years from 1990 to 2015. Each of these 9·3 billion estimates was calculated 1000 times to determine uncertainty intervals. Here, we present key summary findings on global incidence of short duration diseases, global prevalence of long-term disorders, global prevalence of impairments, global numbers and rates of YLDs and changes from 2005 to 2015, global YLL and YLD rates of change, patterns of comorbidity, the expected changes in the composition of YLDs with SDI, and country findings of leading causes of YLDs.

Disorders of less than 3 months duration and injuries with incidence of more than 1 million cases per year in 2015 are listed in table 1. There were two disorders with incidence greater than 1 billion per year: upper respiratory infections (17·2 billion [95% UI 15·4–19·2 billion]), and diarrhoeal diseases (2·39 billion [2·30–2·50 billion]). A further 13 diseases and injuries caused between 100 million and 1 billion incident cases a year and 16 diseases and injuries had incident cases of between 10 million and 100 million per year (table 1).

The disease and injury sequelae with a duration of more than 3 months and a global prevalence of more than 1% in 2015 are presented in table 2, aggregated to the cause level. Prevalence for impairments is presented at the bottom of table 3. Eight out of 56 high-prevalence causes affected more than 10% of the world's population in 2015. A further 48 causes affected between 1% and 10% of the world's population (table 2). Although many of

these causes are not among the dominant causes of YLDs because of comparatively low average disability weights, some causes, such as headaches, gynaecological diseases, oral disorders, and skin diseases, put great demands on health system resources by their sheer numbers.

Anaemia was the most common of our nine impairments, affecting 2.36 billion (2.35-2.37 billion) people in 2015. The next most common impairments were hearing loss of greater than 20 dB (1.33 billion [1.26-1.40 billion]), vision loss (940 million [905–974 million]), developmental intellectual disability (153 million [114–191 million]), infertility (113 million [93·4–136 million]), heart failure (40·0 million [38·6-41·4 million]), and epilepsy (39·2 million [34·3–43·7 million]; table 3; see results appendix (pp 740–48) for the prevalence estimates of the underlying causes of these impairments). Iron deficiency was the cause of anaemia in more than half of all cases. Over 90% of hearing loss was classified as age-related or other hearing loss. The largest number of people with vision loss had uncorrected refraction error. Idiopathic developmental intellectual disability, idiopathic female infertility, and idiopathic epilepsy were the most common causes of their impairments. Ischaemic heart disease was the most common cause of heart failure.

Global causes of disability

Global trends in YLDs 2005 to 2015

GBD 2015 included the assessment of 2619 sequelae at Level 6 of the GBD cause hierarchy, including 1316 sequelae from injuries that contributed to the global burden of disability. Causes at Level 4 of the hierarchy that resulted in 30 million or more YLDs in 2015 included lower back pain, major depressive disorder, age-related and other hearing loss, and neck pain. Figure 2 compares the leading causes of global YLDs in 2005 and 2015, using the cause breakdowns at Level 3 of the GBD cause hierarchy. Among the ten leading causes of YLDs, iron-deficiency anaemia and depressive disorders switched ranks to positions three and four respectively, diabetes rose from the eighth to the sixth position, migraine dropped from position six to seven, and other musculoskeletal disorders dropped from rank seven to eight (figure 2).

Estimates of prevalence and YLDs at the global level for 2005 and 2015 for each cause are presented in table 3 (see results appendix pp 717–40 for full detail at the sequelae level). Prevalence and age-standardised YLDs for 21 Group 1 diseases decreased significantly and by more than 10%, including measles, African trypanosomiasis, diphtheria, lymphatic filariasis, and rabies.

Age-standardised YLD rates for all maternal causes and sequelae combined decreased between 2005 and 2015, whereas overall age-standardised YLDs for neonatal disorders increased by 3 · 11% (-0 · 32-6 · 79%) since 2005 to 144 YLDs per 100 000 (109–185 YLDs per 100 000) in 2015, and age-standardised YLD rates decreased and absolute numbers of cases increased for nutritional deficiencies.

	Incidence (thousand	Percentage change (%)	
	2005	2015	
(Continued from previous pa	ge)		
Typhoid fever	15 341	12538	-18·3
	(13 454-17 417)	(10887-14283)	(-20·8 to -15·6)*
Maternal sepsis and other maternal infections	11 938	11 817	-1·0
	(9457-15 125)	(9300–15 009)	(-4·6 to 2·8)
Appendicitis	10 159	11 619	14·4
	(9549 to 10 840)	(10 918 to 12 407)	(13·5 to 15·2)*
Pancreatitis	7160	8 902	24·3
	(6 660-7 694)	(8 212-9 643)	(21·9 to 26·6)*
Maternal haemorrhage	8438	8740	3.6
	(6614-10620)	(6774–11 061)	(-0.3 to 7.6)
Other sexually transmitted diseases	7569	7656	1·2
	(6511-8741)	(6583-8853)	(-0·1 to 2·4)
Ischaemic heart disease	6308	7287	15·5
	(5918-6716)	(6798-7808)	(14·3 to 16·8)*
Maternal obstructed labour and uterine rupture	6830	6521	-4·5
	(5202-8933)	(4977–8588)	(-9·1 to -0·3)*
Hepatitis C	4609	5394	17·0
	(4316 to 4918)	(5032 to 5778)	(15·7 to 18·4)*
Ischaemic stroke	4651	5385	15·8
	(4397 to 4912)	(5017 to 5726)	(13·4 to 18·4)*
Measles	15 610	4652	-70·2
	(6699-31 357)	(2072–9206)	(-73·6 to -66·0)*
Paratyphoid fever	5580	4587	-17·8
	(4498-6896)	(3738-5601)	(-22·6 to -12·7)*
Haemorrhagic stroke	3144	3583	14·0
	(2973–3307)	(3336-3822)	(11·4 to 16·2)*
Paralytic ileus and intestinal obstruction	2664	3167	18·9
	(2494–2855)	(2946-3406)	(16·9 to 20·7)*
Encephalitis	1489	1603	7·7
	(1380–1608)	(1472–1750)	(5·9 to 9·5)*
Acute glomerulonephritis	1528	1534	0·4
	(1395–1672)	(1394–1685)	(-2·6 to 3·1)

Data in parentheses are 95% UIs. *Percentage changes that are statistically significant.

Table~1: Global incidence~of~short~duration~(less~than~3~months)~sequelae~in~2005~and~2015~for~all~ages~and~both~sexes~combined,~with~percentage~change~between~2005~and~2015~for~level~4~causes~with~incidence~greater~than~1~million~cases~per~year~

Age-standardised YLD rates attributable to hepatitis A, B, and C increased, and decreased for hepatitis E. The number of individuals with chronic hepatitis C infection increased from 121 million (108–133 million) in 2005 to 142 million (127–157 million) in 2015.

$Changes\ in\ age-standardised\ YLDs\ and\ YLLs\ over\ time$

In 2005, non-communicable diseases (NCDs) accounted for 23 of the leading 25 causes of age-standardised YLDs worldwide and 23 of the 25 leading causes in 2015 (figure 2). Although diabetes rose only two ranks in the list of leading cause of YLDs, from position eight to six between 2005 and 2015, the increase in age-standardised rate was 5.4% (3.2–7.5%). Musculoskeletal disorders occupied three of the leading 25 causes of disability in both 2005 and 2015; lower back and neck pain were the single largest cause with little change in their rates.

	Prevalence (thousands)	Percentage change (%)	
	2005	2015	_
Permanent caries	2 045 859	2344628	14·6
	(1 909 845-2 170 355)	(2193751-2488741)	(13·7 to 15·5)*
Tension-type headache	1306390	1505892	15·3
	(1160423-1463889)	(1337310-1681575)	(14·0 to 16·6)*
Iron-deficiency anaemia	1456387	1477531	1·5
	(1449846-1462782)	(1470902-1485322)	(0·9 to 2·1)*
Age-related and other hearing loss	943 504	1210 055	28·3
	(886 325-995 403)	(1140 224-1274 665)	(27·4 to 29·2)*
Migraine	831726	958789	15·3
	(755 991-918 968)	(872109-1055631)	(14·0 to 16·6)*
Genital herpes	716 115	845 826	18·1
	(626 987-817 387)	(736 724 – 968 066)	(16·4 to 19·9)*
Refraction and accommodation disorders	672165	819 307	21·9
	(649969-694024)	(789 917-848 059)	(20·5 to 23·2)*
Ascariasis	854489	761894	-10·8
	(790000-924895)	(682558-861031)	(-22·5 to 2·5)
G6PD trait	663704	728 549	9·8
	(625032-703799)	(676 735-781 534)	(7·9 to 11·4)*
Acne vulgaris	605 008	632741	4·6
	(568 577-642 061)	(595242-671249)	(3·5 to 5·6)*
Other skin and subcutaneous diseases	492 883	605 036	22·8
	(480 852-505 426)	(589 500-619 676)	(22·1 to 23·4)*
Deciduous caries	534122	558 028	4·5
	(449525-635866)	(462 649-669 027)	(2·3 to 6·1)*
Low back pain	460164	539 907	17·3
	(444680-477119)	(521 449–559 556)	(16·5 to 18·2)*
Periodontal diseases	428784	537 506	25·4
	(372 953-498 682)	(465 114-625 889)	(24·1 to 26·5)*
Fungal skin diseases	434 604	492 373	13·3
	(395 512-475 904)	(448 951-538 232)	(12·5 to 14·1)*
Trichuriasis	473 399	463 652	-2·1
	(443 689-505 928)	(426 621–502 939)	(-12·0 to 9·0)
Diabetes	333325	435328	30·6
	(310773–355510)	(404736-468562)	(28·0 to 33·0)*
Premenstrual syndrome	391207	430 697	10·1
	(375009-407896)	(410 841-450 494)	(7·9 to 11·8)*
Hookworm disease	462 111	428 246	-7·3
	(430 885-495 596)	(394 486-468 292)	(-16·9 to 3·5)
Sickle-cell trait	338756	404 566	19·4
	(318736-378582)	(381 223-448 155)	(18·3 to 20·3)*
Asthma	327 097	358 198	9·5
	(296 406-358 060)	(323 134-393 466)	(7·6 to 11·6)*
Neck pain	295 532	358 007	21·1
	(258 878-338 138)	(313 408-409 411)	(19·0 to 23·3)*
Hepatitis B	293745	343 251	16·9
	(284478-303036)	(330 541-357 195)	(15·3 to 18·4)*
Other musculoskeletal disorders	283 317	342 068	20·7
	(254 135-315 519)	(305 431-385 147)	(17·5 to 24·0)*
Urolithiasis	259 567	318763	22·8
	(238 100-281 892)	(290695349154)	(20·8 to 24·9)*
Thalassaemias trait	252798	279 451	10·5
	(247008-259972)	(272 819-287 357)	(10·1 to 11·0)*
Malaria	207773	278 961	34·3
	(186530-230942)	(240 158-320 921)	(26·8 to 42·3)*
Edentulism and severe tooth loss	216 473	275 619	27·3
	(207 563-226 331)	(264 201–288 252)	(26·9 to 27·7)*
Anxiety disorders	232 597	267202	14·9
	(204 165-264 445)	(234064-306318)	(13·0 to 16·8)*
		(Table 2	continues on next page)

However, age-standardised rates of YLDs increased for osteoarthritis 3.90% (3.00–4.83%) by 2015. Depressive disorders were the fourth leading cause of disability in 2005 and the third leading cause of disability in 2015, and age-standardised YLD rates associated with the disorder increased marginally (1.0% [0.5–1.5%]). Age-standardised rates of YLDs from alcohol use disorders decreased after 2005 (4.5% [2.3–6.4%]) whereas disability due to drug use disorders increased by 8.2% (6.2–10.2%).

In contrast with global trends for NCDs, both relative ranks and age-standardised YLD rates decreased for most injuries. Falls, which were the 13th leading cause of disability in 2005, dropped to the 15th rank, and age-standardised YLD rates decreased (8·58% [5·23–12·2%]). Other unintentional injuries decreased in global rank from 27th in 2005 to 34th in 2015, and age-standardiSed YLD rates decreased by 16·7% (15·7–17·9%).

The leading causes of disability varied considerably with age (figure 3). The leading cause in children younger than 5 years was iron-deficiency anaemia followed by skin diseases, protein-energy malnutrition, and diarrhoea. In older children, iron-deficiency anaemia, skin diseases, asthma, and mental health disorders such as conduct, autistic spectrum, and anxiety disorders were top ten causes of disability. In adolescents and young adults (aged between 15 and 39 years), iron-deficiency anaemia, skin diseases, depression, lower back and neck pain, and migraine led the rankings. Other mental health disorders such as anxiety disorders and schizophrenia were in the top ten causes in this age group. In middle-aged adults, musculoskeletal disorders dominated the top rankings followed by mental health disorders, especially depression. Diabetes and sense organ disorders were more prominent causes of disability in middle-age. In older adults (older than 65 years), sense organ disorders were the top-ranked cause of disability. Musculoskeletal disorders remained a dominant source of disability, and chronic obstructive pulmonary disease entered the top ten. In the oldest age groups, ischaemic heart disease and Alzheimer's and other dementias made their first appearance in the top ten.

We examined the trends in YLDs and YLLs in a scatterplot (figure 4). YLLs decreased for the majority of causes. For Group 1 causes and injuries, the decrease in YLLs was accompanied by a decrease in YLDs, albeit at a slower pace. The exceptions were neonatal encephalopathy, haemolytic disease and other neonatal jaundice, leishmaniasis, meningitis, hepatitis, and sexually transmitted diseases with increasing YLD rates between 1990 and 2015. A few Group 1 disorders and injuries had a faster decrease in YLDs than in YLLs: intestinal infections. obstructed labour, and neonatal sepsis. The only NCDs with a faster decrease in YLDs compared with YLLs were epilepsy and cervical cancer. Another small number of NCDs saw an increase in YLDs and YLLs, including drug use disorders, diabetes, and Parkinson's disease. NCDs with decreasing YLLs but increasing YLDs included

cancers of the prostate, testis, uterus, kidney, colorectum, and pancreas, melanoma, and congenital disorders. Some cancers (stomach and Hodgkin's lymphoma), rheumatic heart disease, asthma, chronic obstructive pulmonary disease, acute glomerulonephritis, peptic ulcer disease, gastritis, hernia, and gallbladder disease had decreasing YLLs and YLDs, but with a faster decrease in YLLs than in YLDs. The rate of change in YLDs for the main drivers of non-fatal health loss, musculoskeletal disorders, and mental and substance use disorders was small.

Global distribution of disability weights across individuals

Figure 5 shows the global distribution of individuals from our comorbidity microsimulation by six categories of disability, age, and sex for the highest and lowest SDI quintile. The six categories of disability are no disability, very mild disability (disability weight less than or equal to 0.01), mild disability (from 0.01 to 0.05 inclusive), moderate (from 0.05 to 0.1 inclusive), severe (from 0.1 to 0.3 inclusive), and profound (greater than 0.3). In 2015, most of the world's population experienced mild or greater disability. Having no disability at all was most common in children. After age 25 years, the proportion of the population having no disability became progressively smaller, and by age 55 years in low SDI countries and age 75 years in high SDI countries, nearly everyone had some form of disability. Very mild to moderate disability (ie, individuals with a disability weight of 0.1 or less) was common in childhood and young adults, but was replaced by more severe disability with increasing age. The patterns were similar for both sexes, apart from a much larger amount of disability in women older than 80 years in the top SDI quintiles, reflecting the much higher average age of women in this age category. For policy considerations around the age of retirement in ageing populations, it is noteworthy that from age 60 years onwards more than half of the population had severe or worse disability. The extent to which this loss of health limits or precludes the ability to work depends on the nature of the impairments and the type of employment in older workers with that level of disability.

Expected changes in disease profile with higher Socio-demographic Index

Figure 6 depicts changes in patterns of disability by level of SDI for age-standardised and all-age YLD rates per 100 000. Age-standardised YLD rates gradually decreased with increasing SDI in both sexes (figure 6A). The cause composition of YLDs somewhat varied across levels of SDI; these differences were largely derived from absolute levels of disability due to communicable causes and nutritional deficiencies, and to a lesser extent, maternal and neonatal disorders. Age-standardised YLD rates due to NCDs and injuries were similar at all SDI levels.

Across levels of SDI, mental and substance use disorders, musculoskeletal disorders, and other NCDs were consistently among the leading causes of

	Prevalence (thousands	Percentage change (%)	
	2005	2015	` `
(Continued from previous pa	age)		
Other sense organ diseases	214761	266 346	24·0
	(207 401–222 622)	(257 047-276 383)	(23·1 to 24·9)*
Schistosomiasis	329 773	252 340	-23·5
	(297 093-367 878)	(211 032–321 081)	(-32·5 to -4·4)*
G6PD deficiency	231109	247 074	6·9
	(205067-259769)	(209 307-286 713)	(1·4 to 11·9)*
Dermatitis	215 260	245 291	14·0
	(199 590–230 536)	(227 283-262 752)	(13·2 to 14·8)*
Osteoarthritis	178 665	237 369	32·9
	(173 558-184 053)	(230 336-244 648)	(31·9 to 33·8)*
Major depressive disorder	183 434	216 047	17·8
	(163 947-206 420)	(192 863-243 319)	(16·6 to 19·0)*
Scabies	191482	204 152	6·6
	(166101-223739)	(177 534-237 466)	(4·0 to 9·5)*
Viral skin diseases	161 167	174 843	8·5
	(152 218-170 651)	(165 156-185 072)	(8·0 to 9·0)*
Chronic obstructive pulmonary disease	149 115	174 483	17·0
	(137 380-160 739)	(160 205-188 952)	(15·1 to 19·0)*
Genital prolapse	137 383	161 679	17·7
	(121 623-154 875)	(142 335-182 566)	(15·2 to 20·0)*
Gastritis and duodenitis	135 993	157 060	15·5
	(134 420-137 351)	(154 055-160 141)	(12·9 to 17·9)*
Peripheral vascular disease	115 109	154 651	34·4
	(101 439-131 405)	(136 318-176 211)	(33·4 to 35·2)*
Uterine fibroids	126797	151115	19·2
	(120 900–133 050)	(144147-158477)	(18·8 to 19·5)*
Hepatitis C	120 457	142 123	18⋅0
	(108 080-133 129)	(126 978-157 045)	(16⋅7 to 19⋅2)*
Other mental and substance use disorders	107895	128 178	18·8
	(107213-108449)	(127 512-128 877)	(17·9 to 19·7)*
lodine deficiency	103701	110 920	7·0
	(93441-116438)	(100 337-125 253)	(4·7 to 9·4)*
Ischaemic heart disease	87 511	110 193	25·9
	(80 133-96 170)	(100 332-121 427)	(24·6 to 27·2)*
Benign prostatic	80 684	104 625	29·7
hyperplasia	(70 338-90 853)	(90 730-118 244)	(27·5 to 32·0)*
Dysthymia	86 812	104 106	19·9
	(74 974-99 103)	(90 398-118 969)	(18·4 to 21·5)*
Chronic kidney disease due to diabetes	79 184	100 824	27·3
	(68 481–90 737)	(86 923-115 652)	(24·9 to 29·9)*
Chronic kidney disease due to other causes	74 917	94553	26·2
	(64 012-86 576)	(81142-109371)	(24·2 to 28·3)*
Idiopathic developmental intellectual disability	82 996	92 074	10·9
	(47 588-117 109)	(52 280-130 411)	(9·5 to 11·9)*
Other cardiovascular and circulatory diseases	72772	90 348	24·2
	(68 090-77763)	(84 711-96 659)	(22·7 to 25·5)*
Otitis media	83 022	86 393	4·1
	(73 657-93 266)	(76 374-97 104)	(2·7% to 5·5)*
Psoriasis	67753	79 700	17·6
	(65107-70298)	(76 691-82 804)	(17·0 to 18·3)*
Other haemoglobinopathies and haemolytic anaemias	73 045 (72 600-73 470)	74385 (73898-74862)	1·8 (0·9 to 2·7%)

Data in parentheses are 95% UIs. *Percentage changes that are statistically significant.

Table 2: Global prevalence of longer duration (more than 3 months) sequelae in 2005 and 2015 for all ages and both sexes combined, with percentage change between 2005 and 2015 for Level 4 causes with prevalence greater than 1%

age-standardised YLD rates. Anaemia led to generally higher rates of age-standardised YLDs in women than in men across levels of SDI, but the largest imbalance occurred at SDI levels between $0\cdot10$ and $0\cdot50$. Disability from injuries exacted a larger burden for men than for women, particularly at lower levels of SDI.

Without adjustments for population age structure (figure 6B), the effect of ageing populations and causes of disability that disproportionately affect older individuals become prominent. At all levels of SDI, total YLDs per 100 000 did not notably differ by sex; instead, the cause composition of disability showed greater differences. Below an SDI score of 0.25, communicable causes accounted for 30-45% of total disability, primarily due to nutritional deficiencies, malaria, and neglected tropical diseases. YLDs per 100 000 due to musculoskeletal disorders, particularly lower back and neck pain and other musculoskeletal disorders, increased substantially from low to high SDI, with a more pronounced increase beginning at an SDI score of 0.6. This rise was particularly evident in women.

Trends in age-standardised YLDs per capita

Globally, age-standardised YLDs per capita (an indicator of overall disability experienced per person in a given place) moderately decreased for both sexes between 1990 and 2015 (results appendix pp 714–16). Age-standardised YLDs per capita were consistently higher for women than for men at the global level. For both sexes, YLDs per capita were generally higher for lower levels of SDI. YLDs per capita were noticeably larger for low SDI and low-middle SDI groups than for other SDI levels (ie, high SDI, high-middle SDI, and middle SDI), which were more similar to each other.

Leading causes of YLDs and deviations from expected levels based on Socio-demographic Index

Clear, though varied, patterns emerged across and within GBD regions in comparison of observed levels of YLDs due to leading causes of disability with levels expected on the basis of SDI. Figure 7 displays ratios of observed and expected YLDs for the leading ten causes at level 3 of the GBD hierarchy in 2015, colour coded by the magnitude of differences between observed and expected YLDs.

Globally, lower back and neck pain was the leading cause of disability in 2015. Two mental disorders, major depressive and anxiety disorders, were the third and ninth leading causes of global disability, and diabetes was the sixth leading driver of disability. Iron-deficiency anaemia was the only Group 1 cause among the leading ten causes for global YLDs (ranked fourth). Sensory disorders ranked second and skin diseases ranked fifth. Lower back and neck pain was the leading global cause of disability in 2015 in most countries. The leading cause of disability in 2015 was iron-deficiency anaemia in 27 countries; HIV/AIDS in all six southern sub-Saharan

African countries; depression in five eastern sub-Saharan Africa countries; other neglected tropical diseases in Angola, Democratic Republic of the Congo, and Gabon; sense organ disorders in Comoros and Myanmar; diabetes in Fiji and Marshall Islands; war in Lebanon and Syria; and onchocerciasis in Liberia.

Regional, country, territory, and selected subnational results Lower back and neck pain was the leading cause of disability in all high-income countries in 2015. However, ratios of observed to expected YLDs from lower back and neck pain ranged from 0·60 for Singapore to more than 1·59 in Norway. Most high-income countries experienced higher than expected levels of disability due to depressive disorders. The USA and Australia were the only two high-income countries where drug use disorders were a top ten cause of disability, and observed levels of YLDs were much higher than expected. South Korea's ratio of observed-to-expected levels of YLDs due to diabetes exceeded 1·30, whereas Japan's diabetes-related disability was lower than expected on the basis of SDI. In the UK, observed disability due to asthma was well above expected levels.

In 2015, lower back and neck pain was the leading cause of YLDs for all but two countries in Latin America and the Caribbean; Haiti and Venezuela were the exceptions, with iron-deficiency anaemia as the leading causes of disability for both countries. Disability due to diabetes surpassed expected levels by at least a factor of two for six countries and territories: Antigua, Barbados, Dominica, Puerto Rico, Trinidad and Tobago, and the Virgin Islands. Peru, however, had a ratio of less than $0\cdot 6$ for observed and expected YLDs for diabetes.

In 2015, lower back and neck pain was the leading cause of YLDs for 24 out of 28 countries and territories of southeast Asia, east Asia, and Oceania. Other leading causes of disability were sensory disorders in Myanmar, diabetes in Fiji and the Marshall Islands, and iron-deficiency anaemia in Papua New Guinea. Although lower back and neck pain was the primary cause of disability, many countries had far lower levels of this than expected given their SDI, including Thailand (0.73), Indonesia (0.76), and Malaysia (0.75). Observed YLDs due to depressive disorders were often lower than expected, with 22 countries recording ratios below 0.80. Conversely, numerous geographies recorded YLD ratios exceeding 2.00 for diabetes (eg, 2.29 in Taiwan).

Beyond lower back and neck pain, which was the leading cause of YLDs for three of five countries in south Asia in 2015 (with India and Pakistan being the exception), a mixture of causes accounted for the region's main causes of disability; this heterogeneity probably reflects the diversity of countries in the region and their places along the development spectrum. Iron-deficiency anaemia was the first leading cause and lower back and neck pain was the second leading cause of YLDs in both India and Pakistan, whereas sensory disorders, other musculoskeletal disorders, and iron-deficiency anaemia ranked second for

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change ASR between 2005 and 2015
ll causes Communicable, maternal, neonatal, and nutritional diseases	 4133 822·3 (4108 132·5 to 4161262·6)	 8·0 (7·3 to 8·7)*	 -4·2 (-4·8 to -3·6)*	792004·7 (588538·7-1019955·2) 112501·9 (80347·4-156270·9)	15·1 (14·5 to 15·6)* -0·5 (-2·4 to 2·5)	-2·1 (-2·5 to -1·7)* -9·7 (-11·4 to -7·0)*
HIV/AIDS and tuberculosis	46 005·2	18·6	2·9	6702·3	2·1	-12·2
	(44 985·8 to 47 146·6)	(15·2 to 22·1)*	(-0·0 to 5·9)	(4798·6–8863·5)	(-3·5 to 7·2)	(-17·0 to -7·8)*
Tuberculosis	8861·2	9·2	-7·2	2712·4	9·4	-6·8
	(8076·3 to 9707·2)	(6·3 to 12·2)*	(-9·5 to -4·7)*	(1829·9–3744·0)	(6·4 to 12·5)*	(-9·2 to -4·2)*
HIV/AIDS	37 277·5	21·1	5·6	3989·9	-2·4	-15·6
	(36 279·9 to 38 339·1)	(16·9 to 25·5)*	(2·0 to 9·6)*	(2891·2-5159·5)	(-10·5 to 5·5)	(-23·0 to -8·5)*
HIV/AIDS—tuberculosis	1258·0	-17·9	-28⋅3	471·9	-17·8	-28·1
	(1142·5 to 1386·9)	(-20·7 to -14·9)*	(-30⋅7 to -25⋅7)*	(316·0–645·6)	(-20·7 to -14·7)*	(-30·6 to -25·5)*
HIV/AIDS resulting in other diseases	37543·3	23·6	7·9	3518·0	0·2	-13·6
	(36350·0 to 39028·4)	(20·5 to 26·7)*	(5·1 to 10·6)*	(2 555·4-4 558·9)	(-9·3 to 9·3)	(-22·1 to -5·2)*
HIV aggregate	21149·0	-19·5	-29·7	1543·6	-20·4	-30·4
	(20260·3 to 22292·3)	(-22·5 to -16·3)*	(-32·3 to -26·9)*	(1 107·2-2 025·2)	(-23·4 to -17·2)*	(-33·1 to -27·6)*
AIDS aggregate	16 394·3	299·5	232·9	1974·4	25·5	6·1
	(15 747·0 to 17 154·2)	(251·4 to 355·5)*	(193·5 to 279·0)*	(1432·0–2559·9)	(0·9 to 56·1)*	(-14·6 to 32·4)
Diarrhoea, lower respiratory infections, and other common infectious diseases	402 320·4 (393 746·5 to 408 994·2)	8·0 (7·2 to 8·8)*	-2·9 (-3·7 to -2·3)*	14865·0 (10397·0-20283·4)	6·2 (5·4 to 7·1)*	-4·0 (-4·7 to -3·3)*
Diarrhoeal diseases	35 820·6	6·4	-3·3	5731·7	6·4	-3·1
	(34 342·1 to 37 537·8)	(5·5 to 7·3)*	(-4·1 to -2·5)*	(3 943·3-7 890·5)	(5·4 to 7·5)*	(-4·0 to -2·2)*
Diarrhoea episodes	35 816·3	6·4	-3·3	5730·4	6·4	-3·1
aggregate	(34 337·5 to 37 534·1)	(5·5 to 7·3)*	(-4·1 to -2·5)*	(3 942·2–7 889·0)	(5·4 to 7·5)*	(-4·0 to -2·2)*
Guillain-Barré syndrome	4·2	16·9	-0·2	1·3	16·9	-0·2
due to diarrhoeal diseases	(3·2 to 5·5)	(14·7 to 19·5)*	(-0·9 to 0·4)	(0·8–1·9)	(14·7 to 19·5)*	(-0·9 to 0·4)
Intestinal infectious diseases	1666·8	-23·3	-28·2	220·3	-18·6	-24·2
	(1594·4 to 1732·8)	(-25·6 to -20·6)*	(-30·4 to -25·8)*	(149·3–306·5)	(-22·8 to -13·8)*	(-28·1 to -19·8)*
Typhoid fever	1446·7	-18·3	-23·9	191·0	-17·7	-23·2
	(1256·2 to 1 648·1)	(-20·8 to -15·6)*	(-26·3 to -21·3)*	(129·5–267·8)	(-22·1 to -12·6)*	(-27·4 to -18·5)*
Typhoid fever episodes aggregate	1197-9	-18·3	-23·9	113·3	-18·1	-23·7
	(1041-3 to 1366-6)	(-21·7 to -14·8)*	(-27·1 to -20·5)*	(75·8–162·5)	(-22·6 to -13·4)*	(-27·8 to -19·2)*
Complications of typhoid fever aggregate	248·8	-18·2	-23⋅8	77·8	-17·0	-22·6
	(210·8 to 292·2)	(-28·6 to -5·4)*	(-33⋅5 to -12⋅1)*	(51·6–110·8)	(-28·5 to -3·0)*	(-33·1 to -9·9)*
Paratyphoid fever	529·3	-17·8	-23·8	27·5	-17·5	-23·5
	(431·3 to 646·3)	(-22·6 to -12·7)*	(-28·2 to -19·2)*	(17·5–40·7)	(-24·7 to -9·2)*	(-30·1 to -16·1)*
Paratyphoid fever episodes aggregate	502·2	-17·8	-23·8	24·4	-17·5	-23·5
	(409·6 to 612·7)	(-22·6 to -12·4)*	(-28·2 to -19·0)*	(15·5–36·4)	(-24·9 to -8·7)*	(-30·7 to -15·5)*
Intestinal perforation	27·1	-17·6	-23·7	3·1	-17·5	-23⋅6
due to paratyphoid	(20·9 to 34·5)	(-33·0 to 1·4)	(-37·8 to -6·1)*	(1·9–4·6)	(-32·8 to 1·6)	(-37⋅7 to -5⋅9)*
Other intestinal infectious diseases				1·8 (0·6-4·1)	-67·0 (-78·8 to -45·8)*	-69·3 (-80·2 to -50·1)*
Lower respiratory infections	8986·6	5·1	-8·2	540·4	4·9	-8·1
	(8545·2 to 9393·7)	(4·0 to 6·2)*	(-9·1 to -7·3)*	(365·3–760·3)	(3·5 to 6·3)*	(-9·2 to -7·0)*
Lower respiratory infection episodes aggregate	8982·2	5·1	-8·2	539·1	4·9	-8·1
	(8539·8 to 9387·9)	(4·0 to 6·2)*	(-9·1 to -7·3)*	(364·5-759·5)	(3·5 to 6·3)*	(-9·2 to -7·0)*
Guillain-Barré syndrome due to lower respiratory infections	4·4 (2·5 to 6·9)	16·9 (14·7 to 19·5)*	-0·2 (-0·9 to 0·4)	1·3 (0·7-2·2)	16·9 (14·7 to 19·5)*	-0·2 (-0·9 to 0·4)
Upper respiratory infections	233 470·3	10·2	-1·4	2738·4	10·1	-1·3
	(208 009·7 to 259 076·8)	(8·9 to 11·5)*	(-2·0 to -0·8)*	(1538·6-4644·3)	(8·9 to 11·5)*	(-1·9 to -0·6)*
Upper respiratory infection episodes aggregate	233 458·2	10·2	-1·4	2734·8	10·1	-1·3
	(207 997·9 to 259 065·2)	(8·9 to 11·5)*	(-2·0 to -0·8)*	(1 533·9-4 641·4)	(8·9 to 11·5)*	(-1·9 to -0·6)*
Guillain-Barré syndrome due to upper respiratory infections	12·1 (9·6 to 15·2)	16·9 (14·7 to 19·5)*	-0·2 (-0·9 to 0·4)	3·6 (2·2-5·4)	17·0 (14·7 to 19·5)*	-0·2 (-0·9 to 0·4)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
Continued from previous page)						
Otitis media	112 089·1	5·2	-4·4	3321·5	3·3	–5.6
	(100 504·6 to 123 655·6)	(4·1 to 6·5)*	(-5·5 to -3·4)*	(2 114·2-4 906·3)	(0·6 to 6·0)*	(–8.0 to –3.1)*
Acute otitis media	25 630·8	9·4	1·3	334·9	9·4	1·4
	(21 044·2 to 31 570·6)	(7·5 to 11·2)*	(-0·4 to 3·0)	(169·4-623·7)	(7·5 to 11·4)*	(-0·3 to 3·1)
Chronic otitis media aggregate	86 458·3	4·1	-6·0	2986·6	2·7	-6·3
	(76 403·0 to 97 133·7)	(2·7 to 5·5)*	(-7·2 to -4·8)*	(1904·2-4 371·1)	(-0·2 to 5·6)	(-8·9 to -3·6)*
Meningitis	8733·5	16·9	2·5	1516·9	15·0	1.9
	(8320·6 to 9107·1)	(13·4 to 20·0)*	(-0·6 to 5·1)	(1076·1-1986·0)	(13·7 to 16·5)*	(0.7 to 3.0)*
Pneumococcal meningitis	7297·2	22·2	6·9	728·8	17·6	3·9
	(6228·5 to 8635·3)	(19·7 to 25·1)*	(4·7 to 9·6)*	(513·1–959·9)	(15·5 to 19·7)*	(2·0 to 5·6)*
Acute pneumococcal meningitis	40·3	25·3	12·1	5·3	25·1	11·9
	(33·5 to 48·4)	(17·4 to 33·8)*	(5·2 to 19·5)*	(3·4-7·8)	(15·1 to 37·1)*	(3·2 to 22·3)*
Complications of pneumococcal meningitis aggregate	7256·9	22·2	6·9	723·5	17·6	3·8
	(6191·3 to 8591·7)	(19·7 to 25·1)*	(4·7 to 9·6)*	(509·2–952·4)	(15·4 to 19·6)*	(2·0 to 5·6)*
H influenzae type b	2455·6	-8·3	-18·0	302·1	6·9	-3·3
meningitis	(1966·7 to 3032·6)	(-12·2 to -3·9)*	(-21·5 to -13·9)*	(209·4-407·1)	(3·4 to 10·6)*	(-6·6 to 0·0)
Acute H influenzae type	22·0	-19·2	-26·0	2·9	–18·7	-25·6
b meningitis	(17·2 to 28·7)	(-25·3 to -11·9)*	(-31·8 to −19·2)*	(1·8-4·5)	(–26·1 to –10·8)*	(-32·4 to -18·1)*
Complications of H influenzae type b meningitis aggregate	2433·5	-8·2	-17·9	299·1	7·3	-3·0
	(1947·4 to 3010·8)	(-12·1 to -3·7)*	(-21·5 to -13·7)*	(207·9–402·7)	(3·7 to 11·0)*	(-6·3 to 0·3)
Meningococcal meningitis	1720·8	22·0	7·5	161·6	19·7	5·1
	(1327·7 to 2 180·5)	(17·9 to 26·2)*	(3·8 to 11·3)*	(111·7–215·6)	(17·5 to 22·1)*	(3·2 to 7·4)*
Acute meningococcal meningitis	22·8	24·4	12·7	3·0	24·7	12·9
	(18·9 to 27·7)	(14·4 to 34·9)*	(3·7 to 22·4)*	(1·8-4·5)	(10·9 to 39·4)*	(0·5 to 25·9)*
Complications of meningococcal meningitis aggregate	1698·0 (1305·2 to 2155·2)	22·0 (17·9 to 26·2)*	7·4 (3·8 to 11·2)*	158·6 (109·4–211·9)	19·6 (17·4 to 22·0)*	5·0 (3·1 to 7·3)*
Other meningitis	3015·2	19·0	3·5	324·5	15·3	0·9
	(2438·5 to 3618·0)	(16·6 to 21·6)*	(1·4 to 5·9)*	(225·1-428·1)	(13·3 to 17·4)*	(−0·8 to 2·7)
Other meningitis episodes aggregate	134·5	16·3	5·7	17·8	16·6	5·9
	(120·0 to 151·7)	(10·2 to 22·5)*	(0·1 to 11·4)*	(11·5–25·3)	(8·6 to 24·0)*	(-1·1 to 12·5)
Complications of other meningitis aggregate	2880·4	19·1	3·4	306·7	15·2	0·6
	(2300·7 to 3474·9)	(16·5 to 21·9)*	(1·2 to 5·8)*	(213·3-406·6)	(13·2 to 17·5)*	(–1·1 to 2·6)
Encephalitis	4315·8	4·8	-9·3	457·0	6·7	-6·3
	(3145·8 to 5875·6)	(2·8 to 7·4)*	(-11·3 to -6·7)*	(326·7-594·9)	(4·6 to 9·0)*	(-8·2 to -4·3)*
Acute encephalitis	81·8	7·1	-4·1	10·8	7·5	-3·7
	(74·8 to 89·5)	(5·2 to 9·0)*	(-5·7 to -2·4)*	(7·1–15·5)	(3·8 to 11·1)*	(-6·9 to -0·5)*
Complications of encephalitis aggregate	4233.6	4·8	-9·4	446·2	6·7	-6·4
	(3069.6 to 5787.5)	(2·7 to 7·3)*	(-11·4 to -6·8)*	(318·8-581·4)	(4·6 to 9·0)*	(-8·3 to -4·4)*
Diphtheria	0.6	-59·6	-62·8	0·0	-59·7	-62·8
	(0.3 to 1.2)	(-81·7 to -12·7)*	(-83·1 to -19·5)*	(0·0–0·1)	(-81·7 to -12·7)*	(-83·1 to -19·5)*
Whooping cough	2232·6	-27·4	-32·4	110·3	-27·3	-32·3
	(1725·9 to 2800·7)	(-29·5 to -25·2)*	(-34·3 to -30·3)*	(65·6–167·0)	(-29·6 to -24·8)*	(-34·5 to -30·0)*
Tetanus	209·3	8·4	-2·7	9·1	-8·9	-18·1
	(204·8 to 214·7)	(6·1 to 9·8)*	(-5·0 to -1·4)*	(6·7-12·2)	(-14·5 to -3·7)*	(-23·3 to -13·3)*
Severe tetanus	8·6	-46·3	-52·1	1·1	-45·9	-51·8
	(6·5 to 13·0)	(-54·0 to -36·0)*	(-59·1 to -42·9)*	(0·7–1·9)	(-53·8 to -35·7)*	(-58·9 to -42·6)*
Complications of tetanus aggregate	200·7	13·3	1·8	8·0	1·0	-8·8
	(196·7 to 204·5)	(12·8 to 13·9)*	(1·4 to 2·3)*	(5·8–10·6)	(-3·7 to 5·4)	(-13·0 to -4·9)*
Measles	127·4	-70·2	-72·3	11·5	-70·0	-72·0
	(56·8 to 252·2)	(-73·6 to -66·0)*	(-75·4 to -68·4)*	(4·3–25·3)	(-73·6 to -65·4)*	(-75·4 to -67·7)*
Varicella and herpes zoster	5907·7	15·2	-0·4	207·8	18·5	-0·4
	(5489·4 to 6344·6)	(14·1 to 16·5)*	(-0·8 to -0·0)*	(127·9–316·5)	(16·5 to 20·8)*	(-1·4 to 0·7)
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	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change i ASR between 2005 and 2015
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Neglected tropical diseases and malaria	1 900 062·6 (1 863 148·8 to 1 940 058·5)	-0·4 (-2·5 to 1·9)	-10·7 (-12·7 to -8·7)*	20 763·1 (13 382·4-32 174·5)	-4·7 (-13·2 to 8·9)	-14·5 (-22·2 to -2·4)*
Malaria	295717·3	29·9	20·0	3358·2	16·1	8.5
	(257568·4 to 338 449·0)	(22·5 to 37·2)*	(12·9 to 27·2)*	(2356·9-4703·9)	(12·8 to 19·3)*	(5.4 to 11.4)*
Asymptomatic malaria parasitaemia (PfPR)	206 147-2 (168 670-7 to 246 888-7)	37·3 (27·0 to 48·1)*	26·4 (16·6 to 36·8)*	0·0 (0·0–0·0)	0·0 (0·0 to 0·0)	0·0 (0·0 to 0·0)
Malaria episodes	16756·8	-15·9	-21·1	346·6	-15·8	–20·9
aggregate	(10341·8 to 25965·5)	(-23·6 to -8·4)*	(-28·4 to -14·1)*	(174·4-598·4)	(-23·6 to -8·2)*	(–28·2 to −13·7)*
Complications of malaria aggregate	876·9	28·4	18·4	274·9	24·7	15·0
	(798·1 to 957·9)	(26·5 to 30·5)*	(16·5 to 20·3)*	(209·7–344·8)	(20·5 to 29·1)*	(11·2 to 18·9)*
Anaemia due to malaria parasitaemia (PfPR) aggregate	71 936·4 (70 137·5 to 73 504·2)	26·3 (24·4 to 28·0)*	17·0 (15·3 to 18·6)*	2736·7 (1848·0-3918·8)	21·1 (18·1 to 24·3)*	13·1 (10·5 to 16·1)*
Chagas disease	6653.6	-7·1	-22·1	63·1	-1·8	-21·0
	(5750.5 to 7575.6)	(-9·7 to -4·1)*	(-24·2 to -19·7)*	(42·1–90·8)	(-5·1 to 1·7)	(-23·6 to -18·3)*
Chagas disease episodes aggregate	5634·7	-7·8	-22·2	0·0	-18·7	-27·6
	(4850·5 to 6414·1)	(-10·3 to -4·9)*	(-24·3 to -19·8)*	(0·0–0·1)	(-22·9 to -14·4)*	(-31·1 to -24·1)*
Complications of Chagas disease aggregate	666-0	-5·2	-22·0	44·8	-2·5	-21·0
	(525-5 to 820-6)	(-7·9 to -1·9)*	(-24·3 to -19·6)*	(30·0–62·8)	(-6·0 to 1·2)	(-23·6 to -18·3)*
Heart failure due to	353·0	0·3	-20·9	18·2	0·2	-20·9
Chagas disease aggregate	(221·5 to 503·5)	(-2·9 to 4·0)	(-23·4 to -18·2)*	(9·7–29·4)	(-3·6 to 4·4)	(-23·8 to -17·8)*
Leishmaniasis	3859·3	27·3	11·4	45·8	25·5	10·5
	(3438·4 to 4570·4)	(5·9 to 55·0)*	(-7·0 to 35·1)	(22·8–86·7)	(21·9 to 28·9)*	(7·5 to 13·5)*
Visceral leishmaniasis	60·8	10·6	2·3	4·3	10·8	2·6
	(57·5 to 64·7)	(9·7 to 11·4)*	(1·6 to 3·0)*	(2·9–6·1)	(2·1 to 20·7)*	(-5·4 to 11·5)
Cutaneous and mucocutaneous leishmaniasis	3895·9	27·0	11·0	41·5	27·3	11·4
	(3324·6 to 4767·5)	(23·7 to 30·1)*	(8·3 to 13·8)*	(19·4–80·8)	(23·6 to 30·5)*	(8·3 to 14·5)*
African trypanosomiasis	10·7	-68·7	-72·3	3·0	-67·4	-71·0
	(6·0 to 17·0)	(-70·4 to -67·3)*	(-73·8 to -71·0)*	(1·4-5·3)	(-72·6 to -62·1)*	(-75·5 to -66·6)*
Schistosomiasis	252 339·5	-23·5	-30·9	2472·6	-21·8	-29·1
	(211 032·5 to 321 081·3)	(-32·5 to -4·4)*	(-39·0 to −13·7)*	(1275·0-4521·2)	(-29·3 to -4·0)*	(-36·0 to -13·1)*
Schistosomiasis episodes aggregate	130 285·7	-26·3	-33·1	740·7	-27·3	-34·0
	(114 433·9 to 156 900·0)	(-34·5 to -13·6)*	(-40·5 to -21·3)*	(292·3–1584·0)	(-35·2 to -14·1)*	(-41·0 to -22·0)*
Complications of schistosomiasis aggregate	105 621·9	-19·6	-27·7	1188·8	–18·0	-26·3
	(76 641·0 to 160 339·4)	(-32·6 to 14·1)	(-39·4 to 3·0)	(535·9-2 463·3)	(–30·5 to 15·6)	(-37·5 to 4·3)
Anaemia due to schistosomiasis aggregate	16 331·3	-19·6	-27·0	543·1	-21·5	-27·8
	(16 075·8 to 16 655·1)	(-21·1 to -17·7)*	(-28·3 to -25·3)*	(365·1-781·5)	(-24·0 to −18·6)*	(-30·0 to -25·1)*
Cysticercosis	1931·0	-6·2	-20·8	286·7	-16·3	-29·2
	(1597·8 to 2312·0)	(-10·2 to -2·5)*	(-24·3 to -17·6)*	(194·2-392·8)	(-21·3 to -11·8)*	(-33·3 to -25·3)*
Cystic echinococcosis	1383.0	24·7	6·7	126·8	24·3	6.6
	(1265.9 to 1498.6)	(22·6 to 27·0)*	(5·0 to 8·5)*	(86·7–174·6)	(21·1 to 27·5)*	(4.1 to 9.2)*
Lymphatic filariasis	38 464·1	-44·9	–51·6	2075·0	-16·2	-27·7
	(31 328·2 to 46 783·0)	(-49·9 to -39·9)*	(–56·0 to −47·2)*	(1120·6–3311·2)	(-32·1 to -3·9)*	(-41·5 to -17·1)*
Prevalence of detectable microfiliaria due to lymphatic filariasis	19 707·9 (17 173·8 to 22 270·5)	-58·8 (-61·8 to -55·3)*	-63·5 (-66·1 to -60·3)*	0·0 (0·0–0·0)	0·0 (0·0 to 0·0)	0·0 (0·0 to 0·0)
Complications of lymphatic filariasis aggregate	18756·3 (12420·1 to 26397·1)	-14·8 (-29·9 to -3·5)*	-26·7 (-40·1 to -16·7)*	2075-0 (1120-6-3311-2)	-16·2 (-32·1 to -3·9)*	-27·7 (-41·5 to -17·1)*
Onchocerciasis	15 531·5	-29·1	-36·8	1135·7	-21·2	-31·2
	(11 963·5 to 19 993·8)	(-39·5 to -18·8)*	(-46·0 to -27·5)*	(546·2–2005·4)	(-38·5 to -4·8)*	(-46·7 to −16·2)*
Asymptomatic onchocerciasis	2280·4	-62·4	-66·2	0·0	0·0	0·0
	(1525·0 to 3173·2)	(-66·1 to -59·6)*	(-69·5 to -63·7)*	(0·0–0·0)	(0·0 to 0·0)	(0·0 to 0·0)

	Prevalence (thousands)			YLDs (thousands)		
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Skin disease due to onchocerciasis aggregate	12 223·9	-17·4	-26·5	1055·6	-22·2	-31·6
	(8 474·7 to 16 585·2)	(-31·0 to -2·7)*	(-38·6 to -13·1)*	(467·4-1894·7)	(-41·8 to -3·2)*	(-49·0 to −14·8)*
Vision loss due to onchocerciasis aggregate	1025·4	-2·3	-22·3	80·1	-6·1	-25·4
	(724·8 to 1452·1)	(-15·0 to 11·8)	(-32·3 to -11·0)*	(51·3–117·5)	(-16·9 to 6·0)	(-33·9 to -15·9)*
Trachoma	3557·1	4·6	-19·3	279·2	-1·2	-23·9
	(2940·5 to 4321·8)	(0·4 to 9·2)*	(-22·9 to -15·2)*	(192·8–396·0)	(-5·6 to 3·0)	(-27·5 to -20·3)*
Dengue	4730·0	143·6	119·7	764·1	140·8	117·7
	(2654·1 to 10 254·2)	(-0·3 to 564·7)	(-10·1 to 498·7)	(346·8–1744·2)	(-0·1 to 558·4)	(-9·8 to 494·3)
Dengue episodes	1409·2	134·1	110⋅8	82·1	141·4	117·9
aggregate	(937·6 to 2943·8)	(-3·8 to 521·2)	(-13⋅2 to 459⋅0)	(45·0-183·9)	(0·4 to 549·2)*	(-9·4 to 486·7)
Post-dengue chronic fatigue syndrome	3324·6	143·7	119·8	682·0	140·8	117·6
	(1600·3 to 7347·9)	(-0·3 to 564·7)	(-10·1 to 498·7)	(295·2–1 608·9)	(-0·2 to 558·0)	(-9·8 to 494·1)
Yellow fever	2·8	-25·7	-31·6	0·1	-25·7	-31·6
	(0·8 to 7·7)	(-31·6 to -19·2)*	(-36·9 to -25·8)*	(0·0-0·3)	(-31·6 to -19·2)*	(-36·9 to -25·8)*
Rabies	0·7	-43·4	-50·8	0·1	-43·4	-50·8
	(0·6 to 0·8)	(-51·5 to -33·8)*	(-57·6 to -42·6)*	(0·1-0·1)	(-51·5 to -33·8)*	(-57·6 to -42·6)*
Intestinal nematode infections	1447209·3 (1414995·0 to 1481332·3)	-2·5 (-5·1 to 0·2)	-12·8 (-15·2 to -10·4)*	3173·8 (1861·3–5090·8)	-22·8 (-27·3 to -17·3)*	-30·4 (-34·5 to -25·4)*
Ascariasis	761893.8	-10·8	-20·3	871·0	-37·7	-43·9
	(682557.5 to 861031.5)	(-22·5 to 2·5)	(-30·7 to -8·2)*	(482·5-1 464·3)	(-45·3 to -29·4)*	(-50·8 to -36·4)*
Complications of ascariasis aggregate	47 626·4	-35·6	-42·1	871·0	-37·7	-43·9
	(44 087·5 to 51711·9)	(-41·9 to -28·4)*	(-47·8 to -35·6)*	(482·5-1 464·3)	(-45·3 to -29·4)*	(-50·8 to -36·4)*
Asymptomatic ascariasis	714230·7	-8·5	-18·2	0·0	0·0	0·0
	(634201·9 to 814009·5)	(-20·9 to 6·4)	(-29·3 to -4·9)*	(0·0–0·0)	(0·0 to 0·0)	(0·0 to 0·0)
Trichuriasis	463 652·1	-2·1	-12·2	544·1	−16·7	-24·9
	(426 621·2 to 502 939·4)	(-12·0 to 9·0)	(-21·1 to -2·1)*	(290·7–946·0)	(−30·6 to 3·6)	(-37·5 to -6·8)*
Complications of trichuriasis aggregate	27129·6	-15·6	-24·1	544·1	−16·7	-24·9
	(24188·7 to 31890·8)	(-27·8 to 2·1)	(-35·0 to -8·2)*	(290·7–946·0)	(−30·6 to 3·6)	(-37·5 to -6·8)*
Asymptomatic trichuriasis	436 525·4	-1·1	-11·3	0·0	0·0	0·0
	(400 013·3 to 475 985·5)	(-11·8 to 11·0)	(-21·0 to -0·3)*	(0·0-0·0)	(0·0 to 0·0)	(0·0 to 0·0)
Hookworm disease	428 245·9	-7·3	-17·2	1758·8	-14·6	-22·9
	(394 486·0 to 468 292·4)	(-16·9 to 3·5)	(-25·8 to -7·5)*	(1088·5-2754·9)	(-19·9 to -9·4)*	(-27·7 to -18·2)*
Complications of hookworm disease aggregate	51 012·9 (48 151·1 to 54 346·2)	-14·0 (-21·0 to -6·2)*	-23·2 (-29·5 to -16·3)*	1046·7 (572·8–1732·1)	-15·5 (-24·0 to -6·3)*	-24·4 (-32·1 to -16·3)*
Anaemia due to hookworm disease aggregate	24220·2 (24078·2 to 24365·2)	-6.2 (-7.3 to -5.1)	-15·5 (-16·5 to -14·6)*	712·1 (476·3–1031·1)	-13·2 (-15·6 to -11·1)*	-20·5 (-22·6 to -18·7)*
Asymptomatic	352 948·6	-6·4	-16·4	0·0	0·0	0·0
hookworm disease	(319 143·2 to 392 574·4)	(-18·3 to 7·5)	(-27·2 to -3·9)*	(0·0-0·0)	(0·0 to 0·0)	(0·0 to 0·0)
Food-borne trematodiases	71 095·4	4·0	-9·3	1686·5	3·7	-10·0
	(67 365·5 to 75 246·9)	(1·5 to 6·6)*	(-11·4 to -7·1)*	(857·1–3066·8)	(-0·4 to 10·1)*	(-13·4 to -5·2)*
Asymptomatic food- borne trematodiases aggregate	56 131·7 (45 765·9 to 63 620·3)	3·8 (1·2 to 6·6)*	-9·3 (-11·6 to -6·9)*	0·0 (0·0–0·0)	0·0 (0·0–0·0)	0·0 (0·0-0·0)
Symptomatic food-borne trematodiases aggregate	14963·7	4·8	-9·3	1686·5	3·7	-10·0
	(8632·7 to 25256·6)	(0·5 to 11·4)*	(-12·8 to -4·2)*	(857·1–3066·8)	(-0·4 to 10·1)	(-13·4 to -5·2)*
Leprosy	514·2	-0·1	-19·7	31·0	0·6	-19·2
	(487·0 to 545·7)	(-0·5 to 0·3)	(-20·2 to -19·3)*	(20·9–43·5)	(-1·5 to 2·9)	(-20·8 to -17·4)*
Ebola	2·8 (1·2 to 5·2)	54521·3 (46386·2 to 68346·9)*	49 817·9 (41 865·6 to 62 764·9)*	0·6 (0·2–1·1)	51 908·8 (42 739·0 to 66 884·3)*	47 453·0 (38 609·5 to 61730·3)*
Other neglected tropical diseases	61452·6	0·1	-7·7	5261·1	7·4	-1·2
	(60842·8 to 62054·9)	(-1·0 to 1·3)	(-8·8 to -6·6)*	(2558·5-12 199·0)	(-18·9 to 50·6)	(-25·5 to 38·0)
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	Prevalence (thousands)			YLDs (thousands)			
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change ir ASR between 2005 and 2015	
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Anaemia due to other neglected tropical diseases aggregate	61452·6 (60842·8 to 62054·9)	0·1 (-1·0 to 1·3)	-7·7 (-8·8 to -6·6)*	2158·2 (1446·3–3073·1)	-6.0 $(-8.2 to -3.8)$ *	-12·4 (-14·5 to −10·4)*	
Maternal disorders	11372·9 (10686·6 to 11894·1)	2.1	-7·9 (-13·6 to -1·8)*	898-8	-5·1	-15·1	
Maternal haemorrhage	2304·8 (2231·6 to 2 393·5)	(-4·2 to 9·2) 1·6 (-3·0 to 6·5)	-7·8 (-12·0 to -3·4)*	(642·8-1222·9) 84·7 (57·3-119·5)	(-18·7 to 11·2) 0·1 (-13·5 to 17·2)	(-27·0 to -0·3)* -9·1 (-21·3 to 6·6)	
Maternal hemorrhage episodes aggregate	191-7 (127-8 to 277-5)	2·9 (-36·5 to 66·2)	-6·4 (-42·1 to 49·9)	31·3 (18·1-49·0)	3·3 (-30·6 to 54·9)	-6.0 (-36.3 to 41.1)*	
Anaemia due to maternal haemorrhage aggregate	2113·4 (2082·8 to 2141·5)	1·5 (-0·9 to 3·9)	-7·9 (-10·1 to -5·8)*	53·5 (35·0-78·6)	-1·8 (-5·9 to 2·6)	-10·8 (-14·5 to -6·9)*	
Maternal sepsis and other maternal infections	3140·8 (2658·0 to 3727·7)	3·7 (-15·2 to 27·8)	-7⋅8 (-24⋅2 to 13⋅1)	56·5 (29·5–102·4)	-1·0 (-42·3 to 70·4)	-9·7 (-47·1 to 55·2)	
Maternal sepsis and other maternal infections aggregate	900·0 (486·7 to 1462·3)	-2·4 (-50·3 to 93·7)	-10·2 (-54·4 to 78·1)	45·3 (22·2-82·7)	-2·7 (-49·0 to 91·0)	-10·5 (-53·2 to 73·6)	
Infertility due to puerperal sepsis	2242·1 (2 030·5 to 2483·0)	6·4 (4·4 to 8·9)*	-6⋅8 (-8⋅6 to -4⋅7)*	11·2 (4·2–24·3)	6·7 (4·0 to 9·5)*	-6·6 (-8·9 to -4·1)*	
Maternal hypertensive disorders	4624·4 (3030·8 to 6592·4)	2·7 (-37·9 to 69·8)*	-5⋅8 (-43⋅0 to 55⋅4)*	222·4 (118·4-365·4)	2·6 (-37·3 to 68·5)	-5·9 (-42·4 to 54·4)	
Maternal hypertensive disorders episodes aggregate	4629·2 (3031·5 to 6602·4)	2·8 (-37·8 to 70·0)	-5·7 (-42·9 to 55·7)	219·1 (115·4-363·3)	2·7 (-37·0 to 68·8)	-5·8 (-42·4 to 55·2)	
Eclampsia	5·4 (2·2 to 10·5)	-5·8 (-69·0 to 182·2)	-12·9 (-71·6 to 158·3)	3·3 (1·2-6·3)	-5·7 (-69·0 to 182·2)	-12·8 (-71·6 to 158·3)	
Maternal obstructed labour and uterine rupture	1077·2 (933·3 to 1255·2)	-13·1 (-17·5 to -7·9)*	-23·9 (-27·7 to -19·7)*	352·0 (234·6-499·9)	-12·6 (-17·3 to -7·3)*	-23·4 (-27·4 to -18·9)*	
Obstructed labour, acute event	89·6 (53·0 to 142·9)	-3·6 (-45·9 to 74·7)	-11·4 (-49·7 to 60·1)	27·7 (14·8–47·0)	-3·1 (-44·9 to 72·1)	-10·9 (-49·0 to 58·2)	
Fistula due to maternal obstructed labour and uterine rupture aggregate	987·6 (850·6 to 1156·0)	-13·9 (-16·3 to -11·4)*	-24·9 (-27·0 to -22·8)*	324·3 (216·2-457·4)	-13·3 (-16·6 to -10·1)*	-24·3 (-27·1 to -21·7)*	
Maternal abortion, miscarriage, and ectopic pregnancy	441·3 (285·5 to 630·6)	-0·4 (-37·2 to 58·1)	-9·4 (-42·4 to 44·2)	48·2 (27·2–78·5)	-0·4 (-36·9 to 60·5)	-9·3 (-42·4 to 45·6)	
Other maternal disorders				135·0 (90·6–190·2)	-2·2 (-19·6 to 17·5)	-12·2 (-27·7 to 5·5)	
Neonatal disorders	52 961·9 (50 435·8 to 54 978·4)*	14·7 (11·9 to 17·8)*	3·9 (1·3 to 6·7)*	10710·5 (8113·9-13786·1)	13·3 (9·5 to 17·4)*	3·1 (-0·3 to 6·8)	
Neonatal preterm birth complications	41 855·4 (36 843·4 to 47 188·1)*	10·6 (6·2 to 15·2)*	-0·4 (-4·3 to 3·8)*	5090·9 (3864·5-6555·6)*	4·4 (-1·1 to 10·1)	-5·2 (-10·1 to 0·2)	
Vision loss due to retinopathy of prematurity aggregate	4245·1 (3386·2 to 5265·3)	11·1 (7·2 to 15·1)*	-2·3 (-5·8 to 1·3)	135·5 (83·0–208·8)	13·9 (9·5 to 18·4)	-0.8 (-4.5 to 3.2)	
Complications of preterm birth complications aggregate	37 609·0 (33 086·7 to 42 697·0)	10·5 (5·6 to 15·7)*	-0·1 (-4·6 to 4·5)	4955·4 (3756·9–6394·8)	4·1 (-1·4 to 10·0)	-5·3 (-10·4 to 0·2)	
Neonatal encephalopathy due to birth asphyxia and trauma	13 681·5 (9 797·3 to 19 759·3)	27·3 (24·6 to 30·0)*	16·2 (13·7 to 18·5)*	3769·4 (2451·7–5809·3)	25·1 (22·0 to 28·3)*	14·2 (11·5 to 17·0)*	
Neonatal sepsis and other neonatal infections	139·1 (79·9 to 216·0)	-1·6 (-3·0 to -0·0)*	-6·7 (-8·0 to -5·2)*	17·7 (9·2–31·0)	-1·8 (-3·5 to 0·1)	-6·9 (-8·5 to -5·1)*	
Severe infection due to neonatal sepsis and other neonatal infections	130·1 (71·1 to 207·6)	-2·2 (-3·8 to -0·5)*	-7·3 (-8·8 to -5·6)*	16·6 (8·1–30·0)	-2·1 (-3·7 to -0·3)*	-7·1 (-8·7 to -5·4)*	

	Prevalence (thousands)			YLDs (thousands)		
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Complications of neonatal sepsis and other neonatal infections aggregate	8·9 (7·5 to 10·4)	8·0 (4·3 to 12·7)*	2·4 (-1·1 to 6·9)	1·1 (0·6-1·6)	2·6 (-9·4 to 14·4)	-2·7 (-14·1 to 8·5)
Haemolytic disease and other neonatal jaundice	2042·8 (1825·0 to 2309·0)	20·0 (18·8 to 21·4)*	9·2 (8·0 to 10·3)*	603·0 (451·9–770·8)	17·0 (14·8 to 19·4)*	6·5 (4·4 to 8·6)*
Other neonatal disorders				1229·5 (836·5–1 647·2)	19·9 (0·8 to 41·1)*	8·5 (-8·5 to 27·6)
Nutritional deficiencies	1482655·3 (1477085·9 to	2·7 (2·2-3·2)*	-7·1 (-7·5—6·7)*	54 632·4 (36 772·8-77 894·3)	-3·5 (-4·9—2·1)*	-11·4 (-12·7—10·3)*
Protein-energy malnutrition	22 834·3 (21764·0 to 23 985·1)	-4·8 (-10·2 to 0·8)	-12·1 (-17·1 to -6·9)* -6·0	2823·3 (1820·3–3984·2)	-4·5 (-9·8 to 1·1)	-11·8 (-16·7 to -6·6)*
lodine deficiency	110 919·5 (100 337·2 to 125 252·8)	7·0 (4·7 to 9·4)*	-6·0 (-8·0 to -3·8)*	2386-9 (1508-3-3564-2)	7·7 (5·7 to 9·8)*	-4·4 (-6·4 to -2·4)*
Visible goiter due to iodine deficiency aggregate	108 307·6 (97 705·7 to 122 260·0)	6·9 (4·5 to 9·4)*	-6·2 (-8·2 to -4·0)*	1927·5 (1192·0–3040·1)	6.9 (4.4 to 9.3)*	-6·0 (-8·2 to -3·9)*
Visible goiter with heart failure due to iodine deficiency aggregate	0·3 (0·3 to 0·4)	39·6 (38·3 to 41·0)*	4·2 (3·3 to 5·2)*	0·0 (0·0–0·1)	39·6 (38·3 to 41·0)*	4·2 (3·3 to 5·2)*
Intellectual disability due to iodine deficiency aggregate	2611·6 (1085·6 to 3621·3)	10·4 (6·1 to 12·6)*	2·0 (-2·0 to 4·3)	459·3 (163·4–752·8)	11·0 (6·4 to 13·6)*	2·8 (-1·6 to 5·2)
Vitamin A deficiency	4901·1 (3975·6 to 5921·7)	9·3 (6·2 to 12·5)*	0·1 (-2·7 to 2·7)	232·4 (143·5–345·9)	10·9 (7·5 to 14·6)*	0·2 (-2·7 to 3·3)
Iron-deficiency anaemia	1477530·5 (1470902·3 to 1485322·2)	1·5 (0·9 to 2·1)*	-8·0 (-8·5 to -7·4)*	48 529·2 (32 560·8–69 725·2)	-3·8 (-5·1 to -2·4)*	-11·6 (-12·8 to -10·5)*
Iron-deficiency anaemia without heart failure aggregate	1477 458·4 (1470 824·4 to	39·7 (38·1-41·3)*	6·1 (4·7-7·4)*	8·1 (5·5-11·4)	39·6 (35·2-44·3)*	6·0 (2·5–9·9)*
Iron-deficiency anaemia with heart failure aggregate	72·1 (64·6 to 80·7)	39·7 (38·1 to 41·3)*	6·1 (4·7 to 7·4)*	8·1 (5·5–11·4)	39·6 (35·2 to 44·3)*	6.0 (2.5 to 9.9)*
Other nutritional deficiencies				660·7 (396·0–1026·9)	-16·7 (-36·8 to 5·8)	-22·9 (-41·5 to -2·1)*
Other communicable, maternal, neonatal, and nutritional diseases	1604 919·9 (1563 786·8 to 1636 531·0)	16·1 (15·3 to 17·0)*	-0·4 (-1·1 to 0·4)	3929·7 (2566·9-5786·6)	6·5 (3·9 to 9·2)*	-4·0 (-6·2 to -1·7)*
Sexually transmitted diseases excluding HIV	1145527·5 (1100971·2 to 1176685·1)	16·9 (15·7 to 18·3)*	-1·0 (-2·1 to 0·1)	1573·7 (974·0–2501·8)	16·9 (14·5 to 19·1)*	1·3 (-0·6 to 3·4)
Syphilis	43 604·9 (37 160·3 to 50 658·3)	3·7 (-0·4 to 7·0)	-9·8 (-13·4 to -6·9)*	242·8 (166·1–333·7)	15·3 (12·2 to 18·5)*	-5·6 (-8·0 to -3·0)*
Early syphilis aggregate	42 350·9 (35 898·4 to 49 441·3)	3·3 (-0·9 to 6·8)	-9·9 (-13·6 to -6·9)*	10·3 (3·0-25·3)	3·4 (-1·1 to 7·4)	-9·9 (-13·7 to -6·3)*
Adult tertiary syphilis	1254·0 (1127·1 to 1385·8)	15·8 (13·2 to 18·3)*	–5·6 (–7·8 to –3·6)*	232·6 (155·4-322·8)	15·9 (12·6 to 19·1)*	-5·4 (-7·9 to -2·8)*
Chlamydial infection	82 822·4 (66 426·4 to 104 328·5)	8·3 (6·1 to 10·7)*	-2·9 (-4·8 to -0·9)*	364·6 (210·6–589·5)	10·1 (6·9 to 13·4)*	-1·8 (-4·5 to 0·8)
Chlamydial infection episodes aggregate	81 072·5 (64 735·8 to 102 635·8)	8·3 (6·0 to 10·7)*	-2·9 (-4·8 to -0·9)*	332·7 (185·2–546·0)	10·7 (7·1 to 14·4)*	-1·1 (-4·1 to 2·1)
Pelvic inflammatory diseases due to chlamydial infection aggregate	1-73 (147-8 to 202-0)	1·4 (-0·8 to 3·4)	-11·5 (-13·2 to -9·8)*	23·1 (15·5 to 32·5)	1·8 (-2·2 to 5·7)	-11·1 (-14·5 to 7·8)*
aggregate					(Table 3	continues on next pag

	Prevalence (thousands)			YLDs (thousands)			
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change ir ASR between 2005 and 2015	
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Infertility due to chlamydial infection aggregate	1576-6 (1450-8 to 1703-9)	11·1 (10·0 to 12·2)*	-3·0 (-3·9 to -2·0)	8·8 (3·6 to 18·4)	11·1 (9·0 to 13·2)*	-2·9 (-4·8 to -1·1)*	
Gonococcal infection	47 468·5	25·3	12·2	444·9	26·3	12·6	
	(35 848·1 to 62 099·7)	(18·3 to 31·2)*	(5·9 to 17·7)*	(260·8–691·8)	(20·0 to 31·4)*	(6·7 to 17·3)*	
Gonococcal infection episodes aggregate	46 561·5	25·6	12·5	430·3	27·3	13·5	
	(34 928·2 to 61 227·4)	(18·5 to 31·6)*	(6·1 to 18·1)*	(250·5–671·5)	(20·7 to 32·5)*	(7·4 to 18·3)*	
Pelvic inflammatory diseases due to gonococcal infection aggregate	907·0 (822·0 to 1 016·3)	10·0 (7·6 to 11·8)*	-3·8 (-6·0 to -2·2)*	14·6 (9·6-21·4)	3·5 (-0·8 to 7·1)	-8·7 (-12·4 to -5·7)*	
Trichomoniasis	167 618·6	16·2	0·9	194·3	16·1	1·0	
	(144744·1 to 193 444·0)	(15·3 to 17·2)*	(0·4 to 1·3)*	(78·0–408·3)	(15·0 to 17·2)*	(0·3 to 1·8)*	
Genital herpes	885 169·4 (772 303·8 to 1008 588·1)	18·1 (16·4 to 19·9)*	-1·3 (-2·7 to 0·1)	236·4 (74·3–554·2)	19·5 (17·4 to 23·0)*	0·7 (-1·4 to 5·0)	
Moderate infection due to initial genital herpes episode	389·0 (88·0 to 909·7)	38·1 (33·6 to 43·2)*	29·5 (25·4 to 34·3)*	19·0 (4·3-45·8)	37·6 (30·9 to 45·6)*	29·1 (22·9 to 36·7)*	
Complications of genital herpes aggregate	884780·3 (771914·6 to 1008400·6)	18·1 (16·4 to 19·9)*	-1·3 (-2·7 to 0·0)	217·3 (62·2–534·5)	18·1 (16·3 to 19·9)*	-1·2 (-2·7 to 0·3)	
Other sexually transmitted diseases	4821.8	9·6	-4·7	90·7	3·7	-10·0	
	(4452.8 to 5174.6)	(8·1 to 11·0)*	(-5·9 to -3·6)*	(61·5–129·7)	(1·1 to 6·2)*	(-12·2 to -7·8)*	
Pelvic inflammatory diseases due to other sexually transmitted diseases aggregate	507·2 (437·2 to 591·9)	1·2 (-0·1 to 2·6)	-12·2 (-13·1 to -11·2)*	66·8 (45·4-92·9)	1·3 (-1·2 to 4·2)	-12·0 (-14·3 to -9·8)*	
Infertility due to other sexually transmitted diseases aggregate	4314·5 (3943·8 to 4657·1)	10·7 (9·1 to 12·2)*	-3·8 (-5·1 to -2·4)*	23·9 (9·7–48·8)	10·8 (9·0 to 12·7)*	-3·6 (-5·2 to -2·0)*	
Other sexually transmitted diseases				329·4 (200·1–504·2)	19·5 (14·4 to 24·6)*	6·7 (2·2 to 11·3)*	
Hepatitis	497 901·2	16·7	2·2	406·4	12·6	1·2	
	(490 397·2 to 505 815·2)	(15·8 to 17·6)*	(1·4 to 2·9)*	(267·6–588·8)	(1·7 to 24·1)*	(-8·3 to 11·5)	
Acute hepatitis A	8785·5	4·2	-2·8	172·5	9·5	1·9	
	(7 950·0 to 9 598·2)	(-7·2 to 17·0)	(-13·5 to 9·1)	(111·5–249·7)	(-0·1 to 20·4)	(-6·9 to 12·0)	
Hepatitis B	356 083·4	16·7	2·8	190·2	16·9	1·2	
	(342 942·4 to 370 231·5)	(15·0 to 18·4)*	(1·3 to 4·3)*	(121·1–282·2)	(-4·5 to 40·2)	(-16·9 to 20·7)	
Acute hepatitis B aggregate	12 832·2	13·2	1·0	190·2	16·9	1·2	
	(11 239·6 to 14 567·4)	(-4·3 to 35·2)	(-14·6 to 20·0)	(121·1–282·2)	(-4·5 to 40·2)	(-16·9 to 20·7)	
Chronic hepatitis B	343 251·2 (330 541·3 to 357 194·6)	16·9 (15·3 to 18·4)*	2·8 (1·5 to 4·3)*	0.0	0·0 (0·0 to 0·0)	0·0 (0·0 to 0·0)	
Hepatitis C	142745·3	18·0	1·2	8.7	17·0	3·3	
	(127559·8 to 157704·1)	(16·7 to 19·2)*	(0·1 to 2·2)*	(4·3–16·5)	(12·2 to 21·7)*	(-0·5 to 7·3)	
Acute hepatitis C	622·3	17·0	3·4	8.7	17·0	3·3	
aggregate	(580·4 to 666·6)	(15·7 to 18·4)*	(2·5 to 4·3)*	(4·3–16·5)	(12·2 to 21·7)*	(-0·5 to 7·3)	
Chronic hepatitis C	142 122·9	18·0	1·2	0·0	0.0	0·0	
	(126 977·7 to 157 044·9)	(16·7 to 19·2)*	(0·1 to 2·2)*	(0·0-0·0)	(0.0 to 0.0)	(0·0 to 0·0)	
Acute hepatitis E	1501·9	3·5	-3·2	34·9	4·9	-2·6	
	(1385·4 to 1636·4)	(2·2 to 4·7)*	(-4·1 to -2·3)*	(22·4–51·7)	(-4·4 to 15·5)	(-10·9 to 6·8)	
Other infectious diseases	54835·4	0·4	-8·0	1949·6	-1·6	-8·7	
	(54385·8 to 55273·8)	(−1·6 to 2·3)	(-9·8 to -6·2)*	(1307·1–2797·1)	(-5·1 to 2·0)	(-12·0 to -5·5)*	
Ion-communicable diseases	6 652 153·4	13·7	0·0	638 480·8	18·8	-0·1	
	(6 633 099·8 to	(13·6 to 13·8)*	(-0·0 to 0·1)	(478 716·6-819 498·9)	(18·4 to 19·2)*	(-0·4 to 0·2)	

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Neoplasms	90 497·5	44·4	12·4	8569·3	36.6	6·4
	(89 215·7 to 91 896·1)	(42·1 to 46·7)*	(10·6 to 14·2)*	(6265·5-11 079·2)	(31.8 to 41.2)*	(2·7 to 9·7)*
Lip and oral cavity cancer	2425·1	38·6	7·9	209·1	36·7	6·2
	(2278·7 to 2582·3)	(30·3 to 47·1)*	(1·5 to 14·3)*	(150·2–271·4)	(28·6 to 45·3)*	(-0·1 to 12·7)
Nasopharynx cancer	732·7	18·0	-5·5	69·7	16·8	-6⋅8
	(580·5 to 883·9)	(0·2 to 39·8)*	(-19·2 to 11·1)	(47·2–95·8)	(0·8 to 36·0)*	(-18⋅7 to 7⋅7)
Other pharynx cancer	945·5	32·7	2·2	79·5	30·7	0.6
	(885·9 to 1015·0)	(23·8 to 41·5)*	(-4·5 to 9·0)	(57·8–104·3)	(21·9 to 39·2)*	(-6.2 to 7.2)
Oesophageal cancer	746·0	19·6	-8·2	128·6	9·5	-16·1
	(641·5 to 925·7)	(2·5 to 39·9)*	(-21·2 to 7·1)	(92·0–167·5)	(-3·1 to 24·1)	(-25·4 to -5·2)*
Stomach cancer	3539·4	16⋅3	-9·9	396·5	12·0	-13·3
	(3339·9 to 3776·2)	(8⋅9 to 24⋅4)*	(-15·5 to -3·7)*	(292·4–504·2)	(4·8 to 20·2)*	(-18·8 to -7·1)*
Colon and rectum cancer	9399·0	42·6	8·7	762·8	38·4	5·3
	(9059·3 to 9758·3)	(38·1 to 47·2)*	(5·4 to 12·2)*	(564·2–979·6)	(33·7 to 43·3)*	(1·9 to 8·9)*
Colon and rectum cancer aggregate	9377·0	42·6	8·7	759·8	38·4	5·4
	(9037·3 to 9736·4)	(38·1 to 47·3)*	(5·4 to 12·2)*	(561·9–975·3)	(33·7 to 43·3)*	(1·9 to 9·0)*
Stoma due to colon and rectum cancer	22·0	26·1	-4·9	3·0	26·0	-4·7
	(21·3 to 22·8)	(25·0 to 27·2)*	(-5·7 to -4·1)*	(2·0-4·1)*	(20·7 to 31·6)*	(-9·4 to -0·0)*
Liver cancer	618·7	59·8	23·6	188·2	25·2	-1⋅8
	(550·7 to 688·0)	(38·2 to 82·2)*	(6·8 to 40·6)*	(130·4–245·6)	(9·9 to 45·0)*	(-13⋅5 to 13⋅2)
Liver cancer due to hepatitis B	305·9	20·6	–3·6	59·9	9·5	-12·5
	(235·8 to 418·9)	(-9·1 to 63·2)	(–26·5 to 29·1)	(40·2–80·4)	(-8·3 to 34·1)	(-26·5 to 6·1)
Liver cancer due to hepatitis C	268·7	80·1	38·1	44·3	47·7	13·2
	(228·4 to 320·8)	(53·9 to 113·8)*	(17·1 to 65·1)*	(31·6–57·3)	(34·2 to 65·9)*	(2·7 to 27·7)*
Liver cancer due to alcohol use	273·6	58·8	22·1	53·5	39·3	7·2
	(229·0 to 350·7)	(28·9 to 94·8)*	(-0·3 to 49·1)	(37·3-70·1)	(22·1 to 61·6)*	(-5·8 to 23·5)
Liver cancer due to other causes	151·0	24·9	-2·0	30·5	12·1	-11·9
	(118·8 to 201·0)	(-5·6 to 63·5)	(-26·3 to 29·5)	(20·7–40·6)	(-4·5 to 32·8)	(-24·9 to 4·8)
Gallbladder and biliary tract cancer	149·4	22·0	-7·4	39·5	19·0	-9·5
	(138·5 to 158·9)	(14·4 to 30·1)*	(-13·2 to -1·3)*	(27·7–51·5)	(11·8 to 26·8)*	(-15·1 to -3·7)*
Pancreatic cancer	393·8	46·2	10·9	86.6	38·4	5·1
	(372·9 to 417·1)	(40·7 to 52·1)*	(6·7 to 15·4)*	(61.1–112.2)	(33·1 to 43·8)*	(1·1 to 9·3)*
Larynx cancer	1412·6	26·2	-2·3	147·3	24·0	-3·9
	(1340·0 to 1499·9)	(20·1 to 33·0)*	(-7·0 to 2·9)	(105·7-194·3)	(18·1 to 30·0)*	(-8·4 to 0·7)
Larynx cancer aggregate	820·9	24·4	-4·9	93·6	21·5	-6·8
	(749·1 to 911·1)	(14·2 to 35·8)*	(-12·5 to 3·4)	(67·4–121·5)	(12·7 to 30·8)*	(-13·1 to 0·2)
Laryngectomy due to	591·0	28·7	1·8	53·7	28.6	1·9
larynx cancer	(576·5 to 605·4)	(28·0 to 29·3)*	(1·3 to 2·4)*	(35·8–77·6)	(25.8 to 31.4)*	(-0·2 to 4·1)
Tracheal, bronchus, and lung cancer	3299·7	37·7	5·6	513·6	31·1	0.6
	(3095·1 to 3536·4)	(29·6 to 47·1)*	(-0·5 to 12·6)	(377·9–648·8)	(23·4 to 40·0)*	(–5.2 to 7.1)
Malignant skin melanoma	3082·3	59·0	26·2	180·5	56·3	23·7
	(2475·8 to 3901·7)	(50·7 to 67·0)*	(19·5 to 32·6)*	(120·8–257·6)	(47·9 to 64·0)*	(16·9 to 29·9)*
Non-melanoma skin cancer	2558·2	76·8	34·5	148·0	82·9	40·1
	(2494·8 to 2625·6)	(72·0 to 81·6)*	(30·8 to 38·2)*	(104·7-197·7)	(72·7 to 93·9)*	(31·9 to 48·7)*
Non-melanoma skin cancer (squamous-cell carcinoma)	2155·9 (2019·5 to 2312·2)	93·5 (82·2 to 105·8)*	49·6 (40·5 to 59·2)*	142·0 (100·7–187·1)	86·4 (75·6 to 98·2)*	42·8 (34·2 to 52·1)*
Non-melanoma skin cancer (basal-cell carcinoma)	578·1 (497·7 to 671·7)	27·0 (24·0 to 29·7)*	-3·9 (-6·0 to −1·8)*	6·0 (2·9–11·1)	26·9 (23·5 to 30·1)*	-3·8 (-6·4 to -1·2)*
Breast cancer	21361·8	41·5	9·9	1796·5	36·1	5·3
	(20249·5 to 22266·3)	(33·4 to 49·6)*	(4·1 to 15·7)*	(1270·7–2411·6)	(29·2 to 42·7)*	(0·4 to 10·1)*
Breast cancer aggregate	10718·5	62·7	32·0	989·8	46.6	15·8
	(9581·7 to 11667·2)	(42·9 to 83·3)*	(16·2 to 48·1)*	(719·8–1271·0)	(32.7 to 61.5)*	(5·3 to 27·5)*
Mastectomy due to breast cancer	10 638·5	25·1	-4·7	806·8	25·1	-4·6
	(10 404·8 to 10 894·5)	(24·1 to 26·1)*	(-5·4 to -3·9)*	(520·3–1171·8)	(24·0 to 26·2)*	(-5·5 to -3·8)*
					(Table 3	continues on next pag

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change i ASR between 2005 and 2015
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Cervical cancer	3442-4	-1.6	-20-5	264.0	-1.4	-20.5
Uterine cancer	(3129·9 to 3765·0)	(-11·4 to 9·8)	(-28·2 to -11·5)*	(187·9-342·6)	(-10·9 to 9·8)	(-28·0 to -11·6)*
	3827·7	39·5	8·2	250·2	37·1	6·1
	(3425·6 to 4285·2)	(22·9 to 55·9)*	(-4·0 to 20·0)	(172·4-341·0)	(21·1 to 52·9)*	(-5·4 to 17·6)
Ovarian cancer	1174·2	28·3	0·3	150·7	25·8	-1·7
	(1107·4 to 1250·1)	(20·8 to 36·3)*	(-5·1 to 6·3)	(110·5-192·3)	(17·8 to 33·9)*	(-7·4 to 4·3)
Prostate cancer	14 434·4	70·6	29·7	1150·3	60·5	21·5
	(11 932·2 to 19 785·1)	(61·6 to 81·4)*	(22·5 to 38·4)*	(804·9-1 643·3)	(51·7 to 70·6)*	(14·5 to 29·6)*
Prostate cancer	13 492·3	74·6	32·5	1069·7	63·5	23·5
aggregate	(10 984·7 to 18 837·5)	(64·7 to 86·3)*	(24·6 to 42·1)*	(739·5–1 543·7)	(53·7 to 74·9)*	(15·8 to 32·6)*
Impotence and incontinence due to prostate cancer aggregate	942·0	29·0	-0·9	80·6	28·9	-1·0
	(927·1 to 957·3)	(28·6 to 29·4)*	(-1·2 to -0·5)*	(55·2–111·8)	(27·1 to 30·5)*	(-2·4 to 0·2)
Testicular cancer	685·8	41·9	23·4	42·1	40·3	21·5
	(634·7 to 732·4)	(30·8 to 52·4)*	(13·9 to 32·2)*	(28·4–58·3)	(29·3 to 50·6)*	(11·9 to 30·4)*
Kidney cancer	2870·3	57·9	22·8	202·7	54·4	19·8
	(2728·0 to 3031·5)	(50·0 to 66·0)*	(16·8 to 29·1)*	(145·7-270·5)	(46·9 to 61·8)*	(14·0 to 25·6)*
Bladder cancer	3407·9	35·7	3·3	267·0	32·1	0·7
	(3240·0 to 3603·3)	(28·6 to 43·3)*	(-2·0 to 9·0)	(193·7-349·4)	(25·8 to 38·8)*	(-4·0 to 5·6)
Bladder cancer episodes aggregate	3273·4	36·3	3·7	244·1	33·1	1·2
	(3 105·3 to 3 470·7)	(28·9 to 44·3)*	(-1·8 to 9·6)	(176·2–317·4)	(26·1 to 40·4)*	(-3·9 to 6·7)
Urinary incontinence due to bladder cancer	134·3	22·8	-5·7	22·9	23·0	-5·3
	(129·7 to 139·4)	(22·0 to 23·7)*	(-6·2 to -5·1)*	(15·9–31·2)	(18·7 to 27·1)*	(-8·6 to -2·2)*
Brain and nervous system cancer	1205·1	25·3	4·2	126·1	24·7	2·8
	(1101·7 to 1322·9)	(14·9 to 37·6)*	(-3·9 to 13·8)	(90·3-164·5)	(15·0 to 36·3)*	(-4·7 to 12·0)
Thyroid cancer	3166·5	101·1	60·8	190·0	96·8	56·5
	(2936·7 to 3340·6)	(86·4 to 112·9)*	(48·9 to 70·2)*	(132·5-259·3)	(81·6 to 109·2)*	(44·3 to 66·3)*
Mesothelioma	60·8	41·3	9·5	12·2	39·5	7·9
	(58·1 to 63·6)	(35·1 to 47·7)*	(4·6 to 14·5)*	(8·7–15·8)	(32·1 to 46·8)*	(2·1 to 13·9)*
Hodgkin's lymphoma	574·4	17·8	0·0	48·5	14·6	-3.6
	(519·0 to 672·9)	(10·5 to 25·3)*	(-6·1 to 6·4)	(33·7-65·8)	(7·4 to 21·7)*	(-9.6 to 2.4)*
Non-Hodgkin lymphoma	4292·3	63·5	30·5	312·3	57·9	25·6
	(3741·0 to 4593·6)	(47·5 to 74·3)*	(19·1 to 38·4)*	(221·8–414·2)	(43·8 to 67·8)*	(15·1 to 33·1)*
Multiple myeloma	488·2	54·6	18·0	104·0	49·0	13·6
	(449·4 to 527·7)	(44·8 to 64·5)*	(10·7 to 25·4)*	(73·6−134·8)	(40·0 to 58·5)*	(6·9 to 21·0)*
Leukaemia	2303·6	35·8	10·2	379·4	28·6	4·3
	(2216·1 to 2384·5)	(29·7 to 41·8)*	(5·7 to 14·8)*	(276·2–488·9)	(21·2 to 36·9)*	(-1·1 to 10·4)
Acute lymphoid leukaemia	875·5 (709·0 to 1072·3)	25·4 (7·3 to 46·4)*	10·2 (-5·0 to 27·8)	97·3 (68·0–129·8)	25·7 (11·5 to 40·8)* 27·6	8·1 (-3·1 to 20·0)
Chronic lymphoid leukaemia Acute myeloid	904·0 (833·2 to 974·4)	31·7 (22·2 to 42·9)* 38·4	4·5 (-2·2 to 12·4) 14·2	119·8 (87·2-152·9) 120·9	2/·6 (18·9 to 37·4)* 36·5	0·5 (-6·0 to 7·7) 10·4
leukaemia Chronic myeloid	999·3 (882·5 to 1136·8) 297·7	36·4 (26·6 to 50·7)* 22·2	(5·7 to 23·5)*	(87-3-155-6)	30·5 (27·5 to 45·6)* 17·7	10·4 (3·7 to 17·4)* -7·5
leukaemia Other neoplasms	(271·8 to 324·7)	22·2 (14·3 to 31·0)* 36·1	-3·4 (-9·3 to 3·4) 10·9	41·4 (29·5–53·6)	(10·2 to 26·0)*	-/·5 (-13·3 to -1·2)* 9·0
other neoplasitis	4577·5 (4145·5 to 4922·0)	(27·2 to 44·5)*	(4·1 to 17·6)*	323·2 (228·6–431·8)	34·4 (26·2 to 42·6)*	(2·4 to 15·4)*
Cardiovascular diseases	422 738·4 (415 534·5 to 427 870·8)	24·8 (24·0 to 25·6)*	-1·8 (-2·5 to -1·2)*	25 620·1 (18 401·6-33 656·6)	23·5 (21·7 to 24·5)*	-2·2 (-3·2 to -1·7)*
Rheumatic heart disease	33 438·8	-4·7	-16·3	1654·7	-3·4	-15·4
	(29 725·7 to 43 119·8)	(-11·0 to 0·1)	(-22·5 to -12·1)*	(1041·4-2530·9)	(-9·9 to 1·4)	(-21·5 to -11·1)*
Rheumatic heart disease, without heart failure	32 236·8 (28 520·8 to 41 912·7)	-5·6 (-11·9 to -0·8)*	-16·9 (-23·1 to -12·7)*	1518·6 (941·1–2356·4)	-5·5 (-11·7 to -0·6)*	-16·8 (-22·9 to -12·3)*

	Prevalence (thousands)			YLDs (thousands)			
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	
(Continued from previous page)							
Heart failure due to rheumatic heart disease aggregate	1202·0 (1138·2 to 1269·0)	29·8 (28·5 to 31·2)*	2·2 (1·3 to 3·2)*	136·1 (94·5-187·6)	29·6 (27·2 to 32·1)*	2·4 (0·6 to 4·2)*	
Ischaemic heart disease	110 550·3	25·9	-3·4	7274·7	30·2	-0·3	
	(100 675·9 to 121798·7)	(24·6 to 27·2)*	(-4·2 to -2·6)*	(4958·6–9940·3)	(29·1 to 31·3)*	(-1·0 to 0·3)	
Myocardial infarction episodes aggregate	15 930·1	6·4	-18·1	33·0	15·2	-13·1	
	(14 106·5 to 17 764·0)	(1·9 to 10·6)*	(-21·6 to −14·8)*	(23·0-44·9)	(12·6 to 17·9)*	(-15·1 to -11·1)*	
Angina due to ischaemic	72 344·6	29·3	-0·3	4794·5	29·5	-0·0	
heart disease aggregate	(62 699·8 to 83 460·0)	(27·7 to 30·7)*	(-1·2 to 0·5)	(3100·7-6633·6)	(27·8 to 31·0)*	(-1·0 to 0·8)	
Heart failure due to ischaemic heart disease aggregate	22 275·7 (21 272·0 to 23 320·4)	31·9 (31·1 to 32·7)*	-0·6 (-1·2 to -0·1)*	2447·2 (1736·8–3342·6)	31·9 (31·0 to 32·8)*	-0·5 (-1·1 to 0·1)	
Cerebrovascular disease	42 430·9	21·0	-4·4	6455·2	20·7	-4·2	
	(42 068·2 to 42 767·1)	(20·4 to 21·5)*	(-4·8 to -3·9)*	(4487·0–8609·6)	(19·8 to 21·6)*	(-4·9 to -3·5)*	
Ischaemic stroke	24 929·0	21·8	-5·2	3660·0	22·0	-4·9	
	(24 362·2 to 25 610·0)	(21·0 to 22·6)*	(-5·8 to -4·6)*	(2559·3-4874·9)	(21·0 to 23·0)*	(-5·7 to -4·2)*	
Chronic ischaemic stroke aggregate	24 663·2	21·9	-5·1	3605·1	22·1	-4·9	
	(24 092·1 to 25 345·8)	(21·1 to 22·7)*	(-5·8 to -4·5)*	(2512·6-4815·3)	(21·1 to 23·1)*	(-5·6 to -4·1)*	
Ischaemic stroke	265·8	16·8	-10·6	54·9	17·5	-10·0	
episodes aggregate	(246·6 to 285·1)	(14·2 to 19·7)*	(-12·7 to -8·5)*	(36·3-74·5)	(13·8 to 20·9)*	(-12·9 to -7·2)*	
Haemorrhagic stroke	18 669·6	19·2	-3·3	2795·2	19·1	-3·1	
	(18 258·7 to 19 124·5)	(18·5 to 19·9)*	(-3·8 to -2·7)*	(1945·8–3749·6)	(18·1 to 20·2)*	(-3·9 to -2·3)*	
Chronic haemorrhagic stroke aggregate	18 494·8	19·3	-3·2	2757·3	19·2	-3·0	
	(18 088·5 to 18 959·5)	(18·6 to 19·9)*	(-3·8 to -2·6)*	(1918·2–3700·6)	(18·1 to 20·3)*	(-3·8 to -2·2)*	
Acute haemorrhagic stroke aggregate	174·8	14·3	-9·1	37·9	14·6	-8·8	
	(162·3 to 187·8)	(11·6 to 16·6)*	(-11·1 to -7·3)*	(24·7–51·2)	(11·8 to 17·0)*	(-11·0 to -6·9)*	
Hypertensive heart disease	6086·2	37·1	3·4	670·0	37·2	3·6	
	(5732·7 to 6434·1)	(36·2 to 37·9)*	(2·9 to 3·9)*	(467·1–917·1)	(35·9 to 38·4)*	(2·8 to 4·4)*	
Cardiomyopathy and myocarditis	2536·8	26·7	-1·3	275·1	26·3	-1·3	
	(2404·8 to 2661·4)	(25·6 to 27·7)*	(-2·0 to -0·7)*	(190·3-377·9)	(24·6 to 28·1)*	(-2·5 to -0·2)*	
Acute myocarditis	156·3	21·4	2·2	7·8	21·1	2·1	
	(137·1 to 179·6)	(19·1 to 23·7)*	(1·3 to 3·1)*	(4·8–11·6)	(18·5 to 23·9)*	(0·3 to 3·8)*	
Heart failure due to cardiomyopathy aggregate	2380·4 (2254·3 to 2502·2)	27·0 (25·9 to 28·1)*	-1·5 (-2·2 to -0·9)*	267·3 (185·9–367·9)	26·5 (24·7 to 28·3)*	-1·4 (-2·6 to -0·2)*	
Atrial fibrillation and flutter	33 294·3	28·2	-2·5	2634·6	28·2	-2·4	
	(29 959·8 to 37 202·0)	(27·2 to 29·1)*	(-3·1 to -1·9)*	(1782·6-3637·3)	(27·1 to 29·3)*	(-3·1 to -1·7)*	
Peripheral vascular disease	154650·6	34·4	1·8	572·8	34·5	2·0	
	(136318·0 to 176210·9)	(33·4 to 35·2)*	(1·3 to 2·3)*	(272·1–1056·2)	(32·4 to 36·8)*	(1·2 to 2·8)*	
Endocarditis	115·7	22·7	-5·4	12·3	22·9	-5·2	
	(108·1 to 124·8)	(20·9 to 24·5)*	(-6·7 to -4·1)*	(8·5-17·1)	(18·6 to 27·6)*	(-8·8 to -1·2)*	
Endocarditis episodes	16·9	22·8	-0·7	1·0	22·7	-0·9	
aggregate	(15·7 to 18·2)	(20·3 to 25·3)*	(-2·7 to 1·2)	(0·6-1·4)	(20·0 to 25·3)*	(-3·1 to 1·2)	
Heart failure due to endocarditis aggregate	98·8	22·7	-6·1	11·4	22·9	-5·5	
	(91·1 to 107·6)	(20·7 to 24·7)*	(-7·6 to -4·6)*	(7·9-15·9)	(18·3 to 28·0)*	(-9·3 to -1·2)*	
Heart failure due to other cardiovascular diseases aggregate	1126·5 (1045·9 to 1206·1)	34·4 (33·3 to 35·6)*	1·2 (0·4 to 2·0)*	123·7 (86·0-173·3)	34·4 (32·5 to 36·5)*	1·3 (-0·2 to 2·7)	
Other cardiovascular diseases episodes aggregate	89 221·1 (83 635·8 to 95 470·4)	24·0 (22·6 to 25·4)*	0·4 (-0·2 to 1·0)*	5947·0 (4074·6–8105·9)	23·7 (22·2 to 25·2)*	0·5 (-0·2 to 1·1)	
Chronic respiratory diseases	514 625.8	12·1	-3·3	30 465·9	11·8	-4·6	
	(503 322.3 to 527 617.0)	(11·1 to 13·1)*	(-4·1 to -2·4)*	(23 341·4-38 294·2)	(10·2 to 13·6)*	(-6·2 to -3·1)*	
Chronic obstructive pulmonary disease	174 483·1	17·0	–5·8	12 047·0	16·2	-5·9	
	(160 204·9 to 188 951·7)	(15·1 to 19·0)*	(–7·3 to −4·4)*	(10 206·8-13 725·4)	(13·4 to 18·8)*	(-8·0 to -3·9)*	
					(Table 3	continues on next page)	

	Prevalence (thousands)			YLDs (thousands)		
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tinued from previous page)						
Chronic obstructive pulmonary disease without heart failure aggregate	168 694·6 (154 443·1 to 182 957·1)	16·5 (14·5 to 18·5)*	-6·1 (-7·6 to -4·6)*	9671·6 (8183·6–11 080·9)	12·3 (9·2 to 15·3)*	-8·0 (-10·5 to -5·6)*
Severe chronic obstructive pulmonary disease with heart failure aggregate	5876·4 (5073·7 to 6548·3)	34·9 (31·1 to 37·8)*	2·4 (-0·3 to 4·2)	2375·5 (1890·3–2747·2)	35·1 (31·4 to 38·2)*	2·5 (-0·0 to 4·5)
Pneumoconiosis	2405·8 (2317·2 to 2486·3)	30·3 (27·8 to 32·9)*	8·2 (6·2 to 10·2)*	474·0 (305·6–682·6)	26.6 (23.6 to 29.3)*	6·0 (3·6 to 8·4)*
Silicosis	402·9 (364·9 to 442·2)	18·1 (15·7 to 20·3)*	-4·3 (-6·1 to -2·6)*	65·7 (41·7–95·4)	18·5 (14·5 to 22·4)*	-3·9 (-7·0 to -0·9)*
Silicosis without heart failure aggregate	389·0 (351·0 to 426·8)	17·6 (15·2 to 19·9)*	-4·6 (-6·4 to -2·8)*	59·2 (36·8–87·9)	17·1 (12·7 to 21·3)*	-4·7 (-8·0 to -1·5)*
Severe silicosis with heart failure aggregate	13·9 (12·0 to 15·6)	32·2 (29·1 to 34·8)*	1·5 (-0·5 to 3·3)	6·4 (4·5-8·5)	33·1 (26·9 to 40·5)*	2·3 (-2·7 to 7·9)
Asbestosis	157·3 (144·8 to 170·9)	30·6 (28·8 to 32·6)*	2·7 (1·3 to 4·1)*	24·8 (16·0–35·6)	30·2 (27·0 to 33·6)*	2·5 (-0·1 to 5·2)
Asbestosis without heart failure aggregate	154·5 (141·8 to 168·1)	30·6 (28·7 to 32·5)*	2·7 (1·3 to 4·1)*	23·5 (15·0-34·0)	29·9 (26·4 to 33·4)*	2·5 (-0·2 to 5·4)
Severe asbestosis with heart failure aggregate	2·8 (2·6 to 3·0)	35·6 (34·2 to 37·1)*	1·1 (0·1 to 2·1)*	1·3 (0·9-1·7)*	35·7 (31·3 to 40·0)*	1·3 (-2·2 to 4·7)
Coal workers	84-2	37-0	7.6	13.3	36.3	7.0
pneumoconiosis Coal workers pneumoconiosis without heart failure	(76·7 to 93·6) 82·3 (74·9 to 91·7)	(34·1 to 39·9)* 36·9 (34·0 to 39·8)*	(5·2 to 10·0)* 7·5 (5·2 to 10·0)*	(8·4-19·1)* 12·4 (7·8-17·9)*	(29·4 to 43·4)* 36·0 (28·6 to 43·5)*	(1·3 to 12·9)* 6·9 (0·9 to 13·2)*
aggregate Severe coal workers pneumoconiosis with heart failure aggregate	1·9 (1·6 to 2·1)	41·6 (37·8 to 45·1)*	8·2 (5·4 to 10·9)*	0·9 (0·6-1·2)*	41·6 (37·7 to 45·2)*	8·2 (5·2 to 10·9)*
Other pneumoconiosis	2388·6 (2164·7 to 2629·1)	27·9 (25·3 to 30·7)*	8·3 (6·0 to 10·5)*	370·2 (239·2–533·9)*	27·5 (23·9 to 31·0)*	8·3 (5·3 to 11·2)*
Other pneumoconiosis without heart failure aggregate	2374-2 (2150-3 to 2615-1)	27·8 (25·2 to 30·7)*	8·3 (6·0 to 10·6)*	363·6 (234·0–524·9)*	27·3 (23·6 to 30·8)*	8·4 (5·3 to 11·4)*
Severe other pneumoconiosis with heart failure aggregate	14·4 (13·1 to 15·6)	41·1 (39·0 to 43·0)*	4·5 (3·4 to 5·5)*	6·5 (4·6–8·5)	40·5 (35·1 to 45·7)*	4·3 (0·3 to 8·4)*
Asthma	358 197·9 (323 133·7 to 393 465·6)	9·5 (7·6 to 11·6)*	-2·5 (-4·3 to -0·5)*	15 898·9 (10 371·0-22 344·1)	9·4 (7·4 to 11·5)*	-2·3 (-4·3 to -0·3)*
Asymptomatic asthma	108 461·1 (92 842·5 to 126 250·5)	9·5 (7·6 to 11·6)*	-2·5 (-4·3 to -0·5)*	0·0 (0·0-0·0)	0·0 (0·0 to 0·0)	0·0 (0·0 to 0·0)
Symptomatic asthma aggregate	249736.8 (220469.6 to 278419.7)	9·5 (7·6 to 11·6)*	-2·5 (-4·3 to -0·5)*	15 898·9 (10 371·0-22 344·1)	9·4 (7·4 to 11·5)*	-2·3 (-4·3 to -0·3)*
Interstitial lung disease and pulmonary sarcoidosis	1916·0 (1757·4 to 2075·5)	25·8 (23·8 to 27·7)*	0·6 (-0·9 to 2·1)	234·7 (147·5-334·2)	25·7 (23·5 to 28·1)*	0·7 (-1·1 to 2·6)
Interstitial lung disease and pulmonary sarcoidosis without heart failure aggregate	1718-7 (1565-7 to 1881-0)	25·2 (23·1 to 27·2)*	0·5 (-1·1 to 2·1)	148·5 (88·8–221·4)	22·7 (19·6 to 25·9)*	0·2 (-2·2 to 2·6)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Severe interstitial lung disease and pulmonary sarcoidosis with heart failure aggregate	197·4 (144·1 to 242·7)	31·5 (29·7 to 33·2)*	1·3 (-0·1 to 2·9)	86·2 (54·8–117·6)	31·3 (28·6 to 34·1)*	1·3 (-1·0 to 3·7)
Other chronic respiratory diseases				1811·3 (1483·7–2123·4)	1·7 (-7·3 to 12·2)	-15·8 (-23·5 to -6·9)*
Cirrhosis and other chronic liver diseases	2828·3	34·0	8·2	501·0	31·3	7·3
	(2791·3 to 2862·3)*	(32·4 to 35·8)*	(7·0 to 9·5)*	(352·9-683·1)	(29·8 to 32·9)*	(6·1 to 8·4)*
Cirrhosis and other chronic liver diseases due to hepatitis B	796-5 (737-6 to 857-3)	30·5 (28·4 to 32·7)*	4·7 (3·2 to 6·2)*	130·6 (88·7-178·7)	30·5 (27·0 to 33·8)*	4·8 (2·2 to 7·3)*
Cirrhosis and other chronic liver diseases due to hepatitis C	680-8 (628-3 to 734-2)	35·3 (33·0 to 37·5)*	7·6 (6·0 to 9·1)*	111·4 (78·2–152·6)	35·2 (31·6 to 38·6)*	7·6 (5·0 to 10·3)*
Cirrhosis and other chronic liver diseases due to alcohol use	815·6 (758·7 to 877·2)	37·3 (35·4 to 39·3)*	9.8 (8·5 to 11·2)*	133·9 (92·5–185·1)	37·2 (33·9 to 40·5)*	9·9 (7·5 to 12·4)*
Cirrhosis and other chronic liver diseases due to other causes	750·3 (714·3 to 785·9)	23·8 (22·5 to 25·1)*	6.8 (5.9 to 7.6)*	125·0 (87·5-172·2)	23·3 (20·1 to 26·8)*	6·8 (4·2 to 9·5)*
Digestive diseases	258 248·2 (255 386·4 to 260 843·9)	12·3 (10·9 to 13·7)*	-9·4 (-10·4 to -8·3)*	12142·8 (8492·0-16592·5)	11·4 (9·8 to 13·2)*	-9·3 (-10·6 to -7·9)*
Peptic ulcer disease	72 044-2	4·8	-17·0	2341·7	1·1	-19·5
	(71 302-7 to 72 763-7)	(3·7 to 6·0)*	(-17·8 to -16·0)*	(1605·6–3293·6)	(-0·2 to 2·5)	(-20·6 to -18·4)*
Peptic ulcer disease	4977·7	4·7	-16·7	518·7	4·5	-16·7
symptomatic episodes	(4577·6 to 5379·5)	(2·4 to 6·9)*	(-18·4 to -15·0)*	(357·9–703·2)	(2·2 to 6·8)*	(-18·5 to -14·9)*
Anaemia due to peptic ulcer disease aggregate	67 066-6	4·8	-17·0	1823·1	0·2	-20·3
	(66 465-4 to 67 616-9)	(3·7 to 6·1)*	(-17·9 to -16·0)*	(1225·8–2621·2)	(-1·4 to 1·9)	(-21·6 to -19·0)*
Gastritis and duodenitis	161 001·1	15·5	-6·9	4913·0	11·2	-9·5
	(157 928·3 to 164 071·1)	(13·0 to 17·9)*	(-8·8 to -5·0)*	(3347·3-6967·1)	(8·0 to 14·6)*	(-11·9 to -6·9)*
Gastritis and duodenitis, symptomatic episodes	3941·0	15·4	-6·9	416·0	15·4	-6·7
	(3547·0 to 4358·9)	(10·9 to 17·3)*	(-10·8 to -5·7)*	(277·1–567·9)	(11·2 to 17·5)*	(-10·6 to -5·2)*
Anaemia due to gastritis and duodenitis aggregate	157 060·1 (154 055·1 to 160 141·0)	15·5 (12·9 to 17·9)*	-6·9 (-8·9 to -5·0)*	4497·1 (3050·6–6393·0)	10·8 (7·4 to 14·5)*	-9·8 (-12·4 to -7·0)*
Appendicitis	442·1	14·3	2·8	136·0	14·4	3·0
	(414·6 to 472·3)	(13·4 to 15·2)*	(2·1 to 3·4)*	(92·0–184·6)	(10·6 to 18·0)*	(-0·2 to 6·1)
Paralytic ileus and intestinal obstruction	119·7	18·9	-1·3	37·3	18·6	-1·1
	(110·7 to 130·4)	(16·9 to 20·7)*	(-2·8 to -0·0)*	(25·1–51·1)	(15·9 to 21·2)*	(-3·4 to 1·2)
Inguinal, femoral, and	18 476·7	6·5	-10·6	195·8	6·5	-10·4
abdominal hernia	(16 577·2 to 20 339·8)	(4·2 to 8·6)*	(-12·4 to -9·0)*	(97·0–363·1)	(4·3 to 8·7)*	(-12·2 to -8·8)*
Inflammatory bowel	11 223·5	15·7	-3·7	2387·1	15·6	-3·6
disease	(10 396·8 to 12 000·4)	(15·0 to 16·3)*	(-4·1 to -3·4)*	(1653·2–3 263·5)	(14·7 to 16·4)*	(-4·2 to -3·0)*
Vascular intestinal	35·1	23·3	-2·4	11·0	23·4	-2·0
disorders	(32·4 to 38·1)	(21·0 to 25·8)*	(-4·1 to -0·6)*	(7·4-14·8)	(19·7 to 27·2)*	(-5·2 to 1·2)
Gallbladder and biliary	5656·5	16⋅9	-4·1	601·0	16·8	-3·9
diseases	(5199·3 to 6168·1)	(14⋅5 to 19⋅2)*	(-5·9 to -2·2)*	(409·4–826·4)	(14·2 to 19·3)*	(-5·9 to -1·9)*
Pancreatitis	1018·2	24·2	1·9	300·0	24·0	2·0
	(941·6 to 1 097·4)	(21·7 to 26·5)*	(-0·0 to 3·7)	(203·4–406·7)	(20·3 to 27·4)*	(-0·8 to 4·6)
Other digestive diseases				1219·9 (838·5-1677·5)	21·9 (14·4 to 32·4)*	−1·0 (−7·1 to 7·4)
Neurological disorders	2261316·4 (2204366·6 to 2316282·6)	15·5 (14·6 to 16·4)*	0·5 (-0·3 to 1·2)	59325.6 (40845.0-81083.3)	15·6 (14·3 to 16·9)*	-1·4 (-2·4 to -0·4)*
Alzheimer's disease and other dementias	45 956·3	37·7	1·1	6851·2	38·8	1·1
	(40 178·5 to 52 655·9)	(36·2 to 39·0)*	(0·3 to 1·8)*	(4883·9–9062·8)	(37·1 to 40·5)*	(0·2 to 1·8)*
					(Table 3	continues on next page)

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change i ASR between 2005 and 2015
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Parkinson's disease	6193-3	31.6	0-6	737-8	31.4	0.7
Epilepsy	(5725·7 to 6777·2)	(28·2 to 35·5)*	(-1·9 to 3·5)	(516·1-990·1)	(27·8 to 35·3)*	(-2·1 to 3·7)
	23 414·5	11·3	-1·1	6286·9	-6·4	-16·3
	(21 549·6 to 25 419·4)	(7·0 to 15·5)*	(-5·0 to 2·7)	(4315·8-8250·2)	(-11·5 to -1·2)*	(-20·9 to -11·6)*
Multiple sclerosis	2012·0	19·1	-2·0	667·5	18·8	-2·0
	(1865·7 to 2166·9)	(17·0 to 21·3)*	(-3·9 to -0·3)*	(472·0–855·6)	(16·2 to 21·5)*	(-4·2 to 0·1)
Motor neuron disease	202·4	22·3	-1·3	42·3	22·1	-1·4
	(189·7 to 216·0)	(19·9 to 24·8)*	(-3·3 to 0·7)	(30·5–55·1)	(19·5 to 24·8)*	(-3·5 to 0·8)
Migraine	958 789·2 (872 109·0 to 1 055 630·6)	15·3 (14·0 to 16·6)*	0.6 (-0.3 to 1.5)	32 898·8 (20 303·6-48 883·2)	15·3 (14·0 to 16·6)*	0.8 (-0.1 to 1.8)
Tension-type headache	1505 892·3 (1337 310·0 to 1681575·0)	15·3 (14·0 to 16·6)*	0·5 (-0·1 to 1·1)	2260·5 (1058·1-4169·7)	15·3 (14·0 to 16·7)*	0.6 (-0.1 to 1.3)
Medication overuse headache	58 454·5	19·0	0·4	9164·7	18·9	0·6
	(50 834·9 to 67 363·9)	(15·4 to 22·7)*	(-2·4 to 3·3)	(6094·5-13078·0)	(15·4 to 22·8)*	(-2·3 to 3·5)
Other neurological disorders	11·5	17·0	-0·2	415·9	32·2	-2·1
	(9·1 to 14·6)	(14·8 to 19·6)*	(-0·9 to 0·4)	(301·2–547·5)	(27·5 to 36·3)*	(-5·3 to 0·9)
Mental and substance use disorders	1058903.8 (1038544.9 to 1080413.8)	14·3 (13·6 to 14·9)*	0·3 (-0·3 to 0·9)	149 977·9 (108 716·1-193 130·8)	16·1 (15·5 to 16·8)*	1·1 (0·7 to 1·4)*
Schizophrenia	23 383·0	19·5	0·1	15 020·5	19·5	0·3
	(20 608·0 to 26 418·4)	(18·6 to 20·4)*	(-0·4 to 0·6)	(10 816·1–18 623·2)	(18·5 to 20·4)*	(-0·4 to 0·9)
Alcohol use disorders	63 469·5	11·1	-4·6	6321·3	11·1	-4·5
	(57 507·8 to 69 863·5)	(8·9 to 13·5)*	(-6·5 to -2·4)*	(4205·6-8985·8)*	(9·0 to 13·6)*	(-6·4 to -2·3)*
Alcohol dependence aggregate	62 575·3	11·1	-4·6	6276·2	11·1	-4·5
	(56 621·7 to 68 944·3)	(8·9 to 13·5)*	(-6·6 to -2·5)*	(4167·7-8929·7)	(8·9 to 13·6)*	(-6·5 to -2·3)*
Fetal alcohol syndrome	894·2	11·1	-0·5	45·1	10·7	-0·6
aggregate	(833·3 to 951·9)	(10·3 to 11·7)*	(-1·1 to 0·2)	(29·2–65·1)	(8·2 to 13·1)*	(-2·7 to 1·6)
Drug use disorders	46 388·6	16⋅3	3·9	9849·4	23·6	8·2
	(45 252·4 to 47 446·6)	(15⋅0 to 17⋅6)*	(2·8 to 5·0)*	(6959·0–12775·2)	(21·5 to 25·9)*	(6·2 to 10·2)*
Opioid use disorders	16746·5	23·3	6·4	6969·6	23·3	6·5
	(14659·3 to 19107·5)	(20·7 to 26·0)*	(4·2 to 8·6)*	(4842·5–8999·8)	(20·6 to 26·2)*	(4·2 to 9·0)*
Asymptomatic opioid dependence	2716·8 (1979·9 to 3704·4)	23·3 (20·7 to 26·0)*	6·4 (4·2 to 8·6)*	0.0	0·0 (0·0 to 0·0)	0·0 (0·0 to 0·0)*
Symptomatic opioid dependence aggregate	14 029·7 (12 042·2 to 16 122·8)	23·3 (20·7 to 26·0)*	6·4 (4·2 to 8·6)*	6969·6 (4842·5–8999·8)	23·3 (20·6 to 26·2)*	6·5 (4·2 to 9·0)
Cocaine use disorders	3846·3	31·1	14·1	521·5	30·6	14·0
	(3401·6 to 4309·8)	(28·0 to 34·3)*	(12·0 to 16·4)*	(329·0-733·9)	(27·2 to 34·4)*	(11·4 to 17·0)*
Asymptomatic cocaine dependence	1927·8	31·1	14·1	0·0	0·0	0·0
	(1544·3 to 2357·8)	(28·0 to 34·3)*	(12·0 to 16·4)*	(0·0–0·0)	(0·0 to 0·0)	(0·0 to 0·0)
Symptomatic cocaine dependence aggregate	1918·5	31·1	14·1	521·5	30·6	14·0
	(1542·1 to 2341·9)	(28·0 to 34·3)*	(12·0 to 16·4)*	(329·0-733·9)	(27·2 to 34·4)*	(11·4 to 17·0)*
Amphetamine use disorders	6599·8	30·4	19·2	874·7	30·3	19·1
	(5295·7 to 8024·4)	(27·0 to 34·2)*	(16·1 to 22·5)*	(513·4–1308·7)	(26·1 to 34·8)*	(15·5 to 23·0)*
Asymptomatic amphetamine dependence	3613·7 (2722·6 to 4630·7)	30·4 (27·0 to 34·2)*	19·2 (16·1 to 22·5)*	0·0 (0·0–0·0)	0·0 (0·0 to 0·0)	0·0 (0·0 to 0·0)
Symptomatic amphetamine dependence aggregate	2986·1 (2186·5 to 3 893·0)	30·4 (27·0 to 34·2)*	19·2 (16·1 to 22·5)*	874·7 (513·4-1 308·7)	30·3 (26·1 to 34·8)*	19·1 (15·5 to 23·0)*
Cannabis use disorders	19762·5	5·3	-3·8	577·2	5·3	-3·7
	(17982·4 to 21770·2)	(4·0 to 6·5)*	(-4·7 to -2·9)*	(371·8-816·2)	(3·7 to 7·1)*	(-5·0 to -2·3)*
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	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Asymptomatic cannabis dependence	11 430·9	5·3	-3·8	0·0	0·0	0·0
	(10 159·4 to 12 892·9)	(4·0 to 6·5)*	(-4·7 to -2·9)*	(0·0–0·0)	(0·0 to 0·0)	(0·0 to 0·0)
Symptomatic cannabis dependence aggregate	8331·6 (7244·3 to 9593·1)	5·3 (4·0 to 6·5)*	–3·8 (–4·7 to −2·9)*	577·2 (371·8–816·2)	5·3 (3·7 to 7·1)*	–3·7 (–5·0 to –2·3)*
Other drug use disorders				906·4 (550·1–1 312·1)	30·5 (25·7 to 35·6)*	17·2 (12·9 to 21·7)*
Depressive disorders	311147·6	18·4	0·7	54 255·4	18·2	1·0
	(300016·8 to 320544·3)	(17·2 to 19·5)*	(-0·3 to 1·6)	(37 569·9–72 943·3)	(17·2 to 19·2)*	(0·5 to 1·5)*
Major depressive	216 047·0	17·8	0·8	44 224·4	17·8	1·1
disorder	(192 863·4 to 243 319·4)	(16·6 to 19·0)*	(0·2 to 1·4)*	(29 672·5–60 297·0)	(16·6 to 19·0)*	(0·5 to 1·7)*
Dysthymia	104106·3	19·9	0·6	10 031·0	19·8	0·7
	(90398·1 to 118 968·9)	(18·4 to 21·5)*	(-0·3 to 1·4)	(6 605·9–14 267·1)	(18·3 to 21·5)*	(-0·2 to 1·6)
Bipolar disorder	44 015·8	14·9	0·3	9004·7	14·9	0·5
	(38 150·4 to 50 912·5)	(14·1 to 15·8)*	(0·1 to 0·5)*	(5501·7–13388·0)	(13·9 to 15·9)*	(-0·0 to 0·9)
Anxiety disorders	267 202·4	14·9	0·8	24 643·0	14·8	1·0
	(234 064·3 to 306 318·0)	(13·0 to 16·8)*	(-0·5 to 2·1)	(16 813·7-33 647·8)	(12·8 to 16·6)*	(-0·4 to 2·3)
Eating disorders	6468·4	19·1	9·9	1386·1	19·0	10·0
	(6170·6 to 6753·6)	(16·5 to 21·8)*	(7·5 to 12·4)*	(915·9–1941·2)	(16·9 to 21·2)*	(8·1 to 11·9)*
Anorexia nervosa	2912·1	12·0	5·1	620·5	12·1	5·2
	(2357·7 to 3596·2)	(9·3 to 14·8)*	(2·6 to 7·6)*	(411·3–897·7)	(9·0 to 15·1)*	(2·3 to 8·0)*
Bulimia nervosa	3629·4	25·3	14·2	765·6	25·3	14·2
	(2937·5 to 4406·8)	(23·6 to 27·1)*	(12·5 to 15·8)*	(490·0–1108·5)	(23·2 to 27·6)*	(12·2 to 16·2)*
Autistic spectrum	62 212·4	12·3	0·3	10 051·5	12·3	0·6
disorders	(58 979·0 to 64744·3)	(10·5 to 14·1)*	(-1·2 to 1·9)	(6740·1–13800·3)	(11·9 to 12·8)*	(0·2 to 0·9)*
Autism	24790·1	12·6	0·5	6335·9	12·5	0·7
	(20957·6 to 29393·1)	(12·2 to 12·9)*	(0·3 to 0·7)*	(4112·0-8916·1)	(11·9 to 13·1)*	(0·2 to 1·2)*
Asperger syndrome and other autistic spectrum disorders	37245·4	12·1	0·2	3715·6	12·0	0·3
	(31510·5 to 44883·1)	(11·8 to 12·4)*	(0·1 to 0·3)*	(2479·6–5424·6)	(11·5 to 12·5)*	(-0·1 to 0·7)
Attention-deficit or hyperactivity disorder	51094·3	0·3	-3·6	620·1	0·4	-3·5
	(46162·4 to 57267·7)	(-0·4 to 1·0)	(-4·2 to -3·0)*	(370·1–947·7)	(-0·6 to 1·2)	(-4·4 to -2·8)*
Conduct disorder	48 135.5	0·4	1·3	5770·5	0·5	1·4
	(39 315.2 to 58 151.6)	(-0·1 to 1·1)	(0·9 to 1·6)*	(3485·2-8955·7)	(-0·3 to 1·4)	(0·7 to 2·0)*
Idiopathic developmental intellectual disability	92 074·0	10·9	1·0	3442·1	10·3	0·2
	(52 280·1 to 130 411·2)	(9·5 to 11·9)*	(0·0 to 1·8)*	(1506·4–5996·9)	(8·2 to 11·4)*	(-1·5 to 1·2)
Other mental and substance use disorders	128 178·3	18·8	0·2	9613·4	18·7	0·3
	(127 512·5 to 128 877·2)	(17·9 to 19·7)*	(-0·6 to 0·9)	(6699·7–12 951·9)	(17·7 to 19·8)*	(-0·6 to 1·1)
Diabetes, urogenital, blood, and endocrine diseases	2 944 626.8 (2 925 016·2 to 2 968 511·3)	15·5 (14·7 to 16·1)*	-0·3 (-0·9 to 0·3)	66 092·7 (46 781·5-88 865·8)	22·3 (20·8 to 24·0)*	1·2 (0·1 to 2·3)*
Diabetes	435 328·4	30·6	4·5	33 360·8	32·5	5·4
	(404 736·1 to 468 562·4)	(28·0 to 33·0)*	(2·4 to 6·4)*	(23 043·5-45 530·8)	(29·7 to 35·3)*	(3·2 to 7·5)*
Uncomplicated diabetes	271713·6	27·8	3·2	12599·0	27·7	3·3
	(241414·2 to 302 381·7)	(24·9 to 30·4)*	(1·1 to 5·3)*	(7979·9-18394·2)	(24·8 to 30·3)*	(1·2 to 5·4)*
Neuropathy and other complications of diabetes aggregate	159 067·7	35.6	6·6	20 459·6	35·6	6·7
	(134 666·7 to 187 174·0)	(32.2 to 38.8)*	(4·1 to 9·0)*	(13 572·0–28 749·6)	(32·3 to 38·8)*	(4·2 to 9·0)*
Vision loss due to	4547·1	34·7	5·0	302·3	36·1	5·7
diabetes aggregate	(3823·8 to 5416·0)	(32·4 to 36·9)*	(3·6 to 6·3)*	(210·1–415·5)	(33·6 to 38·7)*	(4·1 to 7·4)*
Acute glomerulonephritis	100·5	-0·9	–14·5	5·0	-0·6	-14·1
	(92·1 to 110·1)	(-4·4 to 1·9)	(–17·3 to –12·6)*	(3·2–7·5)	(-4·6 to 2·8)	(-17·4 to -11·4)*
Chronic kidney disease	322 510·6	26·9	0.9	8172·8	23·8	0·1
	(312718·2 to 330 351·1)	(25·8 to 28·0)*	(-0.0 to 1.8)	(6118·6–10 229·3)	(22·9 to 24·8)*	(-0·4 to 0·6)
Chronic kidney disease	100 823·9	27·3	2·1	2456·1	23·0	0·6
due to diabetes	(86 922·7 to 115 652·3)	(24·9 to 29·9)*	(0·5 to 3·7)*	(1780·4–3151·2)	(20·7 to 25·4)*	(-0·7 to 2·0)
					(Table 3	continues on next page)

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Stage 3 chronic kidney disease due to diabetes aggregate	90 937·7 (77 906·5 to 104 617·0)	27·9 (25·4 to 30·5)*	2·3 (0·6 to 4·0)*	119·6 (79·7–175·5)	11·8 (6·5 to 16·7)*	-6·3 (-10·0 to -2·8)*	
Stage 4 chronic kidney disease due to diabetes aggregate	6441·3 (5521·3 to 7401·3)	21·6 (19·1 to 24·2)*	0·9 (-0·4 to 2·2)	787·7 (541·8–1 106·1)	20·2 (17·6 to 22·7)*	0·4 (-1·0 to 1·8)	
Stage 5 chronic kidney disease untreated due to diabetes	2694·4 (2331·9 to 3091·5)	25.6 (23.0 to 28.2)*	1·3 (-0·0 to 2·8)	1359·3 (920·5–1791·4)	25·3 (22·4 to 28·4)*	1·4 (-0·3 to 3·2)	
End-stage chronic kidney disease due to diabetes aggregate	750·5 (643·3 to 864·2)	19·8 (16·4 to 23·4)*	-3·0 (-5·2 to -0·7)*	189·5 (130·5–252·0)	26·6 (22·3 to 30·9)*	0·7 (-2·1 to 3·7)	
Chronic kidney disease due to hypertension	78 962·3 (67 846·7 to 91 021·2)	26·0 (23·6 to 28·6)*	0·2 (-1·4 to 1·9)	1508·0 (1108·7–1938·1)	28·5 (25·8 to 31·5)*	1·2 (-0·6 to 3·3)	
Stage 3 chronic kidney disease due to hypertension aggregate	72772·4 (61662·5 to 84498·2)	25·9 (23·3 to 28·6)*	0·2 (-1·6 to 2·0)	80·8 (54·2–120·1)	22·1 (17·5 to 26·6)*	-4·4 (-8·0 to -0·9)*	
Stage 4 chronic kidney disease due to hypertension aggregate	4102-2 (3503-5 to 4772-6)	25·9 (22·4 to 29·5)*	0·7 (-1·6 to 2·8)	489·8 (341·4–696·7)	25·2 (22·1 to 28·7)*	0·5 (-1·5 to 2·5)	
Stage 5 chronic kidney disease untreated due to hypertension	1652·2 (1404·5 to 1913·7)	30·7 (27·5 to 34·2)*	1·8 (-0·5 to 4·1)	816·8 (565·2–1060·3)	30·5 (27·0 to 34·5)*	2·0 (-0·5 to 4·7)	
End-stage chronic kidney disease due to hypertension aggregate	435·5 (363·3 to 509·1)	27·5 (22·0 to 33·6)*	0.6 (-3.1 to 4.9)	120·6 (83·1–162·3)	33·2 (27·4 to 39·8)*	3.8 (-0.3 to 8.9)	
Chronic kidney disease due to glomerulonephritis	67348·7 (58198·9 to 77439·5)	28·9 (25·8 to 32·3)*	1·1 (-1·1 to 3·6)	1951·4 (1432·2–2479·2)	22·2 (20·1 to 24·4)*	-0·7 (-2·1 to 0·7)	
Stage 3 chronic kidney disease due to glomerulonephritis aggregate	59790·8 (50844·6 to 69692·7)	29·9 (26·4 to 33·6)*	1·3 (-1·1 to 4·1)	111·3 (73·5-165·2)	17·1 (13·1 to 20·7)*	-5.6 (-8.8 to -2.7)*	
Stage 4 chronic kidney disease due to glomerulonephritis aggregate	4872·4 (4297·2 to 5506·6)	21·3 (18·8 to 23·9)*	-0·5 (-2·2 to 1·6)	602·5 (418·4–845·6)	20·2 (17·7 to 22·9)*	-0.5 (-2·3 to 1·5)	
Stage 5 chronic kidney disease untreated due to glomerulonephritis		23.6 (21·3 to 25·9)*	-0.6 (-2.0 to 0.7)	1122·6 (773·7–1461·3)	23·5 (20·5 to 26·3)*	-0·4 (-2·4 to 1·5)	
End-stage chronic kidney disease due to glomerulonephritis aggregate	457·4 (377·1 to 543·0)	20·5 (16·7 to 24·5)*	-2·6 (-5·1 to -0·0)*	115·0 (80·2–152·9)	26·0 (21·1 to 31·2)*	0·4 (-3·2 to 4·1)	
Chronic kidney disease due to other causes	94553·5 (81141·8 to 109371·3)	26·2 (24·2 to 28·3)*	0·1 (-1·1 to 1·5)	2257·2 (1663·4–2892·2)	23·0 (21·1 to 25·1)*	-0·5 (-1·6 to 0·6)	
Stage 3 chronic kidney disease due to other causes aggregate	85 459·8 (72 471·6 to 99 605·4)	26·6 (24·6 to 28·8)*	0·1 (-1·2 to 1·6)*	132·2 (88·0–194·5)	17·6 (14·0 to 20·8)*	-6·2 (-8·8 to -3·8)*	
Stage 4 chronic kidney disease due to other causes aggregate	6058-4 (5257-9 to 6938-4)	21·3 (18·9 to 23·5)*	-0·2 (-1·1 to 0·8)	739·7 (513·5-1 029·6)	19·4 (17·0 to 21·6)*	-1·0 (-2·2 to 0·2)	
Stage 5 chronic kidney disease untreated due to other causes	2464·6 (2115·1 to 2835·9)	25·7 (23·8 to 27·5)*	0·1 (-0·9 to 1·1)	1233·0 (855·9-1604·1)	25·5 (23·0 to 28·0)*	0·3 (-1·1 to 1·8)	

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(Continued from previous page)						
End-stage chronic kidney disease due to other causes aggregate	570.6 (485.6 to 668.1)	21·6 (18·0 to 25·0)*	-2·2 (-4·2 to -0·2)*	152·4 (107·3–200·8)	26·3 (21·8 to 30·7)*	0·1 (-3·0 to 3·2)
Urinary diseases and male infertility	436 081·7	24·2	0·9	4220·9	28·9	-0·7
	(425 577·7 to 446 450·5)	(22·8 to 25·4)*	(-0·1 to 1·9)	(2699·9-6045·2)	(26·8 to 31·0)*	(-2·1 to 0·7)
Interstitial nephritis and	2901·2	19·3	3·1	96·6	19·2	3·3
urinary tract infections	(2831·9 to 2980·4)	(19·0 to 19·7)*	(2·9 to 3·3)	(59·9-144·0)	(17·5 to 20·9)*	(1·8 to 4·7)*
Urolithiasis	319 600·2	22·8	0·9	89·7	23·4	2·8
Benign prostatic	(291 519·2 to 349 966·1)	(20·8 to 24·9)*	(-0·7 to 2·5)	(60·9–122·5)	(20·0 to 27·0)*	(0·1 to 5·5)*
	104 625·3	29·7	-1·3	3792·7	29·8	−1·3
hyperplasia	(90729.8 to 118244.0)	(27·5 to 32·0)*	(-2·8 to 0·2)	(2423·8–5437·3)	(27·6 to 32·2)*	(-2·8 to 0·3)
Male infertility	29 033·2	22·6	9.6	173·9	22·7	9.8
	(23 972·5 to 34 766·8)	(19·3 to 25·6)*	(6.8 to 12.1)*	(70·1–365·3)*	(19·3 to 25·9)*	(6.9 to 12.5)*
Other urinary diseases				67·9 (45·5–96·5)	17·2 (9·1 to 27·2)*	-0·6 (-7·3 to 7·7)*
Gynaecological diseases	794303·2	13·8	-1·8	10 001·2	10·7	-3·3
	(786497·9 to 801169·7)	(13·2 to 14·5)*	(-2·4 to -1·2)*	(6807·0–14 312·3)	(9·4 to 11·9)*	(-4·3 to -2·5)*
Uterine fibroids	151115·0	19·2	-0·4	2383·5	11·1	-5·8
	(144146·9 to 158 477·5)	(18·8 to 19·5)*	(-0·6 to -0·2)*	(1450·2–3834·4)	(8·6 to 13·2)*	(-7·7 to -4·2)*
Uterine fibroids cases aggregate	108 335·0	23·0	1·7	645·9	26·1	3·4
	(101 618·4 to 115 233·6)	(22·3 to 23·7)*	(1·2 to 2·1)*	(314·9-1193·3)	(25·0 to 27·3)*	(2·7 to 4·3)*
Mild abdominal pain with anaemia due to uterine fibroids aggregate	42780·0 (41634·5 to 43881·4)	10·5 (9·5 to 11·5)*	-5·5 (-6·3 to -4·6)*	1737·5 (1125·7–2625·4)	6·4 (4·5 to 8·1)*	-8·9 (-10·5 to -7·4)*
Polycystic ovarian	60 106·4	10·7	-0·5	532·6	10·8	-0·5
syndrome	(46 139·3 to 75 963·4)	(10·1 to 11·4)*	(-0·9 to -0·0)*	(238·9–1 044·6)	(9·9 to 11·5)*	(-1·1 to 0·0)
Polycystic ovarian syndrome cases aggregate	47 116·8 (35 880·1 to 60 919·9)	10·4 (9·8 to 11·1)*	-0·8 (-1·2 to -0·3)*	408·0 (181·1–797·7)	10·5 (9·7 to 11·3)*	-0·7 (-1·2 to -0·1)*
Hirsutism and infertility due to polycystic ovarian syndrome aggregate	12 989·6 (8 635·6 to 18 975·6)	11·9 (10·9 to 12·7)*	0·4 (-0·4 to 1·1)	124·6 (47·2–278·0)	11·5 (9·6 to 12·9)*	0·1 (-1·5 to 1·3)
Female infertility	61300·1	24·6	11·2	344·5	25·0	11·7
	(45889·9 to 79647·1)	(19·2 to 30·0)*	(6·5 to 15·8)*	(134·3-746·7)	(19·7 to 30·0)*	(7·2 to 16·2)*
Endometriosis	10758·2	11·6	–2·6	996·4	11·7	-2·4
	(9165·5 to 12485·9)	(10·9 to 12·2)*	(–3·0 to –2·2)*	(647·1–1 384·7)	(10·4 to 12·9)*	(-3·5 to -1·5)*
Endometriosis cases aggregate	10 167·5	11·5	-2·7	937·3	11·6	-2·5
	(8 652·4 to 11 828·8)	(10·9 to 12·2)*	(-3·1 to -2·3)*	(610·7-1 312·6)	(10·3 to 12·9)*	(-3·6 to -1·6)*
Abdominal pain and infertility due to endometriosis aggregate	590·7 (415·2 to 787·2)	12·3 (11·2 to 13·3)*	-1·0 (-1·8 to -0·3)*	59·1 (34·2–90·8)	12·3 (9·5 to 15·2)*	-0·9 (-3·4 to 1·5)
Genital prolapse	161679·1	17·7	-6·6	503·0	17·8	-6·5
	(142334·7 to 182566·5)	(15·2 to 20·0)*	(-8·5 to -4·8)*	(242·5–904·3)	(15·4 to 20·3)*	(-8·4 to -4·6)*
Premenstrual syndrome	430 696·9	10·1	-2·1	3621·6	10·2	-2·0
	(410 840·7 to 450 494·4)	(7·9 to 11·8)*	(-3·9 to -0·7)*	(2249·5–5428·5)	(8·0 to 12·0)*	(-3·8 to -0·4)*
Other gynaecological diseases	45 018·8	8·0	-4·2	1619·8	6·2	-5·7
	(44 057·3 to 46 031·2)	(7·3 to 8·6)*	(-4·8 to -3·7)*	(1129·8–2210·0)	(4·9 to 7·6)*	(-6·8 to -4·6)*
Other gynaecological diseases cases aggregate	23 908·1 (22 975·7 to 24 918·0)	13·8 (13·5 to 14·0)*	-0·4 (-0·5 to -0·2)*	936·8 (635·4–1 305·9)	13·8 (12·6 to 15·1)*	-0·2 (-1·2 to 0·8)
Anaemia due to other gynaecological diseases aggregate	21110·8 (20 900·6 to 21328·2)	2·1 (0·9 to 3·4)*	-8·2 (-9·3 to -7·1)*	682·9 (461·0–978·0)	-2·7 (-4·6 to -0·7)*	-12·3 (-14·0 to −10·6)*
					(Table 3	continues on next page)

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Continued from previous page)						
Haemoglobinopathies and haemolytic anaemias	1571610·0 (1545204·5 to 1605099·8)	11·1 (9·5 to 12·6)*	-1·1 (-2·5 to 0·3)	8221·5 (5528·6–11766·6)	4·3 (2·9 to 5·5)*	-4·9 (-6·0 to -3·9)*
Thalassaemias	439·0 (405·7 to 496·3)	1·4 (-0·1 to 3·4)	-6·5 (-7·8 to -4·6)*	31·3 (21·2–44·4)	1·0 (-3·8 to 6·1)	-7·0 (-11·4 to -2·5)*
β-thalassaemia major cases aggregate	229·2 (222·7 to 238·2)	4·4 (3·6 to 5·2)*	-4·2 (-4·9 to -3·4)*	16·1 (11·0–22·9)	4·1 (-2·5 to 11·5)	-4·7 (-10·6 to 1·9)
Haemoglobin E or β-thalassaemia cases aggregate	67·4 (60·1 to 75·2)	5·4 (2·9 to 8·1)*	-2·5 (-4·8 to 0·1)	5·1 (3·4-7·3)	1·8 (-10·0 to 14·2)	-5·9 (-16·7 to 4·9)
Haemoglobin H disease cases aggregate	139·0 (107·5 to 196·4)	-5·3 (-10·0 to 0·8)	-11·8 (-16·2 to -6·3)*	9·7 (6·4-14·1)	-4·9 (-12·3 to 4·9)	-11·4 (-18·2 to -2·4)*
Heart failure due to thalassaemias aggregate	3·4 (3·2 to 3·6)	26·3 (24·3 to 28·4)*	0·1 (-1·1 to 1·4)*	0·4 (0·3-0·6)	26·4 (16·0 to 38·3)*	0.6 (-9.3 to 11.8)
Thalassaemias trait	279 451·4 (272 818·9 to 287 357·4)	10·5 (10·1 to 11·0)*	-2·1 (-2·5 to -1·7)*	3922·2 (2607·9–5653·6)	6·1 (4·8 to 7·4)*	-3·6 (-4·6 to −2·6)*
β-thalassaemia trait cases aggregate	226 029·6 (220 111·5 to 233 332·0)	10·1 (9·6 to 10·5)*	-2·5 (-3·0 to -2·1)*	3661·7 (2437·6–5270·8)	5·9 (4·7 to 7·2)*	-3·8 (-4·9 to -2·8)*
Haemoglobin E trait cases aggregate	53 421·8 (51 122·3 to 55 882·5)	12·6 (11·5 to 13·7)*	-0·1 (-1·0 to 0·8)	260·5 (174·7–382·8)	8·4 (5·2 to 11·3)*	-0·5 (-3·2 to 2·1)
Sickle-cell disorders	4449·9 (4293·7 to 4600·7)	8·2 (5·2 to 11·9)*	2·8 (-0·2 to 6·2)	371·4 (259·0–518·0)	7·7 (3·3 to 12·3)*	2·3 (-1·8 to 6·6)
Homozygous sickle- cell and severe sickle- cell/β-thalassaemia cases aggregate	4035·1 (3876·0 to 4173·7)	6·5 (3·3 to 10·3)*	1·2 (-1·8 to 4·8)	339·4 (237·7-472·1)	6·5 (2·0 to 11·4)*	1·3 (-2·9 to 5·9)
Haemoglobin SC disease cases aggregate	397·6 (368·2 to 427·4)	29·6 (18·3 to 45·6)*	21·5 (11·0 to 36·5)*	30·1 (21·1–42·1)	22·9 (9·6 to 37·4)*	15·2 (2·8 to 28·9)*
Mild sickle-cell/β- thalassaemia cases aggregate	17·2 (15·9 to 18·6)	16·1 (10·1 to 21·8)*	6·4 (0·8 to 11·5)*	1·9 (1·3-2·6)	13·2 (4·7 to 21·9)*	2·7 (-4·9 to 10·5)
Sickle-cell trait	404565·9 (381223·4 to 448154·6)	19·4 (18·3 to 20·3)*	7·5 (6·5 to 8·3)*	1720·3 (1156·5–2459·0)	10·8 (7·3 to 13·4)*	1·4 (-1·8 to 3·6)
G6PD deficiency	247 073·7 (209 306·7 to 286 712·7)	6.9 (1.4 to 11.9)*	-4·4 (-9·3 to 0·1)	28·4 (19·3–39·6)	1⋅8 (-2⋅1 to 6⋅0)	-7·7 (-11·2 to -4·0)*
G6PD cases aggregate	247 068·5 (209 301·6 to 286 707·7)		-4·4 (-9·3 to 0·1)	27·7 (18·9–38·7)	1·2 (-2·8 to 5·5)	-8·1 (-11·6 to -4·3)*
Heart failure due to G6PD deficiency aggregate	5·2 (4·7 to 5·7)	38·1 (36·6 to 39·8)*	7·9 (6·8 to 9·1)*	0·6 (0·4-0·9)	38·1 (34·6 to 41·5)*	7·8 (5·0 to 10·6)*
G6PD trait	728 549·2 (676 734·8 to 781 533·8)	9.8 (7·9 to 11·4)*	-2·7 (-4·3 to -1·2)*	28·0 (19·3–38·9)	4·3 (0·1 to 8·8)*	-5·3 (-9·0 to -1·2)*
Other haemoglobinopathies and haemolytic anaemias	74385·1 (73897·6 to 74861·6)	1·8 (0·9 to 2·7)*	-9·0 (-9·7 to -8·2)*	2119·8 (1423·9–3029·9)	-3·8 (-5·5 to -2·2)*	-12·3 (-13·8 to -11·0)*
Other haemoglobinopathies and haemolytic anaemias cases aggregate	74273·3 (73786·0 to 74751·1)	1.8 (0.9 to 2.7)*	-9·0 (-9·8 to -8·2)*	2067·4 (1384·3-2963·9)	-4·1 (-5·8 to -2·4)*	-12·6 (-14·1 to -11·2)*

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(Continued from previous page)							
Heart failure due to other haemoglobinopathies and haemolytic anaemias aggregate	111·8 (103·8 to 119·6)	34·9 (33·9 to 36·0)*	2·0 (1·3 to 2·7)*	13·1 (8·9–18·2)	34·9 (31·2 to 38·7)*	2·1 (-0·8 to 5·0)	
Endocrine, metabolic, blood, and immune disorders	66 128-9 (65 597-1 to 66 663-8)	4·0 (2·2 to 5·9)*	-7·2 (-8·7 to -5·5)*	2110·5 (1431·5–2993·7)	1·3 (-1·1 to 3·8)	-8·3 (-10·4 to -6·1)*	
Endocrine metabolic blood and immune disorders cases aggregate	7577·2 (7382·6 to 7766·6)	19·8 (19·0 to 20·6)*	-0·6 (-1·2 to 0·0)	292·4 (200·2-401·8)	19·5 (18·1 to 21·0)*	-0·5 (-1·6 to 0·6)	
	58 430·4 (57 936·0–58 946·7)	2·2 (0·3-4·3)*	-8·0 (-9·7—6·2)*	1804·4 (1216·0–2582·3)	-1·3 (-3·9-1·5)	-9·5 (-11·9—7·0)*	
Anaemia due to endocrine metabolic blood and immune disorders aggregate	58 430·4 (57 936·0 to 58 946·7)	2·2 (0·3 to 4·3)*	-8·0 (-9·7 to -6·2)*	1804·4 (1216·0-2582·3)	-1·3 (-3·9 to 1·5)	-9·5 (-11·9 to -7·0)*	
Heart failure due to endocrine metabolic blood and immune disorders aggregate	121·4 (113·3 to 130·0)	30·7 (29·6 to 31·9)*	-1·9 (-2·6 to -1·2)*	13·6 (9·4-19·1)	30·7 (27·4 to 34·0)*	-1·7 (-4·3 to 0·8)	
Musculoskeletal disorders	1304100·4 (1288602·8 to 1316641·4)	20·7 (20·2 to 21·2)*	-0·7 (-1·1 to -0·3)*	146783·8 (106764·7-194473·5)	20·5 (19·6 to 21·5)*	-0·7 (-1·3 to -0·0)*	
Rheumatoid arthritis	24 491·2 (22 552·0 to 26 750·7)	23·8 (21·1 to 26·7)*	0·6 (-1·5 to 2·9)	5777·8 (4016·1-7769·6)	23·6 (20·9 to 26·7)*	0·7 (-1·4 to 3·1)*	
Osteoarthritis	237368-6 (230335-9 to 244648-1)	32·9 (31·9 to 33·8)*	2·2 (1·6 to 2·9)*	12 886·2 (8 999·7–17 540·0)	34·8 (33·6 to 36·0)*	3·9 (3·0 to 4·8)*	
Osteoarthritis of the hip cases aggregate	35 629·2 (32 482·6 to 38 970·1)	33·5 (32·4 to 34·6)*	1·8 (1·0 to 2·6)*	1776·2 (1224·9–2477·0)	35·8 (34·4 to 37·3)*	3·7 (2·6 to 5·0)*	
Osteoarthritis of the knee cases aggregate	201739·4 (195205·3 to 208276·6)	32·7 (31·7 to 33·9)*	2·3 (1·5 to 3·1)*	11110·0 (7742·1–15123·2)	34·6 (33·3 to 35·9)*	3·9 (3·0 to 4·9)*	
Low back and neck pain	820 689.8 (803 467.4 to 837 808.9)		-2·0 (-2·6 to -1·4)*	94 941·5 (67 825·3–128 035·0)	18·6 (17·6 to 19·6)*	-2·1 (-2·7 to -1·4)*	
Low back pain	539 907·4 (521 448·6 to 559 556·0)	17·3 (16·5 to 18·2)*	-2·8 (-3·4 to -2·2)*	60 074·8 (42 721·9-82 343·7)	17·2 (16·4 to 18·1)*	-2·6 (-3·2 to -2·0)*	
Neck pain Gout	358 006·6 (313 408·4 to 409 411·0) 42 214·2	21·1 (19·0 to 23·3)* 26·4	-1·1 (-2·4 to 0·0) 0·5	34866·7 (23362·1–47636·9) 1342·8	21·0 (18·9 to 23·2)* 26·3	-1·1 (-2·3 to 0·1) 0·6	
	(37688·2 to 47495·5)	(25·2 to 27·7)*	(0·1 to 1·0)*	(910-0-1843-8)	(24·6 to 27·9)*	(-0·3 to 1·5)	
Asymptomatic gout	37712·9 (33598·4 to 42467·0)	26·4 (25·2 to 27·7)*	0.5 (0.1 to 1.0)*	0.0 (0.0–0.0)	0·0 (0·0 to 0·0)	0·0 (0·0 to 0·0)	
Other musculoskeletal disorders	342 067·7 (305 430·7 to 385 146·7)	20·7 (17·5 to 24·0)*	1·2 (-1·2 to 3·7)	31835·4 (21489·1–44268·4)	20·5 (17·2 to 23·7)*	1·3 (-1·1 to 3·8)	
Other non-communicable diseases	5316342·1 (5283701·8 to 5355928·3)	14·6 (14·3 to 14·8)*	0·4 (0·2 to 0·6)*	139 001·8 (95 459·0-197 704·3)	20·4 (19·5 to 21·5)*	1·2 (0·6 to 2·0)*	
Congenital anomalies	95706·4 (88133·4 to 102483·4)	22·5 (18·5 to 26·3)*	9·3 (5·8 to 12·8)*	8621·0 (5389·1-12 950·2)	28·5 (20·0 to 37·6)*	14·7 (7·1 to 22·7)*	
Neural tube defects	1449·6 (988·4 to 2091·8)	17·7 (14·5 to 20·3)*	7·2 (4·3 to 9·7)*	499·1 (285·3–823·7)	18·4 (14·1 to 22·2)*	8·1 (4·1 to 11·6)*	
Congenital heart anomalies	48 869·1 (34 325·7 to 72 829·2)	29·8 (25·3 to 34·1)*	15·4 (11·4 to 19·2)*	1688·2 (659·4–3116·9)	29·7 (25·6 to 33·7)*	15·6 (11·7 to 19·1)*	
Less severe heart anomalies cases aggregate	44599·5 (29800·6 to 68498·5)	29·4 (24·5 to 34·0)*	15·2 (10·7 to 19·1)*	1431·0 (532·7–2747·7)	29·3 (24·2 to 34·0)*	15·2 (10·7 to 19·2)*	
					(Table 3	continues on next page)	

	Prevalence (thousands)			YLDs (thousands)			
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change ASR between 2005 and 2015	
ontinued from previous page)							
Severe congenital	4008-6	33·7	18·5	235·0	33·5	18·5	
heart anomalies	(2671-1 to 6011-2)	(27·4 to 39·0)*	(12·9 to 23·3)*	(93·2–467·8)	(27·4 to 38·7)*	(13·2 to 23·2)*	
Critical congenital	155-8	42·0	28·6	9·2	42·0	28·9	
heart anomalies	(90-3 to 252-4)	(27·5 to 52·9)*	(15·5 to 38·5)*	(3·2–18·7)	(26·4 to 54·2)*	(14·9 to 40·1)*	
Heart failure due to congenital heart anomalies aggregate	105·2	11·6	-1·0	12·9	11·6	-1·0	
	(99·2 to 111·4)	(10·9 to 12·4)*	(-1·5 to -0·5)*	(8·8–18·2)	(8·3 to 15·1)*	(-3·9 to 2·0)	
Cleft lip and cleft palate	6882-6	19·5	6·6	79·6	12·2	0·9	
	(4029-2 to 11 483-5)	(7·7 to 27·5)*	(-3·9 to 13·9)	(39·5-143·1)	(2·6 to 19·9)*	(-7·9 to 7·7)	
Down syndrome	5361·8	17·9	6·4	507·0	20·3	6·5	
	(3424·6 to 8192·1)	(13·7 to 21·8)*	(2·3 to 10·0)*	(286·2–837·2)	(15·8 to 24·2)*	(2·2 to 10·2)*	
Turner syndrome	372·4	11·9	0·1	6.6	11·4	-0·1	
	(174·6 to 677·1)	(9·8 to 13·9)*	(-1·7 to 1·9)	(2.4–14.4)	(8·7 to 14·0)*	(-2·4 to 2·2)	
Klinefelter syndrome	220·5	12·8	0·5	1·3	13·5	0·2	
	(115·5 to 395·5)	(10·9 to 14·2)*	(-1·2 to 1·9)	(0·5–3·1)	(10·6 to 15·6)*	(-2·2 to 2·0)	
Other chromosomal abnormalities	4728.6	16·2	4·3	454·1	18·5	4·2	
	(2628.9 to 8303.2)	(11·8 to 19·8)*	(0·1 to 7·7)*	(233·5–852·1)	(13·6 to 22·3)*	(-0·5 to 7·9)	
Other congenital anomalies	32 363·0	14·5	2·0	5385·0	31·3	17·1	
	(17 788·1 to 59 394·1)	(12·3 to 16·8)*	(-0·1 to 4·1)	(3209·4–8562·1)	(18·8 to 45·9)*	(6·1 to 30·3)*	
Skin and subcutaneous diseases	2 239 493·4 (2 222 716·3 to 2 258 252·8)	12·5 (12·1 to 12·8)*	0·7 (0·4 to 1·1)*	44 896·0 (28 943·3–67159·6)	11·7 (11·0 to 12·3)*	0·4 (0·1 to 0·7)*	
Dermatitis	245 290.6	14·0	0·3	8788-0	13·6	1·6	
	(227 283.4 to 262 752.2)	(13·2 to 14·8)*	(0·1 to 0·7)*	(5963-4-12273-5)	(12·9 to 14·4)*	(1·0 to 2·2)*	
Eczema cases	85 585.6	13·0	3·3	5239·2	13·0	3·4	
aggregate	(75 115.2 to 97 031.7)	(12·4 to 13·6)*	(2·8 to 3·8)*	(3515·1 to 7348·6)	(12·2 13·9)*	(2·7 to 4·2)*	
Contact dermatitis cases aggregate	98 015·1	15·2	-1·0	2632·0	15·0	-0·9	
	(85 097·0 to 110 396·7)	(13·3 to 16·9)*	(-1·2 to -0·9)*	(1625·8–3852·4)	(13·1 to 16·8)*	(-1·4 to -0·5)*	
Seborrhoeic dermatitis cases aggregate	61 690·0 (54 642·8 to 68 788·8)	13·3 (12·0 to 14·7)*	-1·3 (-1·7 to -1·0)*	916·8 (521·6-1454·1)	13·2 (11·8 to 14·7)*	-1·3 (-1·8 to -0·8)*	
Psoriasis	79 699·7	17·6	0·4	6438·3	17·5	0·6	
	(76 690·9 to 82 804·5)	(17·0 to 18·3)*	(-0·1 to 0·9)	(4495·6-8734·5)	(16·6 to 18·4)*	(-0·0 to 1·2)	
Cellulitis	959·9	24·0	5·8	69·0	23.6	5.8	
	(887·6 to 1033·6)	(22·1 to 25·7)*	(4·3 to 7·2)*	(45·0–97·8)	(20.7 to 26.1)*	(3.5 to 8.1)*	
Pyoderma	5812·7	15·5	1·4	32·7	15·4	1·5	
	(5581·7 to 6019·0)	(14·7 to 16·3)*	(0·7 to 2·1)*	(13·2-68·4)	(14·0 to 16·8)*	(0·3 to 2·7)*	
Scabies	204151·7	6·6	-2·6	5268·9	6.6	-2·5	
	(177533·7 to 237 466·2)	(4·0 to 9·5)*	(-4·5 to -0·6)*	(2966·9-8605·6)	(3.8 to 9.5)*	(-4·5 to -0·5)*	
Fungal skin diseases	492372.6	13·3	1·6	2783·3	13·3	1·7	
	(448950.9 to 538232.5)	(12·5 to 14·1)*	(1·3 to 1·9)*	(1105·6–5905·3)	(12·4 to 14·1)*	(1·4 to 2·0)*	
Viral skin diseases	174 843·1	8·5	-1·1	5396·9	8·4	-1·0	
	(165 156·3 to 185 072·0)	(8·0 to 9·0)*	(-1·4 to -0·8)*	(3417·1–7959·9)	(7·9 to 9·0)*	(-1·4 to -0·6)*	
Molluscum contagiosum cases aggregate	40 464·2 (35 685·6 to 46 233·4)	5·2 (4·7 to 5·8)*	-0.9 (-1.3 to -0.5)*	1263·0 (783·0–1932·3)	5·3 (4·6 to 6·1)*	-0.8 (-1.5 to -0.2)*	
Viral warts cases	134378·9	9·5	-1·1	4133·9	9·4	-1·0	
aggregate	(125683·9 to 143028·0)	(8·9 to 10·1)*	(-1·4 to -0·7)*	(2628·4-6153·4)	(8·8 to 10·1)*	(-1·5 to -0·6)*	
Acne vulgaris	632741·0	4·6	0.8	6854·0	4·6	0.8	
	(595241·9 to 671248·9)	(3·5 to 5·6)*	(-0.1 to 1.7)	(3275·1-12713·9)	(3·5 to 5·6)*	(-0.1 to 1.8)	
Alopecia areata	20 594·9	14·0	-1·1	695·6	13·9	-1·0	
	(19 607·8 to 21 617·0)	(13·5 to 14·4)*	(-1·3 to -1·0)*	(428·5–1035·6)	(13·1 to 14·7)*	(-1·7 to -0·4)*	
Pruritus	69 582·5	17·6	0·4	741·6	17·5	0·5	
	(62 063·2 to 78 152·6)	(15·7 to 19·5)*	(-0·7 to 1·6)	(360·0–1348·6)	(15·6 to 19·4)*	(-0·7 to 1·7)	
Urticaria	67749.8	10·7	-0·1	4115·7	10·7	-0·0	
	(58743.2 to 77086.4)	(9·6 to 12·0)*	(-0·3 to 0·0)	(2567·2–5833·6)	(9·6 to 12·0)*	(-0·5 to 0·4)	

	Prevalence (thousands)			YLDs (thousands)		
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015
(Continued from previous page)						
Decubitus ulcer	1087·5	35·4	4·7	161·3	34·5	4·7
	(991·2 to 1189·7)	(33·4 to 37·4)*	(3·3 to 6·0)*	(112·0-218·7)	(31·9 to 37·3)*	(2·6 to 6·9)*
Other skin and subcutaneous diseases	605 036·3	22·8	2·4	3550·5	22·6	2·5
	(589 500·0 to 619 675·6)	(22·1 to 23·4)*	(1·9 to 2·9)*	(1706·2–6514·4)	(22·0 to 23·3)*	(2·0 to 2·9)*
Sense organ diseases	1788 125.6 (1771 367.0 to 1808 438.8)	23·7 (23·3 to 24·1)*	0.6 (0.3 to 0.9)*	68515·2 (47798·2–93894·8)	25·2 (24·2 to 26·4)*	0.6 (-0.0 to 1.3)
Glaucoma	5954·5	39·1	4·2	541·3	39·4	4·3
	(5077·5 to 6905·8)	(36·9 to 41·4)*	(2·8 to 5·5)*	(370·0-747·9)	(37·0 to 41·9)*	(2·6 to 5·8)*
Cataract	59727·0	26·9	-3·2	3879·7	25·5	-4·2
	(53633·2 to 66879·2)	(25·2 to 28·5)*	(-4·5 to −2·0)*	(2766·9–5229·2)	(24·1 to 26·9)*	(-5·2 to -3·2)*
Macular degeneration	6188-4	48·5	8·2	462·4	47·7	6·9
	(5250-3 to 7273-2)	(46·1 to 50·8)*	(6·9 to 9·8)*	(327·4-633·0)	(45·2 to 49·9)*	(5·3 to 8·6)*
Refraction and accommodation disorders	819 307·4 (789 917·4 to 848 059·2)	21·9 (20·5 to 23·2)*	-0·4 (-1·3 to 0·6)	14593·8 (9392·9–22901·1)	21·1 (20·2 to 22·1)*	-0·3 (-0·9 to 0·4)
Age-related and other hearing loss	1 210 055·2 (1 140 224·2 to 1 274 664·8)	28·3 (27·4 to 29·2)*	1·4 (1·0 to 1·8)*	40 596·8 (27 898·4–56 075·0)	26·4 (24·6 to 28·3)*	1·1 (0·0 to 2·1)*
Other vision loss	25 797·4	27·1	5·5	1 756·4	29·9	5·6
	(22 675·6 to 28 504·6)	(24·6 to 29·5)*	(4·2 to 6·7)*	(1248·9–2392·7)	(27·8 to 32·0)*	(4·3 to 6·7)*
Other sense organ	267 689.7	24·0	0·8	6684·7	23·8	0·9
diseases	(258 379.8 to 277 711.4)	(23·1 to 24·9)*	(0·1 to 1·4)*	(4185·0-9714·7)	(22·9 to 24·8)*	(0·2 to 1·5)*
Acute other sense organ diseases	1343.8	14·9	0·9	35·4	14·8	0·9
	(1303.5 to 1385.9)	(14·2 to 15·7)*	(0·4 to 1·4)*	(21·6–51·6)	(12·7 to 16·9)*	(-0·9 to 2·7)
Chronic other sense organ diseases	266 345.9	24·0	0·8	6649·3	23·9	0·9
	(257 047.4 to 276 383.0)	(23·1 to 24·9)*	(0·1 to 1·4)*	(4162·3–9662·4)	(23·0 to 24·9)*	(0·2 to 1·5)*
Oral disorders	3 521 901·1 (3 468 088·2 to 3 575 854·4)	14·5 (13·8 to 15·0)*	0·4 (-0·1 to 0·9)	16 969·6 (10 296·5–26 044·5)	22·4 (21·6 to 23·2)*	-0·2 (-0·5 to 0·1)
Deciduous caries	572 694·1	4·5	-2·3	147·2	4·1	-2·7
	(475 089·8 to 686 991·2)	(2·3 to 6·1)*	(-4·2 to -0·7)*	(63·0-292·1)	(1·6 to 5·9)*	(-4·9 to -0·8)*
Permanent caries	2521197.8 (2361418.3 to 2679668.7)	14·5 (13·7 to 15·4)*	0.8 (0.3 to 1.4)*	1743·4 (776·7–3320·9)	13·5 (12·6 to 14·4)*	-0·4 (-1·2 to 0·4)
Periodontal diseases	537506.0	25·4	1·1	3520·7	25·4	1·2
	(465113.9 to 625888.8)	(24·1 to 26·5)*	(0·5 to 1·7)*	(1359·3-7253·7)	(24·1 to 26·5)*	(0·6 to 1·8)*
Edentulism and severe tooth loss	275 619·1	27·3	-0·9	7640·3	27·3	-0·8
	(264 200·8 to 288 252·3)	(26·9 to 27·7)*	(-1·1 to -0·7)*	(5096·6–10 562·3)	(26·9 to 27·7)*	(-1·1 to -0·6)*
Other oral disorders	133719·0 (127499·7 to 139776·2)		-0·1 (-0·2 to 0·1)	3918·1 (2432·7–5879·4)	15·8 (15·3 to 16·4)*	0·0 (-0·2 to 0·3)
Injuries	1019 157·4 (977 405·7 to 1064 704·2)	16·4 (15·5 to 17·4)*	-3·4 (-4·1 to -2·6)*	41 022·0 (29 290·4–55 288·4)	8.0 (4.6 to 11.1)*	-9·9 (-12·5 to -7·5)*
Transport injuries	115 981·0	23·1	-0·0	6444·8	12·2	-8·0
	(110 575·9 to 123 047·1)	(22·5 to 23·7)*	(-0·4 to 0·4)	(4500·6–8733·0)	(8·7 to 15·8)*	(-10·7 to -5·3)*
Road injuries	107722·4	23·4	0·3	5956·1	13·3	-7·1
	(102214·5 to 114850·0)	(22·8 to 24·1)*	(-0·1 to 0·7)	(4130·8–8076·3)	(9·9 to 16·7)*	(-9·7 to -4·6)*
Pedestrian road injuries	14157·8	22·9	-0·7	770·8	9.6	-10·6
	(12664·5 to 16095·9)	(21·7 to 24·1)*	(-1·4 to 0·0)	(532·9–1060·8)	(5·5 to 13·6)*	(-13·7 to -7·6)*
Cyclist road injuries	18 700·6	21·4	-0·4	968·2	9·9	-9·0
	(16 545·4 to 20 951·1)	(19·9 to 22·7)*	(-1·2 to 0·4)	(655·2–1348·5)	(5·5 to 14·2)*	(-12·5 to -5·9)*
Motorcyclist road injuries	26 417·2	31·7	7·3	1397·9	17.6	-3·3
	(23 496·7 to 29 722·0)	(30·3 to 33·0)*	(6·5 to 8·1)*	(962·0–1919·8)	(13.0 to 22.1)*	(-7·0 to 0·2)
Motor vehicle road	44311·4	18·7	-4·1	2585·4	12·2	-8·4
injuries	(40212·6 to 49496·3)	(17·7 to 19·8)*	(-4·7 to -3·4)*	(1811·2–3556·8)	(9·6 to 14·8)*	(-10·4 to -6·6)*
					(Table 3	continues on next page)

	Prevalence (thousands)			YLDs (thousands)				
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change ASR between 2005 and 2015		
Continued from previous page)								
Other road injuries	4135.3	40.2	15.2	233.7	32:1	9.5		
Other transport injuries	(3603·7 to 4692·7)	(38.6 to 42.1)*	(14·4 to 16·1)*	(158·3–320·9)	(28·3 to 35·5)*	(6·8 to 11·9)*		
	8258·5	18.8	-3·2	488·7	0·3	-17·1		
Unintentional injuries	(7469·0 to 9141·6) 790 101·8	(17·9 to 19·7)*	(-3.6 to -2.7)* - 5.2	(348·6-660·5) 30679·5	(-4·0 to 5·2) 6·2	(-20·5 to -13·3)* -11·4		
Unintentional injuries	(759 587·4 to 824 995·0)	(13·8 to 14·8)*	-5·5 to -4·9)*	(21602.9-41953.7)	(3·1 to 9·0)*	(-13·6 to -9·2)*		
Falls	225736·9	25·1	1·7	11770·2	11·3	-8·6		
	(207 914·3 to 245787·2)	(24·4 to 25·8)*	(1·3 to 2·1)*	(8257·6-16166·4)	(6·7 to 15·7)*	(-12·1 to -5·2)*		
Drowning	4455·5	7·0	-13·1	247·3	-6·7	-23·4		
	(4041·2 to 4919·4)	(5·8 to 8·1)*	(-13·5 to -12·7)*	(173·4-335·8)	(-10·3 to -2·4)*	(-26·2 to -20·2)*		
Fire, heat, and hot substances	69 666.6	10·8	-8⋅3	2269·4	1·9	-13·5		
	(60 988.1 to 78 550.1)	(10·1 to 11·7)*	(-8⋅8 to -7⋅9)*	(1571·4–3131·8)	(−1·8 to 5·5)	(-16·1 to -11·2)*		
Poisonings	5722·4	13·8	-2·0	549·4	8.8	-4·5		
	(4829·3 to 6854·1)	(12·7 to 14·9)*	(-3·2 to -0·6)*	(367·8–791·1)	(6.7 to 10.8)*	(-6·6 to -2·6)*		
Exposure to mechanical forces	183 582·4	13·9	-3·5	3580·8	5·9	-9·9		
	(169 658·6 to 197 859·6)	(13·0 to 14·7)*	(-4·0 to -2·9)*	(2485·7–4922·9)	(2·9 to 8·9)*	(-12·1 to -7·6)*		
Unintentional firearm injuries	2333·0	13·4	-6·3	93·2	4·1	-12·6		
	(2074·5 to 2625·1)	(12·4 to 14·4)*	(-7·1 to -5·6)*	(65·4–126·2)	(0·6 to 7·4)*	(-15·0 to -10·1)*		
Unintentional suffocation	9759·3	9·7	-7·7	353·7	11·4	-6.8		
	(8142·5 to 11 916·9)	(5·5 to 13·9)*	(-10·3 to -5·1)*	(245·8–480·3)	(8·6 to 14·0)*	(-8.6 to -5.1)*		
Other exposure to mechanical forces	171 490·1	14·1	-3·2	3133·9	5·4	-10·1		
	(157 821·0 to 185 364·4)	(13·2 to 14·9)*	(-3·7 to -2·6)*	(2172·1-4356·6)	(2·2 to 8·5)*	(-12·6 to -7·7)*		
Adverse effects of medical treatment	11 158·6	3·1	-13·9	1487·4	3·1	-13·9		
	(8773·5 to 13 610·2)	(1·3 to 4·9)*	(-15·4 to -12·4)*	(926·9–2223·0)	(1·3 to 4·9)*	(-15·4 to -12·4)*		
Animal contact	34049·7	9·4	-6.6	1100·9	4.6	-8.8		
	(30774·2 to 37569·0)	(8·0 to 10·6)*	(-7.5 to -5.9)*	(763·8–1486·9)	(3.0 to 6.2)*	(-10.0 to -7.5)*		
Venomous animal contact	14 680·7	13·2	-3·6	820·7	5·3	-7·8		
	(13 173·4 to 16 375·1)	(12·2 to 14·2)*	(-4·4 to -2·8)*	(558·1–1117·5)	(3·7 to 7·1)*	(-9·2 to -6·2)*		
Non-venomous animal contact	19 369·1	6·7	-8·8	280·3	2.5	-11·4		
	(16 794·5 to 22 306·5)	(4·8 to 8·2)*	(-10·1 to -7·7)*	(187·2–410·5)	(0.2 to 4.7)*	(-13·1 to -9·8)*		
Foreign body	32 345·9 (28 877·3 to 36 026·4)	15·7 (14·1 to 17·1)*	-1·9 (-2·9 to -1·1)* -1·6	1263·3 (899·4-1722·5) 690·2	3·7 (0·2 to 7·5)* -0·2	-10·9 (-13·6 to -8·1)*		
Pulmonary aspiration and foreign body in airway	13351·2 (11069·0 to 16798·7)	15·5 (12·5 to 18·4)*	(-3.6 to 0.2)	(477·9–973·1)	-0·2 (-4·1 to 4·5)	-14·0 (-17·2 to -10·4)*		
Foreign body in eyes	1007·2	16·3	1·4	56·2	15·2	0.5		
	(444·0 to 1593·4)	(15·0 to 18·2)*	(0·8 to 1·9)*	(24·0–100·6)	(13·8 to 16·7)*	(-0.9 to 1.3)		
Foreign body in other body part	17 987·4	15·9	-2·3	517·0	8.0	-7·7		
	(15 874·1 to 20 237·9)	(14·9 to 17·0)*	(-3·0 to -1·6)*	(364·4–713·3)	(5.5 to 10.6)*	(-9·6 to -5·8)*		
Environmental heat and cold exposure	61 942·3	15·9	-4·2	2657·1	6·4	-11·0		
	(56 211·9 to 68 761·8)	(15·4 to 16·5)*	(-4·6 to -3·8)*	(1861·7-3636·2)	(3·7 to 9·4)*	(-13·0 to -8·9)*		
Other unintentional injuries	161 441·6	4·7	-13·7	5753.6	0·4	-16·7		
	(147 971·3 to 176 204·3)	(4·0 to 5·3)*	(-14·1 to -13·3)*	(3913.1-8024.1)	(−1·4 to 2·1)	(-17·8 to -15·7)*		
Self-harm and interpersonal violence	24 083·0	15·2	-5·5	1077·5	1·9	-15·1		
	(22 279·7 to 25 744·4)	(14·6 to 15·7)*	(-5·9 to -5·2)*	(762·7-1 448·2)	(-1·6 to 5·8)	(-17·8 to -12·1)*		
Self-harm	6358·6	15·2	-6·2	332·7	0.9	-15·9		
	(5682·1 to 7154·5)	(14·5 to 15·8)*	(-6·7 to -5·8)*	(232·4-446·9)	(-2.1 to 4.5)	(-18·4 to -13·2)*		
Interpersonal violence	17724·4	15·2	-5·3	744·8	2·4	-14·6		
	(16 186·6 to 19 268·4)	(14·5 to 15·9)*	(-5·7 to -4·8)*	(521·9–1007·3)	(-1·4 to 6·4)	(-17·5 to -11·6)*		
Assault by firearm	1043·6	16·1	-4·9	46·5	4·6	-13·2		
	(900·8 to 1213·2)	(14·9 to 17·3)*	(-5·8 to -4·2)*	(32·4–64·9)	(0·7 to 8·1)*	(-16·1 to -10·5)*		
Assault by sharp object	4019·5	12·5	–7·0	123·4	-6·4	-21·1		
	(3330·4 to 4668·1)	(11·5 to 13·6)*	(–7·7 to –6·3)*	(84·8–168·7)	(-11·4 to -0·9)*	(-24·8 to -16·8)*		
Assault by other means	12 661·3	16·0	-4·7	574·8	4·3	-13·3		
	(11 425·5 to 14 018·9)	(15·2 to 16·8)*	(-5·3 to -4·3)*	(404·0–782·2)	(0·8 to 8·0)*	(-15·9 to -10·4)*		
	,	. =	/			continues on next pa		

	Prevalence (thousands)			YLDs (thousands)				
	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015	2015	Percentage change in counts between 2005 and 2015	Percentage change in ASR between 2005 and 2015		
(Continued from previous page)								
Forces of nature, war, and legal intervention	88 991.7	28·8	11·7	2820·2	23·4	7·2		
	(65 131.0 to 116 786.1)	(18·4 to 40·5)*	(2·9 to 21·7)*	(1921·3 to 3850·1)	(1·7 to 47·8)*	(-11·4 to 28·3)		
Exposure to forces of nature	25 293·7	13·5	-1·3	796·9	-27·4	-35·9		
	(16 193·6 to 37 658·7)	(-2·1 to 32·2)	(-14·5 to 14·1)	(541·4 to 1091·4)	(-43·2 to -6·2)*	(-49·6 to -17·8)*		
Collective violence and legal intervention	63 697·9	36·1	18·1	2023·3	70·5	47·1		
	(45 248·7-83 186·4)	(20·9–54·9)*	(5·1–34·4)*	(1303·8–2903·0)	(33·0–118·3)*	(15·3–88·8)*		
Impairments								
Anaemia	2 359 107·9 (2 349 938·3 to 2 369 065·8)	4·0 (3·5 to 4·5)*	-7·1 (-7·6 to −6·7)*	77 876·7 (52 436·9 to 111 360·0)	-0·7 (-2·0 to 0·4)	-10·2 (-11·2 to −9·2)*		
Developmental intellectual disability	152 664·0	12·6	2·0	16 875·1	15·0	3.8		
	(113 511·5 to 190 764·1)	(12·2 to 13·0)*	(1·6 to 2·4)*	(12 072·6 to 22 373·1)	(13·4 to 17·0)*	(2.2 to 5.7)*		
Epilepsy	39 160·5	15·8	3·5	11770·0	5.8	-4·9		
	(34 270·8 to 43 685·8)	(12·6 to 18·9)*	(0·5 to 6·4)*	(8578·8 to 15276·3)	(0.5 to 11.0)*	(-9·8 to -0·1)*		
Guillain-Barré syndrome	35·0	17·0	-0·2	10·4	17·0	-0·2		
	(28·6 to 42·1)	(14·7 to 19·5)*	(-0·9 to 0·4)	(6·6 to 15·0)	(14·7 to 19·5)	(-0·9 to 0·4)		
Hearing loss	1330 902⋅9 (1261 401⋅0 to 1397 100⋅6)	26·0 (25·2 to 26·7)*	0.9 (0.5 to 1.3)*	46183·1 (31566·7 to 63209·9)	23.6 (22.1 to 25.1)*	0.6 (-0.4 to 1.5)		
Heart failure	40 048.6	32·3	0·3	6199·4	33·2	1·1		
	(38 613.2 to 41 418.2)	(31·6 to 33·1)*	(-0·2 to 0·8)	(4700·7 to 7785·5)	(31·7 to 34·6)*	(0·1 to 2·0)*		
Infertility	113113·8	21·1	8·0	752·4	19·9	7·0		
	(93429·4-136497·5)	(17·6–24·5)*	(4·9–10·9)*	(327·7–1543·8)	(16·8–22·7)*	(4·3-9·4)*		
Pelvic inflammatory	754·1	1·1	-12·0	99·9	1·3	-11·7		
disease	(652·8 to 878·7)	(0·1 to 2·1)*	(-12·7 to -11·2)*	(67·8 to 138·6)	(-0·7 to 3·4)	(-13·4 to −10·0)*		
Vision loss	939 580·2	22·4	-0·4	24 462·7	22·4	-0·2		
	(905 009·5 to 974112·3)	(21·2 to 23·6)*	(-1·3 to 0·4)	(16 954·9 to 34 455·1)	(21·6 to 23·3)*	(-0·8 to 0·3)		

Data in parentheses are 95% UIs. *Percentage changes that are statistically significant.

Table 3: Global prevalence and YLDs for 2015, percentage change of counts, and percentage change of age-standardised rates between 2005 and 2015 for all causes, Level 5 sequelae, and nine impairments

disability in Nepal, Bangladesh, and Bhutan, respectively. YLDs due to anxiety were lower than expected in all countries in the region except for Bangladesh.

In the GBD super-region of central Europe, eastern Europe, and central Asia, lower back and neck pain was the leading cause of disability in every geographical region in 2015; Although all countries in eastern and central Europe recorded higher than expected YLD ratios for most of their top ten causes, central Asia mostly had lower than expected or as expected YLD ratios. Sense organ diseases were the second leading causes of disability and depressive disorders were the third leading causes in this super-region.

Sizeable discrepancies occurred for observed and expected YLDs based on SDI throughout north Africa and the Middle East, probably reflecting the uneven achievements in development found in this region. Lower back and neck pain was the primary driver of YLDs in the region. Exceptions to this were Lebanon and Syria, for which war caused the most disability in 2015, and Afghanistan for which iron-deficiency was the leading cause. Depression, sense organ diseases, diabetes ranked second, third, and fourth, respectively, for the

region. Observed YLDs due to diabetes consistently exceeded expected levels, with two countries posting ratios higher than $3\cdot00$ (ie, Qatar $[3\cdot12]$, and the United Arab Emirates $[3\cdot52]$). Iran and Morocco recorded much higher disability due to drug use disorders than expected based on SDI.

In comparison with the rest of the world, ranks of causespecific disability, and their ratios of observed to expected values, were vastly different in sub-Saharan Africa. Of the 46 countries within the super-regions, nine had lower back and neck pain as the leading cause of YLDs in 2015. In southern Sub-Saharan Africa, HIV/AIDS was the leading cause of disability for all countries. Iron-deficiency anaemia ranked as the leading cause of YLDs for 11 countries in western sub-Saharan Africa. For the remaining countries, lower back and neck pain were primarily the leading cause of YLDs; Liberia, the exception, had onchocerciasis as its leading cause of disability. Malaria and various neglected tropical diseases caused more YLDs than expected in most west African countries. For eastern sub-Saharan Africa, sense organ diseases, iron deficiency, depressive disorders, and lower back and neck pain were leading causes of YLDs

in 2015. In central sub-Saharan Africa, skin diseases were consistently among the leading three-to-four causes of disability in this region. HIV/AIDS resulted in far more YLDs than expected based on SDI, with Equatorial Guinea, and Central African Republic recording ratios of observed versus expected YLDs higher than $2\cdot00$.

Discussion

We used 60900 data sources to estimate the incidence and prevalence of 2619 sequelae of 310 causes for 591 geographical regions for the years 1990, 1995, 2000, 2005, 2010, and 2015. After accounting for variation over time and across regions in case definitions, disease assays, survey instruments, and coding practice, and using standardised modelling approaches to deal with missing data, conflicting data, and a range of sampling and nonsampling errors, we characterised the global and national patterns in non-fatal health outcomes. We found that global age-standardised YLDs per capita decreased slightly in the last 25 years from 0.114 (0.085-0.147) in 1990 to 0.110 (0.082-0.141) per capita in 2015, a total decrease of 3.69%(3.12-4.25%) over a generation (appendix pp 714–16). The ageing of the world's population and the general increase in YLDs per capita with age resulted in an increase in global YLDs per capita. Within this overall pattern, 133 out of 310 causes had statistically significant decreases in YLDs per capita between 2005 and 2015 compared with 82 causes with statistically significant increases in YLDs per capita over the same time period. The most important contributors to global YLDs were musculoskeletal disorders (18.5% [16.4-20.9%] of all YLDs in 2015), mental and substance use disorders (18.4% [15.6-21.2%]), and the category of other NCDs (17.9% [15.0-21.7%]), dominated by hearing loss, vision loss, and skin diseases.

Typology of diseases based on YLDs and YLLs

The comparison of trends in age-standardised YLDs and YLLs provides insight into the drivers of differences in rates of increases and decreases for diseases and injuries. Diseases and injuries can be parsed into four groups. The first of these are a small set of disorders including causes such as drug use disorders and skin cancer for which agestandardised rates of both YLDs and YLLs increased at a rate of more than 0.4% per year from 2005 to 2015. A second category of diseases includes those in which YLDs are increasing but YLLs are decreasing, including disorders such as thyroid cancer, cirrhosis, and neonatal encephalopathy. One explanation for this pattern is that risk factors, broadly defined, are causing an increase in incidence, whereas access to effective treatment has improved enough to continue reducing mortality. For a third category of disorders, YLDs are essentially constant, but YLLs are decreasing. This grouping is quite large and includes many of the major causes of disability such as musculoskeletal disorders, various neurological disorders, and a number of cancers. For many infectious causes, decreased are occurring for both YLLs and YLDs but are much faster for YLLs; this pattern is to be expected if access to effective treatment is decreasing the number of deaths but not reducing prevalence or incidence. The last category includes disorders where decreases are equal, if not slightly higher, for YLDs including ischaemic heart disease, falls, cervical cancer, and other injuries. Examination of differential trends by region or country might provide insights into the role of access to treatment in reducing YLLs and, conversely, the role of increasing or decreasing risks and social determinants in driving changes in disease incidence and prevalence. More detailed exploration of these differential trends is warranted in future work.

Disability and retirement age

As life expectancy has steadily increased in most countries, there have been calls in some high SDI countries to extend retirement ages to reflect these changes in survival.²⁸⁻³⁰ A crucial factor in these debates is whether individuals are living for longer and have higher average levels of functional health at each age or not. In this study, we found that age-standardised YLDs per capita decreased, but rather slowly, over the period 2005 to 2015. Comparison of trends in age-standardised YLDs and YLLs by cause shows that the main causes of YLDs are not decreasing and in some settings might be increasing. Other than a somewhat slower rate of improvement for YLDs at oldest age (≥80 years), age discontinuities are not observable in these patterns. Ultimately, the debate on retirement age will hinge on the skill sets required for different types of work, whether work contributes to diminished or improved functional health status, and societal expectations for retirement. Our findings suggest that the burden from mental and substance use disorders, and musculoskeletal disorders, which are frequent causes of early retirement^{31,32} in terms of age-specific prevalence, has not declined much over time; the continued prevalence of these and other disorders associated with increasing age might limit the capacities of an older workforce.

Epidemiological transition

The analysis of YLD rates by Level 2 causes as a function of SDI shows a very different picture than that reported for YLLs. At the lower end of the SDI spectrum, incremental increases in SDI are associated with reductions in both age-standardised YLDs and all-age YLD rates spurred by decreases in disability from neglected tropical diseases and anaemia as well as decreases in tuberculosis, diarrhoea, pneumonia, meningitis, and other infectious diseases. At SDI levels above 0.8 (see methods appendix pp 698–711 for SDI values for each country), age-standardised rates of health loss increase slightly, attributable to higher rates of mental and substance use disorders, musculoskeletal disorders, and neurological disorders. In this phase of the epidemiological transition, this increase in rates, although small, combined with population ageing, results in increases in YLDs per 100000. The rising average YLD per capita rates have implications for health systems; a larger

eading causes 1990.		Leading causes 2005	% change number of YLDs 1990–2005	% change all-age YLD rate 1990–2005	% change age- standardised YLD rate 1990-2005	-	Leading causes 2015	% change number of YLDs 2005-15	% change all-age YLD rate 2005–15	% change age standardised YLD rate 2005-15
1 Lower back and neck pain		1 Lower back and neck pain	34.5	9.4	-1.8		1 Lower back and neck pain	18-6	4.9	-2.1
2 Iron-deficiency anaemia		2 Sense organ diseases	39.4	13.4	2.1		2 Sense organ diseases	25-2	10.8	0.6
3 Sense organ diseases	***	3 Iron-deficiency anaemia	14.8	-6.6	-0.6	٠.,,	3 Depressive disorders	18-2	4.5	1.0
4 Depressive disorders		4 Depressive disorders	32.9	8.0	0.6		4 Iron-deficiency anaemia	-3.8	-14-9	-11.6
5 Skin diseases		5 Skin diseases	21.9	-0.8	0.5		5 Skin diseases	11.7	-1.2	0.4
6 Migraine		6 Migraine	29.7	5.5	-0.3		6 Diabetes	32.5	17-2	5.4
7 Other musculoskeletal disorders		7 Other musculoskeletal disorders	51.8	23.4	13.5	/	7 Migraine	15.3	2.0	0.8
8 Anxiety disorders		8 Diabetes	69-2	37.6	20.7	/	8 Other musculoskeletal disorders	20-5	6.6	1.3
9 Diabetes	····	9 Anxiety disorders	26.1	2.6	-1.5		9 Anxiety disorders	14.8	1.5	1.0
10 Asthma		10 Asthma	2.6	-16.5	-15.5		10 Oral disorders	22-4	8.2	-0.2
11 Oral disorders		11 Oral disorders	33.9	8.9	-1.6	·····	11 Asthma	9.4	-3.3	-2.3
12 Falls	. j	12 Schizophrenia	36.1	10.7	0.7		12 Schizophrenia	19.5	5.7	0.3
13 Schizophrenia	····	13 Falls	13-4	-7.8	-13-9	. ,	13 Osteoarthritis	34.8	19-2	3.9
14 COPD		14 COPD	22-2	-0.6	-9.8	``,	14 COPD	16.2	2.8	-5.9
15 Autistic spectrum	Ţ	15 Osteoarthritis	53.0	24.4	6.3	/ ````.	15 Falls	11.3	-1.5	-8-6
16 Haemoglobinopathies	``	16 Gynaecological diseases	29-1	5.0	-3.4		16 Autistic spectrum	12-3	-0.7	0.6
17 Gynaecological diseases	7	17 Autistic spectrum	23-2	0.2	0.5		17 Gynaecological diseases	10.7	-2.1	-3.3
18 Intestinal nematode	ベ、オ	18 Other mental and substance	32.5	7.8	0.2		18 Drug use disorders	23.6	9.4	8-2
19 Osteoarthritis		19 Drug use disorders	42.1	15.6	11.6		19 Other mental and substance	18.7	5.0	0.3
20 Other mental and substance		20 Haemoglobinopathies	10.8	-9.9	-5.3		20 Medication overuse headache	18-9	5.2	0.6
21 Bipolar disorder	/ 5	21 Bipolar disorder	29.4	5.2	0.1	``	21 Bipolar disorder	14.9	1.6	0.5
22 Epilepsy	/	22 Medication overuse headache	32.6	7.9	-1.5		22 Congenital anomalies	28.5	13.7	14.7
23 Medication overuse headache	7	23 Epilepsy	10.9	-9.8	-7.9		23 Haemoglobinopathies	4.3	-7.7	-4.9
24 Other unintentional		24 Congenital anomalies	48-9	21.1	22.4	\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\	24 Chronic kidney disease	23.8	9.5	0.1
25 Drug use disorders	٠ / ١	25 Chronic kidney disease	35.3	10.1	-2.4		25 Ischaemic heart disease	30.2	15.2	-0.3
26 Diarrhoeal diseases	1. //	26 Conduct disorder	15.8	-5.8	0.7	1 \ /	26 Alzheimer's disease	38.8	22.8	1.1
27 Conduct disorder	1	27 Other unintentional	0.7	-18-1	-23.6	$\mathbb{N}_{+} \times \mathbb{N}_{+}$	27 Cerebrovascular disease	20.7	6.8	-4.2
28 Chronic kidney disease	7. 1	28 Alcohol use disorders	28.2	4.2	-2.5	X	28 Alcohol use disorders	11.1	-1.7	-4.5
29 Congenital anomalies	\$ /	29 Ischaemic heart disease	40.7	14.4	-2.7		29 Epilepsy	-6.4	-17-2	-16-3
30 Alcohol use disorders	/ //	30 Diarrhoeal diseases	-2.2	-20.5	-9.9		30 Other cardiovascular	23.9	9.6	0.5
33 Cerebrovascular disease	- / i •	31 Cerebrovascular disease				//. X	33 Conduct disorder		Communi	
34 Ischaemic heart disease	1	33 Alzheimer's disease				/ X.N	34 Other unintentional			and nutrition
36 Other cardiovascular —	_	34 Other cardiovascular				/ ```.	35 Diarrhoeal diseases		Non-com	
40 Alzheimer's disease	/	39 Intestinal nematode					46 Intestinal nematode		Injuries	Horneable

Figure 2: Leading 30 Level 3 causes of global YLDs for both sexes combined, 1990, 2005, and 2015, with percentage change in number of YLDs, and all-age and age-standardised rates

Causes are connected by lines between time periods. For the time period of 1990 to 2005 and for 2005 to 2015, three measures of change are shown: percent change in the number of YLDs, percent change in the all-age YLD rate, and percent change in the age-standardised YLD rate. YLD=years lived with disability. COPD=chronic obstructive pulmonary disease.

fraction of the population is likely to need care for many disorders. Some of these disorders are currently costly to manage. The increase in health-care costs per individual in the population that occurs once an SDI of 0.8 is exceeded is a predictable component of the epidemiological transition; these costs should be anticipated during the health planning processes of countries in that stage of the transition.

Disease-specific issues

Musculoskeletal disorders

Musculoskeletal disorders continue to be a leading cause of disability worldwide and more so when taking into account that additional musculoskeletal burden from long-term sequelae of fractures and dislocations is classified under injuries in GBD. A key component of healthy ageing is to maintain mobility, and a key public health intervention recommended for improving health outcomes for all chronic diseases is physical activity. Painful musculoskeletal disorders increase with age and are a great threat to mobility, compromising health more broadly.^{34–36} Even if cures for musculoskeletal disorders are not yet available, the clinical goal of preventing disability is attainable.^{37,38}

Mental and substance use disorders

Consistent with the findings of earlier GBD studies, GBD 2015 confirms the large contribution of mental and substance use disorders to global disability. For the first time, in GBD 2015 we found a positive association between conflict and depression and anxiety, albeit with wide uncertainty. This uncertainty is due to sparse and low-quality data for these disorders in post-conflict countries. Going forward, we plan to make separate estimates for post-traumatic stress disorder in GBD, which might add considerably more data from conflict settings and show a stronger association. GBD results have provided an evidence base to support global action such as a stated commitment by the World Bank and WHO to make mental health a global development priority39 and the consideration of mental health and substance use disorders in shaping the Sustainable Development Goals.⁴⁰ Cost-effective interventions are able to reduce the burden imposed by mental and substance use disorders, including in low-income and middle-income countries.41

Increases in deaths attributable to drug use in the USA have resulted in considerable policy and media attention. In our assessment, the ten countries with the

	1	2	3	4	5	6	7	8	9	10
Early neonatal	Iron	NN sepsis	PEM	Haemog	Other inf	Diarrhoea	NN preterm	Congenital	Endocrine	NN enceph
Late neonatal	Iron	PEM	Diarrhoea	Congenital	Haemog	NN preterm	Other nutr	NN enceph	Other inf	Epilepsy
Post neonatal	Iron	Diarrhoea	PEM	Haemog	Skin	Other NTD	Congenital	Other inf	NN preterm	Endocrine
1–4 years	Iron	Skin	PEM	Diarrhoea	Sense	Asthma	Haemog	Other NTD	Congenital	Otitis
5–9 years	Iron	Skin	Asthma	Sense	Haemog	Other NTD	Conduct	Malaria	ASD	Anxiety
10–14 years	Iron	Skin	Conduct	Anxiety	Asthma	Migraine	Sense	Depression	Back & neck	Haemog
15–19 years	Skin	Depression	Iron	Back & neck	Migraine	Anxiety	Sense	Conduct	Other MSK	Asthma
20–24 years	Depression	Back & neck	Skin	Migraine	Iron	Other MSK	Anxiety	Sense	Other mental	Drugs
25–29 years	Back & neck	Depression	Migraine	Skin	Iron	Other MSK	Anxiety	Sense	Drugs	Schiz
30-34 years	Back & neck	Depression	Migraine	Skin	Iron	Sense	Other MSK	Anxiety	Schiz	Gynae
35-39 years	Back & neck	Depression	Migraine	Sense	Other MSK	Skin	Iron	Anxiety	Diabetes	Schiz
40-44 years	Back & neck	Depression	Sense	Migraine	Other MSK	Diabetes	Skin	Iron	Anxiety	Schiz
45-49 years	Back & neck	Depression	Sense	Diabetes	Other MSK	Migraine	Skin	Iron	Anxiety	Schiz
50-54 years	Back & neck	Sense	Depression	Diabetes	Other MSK	Migraine	Skin	Osteoarth	Anxiety	Schiz
55-59 years	Back & neck	Sense	Diabetes	Depression	Other MSK	Migraine	Osteoarth	Skin	Oral	Anxiety
60-64 years	Back & neck	Sense	Diabetes	Depression	Other MSK	Osteoarth	Oral	Skin	Migraine	COPD
65-69 years	Sense	Back & neck	Diabetes	Depression	Other MSK	Osteoarth	Oral	COPD	Skin	IHD
70-74 years	Sense	Back & neck	Diabetes	Depression	Oral	Other MSK	Osteoarth	COPD	IHD	Skin
	Sense	Back & neck	Diabetes	Alzheimer's	Depression	Oral	Osteoarth	Other MSK	COPD	IHD
75-79 years		Alzheimer's	Back & neck	Diabetes	Falls	IHD	Osteoarth	Depression	COPD	Oral

Figure 3: Leading ten Level 3 causes of global age-specific years lived with disability in 2015

Each cause is coloured by the percentage change in age-specific years lived with disability from 2005 to 2015. Alzheimer's=Alzheimer disease and other dementias. ASD=autism. Back & neck=low back and neck pain. Conduct=conduct disorders. Congenital=congenital anomalies. COPD=chronic obstructive pulmonary disease. Drugs=drug use disorders. Endocrine=endocrine, metabolic, blood, and immune disorders. Gyne=gynaecological disorders. Haemog=haemoglobinopathies and haemolytic anaemias. IHD=ischaemic heart disease. Iron=iron-deficiency anaemia. NN enceph=neonatal encephalopathy due to birth asphyxia and trauma. NN preterm=neonatal preterm birth complications. NN sepsis=neonatal sepsis and other neonatal infections. Other NTD=other neglected tropical diseases. Oral=oral disorders. Osteoarth=osteoarthritis. Other inf=other infectious diseases. Other mental and substance use disorders. Other MSK=other musculoskeletal disorders. Other nutr=other nutritional deficiencies. PEM=protein-energy malnutrition. Schiz=schizophrenia. Sense=sense organ disease. Skin=skin and substuaneous diseases.

highest prevalence of opioid dependence in decreasing order in 2015 were Iran, United Arab Emirates, Russia, Morocco, the USA, Australia, Ukraine, Tunisia, Belarus, Canada, and Iraq. Among these countries, we estimated the highest excess mortality from opioid dependence in the eastern European countries at about twice the level of that in the USA, Canada, and the north African and Middle Eastern countries; the lowest rate among these highly prevalent countries was in Australia. Within the USA, prescription opioids have been estimated to account for 37% of drug overdose deaths in 2013.42,43 The availability of overdose response treatments in the form of naloxone kits to laypersons has also accelerated.44 An alternative approach is opioid substitution treatment to reduce the risk of overdose. The intensity at which countries use harm-reduction strategies such as needle exchange and opioid substitution programmes follows an inverse pattern,45 with the most intense programmes in Australia, but very rare use of such strategies in eastern Europe, suggesting that embracing harm reduction is an effective means of reducing drug deaths.

Diabetes

To obtain standard estimates for health loss due to diabetes for GBD 2015 using both fasting plasma glucose (FPG) means and standard deviations as well as diabetes prevalence, we re-extracted all available data from 1990 to 2015. Across studies, we found 20 different case definitions

for diabetes. We also included, wherever possible, studies reporting on mean FPG but not diabetes prevalence, using a regression between mean FPG and diabetes prevalence from studies reporting on both. We have also made the assessment of diabetes prevalence more consistent with cause of death data for diabetes. Our improved efforts at measuring the prevalence of diabetes confirms the increase in global age-standardised incidence, prevalence, and YLDs. Of note, the increase in YLDs at the global scale is greater than the increase in YLLs, which reflects improved access to treatments that lower case fatality. The rise of diabetes prevalence, related to the global increase in body-mass index,46 given the costs of treating the disease47 and the related increases in cardiovascular risks, poses one of the more important challenges to health systems in the coming years. This is particularly the case in regions with high prevalence of diabetes such as central America, north Africa and the Middle East, and Oceania.

Ezzati and colleagues⁴⁸ and the International Diabetes Federation (IDF)⁴⁹ have estimated diabetes prevalence for many countries; the intra-class correlation coefficient for the estimates from Ezzati and colleagues and the GBD for 2015 is 0.74, and for the IDF estimates is 0.65. These large differences appear to stem from the inclusion of cause of death data in the modelling for GBD and also from the inclusion of self-reported diabetes prevalence in the study by Ezzati and colleagues.⁴⁸ In GBD, these sources were excluded from our analyses because of changing patterns in the prevalence of known and

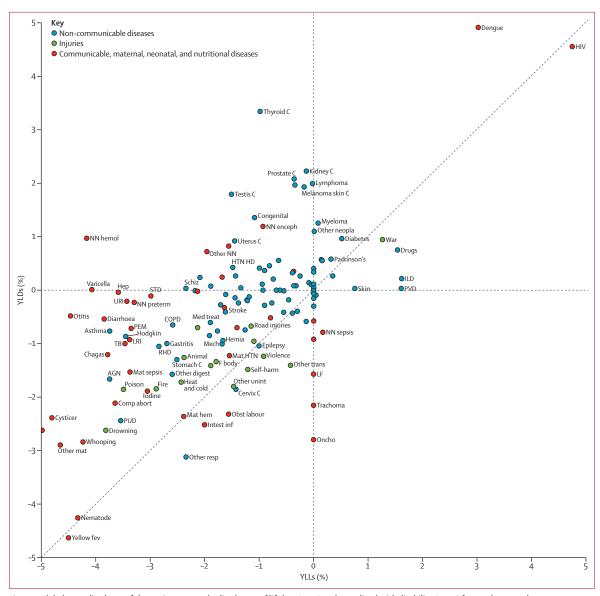


Figure 4: Global annualised rate of change in age-standardised years of life lost (YLLs) and years lived with disability (YLDs) for Level 3 causes between 1990 and 2015

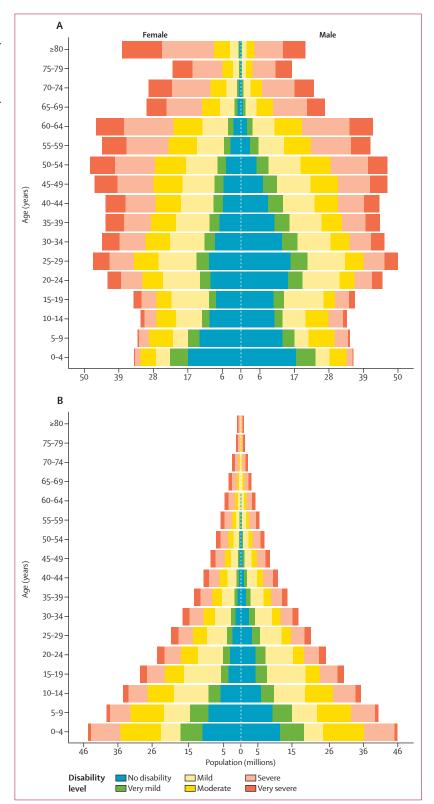
TB=tuberculosis. HIV=HIV/AIDS. Diarrhoea=diarrhoeal diseases. Intest Inf=intestinal infectious diseases. LRI=lower respiratory infections. URI=upper respiratory infections. Otitis=otitis media. Whooping=whooping cough. Varicella=varicella and herpes zoster. Chagas=chagas disease. Cysticer=cysticercosis. LF=lymphatic filariasis. Oncho=onchocerciasis. Trachoma=trachoma. Dengue=dengue. Yellow Fev=yellow fever. Nematode=intestinal nematode infections. Mat hem=maternal haemorrhage. Mat sepsis=maternal sepsis and other maternal infections. Mat HTN=maternal hypertensive disorders. Obst labour=maternal obstructed labour and uterine rupture. Comp abort=maternal abortion, miscarriage, and ectopic pregnancy. Oth mat=other maternal disorders. NN preterm=neonatal preterm birth complications. NN enceph=neonatal encephalopathy due to birth asphyxia and trauma. NN sepsis=neonatal sepsis and other neonatal infections. NN haemol=haemolytic disease and other neonatal jaundice. Oth NN=other neonatal disorders. PEM=protein-energy malnutrition. Iodine=iodine deficiency. Oth nutr=other nutritional deficiencies. STD=sexually transmitted diseases excluding HIV. Hep=hepatitis. Stomach C=stomach cancer. Melanoma=malignant skin melanoma. Skin C=non-melanoma skin cancer. Cervix C=cervical cancer. Uterus C=uterine cancer. Prostate C=prostate cancer. Testis C=testicular cancer. Kidney C=kidney cancer. Thyroid C=thyroid cancer. Hodgkin=Hodgkin lymphoma. Lymphoma=non-Hodgkin lymphoma. Myeloma=multiple myeloma. Oth neopla=Other neoplasms. RHD=rheumatic heart disease. Stroke=cerebrovascular disease. HTN HD=hypertensive heart disease. PVD=peripheral vascular disease. COPD=chronic obstructive pulmonary disease. Asthma=asthma. ILD=interstitial lung disease and pulmonary sarcoidosis. Oth resp=other chronic respiratory diseases. PUD=peptic ulcer disease. Gastritis=gastritis and duodenitis. Hernia=inguinal, femoral, and abdominal hernia. Oth digest=other digestive diseases. Parkinson=Parkinson's disease. Schiz=schizophrenia. Drugs=drug use disorders. AGN=acute glomerulonephritis. Congenital=congenital anomalies. Skin=skin and subcutaneous diseases. Road inj=road injuries. Oth trans=other transport injuries. Drown=drowning. Fire=fire, heat, and hot substances. Poison=poisonings. Mech=exposure to mechanical forces. Med treat=adverse effects of medical treatment. Animal=animal contact. F body=foreign body. Heat & cold=environmental heat and cold exposure. Oth unint=other unintentional injuries. Violence=interpersonal violence. War=collective violence and legal intervention.

unknown diabetes in surveys that vary with geography, making it difficult to crosswalk these studies.

Dementia

Brayne and colleagues50,51 reported that age-specific prevalence of Alzheimer's was decreasing in the UK. Of the four studies in the USA with similar measurements over time that were reviewed, one showed a decrease, whereas no change in prevalence was in the remaining studies. 52,53 Our assessment of age-standardised prevalence showed that Alzheimer's and other dementias decreased slowly in the UK, by 3.69% (2.53-4.85%) from 2005 to 2015, but remained constant in the rest of the world. The decrease might be due to reductions in vascular dementias rather than reductions in Alzheimer's and other dementias. Our assessment also suggests that the number of individuals with Alzheimer's disease increased from 21.7 million (18.9-24.8 million) worldwide in 1990 to 46.0 million $(40 \cdot 2 - 52 \cdot 7 \text{ million})$ in 2015. Although it is useful to know that age-standardised rates might be starting to decrease, if only slightly, the rapid increase in the absolute number of cases points to the major challenge that dementia presents to societies with increasing life expectancy. This number is similar to the most recent estimates from the World Alzheimer Report 2015 that estimated 46.8 million people living with dementia in 2015. However, there is less agreement about new cases per year: The World Alzheimer Report estimated 9.9 million new cases of dementia per year in 2015 compared with 6.44 million (5.52-7.45 million) new cases in 2015 estimated in GBD 2015. The World Alzheimer Report separately analysed prevalence and incidence whereas we use DisMod-MR 2.1 to produce internally consistent prevalence and incidence estimates. We observed a fundamental disagreement in the underlying incidence and prevalence data in many countries and decided to exclude incidence data from our modelling approach, putting greater trust in prevalence studies than incidence studies, arguing that the point of incidence in dementia is more difficult to establish than prevalence when the diagnosis over time becomes more established.

Figure 5: Population pyramids with the number of individuals, by age and sex, grouped by severity of their disability weight (DW) for all comorbid conditions combined into no disability, very mild disability (DW 0-0.01), mild disability (DW 0·01-0·05), moderate disability (DW 0·05-0·1), severe disability (DW 0·1-0·3), and very severe disability (DW >0·3) for geographies of high (A) and low (B) quintiles of Socio-demographic Index in 2015 Disability weights are combined multiplicatively as 1-(1-DW1)(1-DW2)... (1-DWn) for n comorbid sequelae. Socio-demographic Index (SDI) is calculated for each geography as a function of lag-dependent income per capita, average educational attainment in the population aged over 15 years, and the total fertility rate. SDI units are interpretable; a zero represents the lowest level of income per capita, educational attainment, and highest total fertility rate (TFR) observed from 1980 to 2015 and a one represents the highest income per capita, educational attainment, and lowest TFR observed in the same period. Cutoffs on the SDI scale for the quintiles have been selected based on examination of the entire distribution of geographies between 1980 and 2015.



Hepatitis C

The availability of new medical treatments for chronic hepatitis C has driven considerable interest in the prevalence data for this disease. Available treatments are

Males Females 0.75 Socio-demographic Index 0.50 0.25 15000 13000 11000 9000 7000 5000 3000 1000 1000 3000 5000 7000 9000 11000 13000 15000 Age-standardised YLD rate per 100 000 people В 0.75 0.50 Socio-de 0.25 3000 1000 1000 3000 5000 5000 All-age YLD rate per 100 000 people Forces of nature, war, and legal intervention Mental and substance use Other communicable, maternal. neonatal, and nutritional diseases Self-harm and interpersonal violence disorders Unintentional injuries Neurological disorders Nutritional deficiences Transport injuries Digestive diseases Neonatal disorders Other non-communicable diseases Maternal disorders Cirrhosis and other chronic Neglected tropical diseases and malaria Musculoskeletal disorders liver diseases Diabetes, urogenital, blood Chronic respiratory diseases Diarrhoea, lower respiratory, and other common infectious disease Cardiovascular diseases

Neoplasms

HIV/AIDS and tuberculosis

highly effective but costly.⁵⁴ Some programmes have been launched in Egypt and other lower-SDI countries that offer treatment at lower cost. Good data on the number of courses of treatment that have been delivered are not widely available. Given the number of individuals with chronic infection and the potential to be cured, tracking the fraction of people treated each year should be undertaken. Even in high-income countries such as the USA, the fraction of those treated among those who would benefit from treatment is not yet available.

Malaria

Our assessment of malaria prevalence and incidence in high burden sub-Saharan Africa was based on the Malaria Atlas Project spatial analysis of prevalence surveys done across Africa between 2000 and 2015.55,56 The dynamic map of malaria prevalence was updated for GBD 2015 and extended back to 1980. Children younger than 5 years in sub-Saharan Africa showed a remarkable 33.5% (24·8–46·0%) decrease in incidence from the 2005 peak of 0.78 (0.56-0.96) cases per person per year to 0.52(0.33-0.71) in 2015. Yet there were 81.7 million (52.0-111.7 million) incident cases in this age group in 2015. The decrease in children was more rapid than that recorded for older age groups. In adolescents and adults, incidence dropped by 30.0% (17.2–39.2%) and 21.1% (15.3-26.9%), respectively, and resulted in 58.7 million (37.7-91.5 million) and 53.3 million (40.7-67.5 million) incident cases, respectively, in 2015. The observed decreases in malaria incidence underscore the remarkable effect of the scale-up of antimalarial interventions⁵⁷ across Africa in the past decade.⁵⁵ Sustaining the levels of these interventions is crucial to continue to reduce malaria morbidity on the continent, avoid resurgence, and improve on these levels that are vital to global aspirations for malaria eradication.58 Overall, we estimate there were 286 · 9 million (219 · 7–377 · 3 million) cases of malaria worldwide in 2015, whereas the World Malaria Report (WMR) 2015 reports 214 million (149-303 million) cases in 2015.59 Of these cases, the WMR estimates that 88% of cases occurred in the WHO African region (188 million) whereas we estimate 189.4 million (151·7-239·4 million; 66·1%) cases for GBD 2015. Outside of Africa, the WMR reports 26 million cases versus our estimate of 97.3 million (50.2-181.8 million), but both sets of estimates feature identical regional rankings among the six WHO regions with malaria. Increasing convergence in estimates is expected as dynamic maps of

Figure 6: Expected relationship between age-standardised years lived with disability (YLD) rates per 100 000 people for the 21 GBD Level 2 causes and SDI (A) and the expected relationship between all-age years lived with disability (YLD) rates per 100 000 people for the 21 GBD Level 2 causes and SDI (B) by sex These stacked curves represent the average relationship between SDI and each cause of YLDs observed across all geographies over the time period 1990 to 2015. In each figure, the y-axis spans from lowest SDI up to highest SDI. To the left of the midline are male rates, and the female rates are to the right; higher rates are further from the midline. SDI=Socio-demographic Index.

	1	2	3	4	5	6	7	8	9	10
Global	Back & neck (0.96)	Sense (1·0)	Depression (0.93)	lron (1⋅01)	Skin (0·93)	Diabetes (0.98)	Migraine (0.93)	Other MSK (1·29)	Anxiety (0·93)	Oral (0.86)
	Back & neck	Sense	Depression	Skin	Diabetes	Migraine	Other MSK	Anxiety	Oral	Iron
High income ————————————————————————————————————	(1.11)	(0.94)	(1.11)	(0.92)	(1.29)	(1.04)	(1.27)	(1.32)	(0.95)	(0.92)
High-income North America	Back & neck (1.08)	Depression (1·31)	Diabetes (2·3)	Sense (0·89)	Other MSK (1·83)	Skin (0·94)	Anxiety (1·54)	Migraine (0·97)	Drugs (2·94)	lron (0⋅9)
	Back & neck	Sense	Depression	Skin	Other MSK	Diabetes	Migraine	lron	(2·94) Anxiety	Oral
Canada	(1.31)	(0.91)	(0.98)	(1.02)	(1.62)	(1.57)	(1.13)	(1.17)	(1.23)	(0.95)
Greenland	Back & neck	Depression	Other MSK	Skin	Sense	Migraine	Anxiety	Iron	Diabetes	Falls
	(1·1) Back & neck	(0.94) Depression	(2·18) Diabetes	(1·03) Sense	(0.68) Other MSK	(1·03) Skin	(1·28) Anxiety	(0·99) Migraine	(0·72) Drugs	(1·17) Iron
USA	(1·05)	(1.34)	(2.39)	(0.89)	(1·86)	(0.93)	(1.57)	(0·95)	(3.02)	(0.87)
Australasia	Back & neck	Depression	Sense	Other MSK	Skin	Anxiety	Migraine	Diabetes	Asthma	Drugs
Australasia	(1.12)	(1.3)	(0.84)	(1.93)	(0.93)	(1.72)	(1.17)	(1.27)	(1.59)	(2.91)
Australia	Back & neck (1·12)	Depression (1.33)	Other MSK (2·0)	Sense (0·84)	Skin (0.93)	Migraine (1·21)	Anxiety (1·72)	Diabetes (1·22)	Drugs (3·05)	Asthma (1·58)
	Back & neck	Depression	Sense	Anxiety	Skin	Diabetes	Other MSK	Migraine	Asthma	Oral
New Zealand	(1.15)	(1.14)	(0.85)	(1.77)	(0.92)	(1.51)	(1.53)	(1.0)	(1.63)	(1.09)
High-income Asia Pacific	Back & neck	Sense	Depression	Skin	Diabetes	Migraine	Oral	Other MSK	Iron	Falls
•	(0.84) Back & neck	(1.04) Depression	(0.86) Skin	(0.98) Sense	(0.96) Diabetes	(0.89) Iron	(0.96) Migraine	(0.92) Falls	(1·12) Anxiety	(0.89) Asthma
Brunei	(0·82)	(0.9)	(0.93)	(1.05)	(2.39)	(1.06)	(0·84)	(1.35)	(0.85)	(1.13)
lanan	Back & neck	Sense	Depression	Skin	Oral	Diabetes	Migraine	Other MSK	Alzheimer's	Iron
Japan	(0.82)	(1.05)	(0.87)	(1.0)	(0·97)	(0.83)	(0.91)	(0.93)	(1.05)	(1.06)
Singapore	Back & neck (0.6)	Sense (1·01)	Depression (0.95)	Skin (0.93)	Migraine (0·75)	lron (1⋅05)	Oral (0.95)	Other MSK (0.83)	Anxiety (0·85)	Falls (0·92)
	Back & neck	Sense	Depression	Diabetes	Skin	Migraine	lron	Other MSK	Oral	(0.92) Falls
South Korea	(0.9)	(1.02)	(0.83)	(1.37)	(0.92)	(0.83)	(1.28)	(0.9)	(0.94)	(1.06)
Western Europe	Back & neck	Sense	Depression	Migraine	Skin	Anxiety	Diabetes	Oral	Other MSK	Falls
Western Europe	(1.27) Back & neck	(0·94) Sense	(1.05) Depression	(1·17) Skin	(0.87) Migraine	(1·28) Oral	(0.88) Falls	(0.99) Anxiety	(0·94) Diabetes	(1·04) Iron
Andorra	(1.41)	(0·91)	(1·12)	(0.88)	(1·15)	(1.13)	(1.23)	(1·24)	(0·94)	(0.82)
	Back & neck	Sense	Depression	Migraine	Skin	Oral	Anxiety	Diabetes	Falls	Other MSK
Austria	(1.22)	(0.93)	(1.07)	(1.29)	(0.88)	(1.23)	(1.22)	(0.9)	(1.19)	(0.86)
Belgium	Back & neck	Sense	Depression (1.01)	Skin (0·87)	Migraine (1·14)	Diabetes	Oral	Falls	Anxiety	Other MSK
	(1·39) Back & neck	(1·06) Sense	Depression	Migraine	Skin	(1·1) Diabetes	(1·17) Anxiety	(1·25) Oral	(1·17) Asthma	(0.83) Other MSK
Cyprus	(1.4)	(0.96)	(1.07)	(1.14)	(0.88)	(1.27)	(1.22)	(1.09)	(1.24)	(0.79)
Denmark	Back & neck	Sense	Depression	Skin	Migraine	Other MSK	Diabetes	Oral	Anxiety	Iron
Defillark	(1.45)	(0.79)	(1.1)	(0.94)	(1.09)	(1.23)	(1.12)	(1.07)	(1.25)	(0.79)
Finland	Back & neck (1.38)	Sense (0·85)	Depression (1·23)	Skin (0⋅92)	Falls (1·58)	Migraine (1·14)	Diabetes (1.06)	Oral (1·12)	Asthma (1·35)	Iron (1⋅0)
	Back & neck	Sense	Depression	Skin	Anxiety	Migraine	Falls	Oral	Other MSK	Diabetes
France	(1.15)	(0.92)	(1.03)	(0.86)	(1.54)	(1.04)	(1.23)	(1.0)	(1.02)	(0.69)
Germany	Back & neck	Sense	Depression	Migraine	Skin	Anxiety	Diabetes	Oral	Falls	Other MSK
	(1.48) Back & neck	(0.98) Sense	(1.08) Depression	(1·23) Migraine	(0·86) Skin	(1·47) Oral	(1.02) Anxiety	(1.05) Diabetes	(1·02) Falls	(0.82) Alzheimer's
Greece	(1·28)	(0.95)	(1·21)	(1·11)	(0.83)	(1.13)	(1·19)	(0·71)	(0.86)	(1.01)
laste a d	Back & neck	Sense	Depression	Migraine	Skin	Anxiety	Oral	Iron	Diabetes	Falls
Iceland	(1.38)	(0.93)	(0.89)	(1.15)	(0.79)	(1.23)	(1.18)	(0.83)	(0.81)	(0.96)
Ireland	Back & neck (1·25)	Sense (0⋅92)	Depression (1.07)	Skin (0·91)	Migraine (1·14)	Anxiety (1·58)	lron (1⋅0)	Other MSK (1.09)	Oral (1·17)	Asthma (1·36)
	Back & neck	Sense	Depression	Skin	Migraine	Iron	Diabetes	Oral	Other MSK	War
Israel	(1.2)	(0.94)	(1.1)	(0.86)	(1.14)	(1.11)	(1.2)	(1.15)	(0.95)	(6977-72)
Italy	Back & neck	Sense	Depression	Migraine	Diabetes	Skin	Anxiety	Other MSK	Alzheimer's	Falls
reary	(1.31)	(1·09)	(1·06)	(1·39)	(1·01)	(0.86)	(1·24)	(1·0)	(1·22)	(0·94)
Luxembourg	Back & neck (1·42)	Sense (0·92)	Depression (1·11)	Migraine (1·28)	Skin (0·87)	Diabetes (1·3)	Oral (1-21)	Anxiety (1·23)	Falls (1·15)	Asthma (1·42)
	Back & neck	Sense	Depression	Diabetes	Migraine	Skin	Oral	Anxiety	Falls	Asthma
Malta	(1.31)	(0.89)	(1.02)	(1.02)	(1.13)	(0.9)	(1.15)	(1.18)	(1.13)	(1.12)
Netherlands	Back & neck	Sense	Depression	Migraine	Skin	Anxiety	Diabetes	Oral	Other MSK	Falls
	(1.5) Back & neck	(0.9) Sense	(0.99) Depression	(1·21) Anxiety	(0·87) Skin	(1·59) Migraine	(1·23) Diabetes	(1·32) Oral	(1·13) Other MSK	(0.94) Falls
Norway	(1·59)	(0.81)	(1·03)	(1·87)	(0.85)	(1·07)	(1·22)	(1.15)	(0.87)	(1.07)
Dortugal	Back & neck	Sense	Depression	Migraine	Skin	Diabetes	Oral	Anxiety	Other MSK	Asthma
Portugal	(1.34)	(0.83)	(1.09)	(1.13)	(0.88)	(0.78)	(1.09)	(1.13)	(0.94)	(1.18)
Spain	Back & neck	Sense (0.00)	Depression (1.00)	Migraine (1.15)	Skin (0.86)	Diabetes	Iron	Falls	Anxiety	Oral
•	(0.97) Back & neck	(0.99) Sense	(1.09) Depression	(1·15) Diabetes	(0·86) Skin	(0.78) Migraine	(1·25) Anxiety	(1·02) Other MSK	(1·01) Oral	(0·91) Iron
Sweden	(1.25)	(0.78)	(1.05)	(1.36)	(0.9)	(0·91)	(1.22)	(1·02)	(0.9)	(0.89)

	1	2	3	4	5	6	7	8	9	10
Switzerland	Back & neck	Sense	Depression	Skin	Migraine	Oral	Falls	Anxiety	Diabetes	Other MSK
	(1·48) Back & neck	(0.9)	(1.08)	(0.88)	(1.08)	(1.22)	(1·41) Other MSK	(1.25)	(0.89)	(0·74) Iron
UK	(1·17)	Sense (0·79)	Depression (0.96)	Skin (0.88)	Migraine (1·1)	Asthma (1.86)	(1·18)	Anxiety (1.07)	Oral (0⋅91)	(0.92)
	Back & neck	Sense	Depression	Skin	Migraine	Other MSK	Asthma	Anxiety	Oral	Iron
England	(1.17)	(0.79)	(0.95)	(0.87)	(1.04)	(1.22)	(1.79)	(1.08)	(0.91)	(0.95)
Northern Ireland	Back & neck	Sense	Depression	Skin	Migraine	Asthma	Falls	Oral	Anxiety	Diabetes
Northern itelana	(1.11)	(0.73)	(0.97)	(0.88)	(1.03)	(1.41)	(1.21)	(0.91)	(0.89)	(0.63)
Scotland	Back & neck	Migraine	Sense	Depression	Asthma	Skin	Diabetes	Other MSK	Falls	Anxiety
	(1·23) Back & neck	(1.75) Sense	(0·79) Depression	(0.97) Asthma	(2·35) Skin	(0.89) Migraine	(1.07) Diabetes	(1.02) Other MSK	(1·21) Falls	(1·12) Anxiety
Wales	(1·15)	(0.79)	(1.07)	(2·59)	(0.9)	(1·03)	(0·95)	(1.08)	(1.07)	(1.09)
Continue to the Association	Back & neck	Depression	Sense	Skin	Anxiety	Migraine	Iron	Diabetes	Other MSK	Asthma
Southern Latin America	(0.88)	(1.07)	(0.84)	(0.93)	(1.53)	(0.93)	(1.02)	(0.83)	(1.07)	(1.03)
Argentina	Back & neck	Depression	Sense	Skin	Anxiety	Iron	Migraine	Diabetes	Other MSK	Asthma
, rigeriana	(0.87)	(1.07)	(0.84)	(0.92)	(1.52)	(1.12)	(0.91)	(0.82)	(1.09)	(1.01)
Chile	Back & neck	Depression	Sense	Skin	Anxiety	Migraine	Diabetes	Other MSK	Oral	Asthma
	(0.88) Back & neck	(1.1)	(0.86)	(0.96)	(1.57)	(0.95)	(0·9)	(1.05)	(0.98)	(1.08) Other MSK
Uruguay	(0.89)	Sense (0·8)	Depression (1.03)	Skin (0.93)	Anxiety (1·49)	Iron (1·09)	Migraine (0·94)	Diabetes (0.6)	Asthma (1·11)	(0.92)
Central Europe, eastern Europe,	Back & neck	Sense	Depression	Skin	lron	Migraine	Diabetes	Oral	Anxiety	Osteoarth
and central Asia	(1·2)	(1.16)	(1.1)	(0.89)	(1.45)	(0·99)	(1.0)	(0.99)	(0.81)	(1.29)
Forton Forton	Back & neck	Sense	Depression	Iron	Skin	Migraine	Diabetes	Oral	Drugs	Osteoarth
Eastern Europe	(1.21)	(1.25)	(1.17)	(1.81)	(0.89)	(1.01)	(0.93)	(0.96)	(3.26)	(1.33)
Belarus	Back & neck	Sense	Depression	Skin	Migraine	Diabetes	Iron	Falls	Oral	Drugs
Delai US	(1.19)	(1.18)	(1.15)	(1.02)	(1.02)	(0.97)	(1.22)	(1.13)	(0.89)	(2.84)
Estonia	Back & neck	Sense	Depression	Skin	Migraine	Diabetes	Iron	Oral	Osteoarth	Alcohol
	(1·16) Back & neck	(1.08)	(1.25)	(0.88)	(1.03)	(1.05)	(1·16)	(0·93) Oral	(1.38)	(5·28) Anxiety
Latvia	(1·24)	Sense (1·22)	Depression (0.97)	Diabetes (1·32)	Skin (0⋅85)	Migraine (1·03)	Iron (1·18)	(0.89)	Osteoarth (1·35)	(0.77)
	Back & neck	Sense	Depression	Skin	Diabetes	Migraine	Iron	Oral	Osteoarth	IHD
Lithuania	(1.23)	(1.17)	(1.16)	(0.86)	(1.08)	(1.04)	(1.18)	(0.95)	(1.32)	(1.94)
	Back & neck	Sense	Depression	Skin	Migraine	Iron	Diabetes	Oral	Anxiety	Alcohol
Moldova	(1.2)	(0.96)	(1.0)	(0.95)	(1.02)	(1.24)	(0.67)	(0.94)	(0.7)	(3.29)
Russia	Back & neck	Sense	Depression	Iron	Skin	Migraine	Diabetes	Drugs	Oral	Alcohol
Nossia	(1.2)	(1.3)	(1.14)	(2.22)	(0.88)	(1.0)	(0.95)	(3.39)	(0.98)	(6.05)
Ukraine	Back & neck	Sense	Depression	Skin	Migraine	Diabetes	Oral	Osteoarth	Drugs	IHD (1.70)
	(1.26)	(1.17)	(1.29)	(0.89)	(1.03)	(0.86)	(0.92)	(1.24)	(3.26)	(1.76)
Central Europe	Back & neck (1·29)	Sense (1·1)	Depression (1.06)	Diabetes (1·14)	Skin (0⋅9)	Migraine (0.98)	Oral (1.01)	Anxiety (0.93)	Osteoarth (1·29)	lron (0⋅91)
	Back & neck	Sense	Depression	Skin	Iron	Migraine	Anxiety	Oral	Diabetes	Osteoarth
Albania	(1.37)	(0.94)	(0.98)	(0.83)	(1.25)	(0·98)	(0.88)	(0.98)	(0.51)	(0.91)
ъ. :	Back & neck	Sense	Depression	Diabetes	Skin	Migraine	War	Oral	Iron	Anxiety
Bosnia	(1.16)	(0.98)	(0.98)	(1.01)	(0.9)	(0.98)	(1887-24)	(1.06)	(1.01)	(0.88)
Bulgaria	Back & neck	Sense	Depression	Diabetes	Skin	Migraine	Oral	Anxiety	Osteoarth	Iron
Dolgana	(1.28)	(1.04)	(1.06)	(1.18)	(0.9)	(0.98)	(1.01)	(0.96)	(1.21)	(1.12)
Croatia	Back & neck	Sense	Depression	Skin	Diabetes	Migraine	Oral	Iron	Osteoarth	Anxiety
	(1·37) Back & neck	(0·97) Sense	(1·04)	(0.9)	(0.83)	(0·97)	(1·07) Oral	(1·1) Iron	(1·2)	(0.92) Osteoarth
Czech Republic	(1.38)	(1.09)	Depression (1·13)	Diabetes (1·46)	Skin (0.88)	Migraine (0.98)	(1.04)	(1.05)	Anxiety (0.96)	(1.43)
	Back & neck	Sense	Depression	Diabetes	Skin	Migraine	Oral	Anxiety	Osteoarth	Iron
Hungary	(1.38)	(1.05)	(1.07)	(1.62)	(0.96)	(0.98)	(0.94)	(0.95)	(1.35)	(0.85)
Macadonia	Back & neck	Sense	Depression	Diabetes	Skin	Migraine	Oral	Anxiety	Osteoarth	Iron
Macedonia	(1.26)	(0.99)	(1.01)	(1.14)	(0.89)	(0.98)	(0.97)	(0.89)	(1.21)	(0.58)
Montenegro	Back & neck	Sense	Depression	Diabetes	Skin	Migraine	Iron	Oral	Anxiety	Osteoarth
	(1.27)	(1.04)	(1.04)	(1.15)	(0.91)	(0.98)	(1.12)	(0.98)	(0.91)	(1.26)
Poland	Back & neck	Sense	Depression (1.07)	Diabetes	Skin	Migraine	Oral	Anxiety	Osteoarth (1.24)	IHD
	(1·21) Back & neck	(1·27) Sense	(1.07) Depression	(1·25) Skin	(0.88) Migraine	(0.97)	(1·03) Oral	(0·95) Iron	(1·34) Osteoarth	(2·09) Anxiety
Romania	(1·33)	(1.03)	(1·04)	(0·9)	(0.98)	Diabetes (0.75)	(1.04)	(1.18)	(1·28)	(0.89)
c 1:	Back & neck	Sense	Depression	Diabetes	Skin	Migraine	Oral	Anxiety	Osteoarth	Iron
Serbia	(1.31)	(1.0)	(1.02)	(1.22)	(0.92)	(0.98)	(0.98)	(0.91)	(1.23)	(0.8)
Slovakia	Back & neck	Sense	Depression	Skin	Diabetes	Migraine	Iron	Anxiety	Oral	Osteoarth
SIOVANIA	(1.34)	(1.06)	(1.08)	(0.89)	(1.25)	(0.97)	(1.09)	(0.94)	(0.85)	(1.35)
Slovenia	Back & neck	Sense	Depression	Skin	Diabetes	Migraine	Oral	Iron	Osteoarth	Anxiety
**	(1.35)	(0.99)	(1.09)	(0.9)	(1.07)	(0.98)	(1.16)	(1.18)	(1.35)	(0.95)
Central Asia	Back & neck	Sense	Depression	Iron	Skin	Migraine	Diabetes	Anxiety	Oral	Asthma
	(1·01)	(0·98)	(0·97)	(1.36)	(0.87)	(0.98)	(0·98)	(0·8)	(1.06)	(0.69)
Armenia	Back & neck (1.0)	Sense	Depression (1.01)	Diabetes (1.4)	Skin	Migraine	Iron (1·11)	Oral (1.06)	Anxiety (0.81)	Disaster (138-88)
	Back & neck	(1·03) Sense	(1.01) Depression	(1·4) Iron	(0·87) Skin	(0.97)	(1·11) Migraine	Anxiety	(0.81) Oral	Osteoarth
Azerbaijan	(0.98)	(1.08)	(1·02)	(1·55)	(0.88)	Diabetes (1·36)	(0·96)	(0.82)	(1·03)	(1.26)
	Back & neck	Sense	Depression	Diabetes	Skin	Migraine	Iron	Oral	Anxiety	Osteoarth
Georgia	(0·93)	(1.05)	(1.01)	(1.19)	(0.85)	(0·98)	(1.25)	(1.02)	(0.82)	(1.21)

	1	2	3	4	5	6	7	8	9	10
Kazakhstan	Back & neck	Iron	Sense	Depression	Skin	Migraine	Diabetes	Oral	Anxiety	Osteoarth
	(1·04) Back & neck	(2·4) Iron	(1·1) Depression	(1·03) Sense	(0.9) Skin	(0.96) Migraine	(1·17) Anxiety	(1·14) Diabetes	(0·82) Oral	(1·31) Asthma
Kyrgyzstan	(1.04)	(1.3)	(0·91)	(0.85)	(0.86)	(1·0)	(0·76)	(0·52)	(1·05)	(0.69)
Mongolia	Back & neck	Depression	Sense	Skin	Iron	Migraine	Diabetes	Anxiety	Oral	Falls
iviorigolia	(0.98)	(0.94)	(0.94)	(0.87)	(0.97)	(0.98)	(0.79)	(0.78)	(1.02)	(1.37)
Tajikistan	Back & neck (1.06)	Iron (1·03)	Depression (0.87)	Sense (0.8)	Skin (0·85)	Migraine (1·0)	Anxiety	Diabetes (0.66)	Oral (1:06)	Epilepsy (1·54)
	Back & neck	Depression	Sense	Iron	Skin	Migraine	(0.78) Diabetes	Anxiety	Oral	Asthma
Turkmenistan	(0.94)	(1.0)	(1.06)	(1.31)	(0.88)	(0.96)	(1.33)	(0.82)	(1.02)	(0.65)
Uzbekistan	Back & neck	Depression	Sense	Iron	Skin	Migraine	Diabetes	Anxiety	Oral	Asthma
	(1.0) Back & neck	(0.94) Sense	(0.93) Depression	(1·08) Skin	(0.87) Anxiety	(0.98) Iron	(0.84) Diabetes	(0.78)	(1·03) Asthma	(0·79) Other MSK
Latin America and Caribbean	(0·87)	(1.06)	(1.08)	(0.99)	(1.52)	(1.02)	(1.12)	Migraine (1·0)	(1·18)	(1.01)
Central Latin America	Back & neck	Sense	Depression	Diabetes	Skin	Iron	Migraine	Anxiety	Other MSK	Oral
Central Lacin America	(0.86)	(1.12)	(0.94)	(1.44)	(0.95)	(0.93)	(0.9)	(1.01)	(0.89)	(1.08)
Colombia	Back & neck (0.94)	Sense (1·14)	Depression (1.02)	Skin (1.02)	Anxiety	Migraine	Diabetes (0.81)	Asthma	Other MSK	Oral (1·07)
c	Back & neck	Sense	Depression	Skin	(1·35) Iron	(0.89) Migraine	Anxiety	(1·23) Diabetes	(0.93) Asthma	Other MSK
Costa Rica	(0.89)	(1.16)	(1.01)	(0.91)	(1.01)	(0.88)	(1.08)	(0.72)	(1.18)	(0.98)
El Salvador	Back & neck	Sense	Iron	Depression	Skin	Diabetes	Migraine	Anxiety	Asthma	Oral
	(0.94) Back & neck	(0.96)	(1.59)	(0.9) Skin	(0.94)	(0.94)	(0.91) Migraine	(1.03)	(1·24)	(1·07) Oral
Guatemala	(0.96)	Iron (1.06)	Sense (0·94)	Skin (0.96)	Depression (0.85)	Diabetes (1·67)	Migraine (0·9)	Anxiety (1.05)	Asthma (1·02)	(1·18)
Honduras	Back & neck	Sense	Skin	Depression	Iron	Asthma	Migraine	Anxiety	Diabetes	Oral
ionaulas	(0.98)	(0.95)	(1.06)	(0.88)	(0.79)	(1.54)	(0.91)	(1.05)	(0.89)	(1.07)
Mexico	Back & neck	Sense	Diabetes	Depression	Skin	Migraine	Anxiety	Other MSK	Oral	lron
	(0·79) Back & neck	(1·17) Sense	(1.9) Depression	(0·91) Skin	(0·91) Iron	(0·91) Migraine	(0.85) Anxiety	(0.94) Diabetes	(1·08) Asthma	(0·5) Oral
Nicaragua	(0·96)	(0.95)	(0.87)	(0.96)	(0.82)	(0.91)	(1.05)	(0.85)	(0·96)	(1.08)
Panama	Back & neck	Sense	Depression	Skin	Diabetes	Iron	Migraine	Asthma	Anxiety	Oral
	(0.91)	(1.19)	(0.99)	(0.95)	(1.35)	(1.2)	(0.88)	(1.37)	(1.09)	(1.03)
Venezuela	Iron (3.06)	Back & neck	Sense (1·12)	Depression (0.96)	Skin	Diabetes	Migraine (0·86)	Anxiety	Asthma	Other MSK (0.9)
A d I t A	Back & neck	(0.9) Sense	Depression	Iron	(0.98) Skin	(1·17) Anxiety	Migraine	(1.06) Diabetes	(1·01) Asthma	Oral
Andean Latin America	(0.93)	(1.07)	(1.09)	(1.34)	(1.02)	(1.37)	(1.0)	(0.76)	(1.1)	(1.09)
Bolivia	Back & neck	Iron	Sense	Depression	Skin	Anxiety	Migraine	Diabetes	Asthma	Oral
	(1.0) Back & neck	(1·52)	(0.97)	(1·03) Iron	(1·02)	(1.33)	(0.99)	(0.73)	(1.11)	(1·12)
Ecuador	(0·91)	Sense (1-02)	Depression (1.08)	(1.33)	Skin (0·99)	Diabetes (1·15)	Anxiety (1·37)	Migraine (0.87)	Oral (1·09)	Asthma (0.86)
Peru	Back & neck	Sense	Depression	Skin	Iron	Migraine	Anxiety	Asthma	Oral	Diabetes
	(0.92)	(1.14)	(1.11)	(1.03)	(1.29)	(1.07)	(1.38)	(1.22)	(1.09)	(0.57)
Caribbean	Back & neck	Sense	Depression	Iron	Skin	Diabetes	Anxiety	Migraine	Asthma	Other MSK
	(0.89) Back & neck	(1.06) Depression	(1·11) Sense	(1·2) Diabetes	(1·09) Skin	(1·24) Iron	(1·49) Anxiety	(1·0) Migraine	(1.62) Other MSK	(1.05) Asthma
Antigua	(0·82)	(1.2)	(1.21)	(2.2)	(1.05)	(1.43)	(1.49)	(0·94)	(1·37)	(1.31)
Bahamas	Back & neck	Sense	Depression	Diabetes	Skin	lron	Anxiety	Migraine	Other MSK	Asthma
	(0.82)	(1.19)	(1.18)	(1.94)	(1.05)	(1.54)	(1.49)	(0.94)	(1.47)	(1.38)
Barbados	Back & neck (0.81)	Sense (1·07)	Diabetes (2·05)	Depression (1·15)	Skin (0.96)	Other MSK (1·57)	lron (1·49)	Anxiety (1·46)	Migraine (0·94)	Asthma (1·26)
Belize	Back & neck	Depression	Iron	Skin	Sense	Anxiety	Diabetes	Migraine	Asthma	Other MSK
Delize	(0.84)	(1.04)	(1.19)	(1.05)	(0.92)	(1.35)	(1.14)	(0.97)	(1.58)	(1.13)
Bermuda	Back & neck	Depression	Sense	Skin	Iron	Anxiety	Diabetes	Migraine	Other MSK	Asthma
- 1	(0.89) Back & neck	(1·25) Sense	(1·19)	(1·02) Skin	(1·32) Diabetes	(1·53)	(1.87)	(0.96)	(1·41)	(1·46) Asthma
Cuba	(0·82)	(1.11)	Depression (1·14)	(1.08)	(0·91)	Anxiety (1·46)	lron (1⋅36)	Migraine (0.95)	Other MSK (1·11)	(1·31)
Dominica	Back & neck	Diabetes	Sense	Depression	Skin	Iron	Anxiety	Migraine	Asthma	Other MSK
	(0.83)	(2.12)	(1.07)	(1.12)	(1.05)	(1.29)	(1.41)	(0.95)	(1.52)	(1.25)
Dominican Republic	Back & neck	Sense	Depression	Skin	Iron	Anxiety	Migraine	Asthma	Diabetes	Other MSK
Grenada	(0.82) Back & neck	(0.98) Depression	(1·07) Sense	(1·05) Iron	(1·19) Skin	(1·37) Diabetes	(0.97) Other MSK	(1·5) Anxiety	(0·71) Migraine	(0·91) Asthma
Grenaua	(0.8)	(1.11)	(1.06)	(1.52)	(1.05)	(1.78)	(1·73)	(1·41)	(0·94)	(1.43)
Guyana	Back & neck	Iron	Depression	Sense	Diabetes	Skin	Anxiety	Migraine	Asthma	Other MSK
•	(0.84)	(1.41)	(1.02)	(0.91)	(1.36)	(1.04)	(1.33)	(0.97)	(1.58)	(0.85)
Haiti	Iron (1·01)	Back & neck (0.96)	Depression (0.89)	Disaster (1227-89)	Skin (1·02)	Sense (0.8)	Asthma	Anxiety (1.45)	Migraine (1.0)	Diabetes (1·38)
amaica	Back & neck	Sense	Depression	(122/-69) Diabetes	(1·02) Skin	(0⋅8) Iron	(1·97) Anxiety	(1·45) Migraine	(1.0) Other MSK	(1·30) Asthma
parriared	(0.83)	(1.0)	(1·1)	(1.43)	(1.05)	(1.25)	(1.38)	(0·95)	(1·39)	(1.48)
Puerto Rico	Back & neck	Sense	Diabetes	Depression	Skin	Anxiety	Iron	Other MSK	Migraine	Asthma
	(0.88)	(1.2)	(2.37)	(1.23)	(1.02)	(1.53)	(1.38)	(1.3)	(0.95)	(1·41)
Saint Lucia	Back & neck (0.83)	Sense (1.05)	Depression (1·11)	Diabetes (1.48)	Skin (1·05)	Iron (1⋅26)	Anxiety (1·41)	Migraine (0.95)	Asthma	Other MSK (1·25)
Saint Vincent and the Grenadines	Back & neck	Sense	Depression	Diabetes	Skin	(1·20) Iron	Anxiety	(0.95) Migraine	(1·52) Asthma	Other MSK
	(0.82)	(1.05)	(1.1)	(1.84)	(1.05)	(1.23)	(1.41)	(0.95)	(1.45)	(1.21)

	1	2	3	4	5	6	7	8	9	10
Suriname	Back & neck (0.82)	Sense (0.08)	Depression (1.07)	Skin (1.05)	Iron	Diabetes	Anxiety	Migraine	Asthma	Other MSK
	Back & neck	(0.98) Diabetes	(1·07) Sense	(1.05) Depression	(1·25) Skin	(1·21) Iron	(1.38) Anxiety	(0.96) Other MSK	(1·53) Migraine	(1·19) Asthma
Trinidad and Tobago	(0.81)	(3.11)	(1.17)	(1.18)	(1.11)	(1.44)	(1.5)	(1.49)	(0.94)	(1.58)
Virgin Islands, USA	Back & neck	Sense	Diabetes	Depression	Skin	Other MSK	Iron	Anxiety	Migraine	Asthma
	(0.89) Back & neck	(1·19) Depression	(2·14) Sense	(1·24) Anxiety	(1·02) Skin	(1·44) Migraine	(1·38) Iron	(1·54) Diabetes	(0.96) Other MSK	(1·46) Asthma
Tropical Latin America	(0·87)	(1.23)	(0.99)	(2·14)	(1.0)	(1.12)	(1.0)	(0.86)	(1·21)	(1.41)
Brazil	Back & neck	Depression	Sense	Anxiety	Skin	Migraine	Iron	Diabetes	Other MSK	Asthma
biazii	(0.86) Back & neck	(1.23)	(0.99)	(2.15)	(1.0)	(1.12)	(1.0)	(0.86)	(1.22)	(1.43)
Paraguay	(0.97)	Depression (1·24)	Sense (0·97)	Skin (1·14)	Anxiety (1.81)	Iron (1⋅06)	Migraine (1.06)	Diabetes (0.92)	Other MSK (0·9)	Asthma (1·01)
	Back & neck	Sense	Depression	Skin	Diabetes	Other MSK	Iron	Migraine	Schiz	Anxiety
Southeast Asia, east Asia, and Oceania	(0.86)	(0.94)	(0.74)	(1.0)	(0.76)	(1.22)	(0.8)	(0.65)	(1.31)	(0.72)
East Asia	Back & neck (0.89)	Sense (0·91)	Depression (0.72)	Skin	Diabetes (0.64)	Other MSK	Iron	Schiz (1⋅38)	Migraine	Anxiety (0.71)
	Back & neck	Sense	(0.73) Depression	(0·96) Skin	Diabetes	(1·1) Other MSK	(0·79) Iron	Schiz	(0·57) Migraine	Anxiety
China	(0.88)	(0.91)	(0.74)	(0.96)	(0.62)	(1.09)	(0.79)	(1.39)	(0.57)	(0.7)
North Korea	Back & neck	Sense	Skin	Iron	Depression	Other MSK	Diabetes	COPD	Anxiety	Migraine
	(1.05) Back & neck	(0.77)	(0.93)	(0.92)	(0.61)	(1·15) Skin	(0.59)	(2·1)	(0.83)	(0.59)
Taiwan (province of China)	(0.99)	Sense (1.08)	Diabetes (2·29)	Other MSK (1·59)	Depression (0.73)	(0.89)	Anxiety (1.03)	Iron (1.05)	Migraine (0·63)	Osteoarth (1·44)
Southeast Asia	Back & neck	Sense	Skin	Depression	Diabetes	Other MSK	Iron	Migraine	Anxiety	Asthma
Journey Chaid	(0.81)	(1.01)	(1.06)	(0.76)	(1.1)	(1.53)	(0.81)	(0.83)	(0.75)	(1.02)
Cambodia	Back & neck	Iron	Sense	Skin	Depression (0.67)	Migraine (0·82)	Other MSK	Asthma	Anxiety	Diabetes (0.82)
	(0.94) Back & neck	(0·94) Sense	(0·95) Skin	(1.05) Diabetes	(0.67) Depression	Other MSK	(1·11) Iron	(1·14) Migraine	(0·81) Anxiety	Asthma
Indonesia	(0.76)	(0.99)	(1.05)	(1.17)	(0.77)	(1.67)	(0.94)	(0.83)	(0.77)	(0.93)
Laos	Back & neck	Skin	Iron	Sense	Depression	Diabetes	Other MSK	Migraine	Asthma	Anxiety
	(0.9) Back & neck	(1·05) Sense	(0·7)	(0.84)	(0.68)	(1·36) Other MSK	(1·27) Nematode	(0.82)	(1.28)	(0.8) Schiz
Malaysia	(0·75)	(1.0)	Skin (1·05)	Depression (0.84)	Diabetes (1·51)	(1·82)	(4405·93)	Anxiety (1·2)	Migraine (0·79)	(1.18)
Maldives	Back & neck	Sense	Iron	Skin	Depression	Migraine	Other MSK	Anxiety	Haemog	Diabetes
Maidives	(0.82)	(0.98)	(1.35)	(1.08)	(0.75)	(0.84)	(1.17)	(0.76)	(3.61)	(0.56)
Mauritius	Back & neck (0.82)	Diabetes (2·04)	Sense	Depression (0.84)	Skin (1·07)	Other MSK	Migraine (0·82)	Anxiety (0.8)	Asthma (1·03)	Oral (0.84)
	Sense	Back & neck	(1·15) Skin	Depression	Diabetes	(1·77) Iron	Other MSK	Migraine	Asthma	Anxiety
Myanmar	(1.13)	(0.79)	(1.04)	(0.69)	(1.23)	(0.69)	(1.38)	(0.82)	(1.08)	(0.79)
Philippines	Back & neck	Sense	Skin	Diabetes	Iron	Depression	Other MSK	Migraine	Asthma	Anxiety
	(0.95) Back & neck	(1·05) Sense	(1·07) Diabetes	(1·49) Skin	(0.92) Depression	(0·75) Iron	(1.69) Migraine	(0.82) Asthma	(1·29) Anxiety	(0.77) Other MSK
Sri Lanka	(0.81)	(1.14)	(1·44)	(1.06)	(0.81)	(1.04)	(0.82)	(1.14)	(0.79)	(0.86)
Seychelles	Back & neck	Sense	Skin	Depression	Diabetes	Migraine	Other MSK	Anxiety	Asthma	Schiz
,	(0.79)	(1.18)	(1.07)	(0.84)	(1.34)	(0.82)	(1.15)	(0.81)	(0.91)	(1.15)
Thailand	Back & neck (0·73)	Sense (1·04)	Depression (0.82)	Skin (1.06)	Other MSK (1.63)	Diabetes (0.83)	Migraine (0.84)	lron (0⋅82)	Anxiety (0·79)	Asthma (1.08)
Timor-Leste	Back & neck	Sense	Iron	Skin	Depression	Migraine	Asthma	Other MSK	Diabetes	Anxiety
Timor-Leste	(0.87)	(0.97)	(0.66)	(1.07)	· (0·7)	(0.82)	(1.15)	(1.03)	(1.05)	(0.84)
Vietnam	Back & neck	Sense	Skin	Depression	Migraine	Other MSK	Diabetes	Schiz	Other nutr	Asthma
	(0.85) Back & neck	(0·92) Iron	(1·07) Skin	(0.77) Diabetes	(0.85) Sense	(1·18) Depression	(0.6) Asthma	(1·2) Other MSK	(72·36) Nematode	(0.86) Migraine
Oceania	(1.01)	(0.98)	(1.36)	(2.69)	(0.96)	(0·7)	(1.76)	(1·58)	(3.75)	(0·77)
American Samoa	Back & neck	Diabetes	Skin	Sense	Iron	Depression	Other MSK	Asthma	Migraine	Anxiety
	(0·9)	(2.78)	(1.34)	(1.03)	(0.91)	(0.77)	(1.87)	(1.31)	(0.72)	(0.83)
Micronesia	Back & neck (0.91)	Skin (1·31)	Diabetes (1.68)	Sense (0·91)	Depression (0.72)	Other MSK (1.65)	Asthma (1·45)	lron (0.66)	Migraine (0·73)	Anxiety (0.82)
Fiji	Diabetes	Back & neck	Skin	Sense	Iron	Depression	Asthma	Other MSK	Nematode	Migraine
	(3.07)	(0.84)	(1.27)	(1.03)	(1.1)	(0.75)	(1.64)	(1.45)	(58.08)	(0.72)
Guam	Back & neck	Diabetes	Sense	Skin	Other MSK	Depression	Iron	Asthma	Other NTD	Migraine
izi di le	(0.94) Back & neck	(3·17) Diabetes	(1·27) Nematode	(1·24) Skin	(1·96) Sense	(0.89) Iron	(0.97) Depression	(1·47) Asthma	(234964·29) Migraine	(0·72) Other MSK
Kiribati	(0.93)	(3.07)	(7.02)	(1.27)	(0.81)	(0.54)	(0.65)	(1.68)	(0.72)	(1.03)
Marshall	Diabetes	Back & neck	Skin	Iron	Sense	Nematode	Depression	Other MSK	Asthma	Migraine
	(3·31)	(0.88)	(1.29)	(1.23)	(0.87)	(7·21)	(0·69)	(1.7)	(1·41)	(0.72)
Northern Mariana Islands	Back & neck (0.86)	Skin (1·25)	Diabetes (3·3)	Depression (0.82)	Sense (1·28)	Other MSK (1.61)	Iron (1·08)	Migraine (0·71)	Anxiety (0.85)	Asthma (1·16)
Papua New Guinea	Iron	Back & neck	Skin	Sense	Diabetes	Depression	Asthma	Nematode	Other MSK	(1·10) LF
, apparter domen	(0.97)	(0.95)	(1.29)	(0.84)	(2.35)	(0.63)	(1.7)	(3.34)	(1.43)	(100-35)
Samoa	Back & neck	Skin	Sense	Diabetes	Iron	Depression	Other MSK	Asthma	Migraine	Anxiety
	(0.9) Back & neck	(1·33) Skin	(0.92)	(1·5)	(0·84) Iron	(0.74)	(1.6)	(1·16)	(0.73) Migraine	(0.82)
Solomon Islands	(0.95)	(1·25)	Diabetes (2·69)	Sense (0·81)	(0.53)	Depression (0.65)	Other MSK (1·56)	Asthma (1·45)	Migraine (0·73)	Anxiety (0.88)
Tonga	Back & neck	Skin	Diabetes	Sense	Iron	Depression	Other MSK	Asthma	Migraine	Osteoarth
-	(0.92)	(1.34)	(1.49)	(0.83)	(0.97)	(0.73)	(1.69)	(1.68)	(0.73)	(2.01)

	1	2	3	4	5	6	7	8	9	10
Vanuatu	Back & neck (0.91)	Skin (1·31)	Diabetes (1.79)	Sense (0⋅82)	Depression (0.67)	Other MSK (1·55)	Asthma (1·45)	Migraine (0·73)	Nematode (2·91)	Anxiety (0.83)
North Africa and Middle East	Back & neck	Depression	Sense	Diabetes	Iron	Skin	Migraine	Other MSK	Anxiety	Asthma
	(1·07) Back & neck	(0.93) Depression	(0.86) Sense	(1.62) Diabetes	(0·82) Iron	(0.84) Skin	(1·07)	(1·34) Other MSK	(1.06)	(0.94) Asthma
North Africa and Middle East	(1·07)	(0.93)	(0.86)	(1.62)	(0.82)	(0.84)	Migraine (1·07)	(1·34)	Anxiety (1.06)	(0·94)
Afghanistan	Iron	Back & neck	War	Skin	Depression	Sense	Diabetes	Other unint	Migraine	Anxiety
7 tigrium stari	(0.8) Back & neck	(1·15) Diabetes	(163·45) Depression	(0·84) Sense	(0.8) Skin	(0·73) Migraine	(2·72) Other MSK	(5·28) Iron	(1·18) Anxiety	(1·23) Asthma
Algeria	(1.04)	(1.47)	(0.95)	(0.82)	(0.83)	(1·08)	(1·52)	(0.7)	(1.04)	(1.0)
Bahrain	Back & neck	Diabetes	Depression	Sense	Other MSK	Skin	Migraine	Iron	Anxiety	Drugs
	(0.95) Back & neck	(2·9) Iron	(1·11) Diabetes	(1·02) Sense	(1.96) Depression	(0.9) Skin	(1·07) Migraine	(1·43) Other MSK	(1.07) Anxiety	(1.78) Asthma
Egypt	(1.08)	(1.53)	(1.5)	(0.93)	(0.77)	(0.84)	(1.07)	(1.19)	(1.0)	(0.96)
Iran	Back & neck (1·03)	Depression (1.05)	Sense (0·94)	Diabetes (1.48)	Migraine (0.98)	Skin (0·8)	Other MSK (1·44)	Anxiety	Drugs (3·41)	Other cardi
	Back & neck	Diabetes	Depression	(1.40) War	Sense	Skin	lron	(1·07) Migraine	Anxiety	Other MSK
Iraq	(1.08)	(2.19)	(0.93)	(395.43)	(0.79)	(0.75)	(0.65)	(1.06)	(1.14)	(0.92)
Jordan	Back & neck (1.0)	Depression (1.01)	Diabetes (1·99)	Skin (0.92)	Iron (0⋅97)	Sense (0·92)	Migraine (1·05)	Other MSK (1.69)	Anxiety (1.07)	Asthma (1.03)
Kuwait	Back & neck	Depression	Migraine	Skin	Diabetes	Sense	Iron	Anxiety	Other MSK	Asthma
KUWait	(0.88)	(1.18)	(1.06)	(0.89)	(2.46)	(1.09)	(1.18)	(1.12)	(1.32)	(1.01)
Lebanon	War (10689-82)	Back & neck (0.83)	Diabetes (2·0)	Depression (1.02)	Sense (0·9)	Skin (0.93)	Iron (1·35)	Migraine (1·03)	Anxiety (1·27)	Asthma (1·14)
Libus	Back & neck	Depression	Diabetes	Sense	Skin	Migraine	War	Iron	Anxiety	Other MSk
Libya	(1.04)	(1.0)	(1.3)	(0.81)	(0.87)	(1.07)	(464-77)	(0.72)	(1.04)	(1.19)
Morocco	Back & neck (1·14)	Diabetes (2·21)	Sense (0·79)	Depression (0.88)	Skin (0⋅92)	Migraine (1·05)	lron (0⋅56)	Other MSK (1·36)	Anxiety (1.09)	Drugs (4·01)
D.L.	Back & neck	Depression	Skin	Sense	Iron	Migraine	Anxiety	Diabetes	War	Other MSk
Palestine	(1.06)	(1.29)	(0.83)	(0.72)	(0.59)	(1.07)	(1.07)	(1.14)	(198.72)	(1.08)
Oman	Back & neck (0.95)	Depression (1.07)	Diabetes (2·42)	Other Cardio (19·2)	Sense (1·01)	Migraine (1·1)	Skin (0⋅85)	lron (1⋅15)	Other MSK (1·43)	Anxiety (1.05)
0.1	Back & neck	Depression	Diabetes	Migraine	Skin	Sense	Anxiety	Other MSK	Iron	Heat & cold
Qatar	(0.94)	(1.15)	(3.12)	(1.12)	(0.94)	(0.96)	(1.05)	(1.26)	(1.14)	(17-26)
Saudi Arabia	Back & neck (0.99)	Depression (1·11)	Migraine (1·27)	Skin (0.93)	Sense (1.01)	Diabetes (1·51)	Anxiety (1.08)	Other MSK (1·33)	lron (0⋅52)	Asthma (0.8)
	Back & neck	Iron	Depression	Skin	Sense	Migraine	Diabetes	Anxiety	Asthma	Other MSk
Sudan	(1.07)	(0.81)	(0.84)	(0.84)	(0.75)	(1.08)	(1.9)	(1.15)	(1.18)	(1.16)
Syria	War (1612·15)	Back & neck (1·05)	Depression (0.94)	Sense (0.78)	Skin (0.8)	Iron	Migraine (1·07)	Diabetes (1.07)	Anxiety (1.06)	Asthma
	Back & neck	Diabetes	Depression	Sense	Skin	(0.73) Other MSK	Migraine	Anxiety	Iron	(0.99) Asthma
Tunisia	(1.02)	(1.35)	(1.0)	(0.82)	(0.89)	(1.58)	(1.02)	(1.04)	(0.66)	(1.05)
Turkey	Back & neck (1·13)	Sense (0·85)	Depression (0.92)	Diabetes	Other MSK (1.67)	Skin (0·83)	Migraine (1·01)	lron (0.03)	Anxiety	Oral
	Back & neck	Depression	Diabetes	(1·3) Other MSK	Migraine	Sense	Skin	(0.93) Drugs	(0.92) Anxiety	(1·18) Iron
United Arab Emirates	(0.99)	(1.18)	(3.52)	(1.92)	(1.14)	(1.15)	(0.91)	(2.16)	(1.04)	(1.21)
Yemen	Back & neck	War	Depression	Skin	Sense	Iron	Migraine	Diabetes	Anxiety	Other MSk
	(1·11) Iron	(248·22) Back & neck	(0.84) Sense	(0.84) Depression	(0.77) Other MSK	(0·44) Skin	(1·11) Migraine	(1.76) Diabetes	(1·17) Anxiety	(1·34) COPD
South Asia	(1.37)	(0.91)	(1.11)	(0.92)	(1.96)	(0.89)	(1.19)	(0.9)	(0.8)	(1.77)
South Asia	Iron	Back & neck	Sense	Depression	Other MSK	Skin	Migraine	Diabetes	Anxiety	COPD
	(1·37) Back & neck	(0.91) Other MSK	(1·11) Sense	(0.92) Depression	(1·96) Iron	(0·89) Skin	(1·19) Migraine	(0.9) Anxiety	(0.8) Diabetes	(1.77) Epilepsy
Bangladesh	(1.06)	(2.7)	(0.95)	(0.82)	(0.67)	(0.94)	(1.2)	(1.11)	(1.0)	(2.97)
Bhutan	Back & neck	Iron	Sense	Depression	Skin	Other MSK	Migraine	Diabetes	Diarrhoea	Anxiety
	(1·05) Iron	(1·48) Back & neck	(0.98) Sense	(0.83) Depression	(0.95) Other MSK	(1·84) Migraine	(1·19) Skin	(1·17) Diabetes	(3·33) Anxiety	(0.88) COPD
India	(1.58)	(0.89)	(1.16)	(0.94)	(1.93)	(1.2)	(0.86)	(0.83)	(0.74)	(2.01)
Nepal	Back & neck	Sense	Skin	Migraine	Other MSK	Iron	Depression	Disaster	Anxiety	Asthma
n I	(1·13) Iron	(0.89) Back & neck	(0.97) Sense	(1·33) Depression	(1·93) Skin	(0·48) Migraine	(0.63) Other MSK	(661.96) Diabetes	(0·94) Anxiety	(1.07) Asthma
Pakistan	(1.02)	(0.87)	(0.93)	(0.9)	(1.0)	(1.13)	(1.56)	(1.47)	(0.92)	(0.91)
Sub-Saharan Africa	Iron	Back & neck	Depression	Sense	Skin	Other NTD	Migraine	HIV	Asthma	Malaria
control of the second	(0·79) HIV	(0.98) Back & neck	(1-04) Depression	(1·02) Sense	(0·93) Skin	(1·24) Iron	(0.8) Diabetes	(4·74) Migraine	(0.98) Asthma	(2·25) Anxiety
Southern sub-Saharan Africa	(111.59)	(1.03)	(1.05)	(1.06)	(0.95)	(0.96)	(1.54)	(0.78)	(1.29)	(0.79)
Botswana	HIV	Back & neck	Iron	Depression	Sense	Skin	Diabetes	Asthma	Migraine	TB
	(140·61) HIV	(0.98) Back & neck	(1.51) Depression	(1·09) Sense	(0.98) Skin	(0.95) Iron	(1.33) Diabetes	(1·49) Asthma	(0·77) Migraine	(15.89) Anxiety
Lesotho	(73.48)	(1.08)	(1·14)	(0.91)	(0.93)	(0.64)	(1.63)	(1.68)	(0.77)	(0.78)
Namibia	HIV	Back & neck	Iron	Depression	Sense	Skin	Diabetes	Asthma	Migraine	Anxiety
	(77·97) HIV	(0.99) Back & neck	(1·28) Sense	(1·09) Depression	(0.95) Diabetes	(0·96) Skin	(1·07) Iron	(1·42) Migraine	(0.78) Asthma	(0.76) Anxiety
South Africa	(165.51)	(1·04)	(1·12)	(1·04)	(1·65)	(0·95)	(1.01)	(0.78)	(1·27)	(0.81)

	1	2	3	4	5	6	7	8	9	10
Swaziland	HIV	Back & neck	Depression	Skin	Sense	lron	Diabetes	Asthma (1·95)	Nematode (5·73)	Migraine
	(172·35) HIV	(0·94) Back & neck	(1·09) Iron	(0.95) Depression	(0·97) Skin	(0.92) Sense	(1·51) Asthma	Migraine	Diabetes	(0.77) Anxiety
Zimbabwe	(46.19)	(1.04)	(0.82)	(1.04)	(0.94)	(0.88)	(1.18)	(0.78)	(1.12)	(0.75)
Western sub-Saharan Africa	Iron	Back & neck	Depression	Sense	Skin	Malaria	Migraine	Schisto	Anxiety	Asthma
	(0.99) Iron	(1·11) Back & neck	(0.99) Depression	(0.98) Sense	(0.85) Skin	(3·39) Malaria	(0·79) Migraine	(10.95) Anxiety	(0·82) Asthma	(0.82) Schisto
Benin	(0.76)	(1·08)	(0.94)	(0.91)	(0.91)	(5.86)	(0.85)	(0.83)	(0.81)	(43.81)
Burkina Faso	Iron	Back & neck	Depression	Skin	Sense	Malaria	Migraine	Haemog	Anxiety	Diarrhoea
55141141435	(0.73) Back & neck	(1.08)	(0.91)	(0.92)	(0.87)	(1.56)	(0.9)	(1.72)	(0.9)	(0.81)
Cameroon	(1·05)	Depression (0.99)	Iron (0⋅57)	Skin (0·92)	Sense (0·93)	Oncho (654332·73)	HIV (9·44)	Malaria (45·65)	Migraine (0·77)	Anxiety (0.78)
Cara Manda	Back & neck	Depression	Iron	Sense	Skin	Migraine	Diabetes	Other cardio	Anxiety	Asthma
Cape Verde	(1.01)	(1.05)	(1.08)	(0.98)	(0.93)	(0.76)	(0.83)	(9.52)	(0.73)	(0.76)
Chad	lron	Back & neck	Skin	Depression	Sense	Migraine	Diarrhoea	Asthma	Anxiety	Heat & cold
	(1·31) Iron	(1·18) Back & neck	(0·91) Skin	(0·92) Sense	(0.92) Depression	(0.86) Malaria	(1·0) Migraine	(0.85) HIV	(0·87) Asthma	(4·25) Anxiety
Côte d'Ivoire	(0.75)	(1·06)	(1.08)	(0.97)	(0.94)	(15.32)	(0·8)	(4.5)	(0.91)	(0.81)
The Gambia	Iron	Back & neck	Depression	Skin	Sense	Malaria	Migraine	Anxiety	Diarrhoea	Asthma
The Gambia	(0.73)	(1.08)	(1.01)	(0.91)	(0.84)	(3.43)	(0.84)	(0.85)	(0.94)	(0.74)
Ghana	Back & neck	Iron (0.87)	Depression (1.0)	Sense (0.00)	Skin (0.60)	Malaria (150·78)	Migraine	Schisto (6627·12)	Anxiety	Diabetes (0.66)
	(0.92) Back & neck	(0·87) Iron	(1·0) Skin	(0.99) Sense	(0.69) Depression	(150-76) Malaria	(0.76) Schisto	(6627-12) Migraine	(0·75) Asthma	Anxiety
Guinea	(1.1)	(0.68)	(1.06)	(0.93)	(0.92)	(2.14)	(13.79)	(0.87)	(0.95)	(0.86)
Guinea-Bissau	Iron	Back & neck	Depression	Sense	Skin	Migraine	Malaria	HIV	Schisto	Asthma
	(0.77)	(1.11)	(0.92)	(0.92)	(0.91)	(0.85)	(1.77)	(2.78)	(15.56)	(0.9)
Liberia	Oncho (92161·85)	Back & neck (1·1)	Iron (0⋅73)	Sense (0·91)	Skin (0·91)	Depression (0.82)	Schisto (18·67)	Malaria (1·85)	Migraine (0·85)	Anxiety (0.85)
AA 1:	Iron	Back & neck	Sense	Depression	Skin	Malaria	Migraine	Anxiety	Heat & cold	Diarrhoea
Mali	(1.11)	(0.93)	(0.94)	(0.92)	(0.83)	(1.11)	(0.9)	(0.9)	(3.29)	(0.66)
Mauritania	Iron	Back & neck	Sense	Depression	Skin	Migraine	Asthma	Anxiety	Heat & cold	Schisto
	(1·24) Iron	(1·07) Back & neck	(1·06) Sense	(0.96) Skin	(0.91) Depression	(0.79) Migraine	(0.94) Heat & cold	(0∙8) Malaria	(5·99) Diarrhoea	(237·73) Anxiety
Niger	(0.78)	(1·16)	(0.94)	(0.94)	(0·92)	(0.98)	(3·9)	(0·32)	(0·77)	(0.97)
Nigeria	Iron	Back & neck	Depression	Sense	Skin	Schisto	Malaria	Migraine	Anxiety	Asthma
Nigeria	(1.18)	(1.18)	(1.05)	(1.03)	(0.78)	(2926-68)	(58-71)	(0.73)	(0.8)	(0.86)
São Tomé and Príncipe	Back & neck (1.04)	Iron (0⋅65)	Depression (0.99)	Sense (0·97)	Skin (0⋅92)	Other NTD (2·54)	Migraine (0·78)	Malaria (24·3)	Asthma (0.88)	Anxiety (0.79)
6	Iron	Back & neck	Depression	Sense	Skin	Migraine	Anxiety	Diarrhoea	Asthma	Diabetes
Senegal	(1.04)	(0.98)	(0.95)	(0.93)	(0.91)	(0.83)	(0.84)	(0.96)	(0.72)	(1.01)
Sierra Leone	Back & neck	Iron	Depression	Skin	Sense	Malaria	Migraine	Oncho	Asthma	Anxiety
	(1.08) Back & neck	(0.59)	(0.93)	(0.91)	(0.87)	(5.16)	(0.84)	(46362-69)	(0.86)	(0.84)
Togo	(1·1)	lron (0·71)	Depression (0.95)	Skin (0·97)	Sense (0·94)	Malaria (8∙85)	Migraine (0·82)	Asthma (0·93)	Anxiety (0.82)	Diarrhoea (1·04)
Eastern sub-Saharan Africa	Depression	Iron	Sense	Skin	Back & neck	Migraine	Other NTD	Anxiety	HIV	Asthma
Eastern Sub-Sanaran Africa	(1.11)	(0.62)	(0.99)	(0.95)	(0.78)	(0.79)	(0.88)	(0.94)	(3.82)	(0.94)
Burundi	Depression	Back & neck	Skin	Sense	Iron	Migraine	Asthma	Diarrhoea	Anxiety	Malaria
	(1·05) Sense	(0.85) Depression	(0.92) Back & neck	(0·9) Iron	(0·26) Skin	(0.88) Malaria	(1·0) Migraine	(0.99) Asthma	(1·0) Anxiety	(0.79) Diabetes
Comoros	(1.09)	(1.06)	(0·86)	(0.6)	(0.91)	(7.6)	(0·8)	(1.06)	(0.92)	(1.11)
Djibouti	Iron	Depression	Sense	Back & neck	Skin	Other NTD	Heat & cold	Migraine	Diabetes	Anxiety
Djibooti	(0.88)	(1.12)	(1.02)	(0.73)	(0.91)	(2.76)	(9.43)	(0.75)	(1.16)	(0.87)
Eritrea	(0.89)	Depression	Sense	Skin	Back & neck	(102.00)	Schisto	Migraine (0·81)	Asthma	Anxiety
E-1	Depression	(1·06) Sense	(1·0) Skin	(0.91) Back & neck	(0⋅7) Iron	(102·99) Schisto	(42·57) Anxiety	(0.01) Migraine	(0.97) Other NTD	(0.94) Asthma
Ethiopia	(1.13)	(1.0)	(1.02)	(0.8)	(0.32)	(26.64)	(1.0)	(0.76)	(0.66)	(0.79)
Kenya	Iron	Other NTD	Depression	Sense	Skin	Back & neck	HIV	Migraine	Anxiety	Schisto
•	(1·05)	(4·38)	(1·13)	(1·09)	(1·01)	(0.81)	(9.68)	(0.8)	(0.87)	(17-62)
Madagascar	lron (0⋅7)	Sense (1·14)	Depression (1.09)	Back & neck (0.83)	Skin (0·9)	Asthma (1·28)	Schisto (145·77)	Migraine (0·79)	Anxiety (0.92)	Diarrhoea (1·07)
Malawi	Depression	Iron	Back & neck	Sense	Skin	HIV	Malaria	Migraine	Anxiety	Asthma
IVICICITY I	(1.05)	(0.53)	(0.85)	(0.93)	(0.91)	(7-57)	(2.62)	(0.83)	(0.96)	(0.9)
Mozambique	Iron	Depression	Sense	HIV	Back & neck	Skin	Malaria	Migraine	Asthma	Anxiety
	(0.71) Back & neck	(1·04) Sense	(0.99) Depression	(7·43) Iron	(0.83) War	(0⋅89) Skin	(1·88) Asthma	(0·84) Migraine	(0.96) Anxiety	(0.97) Other NTD
Rwanda	(0.8)	(0.94)	(0.88)	(0·5)	(198.09)	(0·77)	(1·25)	(0.79)	(0·92)	(0.82)
Somalia	Iron	Depression	Sense	Skin	Back & neck	Schisto	lodine	Asthma	Migraine	Other NTD
Somula	(1.23)	(1.03)	(0.99)	(0.93)	(0.81)	(1.65)	(3.5)	(1.03)	(0.93)	(0.84)
South Sudan	lron	Depression	Sense	Oncho	Back & neck	Skin	Schisto	Heat & cold	Asthma	Migraine
	(0.81) Iron	(1.05) Depression	(1·02) Skin	(45 415·75) Sense	(0·81) Back & neck	(0.91) Migraine	(11·22) Asthma	(5·89) Anxiety	(1·1) HIV	(0.84) Malaria
Tanzania	(0·85)	(1.08)	(0.93)	(0.86)	(0.65)	(0.72)	(0·97)	(0·9)	(5·38)	(12·32)
Uganda	Depression	Skin	Sense	Iron	Back & neck	Malaria	HIV	Asthma	Migraine	Anxiety
- J	(1.31)	(0.91)	(0.96)	(0.45)	(0.79)	(9.75)	(6.86)	(1.01)	(0.79)	(0.89)

	1	2	3	4	5	6	7	8	9	10
Zambia	Depression	HIV	Skin	Sense	Iron	Back & neck	Migraine	Schisto	Asthma	Anxiety
Zambia	(1.11)	(20.66)	(0.91)	(0.96)	(0.52)	(0.77)	(0.88)	(1984-48)	(0.93)	(0.88)
Central sub-Saharan Africa	Other NTD	Iron	Back & neck	Sense	Skin	Depression	Oncho	Asthma	Malaria	Migraine
Central 300-3anarah Allika	(4.36)	(0.7)	(1.1)	(1.22)	(1.11)	(0.97)	(31767-28)	(1.44)	(1.43)	(0.87)
Angola	Other NTD	Skin	Sense	Back & neck	Depression	Iron	Asthma	Schisto	Malaria	Migraine
Aligola	(4.05)	(1.13)	(1.25)	(1.0)	(1.0)	(0.38)	(1.59)	(567-16)	(15.61)	(0.8)
Central African Republic	Iron	Back & neck	Sense	Skin	Depression	Oncho	Asthma	Other NTD	HIV	Migraine
Central African Republic	(0.84)	(1.12)	(1.17)	(1.1)	(0.95)	(37527-86)	(1.46)	(0.95)	(3.4)	(0.87)
Congo (Brazzaville)	Iron	Sense	Back & neck	Skin	Other NTD	Depression	Asthma	Diabetes	Schisto	Migraine
Congo (Brazzaville)	(1.03)	(1.27)	(1.0)	(1.16)	(5.85)	(1.04)	(1.25)	(1.2)	(10909-69)	(0.77)
DR Congo	Other NTD	Iron	Back & neck	Sense	Skin	Oncho	Depression	Asthma	Malaria	Migraine
DK Congo	(4.56)	(0.76)	(1.15)	(1.21)	(1.1)	(31562-22)	(0.96)	(1.4)	(1.11)	(0.9)
Eguatorial Guinea	Back & neck	Sense	Skin	Depression	Diabetes	Iron	Asthma	Malaria	Migraine	HIV
Equatorial dolliea	(0.95)	(1.23)	(1.14)	(1.11)	(1.22)	(0.75)	(1.72)	(6683-41)	(0.78)	(21.87)
Gabon	Other NTD	Iron	Sense	Back & neck	Depression	Skin	Schisto	Diabetes	Asthma	Migraine
Gabon	(24.82)	(1.58)	(1.31)	(0.94)	(1.13)	(1.1)	(55308.55)	(1.14)	(1.62)	(0.77)
	Colour key									
	0.0-0.81	0.81-0.88	0.88-0.94	0.94-0.99	O-99-1-05					
	1.05-1.13	1.13-1.26	1.26-1.58	■ ≥1.58						

Figure 7: Leading ten causes of years lived with disability (YLDs) with the ratio of observed years lived with disability (YLDs) to years lived with disability (YLDs) expected on the basis of SDI in 2015, by location

Shades of blue represent much lower observed YLDs than expected levels based on SDI, whereas red shows that observed YLDs exceed expected levels. Alzheimer's=Alzheimer's disease and other dementias. Back & neck=low back and neck pain. COPD=chronic obstructive pulmonary disease. Drugs=drug use disorders. Haemog=haemoglobinopathies and haemolytic anaemias. Heat & cold=environmental heat and cold exposure. IHD=ischaemic heart disease. Iron=iron-deficiency anaemia. NTD=neglected tropical diseases. Oncho=onchocerciasis. Oral=oral disorders. Osteoarth=osteoarthritis. Other cardio=other cardiovascular and circulatory diseases. Other MSK=other musculoskeletal disorders. Other nutr=other nutritional deficiencies. Other unintentional injuries. PEM=protein-energy malnutrition. Schitso=schistosomiasis. Schiz=schizophrenia. Sense=sense organ diseases. Skin=skin and subcutaneous diseases. TB=tuberculosis.

malaria prevalence and incidence become available for all malaria-endemic countries outside of Africa.

Tuberculosis

We added new data from tuberculosis prevalence surveys done in Indonesia, Ghana, and several subnational locations in India. Our analysis of tuberculosis relied on prevalence surveys, case notifications and cause-specific mortality estimates. We used the expert judgment values for the case-detection rates (CDRs) from WHO as an initial guide of how much notifications need to be increased to reflect incidence of all tuberculosis, but relied on DisMod-MR 2.1 to find an estimate that is consistent with available prevalence and cause-specific mortality rates. We found that, particularly in older age groups, our estimated CDR often falls below the all-age CDR of WHO. In GBD 2013, we used a relative risk approach to predict the proportion of HIV-infected individuals with tuberculosis. In GBD 2015, we improved our modelling strategy by making use of more abundantly available data on the proportions of HIV infected cases among all tuberculosis cases from the WHO case notifications to separate out combined HIV and tuberculosis from all forms of tuberculosis. In our modelling of tuberculosis, we have not separately estimated the incidence and prevalence related to multidrug-resistant tuberculosis. Given the policy interest in multidrug-resistant tuberculosis, we plan to include estimates for multidrug-resistant tuberculosis in future rounds of the GBD. Our global tuberculosis (allforms) incidence estimate ($10 \cdot 2$ million [$9 \cdot 2 - 11 \cdot 5$ million] cases in 2015) is slightly higher than that of WHO (9.6 million cases in 2014), and we estimate a slightly larger fraction of combined HIV and tuberculosis (13.0%) than WHO (12·0%). Our list of countries with high burden of tuberculosis is consistent with that of WHO, with a few exceptions. Afghanistan and Cambodia are lower in our estimates, and surpassed by Ukraine and Angola.

Cardiovascular diseases

Ageing and population growth have led to a growing number of people living with atherosclerotic vascular disease worldwide, despite the decrease in incident myocardial infarction and ischaemic stroke in highincome regions. Rising levels of obesity and diabetes in many countries makes this an area of major concern for global health. Increased attention will need to be directed toward this highly treatable set of disorders. Health systems will need to improve the delivery of cost-effective treatments such as blood pressure and cholesterollowering drugs while efforts continue to focus on decreasing tobacco smoking, improving diet, and increasing physical activity. Investments in universal primary care and public health campaigns need to be balanced against the need to improve access to emergency care, including the strengthening of pre-hospital systems and expanded access to revascularisation treatments.

Cancer

The International Agency for Research on Cancer last produced cancer estimates by country, age, sex, and cancer site for 2012 for the GLOBOCAN project.⁶⁰ The total estimated cancer incidence from GLOBOCAN for 2012 was 14·1 million individuals. By comparison, GBD 2015 estimated 18·6 million (18·0–19·4 million) new cases of cancer in 2015, which includes cases of non-melanoma skin cancer. GLOBOCAN used nine different

methods to estimate incidence, which precludes the calculation of uncertainty for those estimates. The GBD relies on the strength of the mortality estimates and transformation of the mortality estimates to incidence, taking into account uncertainty associated with both the mortality estimates and with the mortality to incidence ratios. Until high-quality cancer incidence is available in all countries, the GBD approach, which maximises the amount of data used to determine incidence estimates, is highly beneficial.

Vision loss

Globally, 34.3 million (30.7–38.0 million) people are blind, an additional 24.3 million (21.6-27.6 million) have severe vision impairment, 214 million (193-237 million) have moderate vision impairment, and 663 million (638-690 million) have near vision impairment in 2015. Combined, vision loss accounts for 24.5 million (17.0-34.5 million) YLDs and is the third largest impairment after anaemia (77.9 million [52·4–111·4 million] YLDs) and hearing loss (46·2 million [31-6-63-2 million] YLDs). YLDs from vision loss are slightly lower than those for anxiety disorders, which rank ninth worldwide. Largely due to ageing, substantial increases occurred from 2005 to 2015 in the number of people with blindness (increasing by 23.3% [21.9–24.6%]), severe visual impairment (24.3% [22.4-26.0%]), moderate visual impairment (22.0% [20.1-23.7%]), and near vision impairment (22.5% [20.9-24.0%]). Most causes of vision loss can be prevented or cured using cost-effective interventions. 61,62

Chronic kidney disease

Compared with our GBD 2013 estimates, we estimated global prevalence of chronic kidney disease to be lower by a third. Data for chronic kidney disease at younger ages are sparse. We increased the number of datapoints below the age of 30 years from 20 to 51, and that led to considerably lower prevalence estimates for these ages. We have used different strategies for estimating deaths versus YLDs secondary to chronic kidney disease due to other causes. In our cause of death analysis, deaths secondary to many of the causes typically categorised within the chronic kidney disease other category, such as cystic disease and other congenital renal diseases, were already counted under the primary disease following the principles of the ICD, and thus could not be recounted within the chronic kidney disease other category. In comparison, for the YLD analysis we counted chronic kidney disease from these causes in the chronic kidney disease other category. This difference in the classification of morbidity and mortality from chronic kidney disease other is driven by the underlying nature of the data on causes of death and prevalence. In future GBD versions, we plan to make explicit estimates of death and YLDs for polycystic kidney disease, an important component of the other category.

Emerging infectious diseases

Emerging and re-emerging infectious diseases pose a persistent yet dynamic challenge to global health⁶³ and to the GBD study. GBD 2015 has included Ebola virus disease in response to the devastating outbreak in west Africa starting in 2013,⁶⁴ whereas we have not captured the recent expansions of chikungunya⁶⁵ and Zika virus across the Americas,⁶⁶ for instance. Quantifying the effect of newly emerging epidemic diseases is difficult because the formal reporting of the disease events might considerably lag the first presentation of cases. As a result, contemporary events such as the substantial deviation from baseline levels in reported congenital abnormalities in Brazil in 2015,⁶⁷ now established to be associated with Zika virus,⁶⁸ will need to be incorporated into GBD 2016.

Advances in data and analysis

DisMod-MR

For GBD 2015, we improved the estimation for subnational units in DisMod-MR 2.1 compared to GBD 2013 and made substantive efforts to standardise the modelling approach across diseases. Three changes had an important effect on our estimates. First, the cascade was implemented such that national estimates were used to inform subnational estimation (in GBD 2013, each subnational unit was treated as a country within a GBD region). Second, wherever possible, we improved the linkage in the estimation between cause-specific mortality and disease incidence and prevalence by the systematic inclusion of estimated excess mortality data to provide more informative priors for each location in the geographical hierarchy. Third, in GBD 2010, and to a lesser degree in GBD 2013, many DisMod-MR models did not include fixed effects of country covariates for incidence or prevalence. Spatial heterogeneity was largely estimated from the random effects. For GBD 2015, we systematically tested the inclusion of more fixed effects in DisMod-MR 2.1 models including, where appropriate, variables related to the risk factor exposures we estimate for each disease.

Disability weights

For GBD 2015, we have not incorporated any new population-based data on disability weights; we have used the same disability weights as for GBD 2013. The present set of disability weights were based on population surveys in nine countries and an open internet survey. So far, these studies have not found systematic variation in disability weights across populations or within populations as a function of education, income per capita, or other variables. In future, we hope that more population surveys on disability weights will continue to be done to enrich the database for disability weight estimation. Given that we have not been able to collect disability weights in every location included in the GBD, it is possible that disability weights might vary systematically across communities. Even if this were to be confirmed empirically, for global standardised comparisons we would still use the global average disability weights; however, country analyses might use local disability weights. Until there is evidence of systematic variation across communities, this remains a theoretical and not a practical consideration. Given the close correlation between the internet survey and the population-based surveys, we are considering the implementation of an open rolling internet survey to collect more data for disability weights including new or revised health-state descriptions.

GATHER compliance

Providing all documentation for data sources, establishing access to the datasets used in modelling, and posting the code has been a labour-intensive activity; GBD 2015 is compliant with the new GATHER guidelines as a result. Posting code, we believe, will stimulate other researchers to explore the methods used in GBD estimation of nonfatal outcomes and hopefully lead to suggestions for improved estimation. With a steadily growing set of coinvestigators and a widening community interested in global health estimation, the enhanced transparency of GBD will improve the efficacy of the overall effort.

Using claims data

For the USA we used claims data for selected disorders. These data were particularly useful in the estimation of state-by-state prevalence for some disorders. Claims data, however, have important potential biases. Because of exclusion of some disadvantaged groups from health insurance, disease rates might be different for individuals that are covered compared with those from the general population. 69,70 As the claims dataset included information from Medicaid and Medicare, the coverage schemes for low-income citizens and older adults (older than 65 years) in the USA, the bias toward underestimation might not be so great. Indeed, we found that for many diseases, such as asthma, diabetes, or rheumatoid arthritis, the claims data were consistent with high-quality survey estimates. A potential problem of overestimation exists because visits to rule out a diagnosis can be coded to the diagnosis. Others working with claims have counted diagnoses only if they had appeared at least twice during a year. Applying such a restriction would have led to rejection of 44-68% of skin disorders, such as acne, psoriasis, or scabies, where one would expect that even after a single visit a diagnosis can be reliably made. As we tend to adjust the claims data used in DisMod-MR 2.1 if a systematic bias is detected from population-based sources, such as the National Health and Nutrition Examination Survey or other quality surveys, we believe our estimates have not been influenced much by this problem. The potential for greater inclusion of claims data in the GBD analysis is large; in countries with universal health access, claims data might overcome the problems of non-responder bias in survey data, which tend to lead to underestimates of the true population prevalence or incidence.

Self-reported functional health status

We found only modest progress in high-SDI countries in reducing age-standardised YLDs per capita over the last 25 years. There are, however, reports in the literature of substantial improvements based on self-rated health.71,72 In the USA, these improvements seem to be survey programme-specific and have not been reported in all data systems.⁷¹ Nevertheless, more work is needed to understand the different trends between our assessment of YLDs per capita and self-rated health. Item response theory has been used in some fields to identify changing response patterns for single items compared with the pool of all items.73 Item response theory cannot capture systematic changes that affect a set of items, and anchoring vignettes have been proposed as a strategy to deal with these challenges.^{72,74} In the era of increased interest in measuring functional health status, more exploration of the reasons for the divergence from a sequelae-based approach to measuring functional health status used in GBD versus an overall self-rating will be an important avenue of future research.

Crosswalks

Because of the diversity of data sources available, we estimated the statistical association between measurements taken using different case definitions, different assays, different survey items, or different ascertainment methods to a reference category. We used these statistical associations to crosswalk each type of data to the reference category. Crosswalks are a crucial dimension in the GBD analysis; in most cases we estimate these crosswalks from within DisMod-MR 2.1. If crosswalks were believed to vary substantially by age, sex, or location, these crosswalks were estimated outside of DisMod-MR 2.1 and adjusted data were used as inputs. Where data are sparse, the estimated crosswalks can shift substantially when new studies are identified that inform the crosswalk. With each iteration of the GBD, we have recognised the crucial nature of the crosswalk analysis for data processing. We believe that more research is needed in future iterations of GBD to standardise the approaches used for estimating crosswalks across disorders and risk factors, and for propagating uncertainty in crosswalks into the final results.

Data gaps

We have described the availability of data by cause and geography over different time periods, highlighting very large gaps in information availability. The data availability by country ranged from 6·1% in North Korea to 91·3% in the USA (methods appendix p 606). Among the top five causes of YLDs, data availability was rather poor. Lower back pain, iron-deficiency anaemia, major depressive disorder, other hearing loss, and neck pain all have data representativeness indices below 50%. Because of the broad overview provided of major data sources for disease

incidence and prevalence, GBD provides a framework to assist countries in prioritising the collection of new data to inform better monitoring of functional health status. By setting clear reference case definitions and data collection methods (methods appendix pp 4–7), GBD can also provide guidance on how to collect information most relevant to population health measurement.

Future directions for GBD

With each cycle of the GBD we expect, given the present interest in subnational assessments, to report increasingly granular results. These subnational findings will be of important national policy interest, but the discipline of examining the evidence for each community and modelling at the more granular level will, we believe, improve the quality of the national estimation as well. Additions to the cause list will continue to be driven by policy interest, such as the need to incorporate Zika virus, to split diabetes into separate estimates for type 1 and type 2, and by adding diseases to our cause list that are main contributors to large residual categories, such as other cardiovascular disease. other musculoskeletal disorders, and other neurological disorders. We plan to continue to expand our networks of topic-specific and country collaborators to enhance the quality of our estimates through their feedback and by enhancing the amount of data we can bring to bear on GBD estimation. Our country collaborations to generate subnational estimates have shown us that these efforts greatly enhance access to valuable data sources, particularly administrative datasets on inpatient and outpatient episodes and unpublished surveys.

Limitations

The GBD 2015 study has some key limitations. First, although we have sought to capture and include in our estimations many sources of uncertainty, we have not captured all sources. We have not been able to routinely capture uncertainty due to different model specifications in our results. We do not have the ability to reflect the uncertainty of data sources that exist but of which we are not aware. Subnational collaborations in China and Mexico revealed many data sources not captured in previous iterations of the GBD study. Inclusion of these data sources can lead to shifts in estimates that are outside the previously estimated 95% UIs.

Second, within DisMod-MR 2.1 we estimate the average association between different types of studies for reporting on a specific outcome such as the difference between 12-month recall and point prevalence in a survey. These estimated associations from within the DisMod-MR 2.1 likelihood estimation are used to adjust the data to a reference case definition or study design. These estimated adjustment factors are themselves uncertain, which increases the overall uncertainty in our estimation. More standardisation in data collection would be highly desirable and would reduce our dependence on the relatively challenging estimation process involved in crosswalks.

Third, for some disorders the data available to estimate excess mortality by age and sex and to capture how excess mortality changes with development status are very limited. There are probably many unpublished sources of information in some countries that could be usefully brought to bear on the challenge of estimating the age–sex–year–location levels of excess mortality.

Fourth, the microsimulation step of this study assumes that prevalence within an age—sex—location—year group of different sequelae is independent. Although this is clearly known not to be the case for some pairs of disorders, the independence assumption provides reasonable overall adjustments for comorbidity. Progress in incorporating dependent comorbidity has been difficult because information on the correlation structure of prevalence is extremely limited and only available for a minor fraction of all possible pairs of conditions in the GBD study.

Fifth, we estimated separate DisMod-MR 2.1 models for 1990, 1995, 2000, 2005, 2010, and 2015. Independent estimation of each time period implies that the uncertainty intervals for each period are also independent. Furthermore, compositional bias in the data available in different time periods might lead to spurious time trends. A more appealing strategy would be to simultaneously estimate the trends in incidence, excess mortality, and remission by age and sex consistent with all the available data for a geography on prevalence, incidence, remission, excess mortality, and cause-specific mortality by age and sex. DisMod-MR 2.1 does not allow for time-varying trends in incidence, excess mortality, and remission, but a prototype DisMod-MR AT (age and time) has been developed and is being tested. Allowing for all rates to change at different paces over age and time increases the number of parameters that need to be estimated by orders of magnitude.

Sixth, we have emphasised in the reporting of results the changes from calendar year 2005 to calendar year 2015, although we have estimated results for 1990, 1995, 2000, 2005, 2010, and 2015 and interpolated for the years in between. For some disorders, reporting the change from 2005 to 2015 (such as for Ebola virus disease) does not capture the major epidemic in west Africa in 2014. Likewise, disasters and wars are often concentrated in a single year; comparisons of any two years can provide misleading inferences about trends.

Seventh, for a few disorders (eg, tetanus, neonatal sepsis, rabies, and diphtheria) we estimate disease incidence from estimates of mortality and the inverse of the case-fatality rate estimated from available studies. When the case-fatality rate is very low, these estimates of incidence are highly sensitive to very small changes in the estimated case-fatality rate and have large uncertainty intervals.

Finally, although extraordinary efforts have gone into vetting the results by GBD researchers and the collaborator network, there are probably some findings that have not been scrutinised carefully enough. Making all results and underlying data available and having an increasing number of visualisation tools are effective

strategies that we will keep expanding to meet our goal of producing the highest quality global health data to our growing audience of policy makers, researchers, the media, and the general population.

Conclusion

The GBD studies seek to quantify the prevalence and incidence of the major sequelae for a comprehensive list of diseases and injuries; given the diversity of data sources, the range of biases in these sources, and the gaps in availability, this is a challenging task. Despite the limitations, the standardised and comprehensive approach of the GBD studies study provides useful insights. We believe users of the findings, including the wealth of country-specific and cause-specific detail available in the appendix, can usefully examine the broad trends for their country or subnational geography, benchmark against geographies at a similar level of development, and understand the strength or weakness of these estimates. Regular quantification of health is particularly important as the new health-related targets of the Sustainable Development Goals have broadened the health agenda throughout the world. At the same time, development and transformations in the risks to health experienced by different groups in the world are leading to some broad transformations in health. Everyone needs to have access to the timeliest, valid, reliable, and local information possible to enrich debates on how to accelerate health progress in all communities.

GBD 2015 Disease and Injury Incidence and Prevalence Collaborators

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Christopher J L Murray and Theo Vos prepared the first draft. Alan D Lopez and Christopher J L Murray conceived the study and provided overall guidance. All other authors provided data, developed models, reviewed results, initiated modelling infrastructure, and/or reviewed and contributed to the report.

Declaration of interests

Bruce Bartholow Duncan and Maria Inês Schmidt have received additional funding from the Brazilian Ministry of Health (Process No 25000192049/2014-14). Itamar S Santos reports grants from FAPESP (Brazilian public agency), outside the submitted work. Carl Abelardo T Antonio reports grants, personal fees and non-financial support from Johnson & Johnson (Philippines), Inc, outside the submitted work. Cyrus Cooper reports other from Alliance for Better Bone Health, other from Amgen, other from Eli Lilly, other from GSK. other from Medtronic, other from Merck, other from Novartis, other from Pfizer, other from Roche, other from Servier, outside the submitted work. Walter Mendoza is currently employed by the Peru Country Office of the United Nations Population Fund, an institution which does not necessarily endorse this study. Donald S Shepard would like to acknowledge grant support from Sanofi Pasteur. Rafael Tabarés-Seisdedos and Ferrán Catalá-López are supported in part by grant PROMETEOII/2015/021 from Generalitat Valenciana, and Rafael Tabarés-Seisdedos is supported by the national grant PI14/00894 from ISCIII-FEDER. Walter Mendoza is currently employed by the Peru Country Office of the United Nations Population Fund, an institution which does not necessarily endorse this study. Veena J Iyer has received a Public Health Research Initiative Fellowship (2014-17) from the Department of Science and Technology (DST) for a project titled "Relationship between Enteric Fever incidence and Climate in the city of Ahmedabad, 1990-2014". Pablo M Lavados reports grants, personal fees and non-financial support from BAYER, non-financial support from Boehringer Ingelheim, grants and personal fees from AstraZeneca. grants from CONICYT, and grants from The George Institute for Global Health, outside the submitted work. Noela M Prasad reports and is an employee of an international NGO that raises funds from the Australian Public, and holds a honorary position at CERA, which receives Operational Infrastructure Support from the Victorian Government.

Bradford D Gessner reports grants from Crucell, GSK, Hilleman Labs, Novartis, Pfizer, Merck, and Sanofi Pasteur, outside the submitted work. Ai Koyanagi's work is supported by the Miguel Servet contract financed by the CP13/00150 and PI15/00862 projects, integrated into the National R + D + I and funded by the ISCIII - General Branch Evaluation and Promotion of Health Research - and the European Regional Development Fund (ERDF-FEDER). Dorairaj Prabhakaran reports grants from the Bill & Melinda Gates Foundation. Matthias Endres reports that The Center for Stroke Research Berlin has received institutional funding from the German Ministry for Research and Education (BMBF). Katharine J Looker has received funding from the World Health Organization for the HSV-2 seroprevalence review which informs this work; during the study, KJL also received separate funding from the World Health Organization, USAID/PATH, Sexual Health 24 and the National Institute for Health Research (NIHR) Health Protection Research Unit (HPRU) in Evaluation of Interventions at the University of Bristol; these funders had no role in the writing of the manuscript nor the decision to submit it for publication, and the views expressed in this Article do not necessarily represent the views, decisions or policies of the World Health Organization, the NHS, the NIHR, the Department of Health or Public Health England. Donal Bisanzio is supported by Bill and Melinda Gates Foundation (#OPP1068048). Thomas Fürst has received financial support from the Swiss National Science Foundation (SNSF; project no P300P3-154634). Rodrigo Sarmiento-Suarez receives institutional support from Universidad de Ciencias Aplicadas y Ambientales, UDCA, Bogotá Colombia. Ronan A Lyons is supported by two grants: The Farr Institute of Health Informatics Research: Arthritis Research UK, the British Heart Foundation, Cancer Research UK, the Economic and Social Research Council, the Engineering and Physical Sciences Research Council, the Medical Research Council, the National Institute of Health Research, the National Institute for Social Care and Health Research (Welsh Assembly Government), the Chief Scientist Office (Scottish Government Health Directorates), and The Wellcome Trust. Grant No MR/K006525/1, and the National Centre for Population Health and Wellbeing Research. Health and Care Research Wales. Stefanos Tyrovolas's work is supported by the Foundation for Education and European Culture (IPEP), the Sara Borrell postdoctoral programme (reference no CD15/00019 from the Instituto de Salud Carlos III (ISCIII - Spain) and the Fondos Europeo de Desarrollo Regional (FEDER). Manami Inoue is the beneficiary of a financial contribution from the AXA Research fund as chair holder of the AXA Department of Health and Human Security, Graduate School of Medicine, The University of Tokyo from Nov 1, 2012; the AXA Research Fund has no role in this work. Sinead M Langan holds an NIHR Clinician Scientist Fellowship (NIHR/CS/010/014); the views expressed in this publication are those of the authors and not necessarily those of the NHS, the NIHR, or the UK Department of Health. Scott Weichenthal acknowledges financial support from the Cancer Research Society of Canada. Yogeshwar Kalkonde is a Wellcome Trust/ DBT India Alliance Intermediate Fellow in Public Health. John J McGrath received a NHMRC John Cade Fellowship (APP1056929). Sarah Derrett reports grants, personal fees, non-financial support and other from EuroQol Research Foundation, outside the submitted work. Dan J Stein reports personal fees from Lundbeck, personal fees from Novartis, personal fees from AMBRF, grants from NRGF, personal fees from Biocodex, personal fees from Sevier, grants from MRC, personal fees from SUN, and personal fees from CIPLA, outside the submitted work. Tea Lallukka reports funding from The Academy of Finland, grant #287488. Charles D A Wolfe's research was funded and supported by the National Institute for Health Research (NIHR) Biomedical Research Centre based at Guy's and St Thomas' NHS Foundation Trust and King's College London. The views expressed are those of the author(s) and not necessarily those of the NHS, the NIHR, or the Department of Health. The other authors declare no competing interests.

Acknowledgments

We would like to thank the countless individuals who have contributed to the Global Burden of Disease Study 2015 in various capacities. This paper uses data from SHARE Waves 1, 2, 3 (SHARELIFE), 4, and 5 (DOIs: 10.6103/SHARE.w1.500, 10.6103/SHARE.w3.500, 10.6103/SHARE.w4.500, 10.6103/SHARE.w5.500), see Börsch-Supan et al (2013) for methodological details. The SHARE data collection has been

primarily funded by the European Commission through FP5 (QLK6-CT-2001-00360), FP6 (SHARE-I3: RII-CT-2006-062193, COMPARE: CIT5-CT-2005-028857, SHARELIFE: CIT4-CT-2006-028812), and FP7 (SHARE-PREP: N°211909, SHARE-LEAP: N°227822, SHARE M4: N°261982). Additional funding from the German Ministry of Education and Research, the US National Institute on Aging (U01_AG09740-13S2, P01_ AG005842, P01_AG08291, P30_AG12815, R21_AG025169, Y1-AG-4553-01, IAG_BSR06-11, OGHA_04-064) and from various national funding sources is gratefully acknowledged (see www.share-project.org). The data reported here have been supplied by the United States Renal Data System (USRDS). The interpretation and reporting of these data are the responsibility of the author(s) and in no way should be seen as an official policy or interpretation of the US Government. The Palestinian Central Bureau of Statistics granted the researchers access to relevant data in accordance with license no. SLN2014-3-170, after subjecting data to processing aiming to preserve the confidentiality of individual data in accordance with the General Statistics Law—2000. The researchers are solely responsible for the conclusions and inferences drawn upon available data. This study has been realised using the data collectved by the Swiss Household Panel (SHP), which is based at the Swiss Centre of Expertise in the Social Sciences FORS. The project is financed by the Swiss National Science Foundation. The following individuals would like to acknowledge various forms of institutional support: Amanda G Thrift is supported by a fellowship from the National Health and Medical Research Council (GNT1042600). Panniyammakal Jeemon is supported by the Wellcome Trust-DBT India Alliance, Clinical and Public Health, Intermediate Fellowship (2015-20). Amador Goodridge would like to acknowledge funding for me from Sistema Nacional de Investigadores de Panamá-SNI. José das Neves was supported in his contribution to this work by a Fellowship from Fundação para a Ciência e a Tecnologia, Portugal (SFRH/BPD/92934/2013). Boris Bikbov, Monica Cortinovis, Giuseppe Remuzzi, and Norberto Perico would like to acknowledge that their contribution to this paper has been on behalf of the International Society of Nephrology (ISN) as a follow-up of the activities of the GBD 2010 Genitourinary Diseases Expert Group. Lijing L Yan is supported by the National Natural Sciences Foundation of China grants (71233001 and 71490732). Miriam Levi would like to acknowledge the institutional support received from CeRIMP, Regional Centre for Occupational Diseases and Injuries, Tuscany Region, Florence, Italy. Simon I Hay is funded by a Senior Research Fellowship from the Wellcome Trust (#095066), and grants from the Bill & Melinda Gates Foundation (OPP1119467, OPP1093011, OPP1106023 and OPP1132415). No individuals acknowledged received additional compensation for their efforts.

References

- 1 Atun R, de Jongh T, Secci F, Ohiri K, Adeyi O. Integration of targeted health interventions into health systems: a conceptual framework for analysis. Health Policy Plan 2010; 25: 104–11.
- 2 Park J-H, Eum J-H, Bold B, Cheong H-K. Burden of disease due to dementia in the elderly population of Korea: present and future. BMC Public Health 2013; 13: 293.
- 3 George-Carey R, Adeloye D, Chan KY, et al. An estimate of the prevalence of dementia in Africa: a systematic analysis. J Glob Health 2012; 2: 020401.
- 4 Global Burden of Disease 2013 Collaborators. Global, regional, and national incidence, prevalence, and years lived with disability for 301 acute and chronic diseases and injuries in 188 countries, 1990–2013: a systematic analysis for the Global Burden of Disease Study 2013. *Lancet* 2015; 386: 743–800.
- 5 Prince MJ, Wu F, Guo Y, et al. The burden of disease in older people and implications for health policy and practice. *Lancet* 2015; 385: 549–62.
- 6 Hallett TB, Zaba B, Stover J, et al. Embracing different approaches to estimating HIV incidence, prevalence and mortality. AIDS 2014; 28: S523–32.
- 7 Supervie V, Archibald CP, Costagliola D, et al. GBD 2013 and HIV incidence in high income countries. *Lancet* 2015; 385: 1177.
- 8 Cohen J, Vincent J-L, Adhikari NKJ, et al. Sepsis: a roadmap for future research. *Lancet Infect Dis* 2015; 15: 581–614.
- 9 Marshall JC. Understanding the global burden of pediatric sepsis. Am J Respir Crit Care Med 2015; 191: 1096–98.
- 10 Anderson BO, Flanigan J. Novel methods for measuring global cancer burden: implications for global cancer control. JAMA Oncol 2015; 1: 425–27.

- 11 Ton TGN, Mackenzie C, Molyneux DH. The burden of mental health in lymphatic filariasis. *Infect Dis Poverty* 2015; 4: 34.
- Wagner RG, Ibinda F, Tollman S, Lindholm L, Newton CR, Bertram MY. Differing methods and definitions influence DALY estimates: using population-based data to calculate the burden of convulsive epilepsy in rural South Africa. PLoS One 2015; 10: e0145300.
- 13 Alzheimer's Disease International. World Alzheimer Report, 2015. https://www.alz.co.uk/research/WorldAlzheimerReport2015.pdf (accessed May 26, 2016).
- 14 Hausman DM. Health, well-being, and measuring the burden of disease. *Popul Health Metr* 2012; 10: 13.
- Nord E. Uncertainties about disability weights for the Global Burden of Disease study. Lancet Glob Health 2015; 3: e661–62.
- 16 GBD Mortality and Causes of Death Collaborators. Global, regional, and national life expectancy, all-cause and cause-specific mortality for 249 causes of death, 1980–2015: a systematic analysis for the Global Burden of Disease Study 2015. *Lancet* 388: 1459–544.
- 17 Stevens GA, Alkema L, Black RE, et al. Guidelines for Accurate and Transparent Health Estimates Reporting: The GATHER statement. *Lancet* 2016; published online June 28. http://dx.doi.org/10.1016/ S0140-6736(16)30388-9.
- 18 Kiyono P. An integrative metaregression framework for descriptive epidemiology, 1 edn. Seattle: University of Washington Press, 2015.
- 19 Clark DV, Kibuuka H, Millard M, et al. Long-term sequelae after Ebola virus disease in Bundibugyo, Uganda: a retrospective cohort study. Lancet Infect Dis 2015; 15: 905–12.
- 20 Qureshi AI, Chughtai M, Loua TO, et al. Study of Ebola virus disease survivors in Guinea. Clin Infect Dis 2015; 61: 1035–42.
- 21 Rowe AK, Bertolli J, Khan AS, et al. Clinical, virologic, and immunologic follow-up of convalescent Ebola hemorrhagic fever patients and their household contacts, Kikwit, Democratic Republic of the Congo. J Infect Dis 1999; 179: S28–35.
- 22 Bwaka MA, Bonnet M-J, Calain P, et al. Ebola Hemorrhagic Fever in Kikwit, Democratic Republic of the Congo: clinical observations in 103 patients. J Infect Dis 1999; 179: S1–S7.
- 23 PRO-ACT. https://nctu.partners.org/ProACT/Data/Index?Length=0 &LongLength=0&Rank=1&SyncRoot=System.Type%5B%5D&IsRea dOnly=False&IsFixedSize=True&IsSynchronized=False (accessed May 26, 2016).
- 24 Vos T, Flaxman AD, Naghavi M, et al. Years lived with disability (YLDs) for 1160 sequelae of 289 diseases and injuries 1990–2010: a systematic analysis for the Global Burden of Disease Study 2010. *Lancet* 2012; 380: 2163–96.
- 25 Global, regional, and national age-sex specific all-cause and cause-specific mortality for 240 causes of death, 1990–2013: a systematic analysis for the Global Burden of Disease Study 2013. *Lancet* 2015; 385: 117–71.
- 26 Murray CJL, Barber RM, Foreman KJ, et al. Global, regional, and national disability-adjusted life years (DALYs) for 306 diseases and injuries and healthy life expectancy (HALE) for 188 countries, 1990–2013: quantifying the epidemiological transition. *Lancet* 2015; 386: 2145–91.
- 27 United Nations Development Programme. Human development report 2015. http://hdr.undp.org/sites/default/files/2015_human_ development_report.pdf (accessed May 26, 2016).
- 28 Goldman DP, Cutler D, Rowe JW, et al. Substantial health and economic returns from delayed aging may warrant a mew focus for medical research. *Health Aff (Millwood)* 2013; 32: 1698–705.
- 29 Dufouil C, Pereira E, Chêne G, et al. Older age at retirement is associated with decreased risk of dementia. Eur J Epidemiol 2014; 29: 353–61.
- 30 Belbase A, Sanzenbacher G, Gillis CM. Does age-related decline in ability correspond with retirement age? Rochester, NY: Social Science Research Network, 2015 http://papers.ssrn.com/ abstract=2665830 (accessed June 24, 2016).
- 31 van Rijn RM, Robroek SJW, Brouwer S, Burdorf A. Influence of poor health on exit from paid employment: a systematic review. Occup Environ Med 2014; 71: 295–301.
- 32 Lahelma E, Pietiläinen O, Rahkonen O, Lallukka T. Common mental disorders and cause-specific disability retirement. Occup Environ Med 2015; 72: 181–87.

- 33 Begg S, Vos T, Goss J, Mann N. An alternative approach to projecting health expenditure in Australia. Aust Health Rev 2008; 32: 148–55
- 34 Desveaux L, Beauchamp M, Goldstein R, Brooks D. Community-based exercise programs as a strategy to optimize function in chronic disease: a systematic review. Med Care 2014; 52: 216–26.
- 35 Jahanbin I, Hoseini Moghadam M, Nazarinia MA, Ghodsbin F, Bagheri Z, Ashraf AR. The effect of conditioning exercise on the health status and pain in patients with rheumatoid arthritis: a randomized controlled clinical trial. Int J Community Based Nurs Midwifery 2014; 2: 169–76.
- 36 Gay C, Chabaud A, Guilley E, Coudeyre E. Educating patients about the benefits of physical activity and exercise for their hip and knee osteoarthritis. Systematic literature review. Ann Phys Rehabil Med 2016; 59: 174–83.
- 37 McAlindon TE, Bannuru RR, Sullivan MC, et al. OARSI guidelines for the non-surgical management of knee osteoarthritis. Osteoarthris Cartilage 2014; 22: 363–88.
- 38 Heimans L, Akdemir G, Boer KVCW, et al. Two-year results of disease activity score (DAS)-remission-steered treatment strategies aiming at drug-free remission in early arthritis patients (the IMPROVED-study). Arthritis Res Ther 2016; 18: 23.
- 39 Kleinman A, Estrin GL, Usmani S, et al. Time for mental health to come out of the shadows. *Lancet* 2016: 387: 2274–75.
- 40 Izutsu T, Tsutsumi A, Minas H, Thornicroft G, Patel V, Ito A. Mental health and wellbeing in the Sustainable Development Goals. Lancet Psychiatry 2015; 2: 1052–54.
- 41 Patel V, Chisholm D, Parikh R, et al. Addressing the burden of mental, neurological, and substance use disorders: key messages from Disease Control Priorities, 3rd edn. *Lancet* 2016; 387: 1672–85.
- 42 Centers for Disease Control and Prevention. Drug-poisoning deaths involving heroin: United States, 2000–2013. http://www.cdc.gov/ nchs/products/databriefs/db190.htm (accessed June 24, 2016).
- 43 Centers for Disease Control and Prevention. Morbidity and Mortality Weekly Report (MMWR). https://www.cdc.gov/mmwr/ preview/mmwrhtml/mm6401a10.htm (accessed June 24, 2016).
- 44 Centers for Disease Control and Prevention. Opioid overdose prevention programs providing naloxone to laypersons—United States, 2014. http://origin.glb.cdc.gov/mmwr/preview/mmwrhtml/ mm6423a2.htm?s_cid=mm6423a2_w (accessed June 24, 2016).
- 45 Mathers BM, Degenhardt L, Ali H, et al. HIV prevention, treatment, and care services for people who inject drugs: a systematic review of global, regional, and national coverage. *Lancet* 2010; 375: 1014–28.
- 46 GBD 2015 Risk Factors Collaborators. Global, regional, and national comparative risk assessment of 79 behavioural, environmental and occupational, and metabolic risks or clusters of risks, 1990–2015: a systematic analysis for the Global Burden of Disease Study 2015. Lancet 2016; 388: 1659–724.
- 47 Association AD. Economic costs of diabetes in the US in 2012. Diabetes Care 2013; 36: 1033–46.
- 48 NCD Risk Factor Collaboration. Worldwide trends in diabetes since 1980: a pooled analysis of 751 population-based studies with 4-4 million participants. *Lancet* 2016; 387: 1513–30.
- 49 International Diabetes Federation. IDF diabetes atlas. http://www.diabetesatlas.org/ (accessed June 25, 2016).
- 50 Langa KM. Is the risk of Alzheimer's disease and dementia declining? Alzheimers Res Ther 2015; 7: 34.
- Matthews FE, Arthur A, Barnes LE, et al. A two-decade comparison of prevalence of dementia in individuals aged 65 years and older from three geographical areas of England: results of the Cognitive Function and Ageing Study I and II. *Lancet* 2013; 382: 1405–12.

- 52 Rocca WA, Petersen RC, Knopman DS, et al. Trends in the incidence and prevalence of Alzheimer's disease, dementia, and cognitive impairment in the United States. Alzheimers Dement 2011; 7: 80–93.
- 53 Brookmeyer R, Evans DA, Hebert L, et al. National estimates of the prevalence of Alzheimer's disease in the United States. Alzheimers Dement 2011; 7: 61–73.
- 54 Helsper CW, Hellinga HL, van Essen GA, et al. Real-life costs of hepatitis C treatment. Neth J Med 2012; 70: 145–53.
- 55 Bhatt S, Weiss DJ, Cameron E, et al. The effect of malaria control on Plasmodium falciparum in Africa between 2000 and 2015. Nature 2015; 526: 207–11.
- 56 Cameron E, Battle KE, Bhatt S, et al. Defining the relationship between infection prevalence and clinical incidence of *Plasmodium* falciparum malaria. Nat Commun 2015; 6: 8170.
- 57 Bhatt S, Weiss DJ, Mappin B, et al. Coverage and system efficiencies of insecticide-treated nets in Africa from 2000 to 2017. eLife 2015; 4: e09672.
- S8 Aspiration to action. What will it take to end malaria? http:// endmalaria2040.org/assets/Aspiration-to-Action.pdf (accessed May 26, 2016).
- 59 WHO. World Malaria Report 2015. http://www.who.int/malaria/publications/world-malaria-report-2015/report/en/ (accessed Aug 5, 2016).
- 60 Globocan 2012. http://globocan.iarc.fr/Default.aspx (accessed June 24, 2016).
- 61 Pizzarello L, Abiose A, Ffytche T, et al. VISION 2020: The Right to Sight: a global initiative to eliminate avoidable blindness. Arch Ophthalmol 1960 2004; 122: 615–20.
- 62 Baltussen R, Smith A. Cost effectiveness of strategies to combat vision and hearing loss in sub-Saharan Africa and South East Asia: mathematical modelling study. BMJ 2012; 344: e615.
- 63 Currie J, Grenfell B, Farrar J. Beyond Ebola. Science 2016; 351: 815-16.
- 64 Baize S, Pannetier D, Oestereich L, et al. Emergence of Zaire Ebola virus disease in Guinea. *N Engl J Med* 2014; **371**: 1418–25.
- 65 Weaver SC. Arrival of Chikungunya virus in the new world: prospects for spread and impact on public health. PLoS Negl Trop Dis 2014; 8: e2921.
- 66 Fauci AS, Morens DM. Zika Virus in the Americas—yet another arbovirus threat. *N Engl J Med* 2016; **374**: 601–04.
- 67 Microcephaly in Infants, Pernambuco State, Brazil, 2015. Emerg Infect Dis 2016; 22: 1090–93.
- 68 Rasmussen SA, Jamieson DJ, Honein MA, Petersen LR. Zika Virus and birth defects—reviewing the evidence for causality. N Engl J Med 2016; 374: 1981–87.
- 69 Jensen ET, Cook SF, Allen JK, et al. Enrollment factors and bias of disease prevalence estimates in administrative claims data. Ann Evidemiol 2015: 25: 519–25.
- 70 Kottke TE, Baechler CJ, Parker ED. Accuracy of heart disease prevalence estimated from claims data compared with an electronic health record. *Prev Chronic Dis* 2012; 9: E141.
- 71 Salomon JA, Nordhagen S, Oza S, Murray CJL. Are Americans feeling less healthy? The puzzle of trends in self-rated health. Am J Epidemiol 2009; 170: 343–51.
- 72 Salomon JA, Tandon A, Murray CJL. Comparability of self rated health: cross sectional multi-country survey using anchoring vignettes. BMJ 2004; 328: 258.
- 73 Meijer RR. Diagnosing item score patterns on a test using item response theory-based person-fit statistics. *Psychol Methods* 2003; 8: 72–87.
- 74 King G, Wand J. Comparing incomparable survey responses: new tools for anchoring vignettes. *Polit Anal* 2007; 15: 46–66.