# THE LANCET

## Supplementary appendix

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#### **Appendix Text 1.** NCD Risk Factor Collaboration (NCD-RisC)

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#### Appendix Text 2. Data sources

#### Data access

We used data from a population-based database on cardiometabolic risk factors collated by the Non-Communicable Disease Risk Factor Collaboration (NCD-RisC), as detailed previously. Data were obtained from publicly available multi-country and national measurement surveys (e.g., WHO STEPwise approach to Surveillance (STEPS) surveys, and those identified via the Inter-University Consortium for Political and Social Research, UK Data Service, and European Health Interview & Health Examination Surveys Database). With the collaboration of the WHO and its regional and country offices, we identified and accessed population-based survey data from national health and statistical agencies. We searched and reviewed published studies as detailed previously, and invited eligible studies to join NCD-RisC, as we did with data holders from earlier pooled analyses of cardiometabolic risk factors. The NCD-RisC database is continuously updated through all the above routes as well as through periodic requests to NCD-RisC members to suggest additional sources in their countries.

#### Data inclusion criteria and characteristics of included studies

We carefully checked that each data source met our inclusion criteria, described below. All NCD-RisC members were also periodically asked to review the list of sources from their country, to verify that they met the inclusion criteria and were not duplicates. Potential duplicate data sources were first identified by comparing studies from the same country and year, followed by checking with NCD-RisC members who had provided data whether sources from the same country and year, and with similar sample sizes and age ranges, were the same or distinct. If two sources were confirmed as duplicates, one was discarded.

For each data source, we recorded the study population, the sampling approach, the years of measurement and the measurement methods. Only population-based data were included. All data were assessed and classified by whether they covered the whole country, one or more

subnational regions (i.e., one or more provinces or states, more than three cities, or more than five rural communities), or one or a small number of communities (limited geographical scope not meeting above national or subnational criteria). As stated in statistical methods, these study-level attributes were used in the Bayesian hierarchical meta-regression model so that all available data were used, while taking into account the aforementioned differences in the populations from which different studies had sampled. We recorded whether FPG and HbA1c were measured in a laboratory or using a point-of-care portable device. For studies with laboratory measurement, we recorded whether plasma or whole blood was used. For studies that used portable devices, we verified using the device manual, discussion the study investigators, and if relevant the manufacturers, whether the device reported glucose values in whole blood or whether it converted to plasma-equivalent internally, with those reporting glucose in whole blood converted to plasma equivalent as described below. This information was used so that the calculated prevalence for each survey was correctly and consistently calculated using plasma-equivalent glucose. All submitted data were checked by at least two persons independently. Questions and clarifications were discussed with NCD-RisC members and resolved before data were incorporated into the database.

Data were included if the following criteria were met: measured data on FPG and/or HbA1c were available; study participants were 18 years of age and older; data were from population samples at the national, subnational, or community level; and data were from the countries listed in Appendix Table 2.

We excluded all data sources that were solely based on self-reported diagnosis of diabetes or a registry of people with diagnosed diabetes because a substantial proportion of people with diabetes remain undiagnosed, especially in low- and middle-income countries.<sup>7,8</sup> We excluded data sources on population subgroups whose glycaemic levels may be systematically high or low, including studies that had included or excluded participants based on health status; studies whose participants were only from specific educational, occupational,

socioeconomic or ethnic subgroups. We also excluded studies that recruited through health facilities; the exceptions to this exclusion were studies whose sampling frame was health insurance schemes that are not segregated by occupation or socioeconomic status in countries where at least 80% of the population were insured, and studies based on primary care system in high-income and central European countries with universal insurance, as contact with the primary care systems in these countries tends to be as good as or better than response rates for population-based surveys.

We excluded FPG data from studies that had not instructed participants to fast for at least 6 hours before FPG measurement. We excluded FPG or HbA1c data in studies that measured these biomarkers only among participants with previously-diagnosed diabetes or those with high casual glucose levels. We excluded HbA1c data from studies whose decision to measure HbA1c depended on participants' FPG or casual glucose levels, and vice versa. We also excluded HbA1c data from studies whose mid-year was before 2000, before HbA1c assays were standardised. When FPG and/or HbA1c data were missing for more than 10% of participants in a survey, we contacted the data providers and checked the study design documentation to verify that missingness was not based on pre-selected criteria and hence likely at random.

In addition to the studies in the NCD-RisC database, summary statistics such as diabetes prevalence, mean FPG and mean HbA1c for nationally representative studies that could not be accessed via the above routes were extracted from published reports. We also included such summaries from a previous global data pooling study,<sup>4</sup> when individual record data could not be accessed through the above routes, for example because the authors had retired or moved, data had been permanently archived, or data were stored using older storage technologies that could not be easily retrieved.

#### Data cleaning and management

In surveys with data on treatment, we determined whether a person was taking medication for diabetes using survey-specific questions worded as variations of "are you currently taking medication for diabetes or high blood sugar?"; or the combination of "do you currently inject/use insulin for diabetes?" and "are you currently taking any medicines, tablets, or pills for diabetes?"; or using information gathered on types of medicines used by the participant. Fifteen studies (1%) had only collected data on the use of oral hypoglycaemic drugs, but not insulin. For these studies, we used medication data for participants aged 40 years and older, but not for younger participants. We did this because data from studies that had information on both types of treatment showed that using insulin alone was more common for those in younger ages.

We excluded women who were pregnant at the time of examination. We excluded 32,498 participants (<0.1% of all participants) with FPG <2.0 mmol/L or >30.0 mmol/L, or HbA1c <3% or >18%, because these values are likely to reflect measurement or data recording errors. We excluded 254,954 participants with missing medication information (0.2% of all participants); those with and without medication information were similar in their age and BMI (differences of 0.6 years and 0.1 kg/m², respectively). In 74 studies (7%), fasting glucose was reported in capillary whole blood. We converted these measurements to plasma-equivalent values using a linear regression equation that quantified the relationship between the two in studies that had measured both whole-blood-based and plasma-based fasting glucose.<sup>10</sup>

Anonymised individual data from the studies in the NCD-RisC database were reanalysed according to a common protocol. We calculated diabetes prevalence and treatment, as defined in the primary outcomes, by sex and age group for each study. When applicable, we used survey sample weights and accounted for complex survey design in calculating summary statistics.

Some studies had asked questions about diabetes treatment from all participants but measured FPG and HbA1c in a subset of participants by design. The reasons for this practice include the different response rates for questionnaire and laboratory parts of a survey, cost of equipment or laboratory analysis, or the logistics of blood collection including fasting requirement for FPG. For example, the US National Health and Nutrition Examination Surveys (NHANES) divided participants into those who visited the data collection sites in the morning or afternoon; only the morning group was instructed to fast and had FPG data. The ICMR-INDIAB surveys in India measured HbA1c for every fifth of participants who did not have a prior diagnosis of diabetes. Both surveys collected data on diabetes medication from all participants.<sup>11,12</sup> In these studies, untreated diabetes was only calculated among those with biomarker measurement (see also Appendix Text 3).

Some studies in our analysis had data on both FPG and HbA1c; others had only one of these biomarkers (Appendix Table 1). In studies with data on both FPG and HbA1c, untreated diabetes was calculated using both biomarkers, i.e. participants with either FPG ≥7.0 mmol/L or HbA1c ≥6.5%. In studies with data on one biomarker only, participants whose measured biomarker was elevated (i.e., FPG ≥7.0 mmol/L in studies that had measured FPG; HbA1c ≥6.5% in studies that had measured HbA1c) were considered to have diabetes. For the remainder of the sample, who neither used treatment nor had elevated level of the measured biomarker, we used the coefficients of previously-validated regressions8 to estimate the probability of having elevated level of the second (unmeasured) biomarker. These regressions were developed and validated using data from studies that had measured both FPG and HbA1c.8 The model was specified as a logistic regression, and the predictors included the measured biomarker, age, sex, body-mass index, and region.8 For example, in a study that measured only FPG, those who used treatment or had FPG ≥7.0 mmol/L were considered to have diabetes; for those who did not use treatment and whose FPG was <7.0 mmol/L, we estimated the probability of having HbA1c ≥6.5% using the coefficients of the aforementioned regression. The probabilities were then summed across individuals to obtain the equivalent number of people with isolated elevated HbA1c, which was then added to the numerator of prevalence.

Some studies reported diabetes prevalence based on a different definition from the primary definition, e.g., one study used FPG ≥7.8 mmol/L and another used HbA1c ≥10%, or reported mean FPG. Many of these were from a previous global pooling study<sup>4</sup> or extracted from published reports and papers as stated earlier, hence reanalysis of their data was not feasible. This group also included studies with individual participant data that had not collected data on treatment and hence could be used to calculate mean FPG, mean HbA1c or prevalence of people with elevated levels of these markers, but not treated diabetes. We used regressions that converted data from these sources to our primary outcome for diabetes prevalence. The dependent variable in each of these regressions was the primary outcome for diabetes prevalence (prevalence of FPG ≥7.0 mmol/L, HbA1c ≥6.5%, or taking medication for diabetes), and the independent variable was a mean (e.g., mean FPG) or prevalence with a definition that differed from the primary outcome. The coefficients of these regressions were estimated from data sources which could be used to calculate both dependent and independent variables. Details of conversion regressions, and their specification, coefficients and performance, are reported at https://github.com/NCD-RisC/ncdrisc-methods/blob/main/NCD-RisC-conversion-model-for-diabetes-prevalence.pdf.

#### **Appendix Text 3.** Statistical methods

#### Overview

We used a Bayesian hierarchical meta-regression model, fitted using a Markov chain Monte Carlo (MCMC) sampler, with inference made using posterior MCMC samples, to estimate trends in diabetes prevalence and treatment, by sex, age, country, and year. As stated in the main paper, we report trends from 1990 to 2022, a period during which diabetes was recognised as an epidemic and the benefits of treatment were demonstrated in clinical trials. 13,14 Data from 1980 to 1989 were used in the model so that the estimates for 1990 and the subsequent years were informed by as many studies from the early period and from each country as possible. We modelled the two primary outcomes, diabetes prevalence and treatment, separately because their time trends and age associations may be different.

The statistical methods, including its implementation and computation, are detailed in a statistical paper<sup>15</sup> and related substantive papers.<sup>2,16</sup> Model specification is summarised below and further described using statistical notation. This is followed with details of model implementation and computation.

In summary, we organised countries into 20 regions, based on geography and other shared national characteristics, which were further grouped into eight super-regions (Appendix Table 2). The model had a hierarchical structure in which estimates for each country and year were informed by its own data, if available, and by data from other years in the same country and from other countries, especially those in the same region and super-region with data for similar time periods. The extent to which estimates for each country-year were influenced by data from other years and other countries depended on whether the country had data, the sample size of data, whether the available studies were at national, subnational or community level, and the within-country and within-region variability of the available data. Estimates for countries with more national studies, especially with data that were less variable, were informed to a greater degree by its own data than those with fewer studies, especially fewer

national studies, or those with data that varied extensively. At the extreme, for the 25 countries without data, the estimates were informed by data from other countries, especially those in the same region with data for similar time periods, and had larger uncertainty.

The model incorporated non-linear time trends as a combination of linear and second-order random walk terms. Both components were modelled hierarchically. The age association of diabetes was modelled using a cubic spline to allow non-linear age patterns, which might vary across countries. The coefficients of the splines were modelled hierarchically, and the coefficients are allowed to vary over time to reflect changing time trends in diabetes prevalence and treatment across ages, if supported by data.<sup>2,16</sup>

The model accounted for the possibility that diabetes prevalence and treatment in subnational and community studies might systematically differ from, and have larger variation than, nationally representative samples through the inclusion of fixed-effect and random-effect terms. The fixed effects allowed for systematic differences between subnational or community studies and national studies and allowed for these differences to vary over time. The random effects allowed national data to have a larger influence on the estimates than subnational or community data with similar sample sizes. The model also accounted for urban-rural differences in diabetes prevalence and treatment through fixed effect terms for urban-only and rural-only studies, so that our estimates were for the entire population of each country. These urban and rural effects were weighted by the difference between study-level and country-level urbanisation (i.e., proportion of population living in urban areas) in the year when the study was conducted and were also permitted to vary over time.

As stated in the Methods section of the main paper, we analysed diabetes prevalence for people aged 18+ years and treatment for people aged 30+ years. We performed all analyses separately by sex, because levels and trends in diabetes prevalence and treatment may be different by sex.<sup>1</sup>

#### Detailed model specification

As explained in *Overview*, we modelled the two primary outcomes separately. Here we describe the model in detail for diabetes prevalence. The model for diabetes treatment is described in the section *Model specification for treatment coverage*.

For diabetes prevalence, we used a latent variable  $\alpha_{h,i} = \Phi^{-1}(prevalence_{h,i})$  which is the probit-transformed diabetes prevalence  $(prevalence_{h,i})$  for age group h of study i.  $\Phi$  is the cumulative distribution function of the standard normal distribution and referred to as probit transformation. This specification constrains the primary outcome to be between 0 and 1. We modelled  $\alpha_{h,i}$  with a Gaussian distribution

$$\alpha_{h,i} \sim N(\alpha_{i[i]} + b_{i[i]}t[i] + u_{i[i],t[i]} + \gamma_{i[i],t[i]}(z_h) + X_i\beta + e_i, \tau^2 + v_{h,i}^2), \tag{1}$$

where j, the country in which a study was carried out, and t, the study year, are uniquely determined by the study index i; we denote this determination of j and t on i by j[i] and t[i] respectively. The country-specific intercept and linear time slope from country j are denoted  $a_j$  and  $b_j$  respectively, with  $j \in \{1, ..., J\}$ , where J = 200 is the number of countries in our analysis. We describe the hierarchical model used for the linear component of country time trends, a's and b's, in the section Linear component of country time trends.  $u_{j,t}$  captures smooth nonlinear change over time in country j, as described in the section Nonlinear change.  $\gamma_{j,t}(z_h)$  is the age effect for age group h (with mid-age  $z_h$ ) in year t in country j; it is described in detail in the section Age model. The matrix X contains terms describing whether studies were representative at the national, subnational or community level, and whether they were urbanonly, rural-only, or covered both urban and rural populations, and  $\beta$  contains the associated fixed effects. In addition, a random effect  $e_i$  is estimated for each study i. These study-specific terms are described in the section Study-level terms and study-specific random effects. The variance term  $v_{h,i}^2$  in the model accounts for the uncertainty arising from using regression models to estimate diabetes prevalence based on the primary definition. The variance term

 $\tau^2$  captures the variability not accounted for by the study-specific random effects. These are described in the section *Additional age-by-study variability*. Details on model fitting and convergence are given in the section *Model implementation*. Finally, details on how country-level inference was performed are given in the section *Model inference and post-processing*.

The studies used to fit the model provided data on the number (and hence prevalence) of people with diabetes in a sample. Each study contributed up to 13 data points for each sex, with the exact number depending on the age groups represented in the study. If a study collected data on glycaemic markers like FPG or HbA1c and diabetes medication information for all participants, then the number of people with diabetes  $y_{h,i}$ , from age group h of study i, is assumed to be an observation taken from a binomial distribution with sample size  $n_{h,i}$  and prevalence  $prevalence_{h,i}$ .

$$y_{h,i} \sim Bin(n_{h,i}, prevalence_{h,i}),$$
 (2)

with  $prevalence_{h,i}$  linking to the latent variable  $\alpha_{h,i}$  modelled in equation (1).

As stated in Appendix Text 2, some studies collected data on diabetes medication in all participants but measured glycaemic markers only in a subset of participants. This approach to data collection means that the prevalence of treated diabetes was based on the entire sample, whereas untreated diabetes (i.e., people who were not treated but had elevated levels of FPG or HbA1c) was based on the subset with biomarker data, with the two groups having different sample sizes.

With such survey design, the number of participants with diabetes is the sum of two separately observed quantities: treated diabetes  $(y_{h,i}^{treated})$  and untreated diabetes  $(y_{h,i}^{untreated})$ , obtained from their own respective samples. Each is assumed to be an observation taken from a binomial distribution:

$$y_{h,i}^{treated} \sim Bin(n_{h,i}^{treated}, prevalence_{h,i}^{treated}),$$
 (3)

$$y_{h,i}^{untreated} \sim Bin(n_{h,i}^{untreated}, proportion_{h,i}^{untreated}),$$
 (4)

where  $prevalence_{h,i}^{treated}$  is the prevalence of treated diabetes,  $proportion_{h,i}^{untreated}$  is the proportion of participants who do not use treatment with FPG  $\geq$ 7.0 mmol/L or HbA1c  $\geq$ 6.5%. Their respective sample sizes are  $n_{h,i}^{treated}$ , the number of participants with data on diabetes medication, and  $n_{h,i}^{untreated}$ , the number of untreated participants who had data on glycaemic markers. Prevalence of total diabetes can then be calculated as follows:

$$p_{h,i} = prevalence_{h,i}^{treated} + \left(1 - prevalence_{h,i}^{treated}\right) \times proportion_{h,i}^{untreated}. \tag{5}$$

833 (75%) studies (referred to as group A hereafter) had data available for treated and untreated diabetes. Data from all group A studies were used in equations (3) and (4) in the model, to allow for the aforementioned potentially different sample sizes for treated and untreated diabetes. The remaining 275 (25%) studies (group B) had data for prevalence of total diabetes, but not for treated versus untreated diabetes. Data from group B studies were used in equation (2) in the model.

In equation (3), we used the number of participants who answered the questionnaire about medication use as sample size for treated diabetes  $(n_{h,i}^{treated})$ , and in equation (4) we used those participants who had answered "no" to this question and had a biomarker measurement as sample size for untreated diabetes  $(n_{h,i}^{untreated})$ . In studies with complex survey design, effective sample size is smaller than actual sample size because sampling is done in clusters and/or with stratification. We used actual sample size and did not use effective sample size, which was used in previous analyses, 17 because there is no standard way to calculate effective sample size for the untreated group in equation (4), itself because the untreated group relies on data from a subset of participants (those without treatment and with biomarker measurement).

We used an additional latent variable in calculating  $prevalence_{h,i}^{treated}$  and  $proportion_{h,i}^{untreated}$ , namely the probit-transformed treatment coverage  $\zeta_{h,i} = \Phi^{-1}(treatment_{h,i})$  where  $treatment_{h,i}$  is treatment coverage, our second primary outcome, for age group h of study i. This latent variable allows us to consistently model both of our primary outcomes (prevalence and treatment) as stated in the section Model specification for treatment coverage.  $treatment_{h,i}$  has the following relationship with prevalence of diabetes and treated diabetes:

$$prevalence_{h,i}^{treated} = prevalence_{h,i} \times treatment_{h,i}.$$
 (6)

All prevalence terms can be expressed using the two latent variables  $\alpha_{h,i}$  and  $\zeta_{h,i}$ :

$$prevalence_{h,i} = \Phi(\alpha_{h,i}),$$
 (7)

$$prevalence_{h,i}^{treated} = \Phi(\alpha_{h,i}) \Phi(\zeta_{h,i}), \tag{8}$$

$$proportion_{h,i}^{untreated} = \frac{\Phi(\alpha_{h,i}) - \Phi(\alpha_{h,i}) \Phi(\zeta_{h,i})}{1 - \Phi(\alpha_{h,i}) \Phi(\zeta_{h,i})}.$$
 (9)

In the model for diabetes prevalence, we modelled the latent variable  $\alpha_{h,i}$  as described in equation (1) and detailed in the following sections, and we used a normal prior  $\mathcal{N}(0,1)$  for the latent variable  $\zeta_{h,i}$  which is equivalent to an uninformative uniform distribution on [0, 1] in the prevalence scale.

# Linear component of country time trends

The model had a hierarchical structure, whereby studies were nested in countries, which were nested in regions (indexed by l), which were nested in super-regions (indexed by m), which were all nested in the globe (see Appendix Table 2 for a list of countries and territories in each region, and regions in each super-region). This structure allowed the model to share information across units to a greater degree when data were non-existent or weakly

informative (for example, had a small sample size or were not nationally representative) and, to a lesser extent, in data-rich countries and regions.<sup>18</sup>

The a and b terms are country-specific linear intercepts and time slopes with terms at each level of the hierarchy, denoted by the superscripts c, r, s and g, respectively:

$$a_j = a_j^c + a_{l[j]}^r + a_{m[j]}^s + a^g, (10)$$

$$b_j = b_j^c + b_{l[j]}^r + b_{m[j]}^s + b^g, (11)$$

$$a^{x} \sim N(0, \kappa_{a}^{x}), \tag{12}$$

$$b^{x} \sim N(0, \kappa_{b}^{x}), \tag{13}$$

where  $x \in \{c, r, s\}$ . The  $\kappa$  terms were each assigned a flat prior on the standard deviation scale. We also assigned flat priors to  $a^g$  and  $b^g$ .

## Nonlinear change

The prevalence of diabetes may change nonlinearly over time. We captured smooth nonlinear change in time in country j using the vector  $u_j$ . Just as  $a_j$  and  $b_j$  are each defined as the sum of country, region, super-region and global components, we defined

$$u_j = u_j^c + u_{l[j]}^r + u_{m[j]}^s + u^g. (14)$$

To allow the model to differentiate between the degrees of nonlinearity that exist at the country, region, super-region and global levels, we assigned the four components of each u a discrete second-order Gaussian autoregressive prior. In particular, the vectors  $u_j^c$ ,  $j \in \{1, ..., J\}$ ,  $u_l^r$ ,  $l \in \{1, ..., L\}$ ,  $u_m^s$ ,  $m \in \{1, ..., M\}$ , and  $u^g$ , all of length T, are each given a Gaussian prior with mean zero and precision  $\lambda_c P$ ,  $\lambda_r P$ ,  $\lambda_s P$  and  $\lambda_g P$  respectively, where the scaled precision matrix P in the Gaussian autoregressive prior penalizes first and second differences as follows:

$$P = \begin{bmatrix} 1 & 0 & 0 & \cdots & 0 \\ -2 & 1 & 0 & \cdots & 0 \\ 1 & -2 & 1 & \cdots & 0 \\ 0 & 1 & -2 & \cdots & 0 \\ 0 & 0 & 1 & \cdots & 0 \\ \vdots & \vdots & \vdots & \ddots & \vdots \\ 0 & 0 & 0 & \cdots & 1 \end{bmatrix} \begin{bmatrix} 1 & -2 & 1 & 0 & 0 & \cdots & 0 \\ 0 & 1 & -2 & 1 & 0 & \cdots & 0 \\ 0 & 0 & 1 & -2 & 1 & \cdots & 0 \\ \vdots & \vdots \\ 0 & 0 & 0 & 0 & 0 & \cdots & 1 \end{bmatrix}$$

$$= \begin{bmatrix} 1 & -2 & 1 & 0 & 0 & \cdots & 0 \\ -2 & 5 & -4 & 1 & 0 & \cdots & 0 \\ 1 & -4 & 6 & -4 & 1 & \cdots & 0 \\ 0 & 1 & -4 & 6 & -4 & \cdots & 0 \\ 0 & 0 & 1 & -4 & 6 & \cdots & 0 \\ \vdots & \vdots & \vdots & \vdots & \vdots & \ddots & \vdots \\ 0 & 0 & 0 & 0 & 0 & \cdots & 1 \end{bmatrix}.$$

$$(15)$$

P is multiplied by the estimated precision parameters  $\lambda_c$ ,  $\lambda_r$ ,  $\lambda_s$ , and  $\lambda_g$ , thus upweighting or downweighting the strength of its penalties and ultimately determining the degree of smoothing at each level. For each of the four precision parameters, we used a truncated flat prior on the standard deviation scale  $(1/\sqrt{\lambda})$ . We truncated these priors such that  $\log\lambda \leq 20$  for each of the four  $\lambda's$ . This upper bound is enforced as a computational convenience, whereby models with  $\log\lambda > 20$  are treated as equivalent to a model with  $\log\lambda = 20$ , as they essentially have no extra-linear variability in time. In practice, this upper bound had little effect on the parameter estimates. Furthermore, we ordered the  $\lambda's$  a priori as follows:  $\lambda_c < \lambda_r < \lambda_s < \lambda_g$ . This prior constraint conveys the expectation that the global trend in the prevalence of diabetes has less extra-linear variability than the trend of any given super-region, which has less than those of constituent regions, which in turn has less variability than the trends of constituent countries.

The matrix P has rank T-2, corresponding to a flat, improper prior on the mean and the slope of the  $u_j^c$ 's, the  $u_l^r$ 's, the  $u_m^s$ 's and  $u^g$ , and is not invertible. Thus, we had a proper prior in a reduced-dimension space, with the prior expressed as follows:

$$P(u_j^c | \lambda_c) \propto \lambda_c^{\frac{T-2}{2}} \exp\left\{-\frac{\lambda_c}{2} u_j^{c'} P u_j^c\right\}. \tag{16}$$

Note that if  $u_j^c$  had a non-zero mean, this would introduce non-identifiability with respect to  $a_j^c$ . By the same token,  $b_j^c$  would not be identifiable if  $u_j^c$  had a non-zero time slope, and similarly for the other means and slopes. Thus, to achieve identifiability of the a's, b's, and u's, we constrained the mean and slope of  $u^g$  and each of  $u^s$ ,  $u^r$ , and  $u^c$  to be zero. Enforcing orthogonality between the linear and nonlinear portions of the time trends meant that each can be interpreted independently.

For the countries in which there are observations for at least two different time points, this improper prior will not lead to an improper posterior because the data provide information about the mean and slope. In order to enforce the desired orthogonality between the linear and nonlinear portions of the model, we used the Rue and Held correction. For the countries without data (25 for women and 26 for men), we took the Moore-Penrose pseudoinverse of P, setting to infinity those eigenvalues that correspond to the non-identifiability. This effectively constrained the non-identified portions of the model to zero, as the corresponding variances are set to zero; in this case the Rue and Held correction in a country. In this case, the full conditional precision has rank T-1 because the mean but not the linear trend of  $u_j^c$  is identified by the data. We therefore constrained the linear trend of  $u_j^c$  to zero in this case, by taking the generalized inverse of the full conditional precision. We then constrained the mean of  $u_j^c$  to zero using the one-dimensional version of the Rue and Held correction. Computational details are given in the Appendix of a previous paper.

#### Age model

We sought a smooth function that could characterise gradual changes in diabetes over age, as seen in the data, with parsimonious number of parameters to allow robust estimation. To

achieve this, we modelled age using cubic splines with two knots selected based on epidemiological knowledge about diabetes<sup>24</sup> and statistical considerations.

For age group h with mid-age  $z_h$ , in study i, conducted in year t and country j, the age term is given by

$$\gamma_{j[i],t[i]}(z_h) = \gamma_{1,i}z_h + \gamma_{2,i}z_h^2 + \gamma_{3,i}z_h^3 + \gamma_{4,i}(z_h - k_1)_+^3 + \gamma_{5,i}(z_h - k_2)_+^3, \tag{17}$$

where the two knots were placed ages  $(k_1, k_2) = (50, 65)$  years. To reduce dependence among model parameters, we centred the age variable at 55 years.

Each of the spline coefficients was allowed to vary across countries and was modelled hierarchically, and was further allowed to vary across time, in order to reflect different trends in prevalence across age groups. We modelled spline coefficients consistently with previous analysis,  $^{2,16}$  with the  $k^{th}$  age term coefficients for study i given as follows:

$$\gamma_{k,i} = \psi_k^g + \psi_{k,j[i]}^c + \psi_{k,l[i]}^r + \psi_{k,m[i]}^s + \left(\phi_k^g + \phi_{k,j[i]}^c + \phi_{k,l[i]}^r + \phi_{k,m[i]}^s\right)t[i],$$
 (18)

$$\psi_{k,i[i]}^c \sim N(0, \sigma_{\psi,k,c}^2),\tag{19}$$

$$\psi_{k,l[i]}^r \sim N(0, \sigma_{\psi,k,r}^2),\tag{20}$$

$$\psi_{k,m[i]}^s \sim N(0, \sigma_{\psi,k,s}^2),\tag{21}$$

$$\phi_{k,j[i]}^c \sim N(0, \sigma_{\phi,k,c}^2), \tag{22}$$

$$\phi_{k,l[i]}^r \sim N(0, \sigma_{\phi,k,r}^2), \tag{23}$$

$$\phi_{k,m[i]}^{s} \sim N(0, \sigma_{\phi,k,s}^{2}).$$
 (24)

Here  $\psi^g$ ,  $\psi^c$ ,  $\psi^r$ , and  $\psi^s$  are global, country, region, and super-region intercepts, and  $\phi^g$ ,  $\phi^c$ ,  $\phi^r$ , and  $\phi^s$  are global, country, region and super-region time slope parameters. A flat improper prior was placed on each of the  $\sigma_{\psi}$ 's and  $\sigma_{\phi}$ 's.

Study-level terms and study-specific random effects

The prevalence of diabetes as measured in individual studies may differ from the true unobserved country-year prevalence due to study implementation factors such as those associated with sampling, participation and response, and measurement. We included time-varying offsets (referred to above as fixed effects) to help account for potential systematic differences associated with data sources that are representative of subnational or community populations, and data sources that are representative of urban-only or rural-only populations, through the terms in  $X_i\beta$ :

$$X_{i}\boldsymbol{\beta} = \beta_{1}\mathbf{I}\left\{X_{i}^{cvrg} = subnational\right\} + \beta_{2}\mathbf{I}\left\{X_{i}^{cvrg} = subnational\right\}t[i]$$

$$+ \beta_{3}\mathbf{I}\left\{X_{i}^{cvrg} = community\right\} + \beta_{4}\mathbf{I}\left\{X_{i}^{cvrg} = community\right\}t[i]$$

$$+ \beta_{5}X_{j[i],t[i]}^{c.urb}\mathbf{I}\left\{X_{i}^{s.urb} = rural\right\} + \beta_{6}X_{j[i],t[i]}^{c.urb}\mathbf{I}\left\{X_{i}^{s.urb} = rural\right\}t[i]$$

$$+ \beta_{7}\left(1 - X_{j[i],t[i]}^{c.urb}\right)\mathbf{I}\left\{X_{i}^{s.urb} = urban\right\} + \beta_{8}\left(1 - X_{j[i],t[i]}^{c.urb}\right)\mathbf{I}\left\{X_{i}^{s.urb} = urban\right\}t[i],$$

$$(25)$$

where  $X_i^{cvrg}$  is the indicator for whether the coverage of study i, in country j and year t, is subnational or community,  $X_i^{s.urb}$  is the indicator for whether the study i covered rural-only or urban-only populations, and  $X_{j[i],t[i]}^{c.urb}$  is the percentage of the national population of country j in year t living in urban areas, as obtained from the 2018 revision to the United Nation's World Urbanization Prospects. We note that  $\beta_5$  through  $\beta_8$  are all multiplied by zero for studies which are urban-only in countries where all residents lived in urban areas (e.g., Singapore) and for studies which are rural-only in countries where all residents lived in rural areas (e.g., Tokelau), i.e., in such cases the model does not consider studies classified as urban (respectively rural) to have potential systematic differences from the true underlying prevalence in the country.

Even after accounting for sampling variability, national studies may still not reflect the true prevalence of diabetes in a country with perfect accuracy, due to factors related to response and measurement, and subnational and community studies have even larger variability. We include the study-specific random effect  $e_i$  to allow all age groups from the same study to have an unusually high or an unusually low prevalence, after conditioning on the other terms in the

model. Each  $e_i$  is assigned a Gaussian prior with variance dependent on whether study i is representative at the national, subnational or community level. Random effects from national studies were constrained to have smaller variance  $(v_n)$  than random effects of subnational studies  $(v_s)$ , which were in turn constrained to have smaller variance than community studies  $(v_s)$ .

## Additional age-by-study variability

The additional variance term  $v_{h,i}^2$  in the model accounts for the additional variability arising from our use of regression models to estimate diabetes prevalence based on the primary definition for two types of studies described under *Data cleaning and management* section of Appendix Text 2. The first are the studies that only measured one glycaemic marker, for which we used previously-validated conversion regressions to obtain prevalence based on the primary definition. The second are those that only had data for diabetes prevalence based on a different definition or for mean FPG or mean HbA1c, for which we used regression models to convert to diabetes prevalence based on the primary definition. This additional variance term is fixed for each data point and calculated using a simulation approach. Specifically, we sampled 2,000 draws from the joint distribution of the regression coefficients. We used each draw of regression coefficients to repeatedly calculate diabetes prevalence based on the primary definition. We then calculated  $v_{h,i}^2$  term as the variance of the 2,000 probit-transformed diabetes prevalence, by study, sex and age group. The  $v_{h,i}^2$  term is zero when the primary outcome was calculated directly from the data.

Finally, the age patterns across communities within a given country may differ from the overall age pattern of that country. This within-study variability cannot be captured by the  $e_i$  terms, which are equal across age-specific observations in each study, so we included an additional variance component for each study,  $\tau^2$ .

#### Model specification for treatment coverage

As mentioned above, the parameters in the models for diabetes prevalence and diabetes treatment were estimated independently. In the model for treatment coverage, we modelled the latent variable  $\zeta_{h,i}$ , the probit-transformed treatment coverage, in the same way as we modelled the latent variable  $\alpha_{h,i}$  for diabetes prevalence. We modelled  $\zeta_{h,i}$  with a Gaussian distribution:

$$\zeta_{h,i} \sim N(a_{i[i]} + b_{i[i]}t[i] + u_{i[i],t[i]} + \gamma_i(z_h) + X_i\beta + e_i, \tau^2 + \nu_{h,i}^2), \tag{26}$$

with the components having the same definitions as above. Similar to the model for diabetes prevalence, the additional variance term  $v_{h,i}^2$  accounts for the uncertainty from our use of regression models to estimate diabetes prevalence based on the primary definition which appears in the denominator for treatment coverage. We applied the normal prior  $\mathcal{N}(0,1)$  to the other latent variable  $\alpha_{h,i}$ , with this prior again being equivalent to an uninformative uniform distribution on [0, 1] in the prevalence scale. Only group A studies were used to fit this model because group B studies did not contain data on treatment.

## Model implementation

The model was fitted through a bespoke MCMC sampler coded in R, which uses a combination of Metropolis-Hastings and Gibbs updates.<sup>26</sup> Details of the approach for generating starting values were given in a previous paper.<sup>2</sup>

We had a target of eight converged MCMC chains for generating our estimates, which is twice the recommended minimum number to assess convergence using the Rhat diagnostic. 27,28 We ran ten chains for each outcome (prevalence, treatment) and sex combination, with chains ordered by their seeds. The additional two chains per outcome and sex were run to allow for a small number of the first eight chains to be discarded if mixing was slow. In practice, only five of the 32 chains were slow to converge and were replaced. We did not run more chains because the computational and time cost outweighed the gains, if any, in results. We identified,

through visual inspection of hyperparameter trace plots, a burn-in period of 25,000 iterations for diabetes prevalence, and 100,000 for diabetes treatment because treatment models required more iterations for convergence. We took 50,000 post-burn-in iterations from each of the eight target chains, and combined and thinned to obtain a final sample of 5,000 posterior draws for each outcome. Convergence was confirmed through visual inspection as well as through calculated split-Rhat diagnostic for country-year-age outcomes as implemented in the R package 'rstan' v2.26.15.<sup>27,29</sup> The 97.5<sup>th</sup> quantile of split-Rhat for the two primary outcomes and two sexes ranged from 1.005 to 1.066. 99% of country-year-age combinations for the two primary outcomes and two sexes had split-Rhat <1.05.

## Model inference and post-processing

All inference was done for country-year-age combinations, through combining the a, b, u, and  $\gamma$  terms, and setting  $\beta=e_i=0$ . We set  $\beta=0$  as fixed effects associated with study design are not relevant for country-level inference. We set  $e_i=0$  as random effects arising from imperfections and variations in study design and implementation, and from within-country variability of the primary outcomes, are also not relevant for country-level inference.

Posterior estimates were made in five-year age groups. For presentation, we summarised the age-specific results as age-standardised results. Age-standardisation puts the population for each country-year on the same (standard) age distribution, and hence enables comparisons to be made over time and across countries. Age-standardisation was performed by taking the weighted means of age-sex-specific estimates, using age weights from the WHO standard population.<sup>30</sup> Estimates for regions and the world were calculated as population-weighted averages of the constituent country estimates by sex and age group, using population data obtained from the United Nations' World Population Prospects (2024 revision).<sup>31</sup> Consistent with analysis of hypertension treatment,<sup>16</sup> when calculating age-standardised treatment we also accounted for the age pattern of diabetes prevalence, because the denominator of treatment is only people with diabetes, by multiplying the WHO standard population weights

with age-specific diabetes prevalence in each country and year. The number of adults who had diabetes or untreated diabetes were calculated by multiplying the corresponding age-specific prevalence by the age-specific population by sex, country, and year.

The uncertainties of our estimates, represented by their posterior distributions, include the following sources: uncertainty due to sampling in each data source; uncertainty associated with the variability of national data beyond what is accounted for by sampling; uncertainty associated with subnational and community data, which are more variable than national data; uncertainty associated with using regression equations to estimate the primary outcomes; and uncertainty due to making estimates by country, year, and age when data were missing, scarce or weakly informative. The reported credible intervals (CrI) represent the 2.5th to 97.5th percentiles of the posterior distributions, which contain the true estimates with 95% probability. We obtained the posterior probability (PP) that an estimated change in diabetes prevalence or treatment coverage represented a true increase as the proportion of draws from the posterior distribution that indicated an increase, i.e., a positive change.

**Appendix Text 4.** Decomposition of change in the number of people with untreated diabetes

The number of people with untreated diabetes in 1990 can be written as:

$$N_{1990} = P_{1990} \times DP_{1990} \times (1 - TC_{1990}),$$

where  $N_{1990}$  is the number of people with untreated diabetes in 1990,  $P_{1990}$  is the population in 1990,  $DP_{1990}$  is diabetes prevalence in 1990, and  $TC_{1990}$  is treatment coverage in 1990. Similarly for a later year *year2*:

$$N_{year2} = P_{year2} \times DP_{year2} \times (1 - TC_{year2}).$$

Total change in N from 1990 to year2 is the subtraction of the above two numbers, as below:

Total change in 
$$N = N_{year2} - N_{1990}$$

= 
$$P_{year2} \times DP_{year2} \times (1 - TC_{year2}) - P_{1990} \times DP_{1990} \times (1 - TC_{1990})$$
.

Algebraically, *Total change in N* can be written as the sum of the *Contribution attributed to change in P*, the *Contribution attributed to change in DP*, the *Contribution attributed to change in TC*, and a fourth term that represents the residual change after accounting for the three named contributions. The first three terms are defined as below, with the residual being the difference between the total change and the sum of the first three:

Contribution attributed to change in P

$$= (P_{\text{vear}2} - P_{1990}) \times DP_{1990} \times (1 - TC_{1990}),$$

Contribution attributed to change in DP

$$= P_{1990} \times (DP_{year2} - DP_{1990}) \times (1 - TC_{1990}),$$

Contribution attributed to change in TC

$$= P_{1990} \times DP_{1990} \times [(1 - TC_{year2}) - (1 - TC_{1990})],$$

Residual term

$$= P_{year2} \times DP_{year2} \times (1 - TC_{year2}) - P_{year2} \times DP_{1990} \times (1 - TC_{1990})$$
$$- P_{1990} \times DP_{year2} \times (1 - TC_{1990}) - P_{1990} \times DP_{1990} \times (1 - TC_{year2})$$

$$+ 2 \times P_{1990} \times DP_{1990} \times (1 - TC_{1990}).$$

**Appendix Table 1.** Data sources used in the analysis.

	Country	Data years	Survey/Study name/Citation	Level of representative-	Rural, urban, or		ge as used al analysis		size used I analysis	Glycaemic markers available	Information available on diabetes	Whether me laboratory or by portable	a point-of-care	Notes
				ness	both	Female	Male	Female	Male		treatment	Glucose	HbA1c	
	Afghanistan	2018	STEPS	National	both	18-69	18-69	1,727	1,993	Fasting glucose	Yes	Portable		
	Algeria	2001	Temmar et al., J Hpertens 25:2218-26, 2007	Community	rural	35+	35+	655	561	Fasting glucose	No	Unknown		ļ
	Algeria	2003 2005	STEPS Transition and Health Impact in North Africa	Subnational	both	25-64 35-70	25-64 35-70	2,451 2,770	1,613 2,002	Fasting glucose	Yes	Unknown		<del></del>
	Algeria Algeria	2005	The ISOR (InSulino-resistance in ORan) Study	National Community	both urban	30-64	30-64	408	375	Fasting glucose Fasting glucose	Yes Yes	Portable Lab		<del>                                     </del>
6	Algeria	2016-2017	STEPS	National	both	18-69	18-69	3,690	3,022	Fasting glucose	Yes	Portable		<b></b>
	American Samoa	1994	McGarvey, Pac Health Dialog 8(1):157-62, 2001	National	both	29+	29+	247	165	Fasting glucose	Yes	Lab		
8	American Samoa	2004	STEPS	National	both	25-64	25-64	1,061	945	Fasting glucose	Yes	Portable		
9	Angola	2013-2014	CardioBengo - Population based cardiovascular longitudinal study in Bengo Province, Angola	Community	both	18-65	18-65	1,348	768	Fasting glucose	Yes	Portable		
	Argentina	2004-2005	CArdiovascular Risk factors Multiple Evaluation in Latin America (CARMELA)	Community	urban	25-64	25-64	748	734	Fasting glucose	Yes	Lab		
	Argentina	2011-2012	CESCAS Study	Community	urban	35-74	35-74	2,362	1,576	Fasting glucose	Yes	Lab		
	Argentina	2018	Encuesta Nacional de Factores de Riesgo 2018	National	both	18+	18+	15,406	11,925	Fasting glucose	Yes	Unknown		<b>↓</b>
	Armenia	2016	STEPS ADDROG Busseller	National	both	18-69	18-69	1,582	736	Fasting glucose	Yes	Portable		<del></del>
14 15	Australia Australia	1981 1981	APCSC-Busselton Glatthaar et al., Med J Aust 143:436-40, 1985	Community	urban both	25+ 25+	25+ 25+	701 1,739	608 1,457	Fasting glucose Fasting glucose	No No	Unknown Unknown		<b>—</b>
	Australia	1983	Risk Factor Prevalence Study	National	urban	25-64	25-64	3,811	3,731	Fasting glucose	Yes	Unknown		
17	Australia	1988-1989	Dubbo Study of Australian Elderly	Community	urban	59+	59+	1,222	878	Fasting glucose	Yes	Lab		
	Australia	1999-2000	The Australian Diabetes, Obesity and Lifestyle Study 1999-2000	National	both	25+	25+	6,138	5,043	Fasting glucose, HbA1c	Yes	Lab	Lab	
19	Australia	1999-2003	North West Adelaide Health Study	Community	urban	18+	18+	2,089	1,891	Fasting glucose, HbA1c	No	Lab	Lab	
	Australia	2004-2005	The Australian Diabetes, Obesity and Lifestyle Study 2004-2005	National	both	30+	30+	3,438	2,890	Fasting glucose, HbA1c	Yes	Lab	Lab	
	Australia	2004-2006	North West Adelaide Health Study	Community	urban	20+	20+	1,665	1,498	Fasting glucose, HbA1c	No	Lab	Lab	$\longmapsto$
22	Australia	2008-2010	North West Adelaide Health Study	Community	urban	24+	24+	1,277	1,142	Fasting glucose, HbA1c	No	Lab	Lab	ļ
23	Australia	2012	The Australian Diabetes, Obesity and Lifestyle Study 2012	National	both	37+	37+	2,480	2,029	Fasting glucose, HbA1c	Yes	Lab	Lab	<del></del>
24 25	Australia Austria	2011-2012 1985	National Health Measure Survey VHM&PP Ulmer et al., J Intern Med 261:566-76, 2007	National Subnational	both both	18+ 20+	18+ 20+	4,153 42,176	3,320 32,600	Fasting glucose, HbA1c Fasting glucose	No No	Lab Unknown	Lab	<del>                                     </del>
26	Austria	1991	CINDI survey Vorarlberg/Austria	Subnational	both	25-64	25-64	736	695	Fasting glucose	Yes	Lab		$\vdash$
	Austria	1992	Vorariberg Health Monitoring and Promotion Programme (VHM&PP)	Subnational	both	18+	18+	18,769	14,104	Fasting glucose	No	Lab		
-	Austria	1998	Vorariberg Health Monitoring and Promotion Programme (VHM&PP)	Subnational	both	18+	18+	20,902	16,140	Fasting glucose	No	Lab		
29	Austria	1998-1999	CINDI survey Vorarlberg/Austria	Subnational	both	25-64	25-64	88	86	Fasting glucose	Yes	Lab		
30	Austria	2004	Vorarlberg Health Monitoring and Promotion Programme (VHM&PP)	Subnational	both	18+	18+	23,890	20,159	Fasting glucose	No	Lab		
31	Austria	2010-2012	Austrian Study on Nutritional Status 2012	National	both	18-80	18-80	278	176	Fasting glucose, HbA1c	No	Lab	Lab	
	Azerbaijan	2017	STEPS	National	both	18-69	18-69	1,642	1,136	Fasting glucose	Yes	Portable		
33	Bahamas	2019	STEPS	National	both	18-69	18-69	1,403	932	Fasting glucose	Yes	Portable		<b>↓</b>
34 35	Bahrain Bangladesh	2007 2002	STEPS Hussain et al., Diabet Med 22:931-6, 2005	National Community	both rural	20-64 20+	20-64 20+	906 2,720	863 2,037	Fasting glucose Fasting glucose	No No	Lab Unknown		<del>                                     </del>
	Bangladesh	2002	Hussain et al., Diabet Med 22:931-6, 2005	Community	urban	20+	20+	824	731	Fasting glucose	No	Unknown		<b></b>
-	Bangladesh	2006	Urban Health Survey	Subnational	urban	35-59	35-59	1,272	1,520	Fasting glucose	Yes	Portable		
	Bangladesh	2011	Demographic and Health Survey Bangladesh 2011	National	both	35+	35+	3,572	3,753	Fasting glucose	Yes	Portable		
39	Bangladesh	2011-2012	Chronic Disease Risk Factor Study	Community	rural	18+	18+	427	292	Fasting glucose	Yes	Lab		
40	Bangladesh	2016	Diabetes Mellitus: Action through community Groups or Health Information for better Control of population blood glucose, risk factors, knowledge and care seeking (DMagic)	Subnational	rural	30+	30+	6,414	5,630	Fasting glucose	Yes	Portable		
	Bangladesh	2017-2018	Demographic and Health Survey Bangladesh 2017-2018	National	both	18-49		4,858		Fasting glucose	Yes	Portable		
	Bangladesh	2018	STEPS	National	both	18-69	18-69	3,702	3,247	Fasting glucose	No	Lab		<u> </u>
	Barbados	1992	Foster et al., Ethn Dis 3:404-12, 1993	Community	both	40+ 40-59	40+ 40-59	272	188	Fasting glucose	No No	Unknown	University	<del></del>
44 45	Barbados Barbados	1997-2002 2011-2013	The Barbados Incidence Studies of Eye Diseases II Health of the Nation (HotN)	National National	both both	40-59 25+	40-59 25+	840 741	606 469	HbA1c Fasting glucose, HbA1c	No Yes	Lab	Unknown Portable	<del>                                     </del>
46	Belarus	2016-2017	STEPS	National	both	18-69	18-69	2,897	2,089	Fasting glucose, HDATC	Yes	Portable	i ortable	$\vdash$
	Belarus	2020	STEPS	National	both	18-69	18-69	2,990	2,280	Fasting glucose	Yes	Portable		
48	Belgium	1991-1994	Flemish Study on Environment, Genes and Health Outcomes	Community	rural	26+	26+	412	397	Fasting glucose	Yes	Lab		
-	Belgium	2003	The European Male Ageing Study	Community	both		40+		447	Fasting glucose	Yes	Lab		
	Belgium	2005-2008	Flemish Study on Environment, Genes and Health Outcomes	Community	rural	18+	18+	357	346	Fasting glucose	Yes	Lab		igspace
-	Belgium	2006-2008	Flemish Study on Environment, Genes and Health Outcomes	Community	rural	18+	18+	83	97	Fasting glucose	Yes	Lab		<u> </u>
-	Belgium	2008	The European Male Ageing Study	Community	both	20.	45+	205	372	Fasting glucose	Yes	Lab		$\vdash$
53 54	Belgium Belgium	2009-2013 2010-2015	Flemish Study on Environment, Genes and Health Outcomes Flemish Study on Environment, Genes and Health Outcomes	Community	rural rural	20+ 18+	20+ 18+	335 410	330 389	Fasting glucose Fasting glucose	Yes Yes	Lab Lab		$\vdash \!$
55	Belgium	2010-2019	Belgian Health Examination Survey	National	both	18+	18+	614	557	Fasting glucose, HbA1c	Yes	Lab	Lab	1
-	Belize	2005-2006	CAMDI	National	both	20+	20+	1,021	600	Fasting glucose	Yes	Lab		$\vdash$
	Benin	2007	STEPS	Community	urban	25-64	25-64	1,508	955	Fasting glucose	Yes	Portable		
58	Benin	2008	STEPS	National	both	25-64	25-64	3,391	3,442	Fasting glucose	Yes	Portable		
	Benin	2015	STEP\$	National	both	18-69	18-69	2,547	2,307	Fasting glucose	Yes	Portable		igsquare
-	Bhutan	2007	STEPS	Community	urban	25-74	25-74	1,330	1,132	Fasting glucose	Yes	Lab		<b>  </b>
-	Bhutan	2014	STEPS	National	both	18-69	18-69	1,682	1,072	Fasting glucose	Yes	Portable		$\longmapsto$
	Bhutan Bolivia	2019 2005-2007	STEPS  Cardiovascular and metabolic syndrome risk assessment of Bolivian school children and adolescents - Relationships to obesity, diabetes, income, food intake and physical activity	National National	both both	18-69 18	18-69 18	3,280 139	2,099 144	Fasting glucose Fasting glucose	Yes Yes	Portable Lab		
64	Bosnia and Herzegovina	2012	Non-communicable disease risk factor survey, Federation of B&H	Subnational	rural	18+	18+	1,288	1,201	Fasting glucose	Yes	Portable		

March   September   Septembe		Country	Data years	Survey/Study name/Citation	Level of representative-	Rural, urban, or		e as used I analysis		size used Il analysis	Glycaemic markers available	Information available on diabetes		easured in a y a point-of-care e device	Notes
Processor   Proceedings					ness	both	Female	Male	Female	Male		treatment	Glucose	HbA1c	
Fig.   State   1921   Section of the Principle of the P			2012	Non-communicable disease risk factor survey, Federation of B&H	Subnational	urban	18+	18+	716	594	Fasting glucose	Yes	Portable		
Description   1985   PERSON Review of a Line Study Principle   1985													Portable	<b></b>	
					,								Unknown	<b></b>	
17   1962   170					,								Unknown	<b></b>	
77   1982   2000   Ministry of the Control of the													Lab Lab		
To   Read   Sept.   Colorability   Community   Commu													Unknown		+-
The content of the					,								Lab		+
Annal				,	,								Lab		+1
Part   200-200   Rental of Based   Section					,								Lab		-
Part   March   March					,								Portable		
Part   State   Petrop places   Petrop   Petrop   State   State   Petrop places   Petrop   Petrop   State   Petrop places   Petrop   Petr		Brazil			Community	urban					Fasting glucose		Lab		
Proceedings	77 E	Brazil	2008		Community	rural	18+	18+			Fasting glucose		Lab		
Big   Big	78 E	Brazil	2010	San Pedro	Community	rural							Lab		
Bit Road	79 E	Brazil	2011-2012	The 1993 Pelotas (Brazil) Birth Cohort: 18 years follow-up	Community	urban	18-19	18-19	1,963	1,883	HbA1c	Yes		Lab	
Botal	80 E	Brazil	2010-2015		Community	rural	18+	18+	1,407	1,057	Fasting glucose, HbA1c	Yes	Lab	Lab	
201   201   2014   20	81 E	Brazil	2012-2013	The 1982 Pelotas (Brazil) Birth Cohort: 30 years follow-up	Community	urban	30	30	1,737	1,684	HbA1c	No		Lab	
50   State   2014-2015   Depoptions of Sales de Novege (in Neuron Sales de Alegore   10   10   10   10   10   10   10   1	82 E	Brazil	2013	Pesquisas Nacional de Saude	National	both	18+	18+	32,671	25,141	HbA1c	Yes		Unknown	أتسا
Bit Brief   2014-2015   PhiPropa Admit Control Blody of Spages   Version Study   Community   Communi		Brazil				urban		20+		362			Lab		التسا
80   2014-2015 (Epitroge Anabite Color Distry   Community (use)   26-55   26-55   410   288   Fating glacore   Yes   2015-2016   Petrog Patros District   Yes   Stories   Petrog Patros District   Yes   Stories   Petrog District   Yes   Petrog D		Brazil			Subnational	both			3,146			Yes	Portable		
Broad   2015-2016   The Count Principle of the Count Principle (State Strategy (PACASS)   National   164   164   164   167   530   169   Fastrg glacose, N-AAT   Yes   160   164   164   167														Lab	
Brune   Description   Descri													Lab	Lab	
Description	87 E	Brazil	2015-2016	The Ouro Preto Study	Community	rural	18+	18+	330	186	Fasting glucose	Yes	Lab	<u> </u>	
Description	88 [	Darussalam	2010-2011	National Health And Nutritional Status Survey (NHANSS)	National	both	20-75	20-75	807	675	Fasting glucose, HbA1c	Yes	Lab	Lab	
Sec   Durkmar Faso   2021   STEPS   National   50th   18-69   2.026   1.8-61   664   Fasting glucose   Vec   2020   STEPS   National   50th   18-69	o9 [	Darussalam											Lab		
Section   Sect													Portable		+
Section   Sect													Portable		+
Description   Section													Portable Portable		+
Secure continued   1986   1996   19													Portable		+
													Portable		2
98													Portable		+
Sameron   2000   Defining the relationship between powerly and non-communicable disease burden in Cameroon: Preliminary report. Infobase   Subnational   Urban   18+   18+   2,028   1,641   Fasting glucose   No   100   Cameron   2017   Cameron   2017   Cameron   2017   Preventines and determinants of chronic kidney disease in rural and urban Cameroonians. A cross-sectional study   Community   both   20+   20+   20+   20+   389   520   Fasting glucose   Yes   1012   Cameroon   2014 2015   Cardiovascular risk factors screening in urban and rural areas in the Far-North Region Cameroon   Subnational   both   20+				, , , ,	,								Portable		+-
101   Cameroon   2014   Prevalence and determinants of chronic kidney disease in rural and urban Cameroon   Subnational   2014   204   225   250   Fasting glucose   Yes   102   Cameroon   2014-2015   Cardiovascular risk factor's screening in urban and rural areas in the Far-North Region Cameroon   Subnational   both   204   204   209   209   Fasting glucose   Yes   200   Pasting glucose   Ye				Defining the relationship between poverty and non-communicable disease burden in Cameroon: Preliminary report; Infobase							1		Unknown		
101   Cameroon   2014-2015   Cardiovascular risk factors screening in urban and rural areas in the Far-North Region Cameroon   Subnational   both   20+   20+   369   520   Fasting glucose   Yes   103   Caradia   1993-1995   Kinska et al., Diabetes Cher 24:1787-42; 2011   Caradia   1993-1995   Kinska et al., Diabetes Cher 24:1787-42; 2011   Caradia   1995-1997   Caradian Multicentre Osteoporosis Study (GalMos) - Adult Baseline   Community   rural   18-35	99 C	Cameroon	2007	Cameroon Burden of Diabetes - Second Survey	Subnational	urban	18+	18+	4,581	3,305	Fasting glucose	Yes	Portable		
102   Cameroon   2018   Prevalence and determinants of chronic kidney disease in urban adults' populations of northern Cameroon   Community   urban   20+ 20+ 220   209   Fasting glucose   No   103   Canada   1995-1997   Canadan   1995-1997	100 C	Cameroon	2014	Prevalence and determinants of chronic kidney disease in rural and urban Cameroonians: A cross-sectional study	Community	both	20+	20+	253	183	Fasting glucose	Yes	Unknown		
193   Canada   1993-1995   Kinska et al. Diabetes Care 24.1787-92.2001   Community rural   18-35   18-35   180   136   Fasting glucose   No   104   Canada   1998-1997   Canadam Multicentre Osteoporosis Study (CaMos) - Adult Baseline   Community   both   25+   25+   6.539   2.884   Fasting glucose   Yes   105   Canada   2007-2006   Canadam Multicentre Osteoporosis Study (CaMos) - Adult Year 10 follow-up   Subnational   both   35+   35+   3.593   1.570   Fasting glucose   Yes   106   Canada   2007-2006   Canadam Multicentre Osteoporosis Study (CaMos) - Adult Year 10 follow-up   Subnational   both   18-79   18-79   869   789   Fasting glucose   Hoh 1   Yes   107   Canada   2007-2006   Canadam Health Measures Survey, Cycle 2   National   both   18-79   18-79   955   5825   Fasting glucose, Hoh 1   Yes   108   Canada   2012-2013   Canadam Health Measures Survey, Cycle 3   National   both   18-79   18-79   763   798   Fasting glucose, Hoh 1   Yes   108   Canada   2012-2015   Canadam Health Measures Survey, Cycle 4   National   both   18-79   18-79   763   798   Fasting glucose, Hoh 1   Yes   109   Canadam Health Measures Survey, Cycle 5   National   both   18-79   18-79   769   787   Fasting glucose, Hoh 1   Yes   109   Canadam Health Measures Survey, Cycle 6   National   both   18-79   18-79   759   787   Fasting glucose, Hoh 1   Yes   109   Canadam Health Measures Survey, Cycle 6   National   both   18-79   18-79   759   787   Fasting glucose, Hoh 1   Yes   109   Canadam Health Measures Survey, Cycle 6   National   both   18-79   18-79   750   809   Fasting glucose, Hoh 1   Yes   110   Canada   2018-2017   Canadam Health Measures Survey, Cycle 6   National   both   18-79   18-79   750   809   Fasting glucose, Hoh 1   Yes   110   Canadam Health Measures Survey, Cycle 6   National   both   18-79   18-79   750   809   Fasting glucose, Hoh 1   Yes   110   Canadam Health Measures Survey, Cycle 6   National   both   18-79   18-79   750   809   Fasting glucose, Hoh 1   Yes   110   Canadam Health Measures Survey, Cycle 6	101 C	Cameroon	2014-2015	Cardiovascular risk factors screening in urban and rural areas in the Far-North Region Cameroon	Subnational	both	20+	20+	369	520	Fasting glucose	Yes	Portable	1	
196   Canada   1995-1997   Canadian Multicentre Osteoprosis Study (CaMos) - Adult Baseline   Community   both   25+   25+   6,539   2,884   Fasting glucose   Yes	102 C	Cameroon		Prevalence and determinants of chronic kidney disease in urban adults' populations of northern Cameroon	Community	urban	20+	20+	220	209	Fasting glucose	No	Unknown		
105   Canada   2005-2008   Canadian Multicentre Osteoprosis Study (CaMos) - Adult Year 10 follow-up   Subnational   both   35+   35+   3,993   1,570   Fasting glucose   Yes	103 C	Canada	1993-1995	Kriska et al., Diabetes Care 24:1787-92, 2001	Community	rural	18-35	18-35	180	136	Fasting glucose	No	Lab	ı	
Canada   Control   Canada   Ca													Lab		
107   Canada   2009-2011   Canadian Health Measures Survey, Cycle 2   National   both   18-79   18-79   955   825   Fasting glucose, HbA1c   Yes   108   Canada   2012-2013   Canadian Health Measures Survey, Cycle 3   National   both   18-79   18-79   763   788   Fasting glucose, HbA1c   Yes   109   Canada   2014-2015   Canadian Health Measures Survey, Cycle 4   National   both   18-79   764   736   Fasting glucose, HbA1c   Yes   110   Canada   2016-2017   Canadian Health Measures Survey, Cycle 5   National   both   18-79   18-79   759   787   Fasting glucose, HbA1c   Yes   110   Canada   2018-2019   Canadian Health Measures Survey, Cycle 6   National   both   18-79   18-79   750   809   Fasting glucose, HbA1c   Yes   111   Canada   2018-2019   Canadian Health Measures Survey, Cycle 6   National   both   18-79   18-79   750   809   Fasting glucose, HbA1c   Yes   111   Canada   2018-2019   Canadian Health Measures Survey, Cycle 6   National   both   18-79   18-79   750   809   Fasting glucose, HbA1c   Yes   112   Canadian Health Measures Survey, Cycle 6   National   both   18-79   18-79   750   809   Fasting glucose, HbA1c   Yes   112   Canadian Health Measures Survey, Cycle 6   National   both   18-79   18-79   750   809   Fasting glucose, HbA1c   Yes   112   Canadian Health Measures Survey, Cycle 6   National   both   18-79   18-79   750   809   Fasting glucose, HbA1c   Yes   112   Canadian Health Measures Survey, Cycle 6   National   both   18-79   18-79   750   809   Fasting glucose   Yes   113   Chile   1988   Chilean Health Measures Survey, Cycle 6   National   both   18-79   18-79   750   809   Fasting glucose   No   115   Chile   1988   Chilean Health Measures Survey, Cycle 5   National   both   18+   1											- 00		Lab	ļ	
108   Canada   2012-2013   Canadian Health Measures Survey, Cycle 3   National   both   18-79   18-79   763   798   Fasting glucose, HbA1c   Yes													Lab	Lab	$\perp$
109   Canada   2014-2015   Canadian Health Measures Survey, Cycle 4   National   both   18-79   18-79   764   736   Fasting glucose, HbA1c   Yes				7. 7									Lab	Lab	+
110   Canada   2016-2017   Canadian Health Measures Survey, Cycle 5   National   both   18-79   18-79   759   787   Fasting glucose, HbA1c   Yes													Lab	Lab	+-
111   Canada   2018-2019   Canadian Health Measures Survey, Cycle 6   National   both   18-79   18-79   750   809   Fasting glucose, HbA1c   Yes													Lab	Lab	+
112   Central African Republic   2010   STEPS   Subnational   both   25-64   25-64   1,998   1,882   Fasting glucose   Yes													Lab	Lab Lab	+
112   Republic   2010   STEPS   Subnational   25-64   25-64   1,996   1,882   Fasting glucose   Yes	_		2018-2019		inational	noon					rasting glucose, HbA1c	res	Lab	Lab	+
114   Chile   1992-1993   Miquel et al., Gastroenterology 115(4):937-46, 1998   Community   urban   18+   18+   1,032   657   Fasting glucose   No	112 F	Republic							· ·	·			Portable		
115													Lab		+
116   Chile   2003   Encuesta Nacional de Salud   Sa													Lab		+
117         Chile         2004-2005         CArdiovascular Risk factors Multiple Evaluation in Latin America (CARMELA)         Community         urban         25-64         25-64         872         783         Fasting glucose         Yes           118         Chile         2009-2010         Encuesta Nacional de Salud         National         both         18+         18+         2,791         1,841         Fasting glucose         Yes           119         Chile         2011-2012         CESCAS Study         Community         urban         35-74         399         916         Fasting glucose         Yes           120         Chile         2016-2017         Encuesta Nacional de Salud         National         both         18+         18+         3,796         2,199         Fasting glucose         Yes           121         China         1992-1993         Anzhen OZ Cohort Study         Community         urban         34-65         34-65         2,199         Fasting glucose         No           122         China         1991-1992         Fangshan Cohort Study         Community         urban         34-86         34-86         555         266         Fasting glucose         No           123         China         1995-1996         Hong Kong Cardiovas													Lab		$\vdash$
118         Chile         2009-2010         Encuesta Nacional de Salud         National         both         18+         18+         2,791         1,841         Fasting glucose         Yes           119         Chile         2011-2012         CESCAS Study         Community         urban         35-74         35-74         399         916         Fasting glucose         Yes           120         Chile         2016-2017         Encuesta Nacional de Salud         National         both         18+         18+         3,796         2,199         Fasting glucose         Yes           121         China         1992-1993         Anzhen 02 Cohort Study         Community         urban         34-65         2,112         2,030         Fasting glucose         No           122         China         1991-1992         Fangshan Cohort Study         Community         urban         34-86         34-86         555         266         Fasting glucose         No           123         China         1995-1996         Hong Kong Cardiovascular Risk Factor Prevalence Study 1995-1996         Community         urban         25-74         25-74         1,483         1,412         Fasting glucose         Yes													Lab Lab		+-1
119   Chile   2011-2012   CESCAS Study   Community   urban   35-74   35-74   999   916   Fasting glucose   Yes				, ,									Lab		+
120         Chile         2016-2017         Encuesta Nacional de Salud         National         both         18+         18+         3,796         2,199         Fasting glucose         Yes           121         China         1992-1993         Anzhen 02 Cohort Study         Community         urban         34-65         34-65         2,112         2,030         Fasting glucose         No           122         China         1991-1992         Fangshan Cohort Study         Community         urban         34-86         34-86         555         266         Fasting glucose         No           123         China         1995-1996         Hong Kong Cardiovascular Risk Factor Prevalence Study 1995-1996         Community         urban         25-74         25-74         1,483         1,412         Fasting glucose         Yes													Lab	<del></del>	$\vdash$
121         China         1992-1993         Anzhen 02 Cohort Study         Community         urban         34-65         2,112         2,030         Fasting glucose         No           122         China         1991-1992         Fangshan Cohort Study         Community         urban         34-86         34-86         555         266         Fasting glucose         No           123         China         1995-1996         Hong Kong Cardiovascular Risk Factor Prevalence Study 1995-1996         Community         urban         25-74         25-74         1,483         1,412         Fasting glucose         Yes				,									Lab	<del></del>	$\vdash$
122         China         1991-1992         Fangshan Cohort Study         Community         urban         34-86         34-86         555         266         Fasting glucose         No           123         China         1995-1996         Hong Kong Cardiovascular Risk Factor Prevalence Study 1995-1996         Community         urban         25-74         25-74         1,483         1,412         Fasting glucose         Yes													Unknown	<del></del>	
123         China         1995-1996         Hong Kong Cardiovascular Risk Factor Prevalence Study 1995-1996         Community         urban         25-74         1,483         1,412         Fasting glucose         Yes													Unknown	<del></del>	
													Lab	<del></del>	
124   China   1997   DECODA; DECODA Study Group, Diabetes Care 26:1770-80, 2003   Community   urban   30-89   30-89   2,571   1,577   Fasting glucose   No													Unknown	<sub>[</sub>	
125 China 2000-2001 The International Collaborative Study of Cardiovascular Disease in Asia National both 35-74 35-74 7,828 7,327 Fasting glucose No			2000-2001					35-74					Lab	1	
126 China 2003 Fan et al., World J Gastroenterol 14:2418-24, 2008 Community both 25+ 25+ 7,770 5,529 Fasting glucose No													Unknown	1	

	Country	Data years	Survey/Study name/Citation	Level of representative-	Rural, urban, or		e as used I analysis		size used al analysis	Glycaemic markers available	Information available on diabetes		easured in a y a point-of-care e device	Notes
				ness	both	Female	Male	Female	Male		treatment	Glucose	HbA1c	
	China	2003	Wu, et al., Prev Med 51:412-5, 2010	Community	both	15+	15+	8,260	6,123	Fasting glucose	No	Unknown		3
	China	2004	Tian et al., Diabets Res Clin Pract 84:273-8, 2009	Community	rural	35+	35+	405,011	364,781	Fasting glucose	No	Unknown		<u> </u>
	China China	2005 2004-2005	Zhi et al., Chin Med Sci J 23:249-52, 2008 Xiniiang Children and Adolescent Survey	Community	both urban	18-69 18	18-69 18	10,716 68	9,943 55	Fasting glucose Fasting glucose	No No	Unknown Portable	$\overline{}$	-
	China	2004-2003	Beijing Eye Study	Community	both	45+	45+	1,827	1,393	Fasting glucose	Yes	Lab		<b></b>
	China	2006	Qingdao Diabetes Cohort Study	Community	both	35-74	35-74	2,310	1,536	Fasting glucose, HbA1c	Yes	Lab	Lab	
133 C	China	2006-2007	Fu et al., BMC Public Health 11:862, 2011	Community	rural	18-64	18-64	2,582	1,815	Fasting glucose	No	Lab	i	
	China	2006-2007	Handan Eye Study	Community	rural	30+	30+	3,430	2,998	Fasting glucose	Yes	Lab		<u> </u>
	China	2008-2009	Chinese Longitudinal Healthy Longevity Survey	Subnational	both	65+	65+	753	477	Fasting glucose	No	Lab	<del></del>	4
	China China	2009 2009	China Health and Nutrition Study  Qingdao Diabetes Cohort Study	National Community	both both	18+ 37-78	18+ 37-78	5,163 1,528	4,811 979	Fasting glucose, HbA1c Fasting glucose, HbA1c	Yes Yes	Lab Lab	Lab Lab	5
	China	2009-2010	China National Survey of Chronic Kidney Disease	National	both	18+	18+	23.341	16,611	Fasting glucose, HDATC	Yes	Lab	Lab	<del>                                     </del>
	China	2010	China Noncommunicable Disease Surveillance	National	rural	18+	18+	31,297	27,369	Fasting glucose, HbA1c	Yes	Lab	Lab	
140 C	China	2010	China Noncommunicable Disease Surveillance	National	urban	18+	18+	21,279	16,973	Fasting glucose, HbA1c	Yes	Lab	Lab	
141 C	China	2011	Beijing Eye Study	Community	both	50+	50+	1,963	1,505	Fasting glucose	Yes	Lab		
	China	2011-2012	China Health and Retirement Longitudinal Study (CHARLS), baseline survey	National	both	45+	45+	7,083	6,425	Fasting glucose, HbA1c	Yes	Lab	Lab	<u> </u>
	China	2011-2012	Chinese Longitudinal Healthy Longevity Survey China National Nutrition and Health Survey	Subnational National	both	65+ 18+	65+	1,123 65,491	944 50,394	Fasting glucose	No No	Lab		4
	China China	2010-2013 2014	Chinese Longitudinal Healthy Longevity Survey	Subnational	both both	65+	18+ 65+	1,219	1,022	Fasting glucose Fasting glucose	No No	Lab Lab		4
	China	2015	China Adult Chronic Disease and Nutrition Surveillance	National	both	18+	18+	64,059	56,354	Fasting glucose, HbA1c	No	Lab	Lab	-
	China	2013-2017	Children of 1997 Birth Cohort- Biobank Clinical Follow-up	Community	both	18-20	18-20	250	255	Fasting glucose, HbA1c	No	Lab	Lab	
148 C	China	2015-2017	Henan Rural Cohort	Subnational	rural	18-79	18-79	23,721	15,488	Fasting glucose	Yes	Lab	i	
	China	2015-2016	INTERMAP China Prospective (ICP) Study	Subnational	rural	40-79	40-79	433	348	Fasting glucose	Yes	Unknown		
	China	2016-2018	The FAMILY Cohort	Community	urban	18+	18+	1,118	842	HbA1c	Yes		Lab	ļ
	China	2018	Chinese Longitudinal Healthy Longevity Survey	Subnational	both	65+	65+	7,771	6,191	Fasting glucose	Yes	Lab	<del></del>	6
	Colombia Colombia	2001 2004-2005	CINDI/CARMEN-Bucaramaga; Bautista et al., Eur J Cardiovasc Prev Rehabil 13:769-75, 2006  CArdiovascular Risk factors Multiple Evaluation in Latin America (CARMELA)	Community	urban urban	25-64 25-64	25-64 25-64	1,217 815	622 738	Fasting glucose Fasting glucose	No Yes	Unknown Lab	$\overline{}$	-
	Colombia	2004-2003	Encuesta Nacional de Salud	National	both	18-69	18-69	7,637	5,348	Fasting glucose	Yes	Portable		<b></b>
	Colombia	2010	STEPS	Subnational	urban	18-64	18-64	1,239	924	Fasting glucose	Yes	Portable		
156 C	Colombia	2015	STEPS	Subnational	both	18-64	18-64	1,142	868	Fasting glucose	Yes	Portable	i	
157 C	Colombia	2016	The Survey on Health, Well-Being, and Aging in Latin America and the Caribbean (SABE)	National	both	60+	60+	13,582	10,112	Fasting glucose	Yes	Lab		ļ'
	Comoros	2011	STEPS	National	both	25-64	25-64	3,639	1,584	Fasting glucose	Yes	Unknown		<u> </u>
	Congo	2019 2013-2015	Diabetes prevalence and risk factors STEPS	Community National	rural	19+ 18-64	19+ 18-64	797 630	709 622	Fasting glucose	Yes	Portable Portable		<u> </u>
	Cook Islands Cook Islands	2013-2015	STEPS	National	both both	18-69	18-69	718	692	Fasting glucose Fasting glucose	Yes Yes	Portable		
	Costa Rica	1988	Campos et al., Circulation 85:648-58, 1992	Community	rural	20-65	20-65	123	111	Fasting glucose	No	Unknown		†
163 C	Costa Rica	1988	Campos et al., Circulation 85:648-58, 1992	Community	urban	20-65	20-65	120	111	Fasting glucose	No	Unknown		
164 C	Costa Rica	2000	Ministerio de Salud, 2003	Community	urban	25-64	25-64	636	330	Fasting glucose	No	Unknown		
	Costa Rica	2004	CAMDI	Community	urban	20+	20+	756	390	Fasting glucose	Yes	Lab		ļ
	Costa Rica	2004-2006	Costa Rican Longevity and Healthy Aging Study Pre-1945 Cohort Wave 1	National	both	60+	60+	1,534	1,293	Fasting glucose, HbA1c	Yes	Lab	Lab	<u> </u>
	Costa Rica Costa Rica	2006-2008 2010	Costa Rican Longevity and Healthy Aging Study Pre-1945 Cohort Wave 2  Costa Rican National Cardiovascular Risk Factors Survey, 2010	National National	both both	62+ 20+	62+ 20+	1,286 2,571	1,075 992	Fasting glucose, HbA1c Fasting glucose	Yes Yes	Lab Lab	Lab	<del>                                     </del>
	Costa Rica	2010-2011	Costa Rican Longevity and Healthy Aging Study 1945-1955 Cohort Wave 1	National	both	54-66	54-66	1,688	1,077	HbA1c	Yes	Lab	Lab	
	Costa Rica	2014	Costa Rican National Cardiovascular Risk Factors Survey, 2014	National	both	20+	20+	2,126	954	Fasting glucose	Yes	Lab		
	Costa Rica	2018	Costa Rican National Cardiovascular Risk Factors Survey, 2018	National	both	20+	20+	2,291	1,286	Fasting glucose	Yes	Lab		
	Croatia	2008	Endemic Nephropathy and Arterial hypertension (ENAH)	Subnational	rural	18+	18+	649	502	Fasting glucose	Yes	Lab		ļ
	Croatia	2010	Endemic Nephropathy and Arterial hypertension (ENAH)	Subnational	rural	18+	18+	401	299	Fasting glucose	Yes	Lab		<del>                                     </del>
	Croatia Croatia	2015 2018-2021	Endemic Nephropathy and Arterial hypertension (ENAH) Follow-up Study  Epidemiology of arterial hypertension in Croatia (EH-UH)	Subnational National	rural both	18+	18+ 18+	464 738	225 474	Fasting glucose Fasting glucose	Yes Yes	Lab Lab		<del>                                     </del>
	Cuba	2010-2021	National Survey on Risk Factors and Chronic Diseases (NSRFCD)	National	both	18+	18+	4,036	3,558	Fasting glucose	Yes	Lab		$\vdash$
	Cuba	2010-2011	Noncommunicable disease risk factors in Cienfuegos	Community	urban	18-74	18-74	841	587	Fasting glucose	Yes	Lab		
	Cuba	2018-2020	Encuesta nacional de salud Cuba 2018-2022 (ENS)	National	both	18+	18+	3,219	2,057	Fasting glucose	Yes	Lab		
	Czechia	1981	Machova et al., Cas Lek Cesk 143:90-3, 2004; Site 1	Subnational	rural	25+	25+	11,004	9,189	Fasting glucose	No	Unknown		
	Czechia	1981	Machova et al., Cas Lek Cesk 143:90-3, 2004; Site 2	Subnational	rural	25+	25+	11,004	9,189	Fasting glucose	No	Unknown		<b> </b>
400 0	Czechia Czechia	1997-1998	Czech post-MONICA	National	both	25-64	25-64	1,664	1,529	Fasting glucose	Yes	Lab		<del>                                     </del>
	Ozechia Ozechia	2002-2005	Health, Alcohol and Psychosocial Factors In Eastern Europe	Subnational	urban	25-64 45-70	25-64 45-70	1,755 4,665	1,686 4,060	Fasting glucose Fasting glucose, HbA1c	Yes	Lab	Lab	<del>                                     </del>
	Ozechia	2006-2009	Czech post-MONICA	National	both	25-64	25-64	1,840	1,679	Fasting glucose	Yes	Lab		
	Czechia	2014-2015	European Heath Examination Survey	National	both	25-64	25-64	691	473	HbA1c	Yes		Lab	
	Czechia	2015-2018	Czech post-MONICA	National	both	25-64	25-64	1,338	1,215	Fasting glucose, HbA1c	Yes	Lab	Lab	
	Czechia	2019-2020	European Heath Examination Survey	National	both	25-64	25-64	628	426	HbA1c	Yes	ļ	Lab	<u> </u>
	Czechia	2019-2022	CELSPAC: YA (The Central European Longitudinal Studies of Parents and Children: Young Adults)	Community	both	27-30	27-30	151	138	Fasting glucose	No	Lab		<u> </u>
	Denmark Denmark	2007-2008	The Danish Health Examination Survey 2007-2008  European Youth Heart Study	National Community	both both	18+ 18-28	18+ 18-28	10,334 333	7,089 305	HbA1c Fasting glucose	Yes Yes	Lab	Lab	<del>                                     </del>
	Denmark Denmark	2009-2010	Copenhagen General Population Study	Subnational	urban	20-90	20-90	5,031	4,148	HbA1c	Yes	Lau	Lab	<del>                                     </del>
	Denmark	2017	Copenhagen General Population Study  Copenhagen General Population Study	Subnational	urban	20-90	20-90	4,519	3,282	HbA1c	Yes	<u> </u>	Lab	$\vdash$

	Country	Data years	Survey/Study name/Citation	Level of representative-	Rural, urban, or both		ge as used al analysis		size used Il analysis	Glycaemic markers available	Information available on diabetes		easured in a a point-of-care device	Notes
				ness	Dotti	Female	Male	Female	Male		treatment	Glucose	HbA1c	
	Denmark	2018	Copenhagen General Population Study	Subnational	urban	20-90	20-90	4,018	3,042	HbA1c	Yes		Lab	
194	Denmark	2019	Copenhagen General Population Study	Subnational	urban	20-90	20-90	1,635	1,324	HbA1c	Yes		Lab	<b>↓</b>
	Denmark Denmark	2020-2021 2022-2023	Copenhagen General Population Study Copenhagen General Population Study	Subnational Subnational	urban urban	20-90 20-90	20-90 20-90	1,205 4,047	1,066 3,551	HbA1c HbA1c	Yes Yes		Lab Lab	<del>                                     </del>
	Dominica	2007-2008	STEPS	National	both	18-64	18-64	531	430	Fasting glucose	Yes	Unknown	Lau	$\vdash$
198	Dominican Republic	1996-1998	Estudio factores de riesgo cardiovascular y sindrome metabolico en la Republica Dominicana I (EFRICARD I)	National	both	18-75	18-75	4,097	2,087	Fasting glucose	Yes	Lab		
199	Dominican Republic	2010-2012	Estudio factores de riesgo cardiovascular y sindrome metabolico en la Republica Dominicana II (EFRICARD II)	National	both	18-75	18-75	3,318	1,658	Fasting glucose	Yes	Lab		
200	DR Congo	2007	Diabetes and intermediate hyperglycaemia in Kisantu, DR Congo: a cross-sectional prevalence study	Community	urban	20+	20+	1,197	656	Fasting glucose	Yes	Portable		
201	DR Congo	2016-2017	Prevalence and Risk Factors of CKD in South Kivu, Democratic Republic of Congo: A Large-Scale Population Study	Subnational	both	18+	18+	802	515	Fasting glucose	Yes	Portable		
	Ecuador	2004-2005	CArdiovascular Risk factors Multiple Evaluation in Latin America (CARMELA)	Community	urban	25-64	25-64	825	813	Fasting glucose	Yes	Lab		لـــــــا
	Ecuador	2009-2010	The Survey on Health, Well-Being, and Aging in Latin America and the Caribbean (SABE)	National	both	60+	60+	2,706	2,426	Fasting glucose	Yes	Unknown		igsquare
	Ecuador	2011-2013	Encuesta Nacional de Salud y Nutrición (ENSANUT)	National	both	18-59	18-59	8,068	3,690	Fasting glucose	No	Lab		$\vdash$
	Ecuador Egypt	2018 1995	STEPS Herman et al., Diabet Med 12:1126-31, 1995	National Community	both urban	18-69 20-79	18-69 20-79	2,632 425	1,944 604	Fasting glucose Fasting glucose	Yes No	Portable Unknown		$\vdash \!$
	Egypt	2003-2004	Marzouk et al., Gut 56(8):1105-10, 2007	Community	rural	25+	25+	455	321	Fasting glucose	Yes	Lab		$\vdash$
	Egypt	2005	STEPS	National	both	18-65	18-65	3,923	4,164	Fasting glucose	Yes	Lab		
209	Egypt	2007-2009	Mostafa et al., Gut 59(8):1135-40, 2010	Community	rural	35+	35+	846	642	Fasting glucose	Yes	Lab		
210	Egypt	2011	STEPS STEPS	National	both	18-65	18-65	2,959	1,740	Fasting glucose	Yes	Unknown		
	Egypt	2017	STEPS	National	both	18-69	18-69	3,669	2,094	Fasting glucose	Yes	Portable		
	El Salvador	2004	CAMDI	Community	urban	20+	20+	822	405	Fasting glucose	Yes	Lab		igspace
	El Salvador	2014-2015	Encuesta Nacional de Enfermedades Crónicas (ENECA-ELS)	National	both	20+	20+	2,964	1,703	Fasting glucose	Yes	Lab		igwdown
214	Eritrea	2010	STEPS  CHURCHONIA Inhance and I linear Mad STOUTER CO. 2002	National	both	25-74	25-74	4,309	1,725	Fasting glucose	Yes	Portable		$\vdash$
215 216	Estonia Estonia	1997 2003	SWESTONIA; Johansson et al., J Intern Med 252:551-60, 2002 The European Male Ageing Study	Community Community	urban both	35-55	35-55 40+	133	144 428	Fasting glucose Fasting glucose	No Yes	Unknown Lab		$\vdash$
	Estonia	2003	The European Male Ageing Study  The European Male Ageing Study	Community	both		45+		327	Fasting glucose	Yes	Lab		<del>                                     </del>
218	Eswatini	2014	STEPS	National	both	18-69	18-69	1,925	1,016	Fasting glucose	Yes	Portable		
219	Ethiopia	2015	STEPS	National	both	18-69	18-69	5,127	3,752	Fasting glucose	Yes	Portable		
220	Fiji	1980	National Cardiovascular and Diabetes Survey (NCVDS)	Subnational	both	20+	20+	1,523	1,449	Fasting glucose	Yes	Lab		
221	Fiji	2002	STEPS	National	both	25-64	25-64	2,985	2,055	Fasting glucose	Yes	Portable		
	Fiji	2009	Fiji Eye Health Survey 2009	National	both	40+	40+	787	590	HbA1c	Yes		Portable	
	Fiji	2011	STEPS	National	both	25-64	25-64	1,394	1,096	Fasting glucose	Yes	Portable		<b> </b>
224 225	Finland Finland	1984 1984	Tuomilehto et al., Diabetologia 29:611-5, 1986; Site 1 Tuomilehto et al., Diabetologia 29:611-5, 1986; Site 2	Subnational Subnational	both both		65-84 65-84		296 367	Fasting glucose Fasting glucose	No No	Unknown Unknown		$\vdash \!$
	Finland	1984	Finland, Italy, Netherlands, Elderly (Fine-Finland)	Community	rural		65-84		715	Fasting glucose	Yes	Lab		$\vdash$
	Finland	1986	Young Finns Study 1986	National	rural	18-24	18-24	230	200	Fasting glucose	No	Lab		
	Finland	1986	Young Finns Study 1986	National	urban	18-24	18-24	326	253	Fasting glucose	No	Lab		
229	Finland	1984-1989	Kuopio Ischaemic Heart Disease Risk Factor Study	Subnational	both		42-61		2,682	Fasting glucose	Yes	Lab		
	Finland	1989	Finland, Italy, Netherlands, Elderly (Fine-Finland)	Community	rural		70-89		450	Fasting glucose	No	Lab		
	Finland	1990-1992	Oulu 35 Study	Community	urban	56-57	56-57	327	231	Fasting glucose	Yes	Lab		igsquare
232	Finland	1991-1993	Kuopio Ischaemic Heart Disease Risk Factor Study	Subnational	both	00.04	46-65	050	1,038	Fasting glucose	Yes	Lab		igwdown
233 234	Finland Finland	1997 1996-1998	Northern Finland Birth Cohort 1966 Oulu 35 Study	Community	both urban	30-31 60-63	30-31 60-63	256 346	2,631 244	Fasting glucose Fasting glucose	Yes Yes	Lab Portable + Lab		
235	Finland	1998-2001	Kuopio Ischaemic Heart Disease Risk Factor Study	Subnational	both	53-73	53-73	919	834	Fasting glucose	Yes	Lab		
	Finland	2000-2001	Health 2000 Survey	National	both	30+	30+	3,889	3,159	Fasting glucose, HbA1c	Yes	Lab	Lab	
	Finland	2001	Young Finns Study 2001	National	rural	24-39	24-39	395	344	Fasting glucose	No	Lab		
	Finland	2001	Young Finns Study 2001	National	urban	24-39	24-39	770	660	Fasting glucose	No	Lab		
	Finland	2001-2003	Oulu 45 Study	Community	urban	55-58	55-58	550	428	Fasting glucose	Yes	Lab		igsquare
	Finland	2001-2004	Helsinki Birth Cohort Study	Community	urban	56-69	56-69	1,075	928	Fasting glucose	Yes	Lab		igwdown
241 242	Finland Finland	2005	Mantyselka et al., Rheumatology (Oxford) 47:1235-8, 2008	Community	rural	30-65 60-81	30-65 60-81	250 634	229 1,241	Fasting glucose	No Vos	Unknown		$\vdash$
242	Finland	2005-2008 2007	Kuopio Ischaemic Heart Disease Risk Factor Study Oulu 35 Study	Subnational Community	both urban	71-73	71-73	272	1,241	Fasting glucose Fasting glucose	Yes Yes	Lab Lab		$\vdash$
	Finland	2007	Young Finns Study 2007	National	rural	30-45	30-45	448	384	Fasting glucose	Yes	Lab		
	Finland	2007	Young Finns Study 2007	National	urban	30-45	30-45	728	603	Fasting glucose	Yes	Lab		
246	Finland	2008	Control group for Finnish male former elite athletes	National	both		61+		207	Fasting glucose	Yes	Lab		
	Finland	2007-2008	Savitaipale Study, 10-year Follow-up	Community	rural	51-75	51-75	358	259	Fasting glucose	Yes	Portable		
	Finland	2011	Young Finns Study 2011	National	rural	34-49	34-49	436	368	Fasting glucose, HbA1c	Yes	Lab	Lab	igsquare
	Finland	2011	Young Finns Study 2011	National	urban	34-49	34-49	650	513	Fasting glucose, HbA1c	Yes	Lab	Lab	<b>_</b>
	Finland	2011-2012	Health 2011 Survey	National	both	30+	30+	2,819	2,288	Fasting glucose, HbA1c	Yes	Lab	Lab	
	Finland Finland	2012 2017	Northern Finland Birth Cohort 1966 The FinHealth Survey	Community National	both both	45-47 18+	45-47 18+	2,843 3,627	2,206 3,209	Fasting glucose, HbA1c Fasting glucose, HbA1c	Yes Yes	Lab Lab	Lab Lab	<del>                                     </del>
	Finland	2017	Savitaipale Study, 22-year Follow-up	Community	rural	62-86	62-86	320	243	Fasting glucose, HbA1c	Yes	Lab	Lab	
	Finland	2018-2020	Young Finns Study: Follow-up	National	both	18+	18+	2,953	2,138	Fasting glucose, HbA1c	Yes	Lab	Lab	
	Finland	2019-2020	Northern Finland Birth Cohort 1986	Community	both	33-35	33-35	918	581	Fasting glucose, HbA1c	Yes	Lab	Lab	
	France	1996	POLA Study; Defay et al., Int J Obes Relat Metab Disord 25:512-8, 2001	Community	both	60+	60+	1,419	1,113	Fasting glucose	No	Unknown		
257	France	1996	Asmar et al., J Hypertens 19:1727-32, 2001	Subnational	both	18+	18+	31,416	29,692	Fasting glucose	No	Unknown		

	Country	Data years	Survey/Study name/Citation	Level of representative-	Rural, urban, or		ge as used al analysis		size used al analysis	Glycaemic markers available	Information available on diabetes	laboratory or by	easured in a y a point-of-care le device	Notes
				ness	both	Female	Male	Female	Male		treatment	Glucose	HbA1c	
	France	1999-2001	The Three City Study	Community	urban	65+	65+	5,644	3,650	Fasting glucose	Yes	Lab		<u> </u>
	France	2003-2005	The Three City Study	Community	urban	68+	68+	4,435	2,712	Fasting glucose	Yes	Lab		<u> </u>
	France	2006-2007	Etude Nationale Nutrition Santé	National	both	18-74	18-74	1,379	824	Fasting glucose	Yes	Lab	<del>  </del>	7
	France France	2008-2010	The Three City Study	Community	urban	73+ 40-64	73+ 40-64	3,024 779	1,666 751	Fasting glucose, HbA1c	Yes Yes	Lab Lab	Lab Lab	
	France France	2011-2013	Enquête Littorale Souffle Air Biologie EnvironnemenT (ELISABET) Dunkerque  Enquête Littorale Souffle Air Biologie EnvironnemenT (ELISABET) Lille	Community	urban urban	40-64	40-64	838	751	Fasting glucose, HbA1c Fasting glucose, HbA1c	Yes	Lab	Lab	
	France	2012-2014	Cohorte des consultants des Centres d'examens de santé (CONSTANCES)	National	urban	18-69	18-69	25,538	22,367	Fasting glucose	Yes	Lab	Lab	<del>                                     </del>
	France	2014-2016	L'Etude de Sante sur l'Environnement, la Biosurveillance, l'Activite physique et la Nutrition (Etude Esteban)	National	both	18-74	18-74	1,268	992	Fasting glucose	Yes	Lab		8
	France	2015-2017	Cohorte des consultants des Centres d'examens de santé (CONSTANCES)	National	urban	18-69	18-69	50,369	44,390	Fasting glucose	Yes	Lab		
267	France	2017-2019	Cohorte des consultants des Centres d'examens de santé (CONSTANCES)	Subnational	urban	22-76	22-76	8,552	7,703	Fasting glucose	Yes	Lab		
268	France	2018-2019	Cohorte des consultants des Centres d'examens de santé (CONSTANCES)	National	urban	18-69	18-69	26,699	23,288	Fasting glucose	Yes	Lab		
269	France	2020-2021	Cohorte des consultants des Centres d'examens de santé (CONSTANCES)	Subnational	urban	18-69	18-69	1,635	1,344	Fasting glucose	Yes	Lab		
270	France	2020-2022	Cohorte des consultants des Centres d'examens de santé (CONSTANCES)	Subnational	urban	22-80	22-80	19,004	17,393	Fasting glucose	Yes	Lab		
	French Polynesia	2010	STEPS	National	both	18-64	18-64	1,267	950	Fasting glucose	No	Portable		<u> </u>
	Gambia	2018	The Gambia Micronutrient Survey (GMNS)	National	both	18-49		1,192		HbA1c	No	ļ	Portable	<u> </u>
273	Georgia	2010	STEPS	National	both	18-64	18-64	4,499	1,870	Fasting glucose	Yes	Portable		<u> </u>
274	Georgia	2016	STEPS	National	both	18-69	18-69	2,887	1,271	Fasting glucose	Yes	Portable	+	<b></b>
275	Germany	2000-2002	Epidemiological study of the chances of prevention, early recognition and optimal treatment of chronic diseases in an elderly population (ESTHER)	Subnational	both	50-75	50-75	5,418	4,436	Fasting glucose, HbA1c	Yes	Lab	Lab	<u> </u>
276	Germany	2002	Echinoccoccus Multilocularis and Internal Diseases in Leutkirch	Community	urban	18-65	18-65	964	875	HbA1c	Yes		Lab	
277	Germany	2000-2003	Heinz Nixdorf Recall Study	Subnational	urban	45-75	45-75	2,273	2,223	Fasting glucose, HbA1c	Yes	Lab	Lab	9
278	Germany	2002-2006	Study of Health in Pomerania (SHIP-START-1) 5-year follow-up	Subnational	both	25-85	25-85	1,674	1,568	HbA1c	No	1 - 1	Lab	10 9
279	Germany	2005-2008	Heinz Nixdorf Recall Study  Epidemiological study of the chances of prevention, early recognition and optimal treatment of chronic diseases in an elderly	Subnational	both	50-80	50-80	2,103	2,054	Fasting glucose, HbA1c	Yes	Lab	Lab	9
280	Germany	2008-2011	population (ESTHER)	Subnational	both	58-84	58-84	3,267	2,655	HbA1c	Yes	<u> </u>	Lab	<u> </u>
281	Germany	2008-2011	German Health Interview and Examination Survey for Adults 2008-11 (DEGS1)	National	both	18-79	18-79	3,549	3,280	HbA1c	Yes	+	Lab	
282	Germany	2008-2012	Study of Health in Pomerania (SHIP-START-2) 11-year follow-up	Subnational	both	31-81	31-81	1,198 2,229	1,052 2,096	HbA1c	No	1 - 1	Lab Lab	10 10
283 284	Germany Germany	2008-2012 2011-2014	Study of Health in Pomerania, second cohort (SHIP-TREND-0)  Heinz Nixdorf Recall Study	Subnational Subnational	both both	20-79 56-85	20-79 56-85	1,573	1,504	Fasting glucose, HbA1c Fasting glucose, HbA1c	Yes	Lab	Lab	9
	Germany	2011-2014	Study of Health in Pomerania (SHIP-START-3) 16-year follow-up	Subnational	both	37-87	37-87	908	776	HbA1c	Yes Yes	Lab	Lab	10
	Germany	2014-2010	Study of Health in Pomerania, second cohort (SHIP-TREND-1) 8-year follow-up	Subnational	both	28-90	28-90	1,276	1,202	Fasting glucose, HbA1c	Yes	Lab	Lab	10
	Ghana	2003	Women's Health Study of Accra (WHSA-I)	Community	urban	18+	20 30	3,004	1,202	Fasting glucose	Yes	Lab	Lab	10
	Ghana	2006	STEPS	Community	urban	25+	25+	1,706	887	Fasting glucose	Yes	Portable		<del>                                     </del>
	Ghana	2012-2014	Research on Obesity and Diabetes among African Migrants (RODAM), control group	Subnational	rural	25+	25+	679	432	Fasting glucose, HbA1c	Yes	Lab	Lab	
290	Ghana	2012-2014	Research on Obesity and Diabetes among African Migrants (RODAM), control group	Subnational	urban	25+	25+	1,033	419	Fasting glucose, HbA1c	Yes	Lab	Lab	
291	Ghana	2023	STEPS	National	both	18-69	18-69	3,251	2,022	Fasting glucose	Yes	Portable		2
292	Greece	2001	Karalis et al., BMC Public Health 25:1330-6, 2007	Community	rural	25+	25+	103	91	Fasting glucose	No	Unknown		
	Greece	2001-2002	The ATTICA study	Community	urban	18+	18+	1,525	1,505	Fasting glucose	Yes	Lab		<u> </u>
	Greece	2006	Paliouri Study	Community	rural	65-94	65-94	71	95	Fasting glucose	Yes	Lab		
	Greece	2013-2015	Hellenic National Nutrition and Health Survey (HNNHS)	Subnational	urban	18+	18+	2,265	1,559	Fasting glucose	Yes	Lab		<u> </u>
	Greece	2013-2016	National Survey of Morbidity and Risk Factors (EMENO)	National	both	18+	18+	3,400	2,519	Fasting glucose, HbA1c	Yes	Lab	Lab	<u> </u>
	Greenland	2005-2010	Population Health Survey in Greenland	National	both	18+	18+	1,727	1,356	Fasting glucose, HbA1c	Yes	Lab	Lab	
298 299	Greenland Guatemala	2016-2019	Population Health Survey in Greenland  CAMDI	National Community	both urban	18+ 20+	18+ 20+	1,053 683	841 349	Fasting glucose, HbA1c	Yes No	Lab Lab	Lab	
	Guatemala	2001-2002	The Institute of Nutrition of Central America and Panama Nutrition Supplementation Trial Cohort	Community	both	25-41	25-41	293	242	Fasting glucose Fasting glucose	Yes	Portable		<del>                                     </del>
	Guatemala	2015	STEPS	Subnational	urban	18+	18+	1,551	458	Fasting glucose	Yes	Portable	<del>                                     </del>	<del>                                     </del>
	Guatemala	2015-2017	Nutrition on early childhood and metabolomic and cardiometabolic profile on adulthood (META)	Community	both	37-55	37-55	302	207	Fasting glucose, HbA1c	No	Lab	Lab	†
	Guatemala	2016	Sistema de vigilancia Epidemiológica de Salud y Nutrición (SIVESNU)	National	both	18-49		1,341	1	HbA1c	No		Portable	
	Guatemala	2017-2018	Sistema de vigilancia Epidemiológica de Salud y Nutrición (SIVESNU)	National	both	18-49	1	1,344	1	HbA1c	No	1	Lab	
305	Guatemala	2018-2019	Population-Based Survey of Chronic Kidney Disease in Guatemala	Community	rural	18+	18+	508	263	HbA1c	No		Lab	
306	Guatemala	2018-2019	Sistema de vigilancia Epidemiológica de Salud y Nutrición (SIVESNU)	National	both	18-49		1,489		HbA1c	Yes		Lab	
	Guinea	2009	STEPS	Subnational	both	18-64	18-64	1,131	1,056	Fasting glucose	Yes	Portable		
308	Guyana	2016	STEPS	National	both	18-69	18-69	1,585	1,068	Fasting glucose, HbA1c	Yes	Portable	Lab	<b></b> '
	Haiti	2015-2016	Haiti Health Study (Carrefour)	Community	urban	25-65	25-65	685	474	HbA1c	No	<b></b> '	Portable	<u> </u>
	Haiti	2015-2016	Haiti Health Study (Thomonde)	Community	rural	25-65	25-65	423	258	HbA1c	Yes	<del>                                     </del>	Portable	$\vdash$
	Honduras	2003-2004	CAMDI Cimpu et al. Public Health 440:427 44, 2005	Community	urban	20+	20+	786	435	Fasting glucose	Yes	Lab	₩	<b></b>
	Hungary Hungary	1990-1998 2003	Simay et al., Public Health 119:437-41, 2005 The European Male Ageing Study	Community Community	both both	18+	18+ 40+	13,647	10,651 421	Fasting glucose Fasting glucose	No Yes	Lab Lab	+ +	$\vdash$
	Hungary	2008	The European Male Ageing Study  The European Male Ageing Study	Community	both		45+	+	343	Fasting glucose	Yes	Lab	<del>                                     </del>	$\vdash$
	Iceland	2002-2006	AGES	Subnational	urban	66-96	66-96	3,324	2,437	Fasting glucose	Yes	Lab		
	Iceland	2007-2011	AGESII	Subnational	urban	71-98	71-98	1,934	1,382	Fasting glucose	Yes	Lab		
	India	1988-1989	Ramachandran et al., Diabetes Res Clin Pract 58(1):55-60, 2002	Community	urban	20-74	20-74	408	438	Fasting glucose	No	Portable		
	India	1995	Shobana et al., Diabetes Res Clin Pract 42(3):18186, 1998	Community	urban	20-74	20-74	710	741	Fasting glucose	No	Portable		
	India	1996-1999	Chennai Urban Population Study	Community	urban	20+	20+	698	542	Fasting glucose	Yes	Lab		
	India	1999	DECODA; DECODA Study Group, Diabetes Care 26:1770-80, 2003	Community	urban	30-79	30-79	1,322	1,297	Fasting glucose	No	Unknown		
320	IIIdia											0.110.10111		
	India	2000	Ramachandran et al., Diabet Med 20(3):220-24, 2003	Subnational	urban	20-75	20-75	5,267	4,644	Fasting glucose	No	Portable		

	Country	Data years	Survey/Study name/Citation	Level of representative-	Rural, urban, or		ge as used al analysis		size used I analysis	Glycaemic markers available	Information available on diabetes	Whether me laboratory or by portable	a point-of-care	Notes
				ness	both	Female	Male	Female	Male		treatment	Glucose	HbA1c	
	India	1998-2002	Vellore Birth Cohort	Subnational	both	25-31	25-31	1,055	1,163	Fasting glucose	Yes	Lab		
324	India	1999-2002	New Delhi Birth Cohort	Community	urban	26-33	26-33	636	881	Fasting glucose	Yes	Lab		
325	India	2003	Study in Chennai	Community	rural	20-79	20-79	575	346	Fasting glucose	No	Unknown		
326	India	2001-2004	Chennai Urban Rural Epidemiology Study	Community	urban	20+	20+	1,254	1,096	Fasting glucose, HbA1c	Yes	Lab	Lab	$\vdash$
327	India	2003-2004	Jaipur Heart Watch 3	Community	urban	20-75	20-75	228	212	Fasting glucose	Yes	Lab		$\vdash$
328 329	India India	2006 2005-2006	Ramachandran et al., Diabetes Care 31(5):893-98, 2008  Risk factor profile for chronic non-communicable diseases: Results of a community-based study in Kerala, India	Community Community	both both	20+ 18-64	20+ 18-64	3,745 2,810	3,321 2,615	Fasting glucose Fasting glucose	Yes Yes	Lab Unknown		_
330	India	2005-2006	National Nutrition Monitoring Bureau rural survey	Subnational	rural	20+	20+	13,671	11,901	Fasting glucose	Yes	Portable		$\vdash$
331	India	2006-2008	Central India Eye and Medical Study	Community	rural	30+	30+	2,518	2,191	HbA1c	Yes	Foliable	Lab	-
332	India	2006-2007	Jaipur Heart Watch 4	Community	urban	20-75	20-75	536	502	Fasting glucose	Yes	Lab		
333	India	2006-2008	Kashmiri Young Adults	Subnational	both	20-40	20-40	912	2,120	Fasting glucose	No	Portable		
334	India	2007-2008	Urban population in Hyderabad	Community	urban	20-60	20-60	1,552	1,511	Fasting glucose, HbA1c	Yes	Portable	Portable	
335	India	2006-2009	New Delhi Birth Cohort	Community	urban	33-38	33-38	448	652	Fasting glucose	Yes	Lab		
336	India	2008-2010	ICMR-India Diabetes (INDIAB) Study, Phase I	National	both	20+	20+	6,944	7,110	Fasting glucose, HbA1c	Yes	Portable	Lab	
337	India	2009-2010	Jaipur Heart Watch 5	Community	urban	20-75	20-75	274	429	Fasting glucose	Yes	Lab		
338	India	2010-2012	Centre for cArdiometabolic Risk Reduction in South-Asia (CARRS) - Surveillance Study	Community	urban	20+	20+	6,402	5,867	Fasting glucose, HbA1c	Yes	Lab	Lab	11
339	India	2011-2012	National Nutrition Monitoring Bureau rural survey	National	rural	18+	18+	27,080	21,937	Fasting glucose	Yes	Lab		
340	India	2012-2013	ICMR-India Diabetes (INDIAB) Study, Phase II	Subnational	both	20+	20+	10,866	8,252	Fasting glucose, HbA1c	Yes	Portable	Lab	igsquare
341	India	2012-2014	Jaipur Heart Watch 6	Community	urban	20-75	20-75	353	516	Fasting glucose	No	Lab		igsquare
342	India	2012-2013	Processed and non-processed foods - Rural sample	National	rural	18+	18+	2,093	1,855	Fasting glucose	No	Portable		Щ.
343	India	2014	Annual Health Survey: Clinical, Anthropometric and Bio-chemical	National	both	18+	18+	449,906	407,330	Fasting glucose	No	Lab		
344	India	2012-2015	ICMR-India Diabetes (INDIAB) Study, North East Phase	Subnational	both	20+	20+	16,682	14,260	Fasting glucose, HbA1c	Yes	Portable	Lab	
345	India	2013-2014	Vellore Birth Cohort	Subnational	both	39-44	39-44	499	581	Fasting glucose, HbA1c	Yes	Lab	Lab	
346	India	2014-2015	Control of Hypertension In Rural India (CHIRI) - Rishi Valley	Community	rural	18+	18+	3,551	2,616	Fasting glucose, HbA1c	Yes	Portable	Portable	
347	India	2015-2016	Diet and nutritional status of urban population and prevalence of hypertension	National	urban	18+	18+	53,527	39,397	Fasting glucose	Yes	Portable		-
348	India	2017-2018	National Noncommunicable Disease Monitoring Survey (NNMS)	National	both	18-69	18-69	4,666	5,011	Fasting glucose	Yes	Portable		$\vdash$
349	India	2016-2017	Secular TRends in DiabEtes in India (STRiDE-I) -Change in Prevalence in Ten Years among Urban and Rural Populations in Tamil Nadu	Community	both	20+	20+	5,319	4,527	Fasting glucose	Yes	Portable		
350	India	2017-2018	ICMR-India Diabetes (INDIAB) Study, Phase III	Subnational	both	20+	20+	7,693	7,282	Fasting glucose, HbA1c	Yes	Portable	Lab	
351	India	2016-2019	Vellore Birth Cohort	Subnational	both	43-48	43-48	758	843	Fasting glucose, HbA1c	Yes	Lab	Lab	
352	India	2018-2019	ICMR-India Diabetes (INDIAB) Study, Phase IV	Subnational	both	20+	20+	10,320	9,026	Fasting glucose, HbA1c	Yes	Portable	Lab	
353	India	2019-2020	ICMR-India Diabetes (INDIAB) Study, Phase V STEPS. Mumbai	Subnational	both	20+	20+	6,701	6,670	Fasting glucose, HbA1c	Yes	Portable	Lab	-
354 355	India Indonesia	2021 2001		Community	urban both	18-69	18-69	2,575 2.186	2,601	Fasting glucose	Yes	Portable	<del></del>	$\vdash$
356	Indonesia	2001	STEPS/SURKESNAS  A genetic-ecological study of the risk factors for lifestyle-related diseases in Oceanian populations, Study A	Subnational Community	rural	25+ 18-79	25+ 18-79	103	1,895 100	Fasting glucose Fasting glucose	No Yes	Unknown Lab		$\vdash$
357	Indonesia	2003	A genetic-ecological study of the risk factors for lifestyle-related diseases in Oceanian populations, Study B	Community	rural	18-79	18-79	140	100	Fasting glucose	Yes	Lab		-
358	Indonesia	2005	NCD RFS; Soebardi et al., Acta Med Indones 41:186-90, 2009	Community	urban	25-64	25-64	950	641	Fasting glucose	No	Unknown		_
359	Indonesia	2018	Indonesian Basic Health Survey (RISKESDAS) 2018	National	both	18+	18+	20,614	14,493	Fasting glucose	Yes	Portable		12
360	Iran	1994	Sarraf-Zadegan et al., Acta Cardiol 54:257-63, 1999	Community	urban	20-69	20-69	1,069	1,000	Fasting glucose	No	Unknown		
361	Iran	1999-2000	National Health Survey II	Subnational	both	18+	18+	598	498	Fasting glucose	No	Lab		
362	Iran	2001	Report on the pilot project for community-based primary prevention of the major noncommunicable diseases in Qazvin & Abhar cities 2001; 1; Infobase 101013a2	Community	urban	25+	25+	496	471	Fasting glucose	No	Unknown		
363	Iran	2001	Report on the pilot project for community-based primary prevention of the major noncommunicable diseases in Qazvin & Abhar cities 2001; Infobase 101013a1	Community	urban	25+	25+	495	489	Fasting glucose	No	Unknown		
364	Iran	2001	Isfahan Healthy Heart Programme (IHHP), Arak	Community	rural	19+	19+	1,091	1,028	Fasting glucose	Yes	Lab		
365	Iran	2001	Isfahan Healthy Heart Programme (IHHP), Arak	Community	urban	19+	19+	2,131	2,089	Fasting glucose	Yes	Lab		
366	Iran	2001	Isfahan Healthy Heart Programme (IHHP), Isfahan	Community	rural	19+	19+	238	234	Fasting glucose	Yes	Lab		$\overline{}$
367	Iran	2001	Isfahan Healthy Heart Programme (IHHP), Isfahan	Community	urban	19+	19+	1,932	1,782	Fasting glucose	Yes	Lab		
368	Iran	2001	Isfahan Healthy Heart Programme (IHHP), Najaf Abad	Community	rural	19+	19+	419	409	Fasting glucose	Yes	Lab		
369	Iran	2001	Isfahan Healthy Heart Programme (IHHP), Najaf Abad	Community	urban	19+	19+	578	581	Fasting glucose	Yes	Lab		
370	Iran	2004	Azimi-Nezhad et al., Singapore Med J 49:571-6, 2008	Community	both	15+	15+	1,675	1,585	Fasting glucose	No	Lab		3
371	Iran	2003-2004	Childhood and Adolescence Surveillance and Prevention of Adult Noncommunicable Disease (CASPIAN)	National	both	18	18	368	373	Fasting glucose	Yes	Lab		
372	Iran	2003-2004	The Persian Gulf Healthy Heart Study	Subnational	urban	25-75	25-75	1,974	1,741	Fasting glucose	Yes	Lab		
373	Iran	2005	STEPS	National	both	25-64	25-64	31,782	32,723	Fasting glucose	Yes	Lab		ldot
374	Iran	2007	Isfahan Healthy Heart Programme (IHHP), Arak	Community	rural	19+	19+	1,028	1,030	Fasting glucose	Yes	Lab		ш
375	Iran	2007	Isfahan Healthy Heart Programme (IHHP), Arak	Community	urban	19+	19+	1,366	1,429	Fasting glucose	Yes	Lab		-
	Iran	2007	Isfahan Healthy Heart Programme (IHHP), Isfahan	Community	rural	19+	19+	153	158	Fasting glucose	Yes	Lab		
	Iran	2007	Isfahan Healthy Heart Programme (IHHP), Isfahan	Community	urban	19+	19+	1,435	1,415	Fasting glucose	Yes	Lab		
	Iran	2007	Isfahan Healthy Heart Programme (IHHP), Najaf Abad	Community	rural	19+	19+	254	254	Fasting glucose	Yes	Lab		
	Iran	2007	Isfahan Healthy Heart Programme (IHHP), Najaf Abad	Community	urban	19+	19+	544	498	Fasting glucose	Yes	Lab		$\vdash$
	Iran	2007	Isfahan Healthy Heart Programme (IHHP) Students, Arak Isfahan Healthy Heart Programme (IHHP) Students. Arak	Community	rural	18	18	4	8	Fasting glucose	No No	Lab		
381 382	Iran Iran	2007 2007	Isfahan Healthy Heart Programme (IHHP) Students, Arak Isfahan Healthy Heart Programme (IHHP) Students, Isfahan	Community	urban urban	18 18	18 18	6	17 9	Fasting glucose	No No	Lab Lab		
	Iran	2007	Isranan Healthy Heart Programme (IHHP) Students, Isranan Israhan Healthy Heart Programme (IHHP) Students, Najaf Abad	Community	rural	10	18	· ·	3	Fasting glucose				$\vdash$
	Iran Iran	2007	STEPS - National	National	both	25-64	18 25-64	1,886	1,912	Fasting glucose Fasting glucose	No Yes	Lab Lab	<del></del>	
	Iran	2007	STEPS - Provincial	National	both	25-64	25-64	11,705	11,912	Fasting glucose	Yes	Lab	<del></del>	
	Iran	2007	Childhood and Adolescence Surveillance and Prevention of Adult Noncommunicable Disease (CASPIAN)	National	both	18	18	532	497	Fasting glucose	Yes	Lab	<del></del>	-
386	ııaıl	2009-2010	Control to the Advisage Disease (CASPIAN)	เงสแบทสเ	DUIN	Iδ	18	<b>03</b> 2	497	rasung glucose	168	Lab		

	Country	Data years	Survey/Study name/Citation	Level of representative-	Rural, urban, or		ge as used al analysis		size used I analysis	Glycaemic markers available	Information available on diabetes		easured in a y a point-of-care e device	Notes
				ness	both	Female	Male	Female	Male		treatment	Glucose	HbA1c	
	Iran	2009-2010	The Persian Gulf Healthy Heart Study	Subnational	urban	31-79	31-79	1,014	833	Fasting glucose	Yes	Lab		
388	Iran	2008-2011	Tehran Lipid and Glucose Study	Community	urban	20+	20+	6,000	4,704	Fasting glucose	Yes	Lab	ļ	
389	Iran	2010-2012	Golestan Cohort Study Second Phase	Subnational	rural	43-82	43-82	4,921	4,325	Fasting glucose	Yes	Lab		
390 391	Iran	2010-2012 2011	Golestan Cohort Study Second Phase STEPS	Community National	urban both	43-82 25-69	43-82 25-69	1,061 4,854	1,091 3,313	Fasting glucose Fasting glucose	Yes Yes	Lab Lab		-
392	Iran	2011-2012	Amol county study	Community	rural	18+	18+	1,024	1,685	Fasting glucose	Yes	Lab		
393	Iran	2011-2012	Amol county study	Community	urban	18+	18+	1,476	1,522	Fasting glucose	Yes	Lab		
394	Iran	2012-2013	Tehran City	Community	urban	18-90	18-90	519	395	Fasting glucose	Yes	Lab		
395	Iran	2012-2014	Pars Cohort Study	Community	rural	40-90	40-90	4,988	4,276	Fasting glucose	Yes	Lab	ı	
396	Iran	2012-2013	Zahedan City	Community	urban	18-90	18-90	1,072	1,133	Fasting glucose	Yes	Lab		
397	Iran	2013-2014	Bushehr Elderly Health Program (BEH)	Community	urban	60+	60+	1,545	1,455	Fasting glucose	Yes	Lab	<b></b>	
398 399	Iran	2013-2014 2014-2015	Gilan Eye Study  Childhood and Adolescence Surveillance and Prevention of Adult Noncommunicable Disease (CASPIAN)	Subnational	both	50+ 18	50+ 18	1,159 388	839 330	Fasting glucose, HbA1c	No	Lab Lab	Lab	
400	Iran	2014-2015	The PERSIAN Fasa Cohort Study	National Community	both rural	35-70	35-70	4,584	3,752	Fasting glucose Fasting glucose	Yes Yes	Lab		
401	Iran	2014-2016	The PERSIAN Fasa Cohort Study	Community	urban	35-70	35-70	812	722	Fasting glucose	Yes	Lab		-
402	Iran	2014-2016	The PERSIAN Guilan Cohort Study	Community	rural	35-70	35-70	3,261	2,646	Fasting glucose	Yes	Lab		
403	Iran	2014-2016	The PERSIAN Guilan Cohort Study	Community	urban	35-70	35-70	2,350	2,236	Fasting glucose	Yes	Lab		
404	Iran	2014-2016	The PERSIAN Kermanshah Cohort Study	Community	rural	35-70	35-70	2,198	1,809	Fasting glucose	Yes	Lab		$oxed{oxed}$
405	Iran	2014-2016	The PERSIAN Kermanshah Cohort Study	Community	urban	35-70	35-70	2,976	2,948	Fasting glucose	Yes	Lab	ļ	$\sqcup$
406	Iran	2014-2016	The PERSIAN Kharameh Cohort Study	Community	rural	35-70	35-70	3,861	2,914	Fasting glucose	Yes	Lab	<b></b>	
407 408	Iran Iran	2014-2016 2014-2016	The PERSIAN Kharameh Cohort Study The PERSIAN Tabriz Cohort Study	Community Community	urban rural	35-70 35-70	35-70 35-70	2,001 2,549	1,795 1,974	Fasting glucose Fasting glucose	Yes Yes	Lab Lab		-
409	Iran	2014-2016	The PERSIAN Tabriz Cohort Study	Community	urban	35-70	35-70	5,595	4,671	Fasting glucose	Yes	Lab		_
410	Iran	2015-2017	The PERSIAN Mazandaran Cohort Study	Community	rural	35-70	35-70	1,608	936	Fasting glucose	Yes	Lab		
411	Iran	2015-2017	The PERSIAN Mazandaran Cohort Study	Community	urban	35-70	35-70	4,421	3,179	Fasting glucose	Yes	Lab		
412	Iran	2015-2017	The PERSIAN Rafsanjan Cohort Study	Community	rural	35-70	35-70	1,030	1,578	Fasting glucose	Yes	Lab	ı	
413	Iran	2015-2017	The PERSIAN Rafsanjan Cohort Study	Community	urban	35-70	35-70	4,280	3,593	Fasting glucose	Yes	Lab	<b></b>	
414	Iran	2016	Iran STEPS 2016	National	both	25+	25+	10,015	8,704	Fasting glucose, HbA1c	No	Lab	Lab	
415 416	Iran Iran	2015-2017 2016-2018	The PERSIAN Yazd Cohort Study The PERSIAN Ahvaz Cohort Study	Community Community	urban rural	30-70 35-70	30-70 35-70	4,861 2,282	5,002 1,479	Fasting glucose Fasting glucose	Yes Yes	Lab Lab		-
417	Iran	2016-2018	The PERSIAN Ahvaz Cohort Study	Community	urban	35-70	35-70	3,554	2,512	Fasting glucose	Yes	Lab		-
418	Iran	2016-2018	The PERSIAN BandarKong Cohort Study	Community	rural	35-70	35-70	366	234	Fasting glucose	Yes	Lab		
419	Iran	2016-2018	The PERSIAN BandarKong Cohort Study	Community	urban	35-70	35-70	1,902	1,485	Fasting glucose	Yes	Lab		
420	Iran	2017	Northwest Iran - population based blood sample programme	Community	urban	21+	21+	317	176	Fasting glucose, HbA1c	No	Lab	Lab	
421	Iran	2016-2017	IraPEN Study	Community	rural	30+	30+	3,173	2,919	Fasting glucose	No	Portable	<b></b>	
422	Iran	2016-2017	Iranian Children and Adolescents Psychiatric Disorders (IRCAP) Survey	Subnational	both	18	18	16	15	Fasting glucose	No	Lab	<u>_</u>	
423 424	Iran	2016-2018 2016-2018	The PERSIAN Urmia Cohort Study The PERSIAN Urmia Cohort Study	Community Community	rural urban	35-70 35-70	35-70 35-70	2,379 484	1,821 423	Fasting glucose Fasting glucose	Yes Yes	Lab Lab		
425	Iran	2015-2018	The PERSIAN Zahedan Cohort Study	Community	urban	35-70	35-70	6,024	3,909	Fasting glucose	Yes	Lab		-
426	Iran	2016-2020	The PERSIAN Ardabil Cohort Study	Community	urban	35-70	35-70	11,250	9,552	Fasting glucose	Yes	Lab		
427	Iran	2017-2019	The PERSIAN Dena (Yasouj) Cohort Study	Community	rural	35-70	35-70	920	610	Fasting glucose	Yes	Lab		1
428	Iran	2017-2019	The PERSIAN Dena (Yasouj) Cohort Study	Community	urban	35-70	35-70	1,126	960	Fasting glucose	Yes	Lab		
429	Iran	2018-2019	Prevalence of risk factors for cardiovascular disease among a rural population in eastern Iran	Community	rural	18-69	18-69	146	152	Fasting glucose	Yes	Lab		
430	Iran .	2017-2018	The PERSIAN Kavar Cohort Study	Community	urban	35-70	35-70	2,540	2,418	Fasting glucose	Yes	Lab		
431 432	Iran	2016-2019 2017-2018	The Khuzestan comprehensive health study: A platform for NCDs, blood borne and mental diseases research PERSIAN Elderly Component-Iranian Longitudinal Study on Ageing	Subnational Subnational	both urban	20-65 50-95	20-65 50-95	18,413 3,943	10,216 3,497	Fasting glucose Fasting glucose	No Yes	Lab Lab		
433	Iran	2017-2018	The PERSIAN Sabzevar Cohort Study	Community	urban	35-70	35-70	2,341	1,898	Fasting glucose	Yes	Lab		$\vdash$
434	Iran	2016-2019	The PERSIAN Shahrekord Cohort Study	Community	rural	35-70	35-70	1,791	1,233	Fasting glucose	Yes	Lab		$\vdash$
435	Iran	2016-2019	The PERSIAN Shahrekord Cohort Study	Community	urban	35-70	35-70	3,495	3,499	Fasting glucose	Yes	Lab		
436	Iran	2018-2020	Bushehr Elderly Health program Phase II	Community	urban	50-95	50-95	1,120	835	Fasting glucose, HbA1c	Yes	Lab	Lab	$oxedsymbol{oxedsymbol{oxed}}$
437	Iran .	2018-2019	The PERSIAN Dehgolan (Kordistan) Cohort Study	Community	urban	35-70	35-70	2,206	1,748	Fasting glucose	Yes	Lab		$\vdash$
438	Iran	2020-2021	STEPS	National	both	25+	25+	13,979	11,202	Fasting glucose, HbA1c	Yes	Lab	Lab	$\vdash$
439 440	Iraq Iraq	2006 2015	STEPS STEPS	National National	both both	25-65 18+	25-65 18+	2,379 1,530	1,817 959	Fasting glucose Fasting glucose	No Yes	Lab Lab		$\vdash$
441	Ireland	2006-2007	Survey of Lifestyle, Attitudes and Nutritional in Ireland 2006-2007	National	both	45+	45+	680	526	HbA1c	Yes	Lab	Lab	$\vdash$
442	Ireland	2008-2010	National Adult Nutrition Survey	National	both	18+	18+	440	445	Fasting glucose	No	Lab		
443	Israel	1990-1991	The Jerusalem Longitudinal Cohort Study	Community	urban	69-70	69-70	207	249	Fasting glucose	Yes	Lab		
444	Israel	1997-1998	The Jerusalem Longitudinal Cohort Study	Community	urban	76-77	76-77	454	446	Fasting glucose	Yes	Lab	,	$ldsymbol{ldsymbol{eta}}$
445	Israel	1999-2005	The Israel Glucose Intolerance, Obesity and Hypertension Study (GOH)	National	urban	58-93	58-93	607	536	Fasting glucose	Yes	Lab	<b></b>	$\vdash$
446 447	Israel	2002-2008	The Hadera District Study (HDS) The Javandom Length displ Cahort Study	Subnational	urban	25-78	25-78	551	550	Fasting glucose	Yes	Lab		$\vdash$
	Israel	2005-2006 2010-2011	The Jerusalem Longitudinal Cohort Study The Jerusalem Longitudinal Cohort Study	Community Community	urban urban	83-85 89-92	83-85 89-92	635 347	522 259	Fasting glucose Fasting glucose	Yes Yes	Lab Lab		$\vdash$
	Italy	1982	Verrillo et al, Diabetes Res 2:301-6, 1985	Community	rural	18+	18+	476	410	Fasting glucose	No	Unknown	,—————————————————————————————————————	$\vdash$
	Italy	1983-1985	Gubbio Study	Community	both	18+	18+	2,512	2,040	Fasting glucose	Yes	Lab		
	Italy	1983-1984	Malattie cardiovascolari ATerosclerotiche Istituto Superiore di Sanità (MATISS)	Community	rural	19-69	19-69	1,903	1,700	Fasting glucose	Yes	Lab	<u> </u>	
452	Italy	1986-1987	Malattie cardiovascolari ATerosclerotiche Istituto Superiore di Sanità (MATISS)	Community	rural	19-72	19-72	1,479	1,209	Fasting glucose	Yes	Lab		

150   No.   1500   15	Fasting gluce	787 500 469 785 804	Fasting glucose	treatment			Notes
150   190	Fasting gluce	500 469 785 804	Fasting glucose		Glucose	HbA1c	
150	Fasting gluco Fasting gluco Fasting gluco Fasting gluco Fasting gluco Fasting gluco Fasting gluco	469 785 804		No	Lab		
509   1891   1992   1	Fasting gluco Fasting gluco Fasting gluco Fasting gluco Fasting gluco Fasting gluco	785 804		Yes	Lab		
457   1997	Fasting gluco Fasting gluco Fasting gluco Fasting gluco Fasting gluco	804		Yes	Lab	+	_
450 bigs	Fasting gluce Fasting gluce Fasting gluce Fasting gluce			No	Lab	+	+
1909   1999-1992   Galebo Starly   1992-1993   Galebo Starly   1992-1993   State Longspilled Stayly on Aging   National Longspilled Stayly on Aging   National Longspilled Stayly on Aging   National Longspilled Stayly on Aging (Stayled Longspilled Stayled Longspilled Stayled Longspilled Stayled Longspilled Stayled Longspilled Stayled Longspilled Stayled Longspilled Longspilled Stayled Longspilled Longspilled Stayled Longspilled Longspilled Longspilled Stayled Longspilled Longspill	Fasting gluce Fasting gluce Fasting gluce	423		No	Unknown	+	+
1400   1897   1993   1994   1993   1994   1995	Fasting gluce Fasting gluce			Yes Yes	Lab Lab	+	+
448	Fasting gluce			Yes	Lab	+	+
May   1999-1994   MONICO, Ramora   Month   25-64   2				Yes	Lab	+	+
Hard   1986   Volamo Stay, Masean et al, Biod Press 15 14-0, 2006   Community   1986   1986   September				Yes	Lab	+	+ -
465   Bay   1995   Sunten Study				No	Unknown	+	+
May				Yes	Lab	+	+
Hay				Yes	Lab		1
May	Fasting gluo	1,243	Fasting glucose	Yes	Lab		
490	Fasting gluc	582	Fasting glucose	No	Unknown	1	
A	Fasting gluc	599	Fasting glucose	Yes	Lab	1	
471	Fasting glucose,		Fasting glucose, HbA1	c Yes	Lab	Lab	
473   ally   2000-2001   falsen Longiturien Study on Aging   Amount   Study   2002   Victorian Study, Meason at all Blood Press 15.14-0, 2000   Community   both   CF+   1.377   B13   Fasting glucose   Aging   Study   2002-2003   ROpetito Venico Accion (PROVA)   Submitished   Subm	Fasting gluc		Fasting glucose	Yes	Lab		
474   394				Yes	Lab		$oldsymbol{oldsymbol{\perp}}$
##   ##   ##   ##   ##   ##   ##   #	Fasting gluce	941	Fasting glucose	Yes	Lab		
476   Hay   2011-2004   Scutler et al., Nutr Metab Cardrovasc Dis 19-532-41, 2009   The European Mate Apping Study   Community   Loth   19+   19+   2.265   1.687   Fasting glucose   476   Hay   2011-2007   Community   Loth   20+   26+   1.482   1.191   Fasting glucose   478   Hay   2012-2005   PROpetto Verend Anzieni (PROVA)   Subnational   Loth   68+   68+   1.147   629   Fasting glucose   479   Hay   2002-2005   PROpetto Verend Anzieni (PROVA)   Subnational   Loth   68+   68+   1.147   629   Fasting glucose   479   Hay   2004-2005   Islain Project on the Epidemiology of Alzheime's Disease   National   Loth   65-64   65-84   1.444   1.599   Fasting glucose   481   Hay   2004-2005   Vedemas study   Community   Loth   2004-2005   Vedemas study   Pasting glucose   482   Hay   2005-2007   Robin Study   Subnational   Loth   18+   270   276   Fasting glucose   483   Hay   2005-2007   Robin Study   Subnational   Loth   18+   270   276   Fasting glucose   484   Hay   2005-2007   Robin Study   Subnational   Loth   36+   36+   1.2366   11.495   Fasting glucose   485   Hay   2005-2007   Robin Study   Subnational   Loth   36+   36+   1.2366   11.495   Fasting glucose   486   Hay   2005-2010   Robin Study   Subnational   Loth   36+   36+   1.2366   11.495   Fasting glucose   486   Hay   2005-2010   Robin Study   Subnational   Loth   36+   36+   1.2366   11.495   Fasting glucose   486   Hay   2005-2010   Secretaria Study   Subnational   Loth   36+   36+   1.2366   11.495   Fasting glucose   488   Hay   2005-2010   Secretaria Study   Subnational   Loth   36+   36+   1.2366   11.495   Fasting glucose   488   Hay   2005-2010   Secretaria Study   Subnational   Loth   19+   2005-2010   Fasting glucose   488   Hay   2005-2010   Secretaria Study   Subnational   Loth   19+   2005-2010   Robin Study   Subnational   Loth	Fasting gluce		Fasting glucose	No	Unknown		
476   Italy   2003   The European Male Ageing Study   Community   both   40+   428   Fasting glucose   478   Italy   2001-2007 (Subbs Study   Community   Submational   both   68+   68+   1,147   629   Fasting glucose   478   Italy   2002-2005   PRCgetto Veneto Anziani (PROVA)   Submational   both   68+   68+   1,147   629   Fasting glucose   479   Italy   2005-2005   Balan Project on the Epidemiology of Alzheimer's Disease   National   both   68+   68+   1,147   629   Fasting glucose   400   Italy   2004-2005   Italian Project on the Epidemiology of Alzheimer's Disease   National   both   68+   68+   1,147   629   Fasting glucose   400   Italian Project on the Epidemiology of Alzheimer's Disease   National   both   68+   68+   1,147   1,689   Fasting glucose   481   Italy   2004-2005   Vobarno study   Community   rural   55-74   55-74   113   99   Fasting glucose   482   Italy   2006-2007   Molf-amis Study   Disease   Submational   both   18+   18+   270   216   Fasting glucose   483   Italy   2006-2007   Molf-amis Study   Community   both   45+   45+   344   Fasting glucose   480   Italy   2005-2010   Molf-amis Study   Community   both   45+   45+   344   Fasting glucose   480   Italy   2005-2010   Molf-amis Study   Community   both   25-74   25-74   600   600   Fasting glucose   480   Italy   2005-2010   Molf-amis Study   Community   Land	Fasting gluce		Fasting glucose	Yes	Lab		
477		·		No	Lab		
479   Baby   2002-2005   PROgetto Veneto Anziani (PROVA)   Subnational both   68+   68+   68+   1.147   629   Fasting glucose   149   189   2005   Brusses Study   Community   rural   55-93   55-93   5307   253   Fasting glucose   149   189   2004-2005   189   20	- 00			Yes	Lab		
480   Baly   2005   Stuneck Study   Community   rural   55-93   357   283   Fasting glucose, HbA1				Yes	Lab		
480   Italy				Yes	Lab		
481   Italy   2004-2005   Vobamo study   Community   rural   55-74   55-74   113   99   Fasting glucose				_	Lab	Lab	_
482   Italy   2005-2007   Moli-family Study   Subnational   both   18+   18+   270   216   Fasting glucose				Yes	Lab		+
1				Yes	Lab	+	+
484   Italy   2005-2010   Moli-sani Study   Subnational   both   35+   35+   12,366   11,495   Fasting glucose				Yes	Lab	+	+
485   Italy   2008-2009   Progetto VIP   Community   both   25-74   25-74   600   600   Fasting glucose   486   Italy   2010   Bruneck Study   Bruneck Study   Community   nural   60+   60+   259   225   Fasting glucose   A87   Italy   2009-2010   Factors associated with metabolic syndrome in a mediterranean population: role of caffeinated beverages   Community   both   19-88   1-119   752   Fasting glucose   A88   Italy   2008-2012   Osservatorio Epidemiologico Cardiovascolare/Health Examination Survey (DEC/HES)   National   both   35-80   35-80   4.330   4.389   Fasting glucose   489   Italy   2011-2012   CArdiovascular risk MEtabolic syndrome Liver and Autoimmunity diseases (CA.ME.LLA)   Community   both   18-75   18-75   506   466   Fasting glucose   490   Italy   2011-2012   Vobamo study   Vobamo study   Vobamo study   Community   rural   49-62   49-62   143   107   Fasting glucose   491   Italy   2015   Bruneck Study   Community   Community   rural   65+   65+   169   171   Fasting glucose   Habit   49-92   Italy   2016   The Tyrolean Early Vascular Ageing-study (EVA-Tyrol) - South-Tyrol   Subnational   both   18   2   Fasting glucose   495   Jamaica   1995   MacFarlane-Anderson et al., Metabolism 47:67:-21, 1998   Community   both   25-74   25-74   598   600   Fasting glucose   496   Jamaica   1995   MacFarlane-Anderson et al., Metabolism 47:67:-21, 1998   Community   urban   25+   25+   329   233   Fasting glucose   497   Jamaica   2006-2007   Jamaica Health and Lifestyle Survey   National   both   18-74   18-74   1,249   622   Fasting glucose   498   Jamaica   2007-2000   Jamaica Health and Lifestyle Survey   National   both   18-74   18-74   1,249   622   Fasting glucose   499   Jamaica   2007-2001   Jamaica Health and Lifestyle Survey   National   both   60+   60+   60+   208   156   HbA1c   500   Jamaica   2015-2017   Jamaica   2015-2017   Jamaica Health and Lifestyle Survey   National   both   18-74   18-74   1,249   622   Fasting glucose   490   Jamaica   2016-2017   Jamaica Health and Lifestyle				Yes Yes	Lab Lab	+	+
486   Italy   2010   Bruneck Study   2010   Bruneck Study   2009-2010   Factors associated with metabolic syndrome in a mediterranean population: role of caffeinated beverages   Community   both   19-88   19-88   1,119   752   Fasting glucose   A88   Italy   2009-2012   Osservatorio Epidemiologico Cardiovassolare/Health Examination Survey (OEC/HES)   National   both   35-80   35-80   4,330   4,369   Fasting glucose   489   Italy   2010-2012   CArdiovascular risk MEtabolic syndrome Liver and Autoimmunity diseases (CA.ME.LLA)   Community   both   18-75   18-75   506   466   Fasting glucose   490   Italy   2011-2012   Vobano study   Community   rural   49-62   49-62   143   107   Fasting glucose   HaAlt   490   Italy   2015   Bruneck Study   Subnational   both   18   20   Fasting glucose   HaAlt   490   Italy   2015   Bruneck Study   Community   rural   49-62   49-62   143   107   Fasting glucose   HaAlt   490   Italy   2016   The Tyrotiean Early Vascular Ageing-study (EVA-Tyrol) - South-Tyrol   Subnational   both   18   2   Fasting glucose   HaAlt   493   Italy   2016-2019   Progetto VIP   Community   both   25-74   25-74   598   600   Fasting glucose   494   Italy   2017-2020   Moli-sani Study   Subnational   both   47-94   47-94   1,309   1,081   Fasting glucose   496   Jamaica   1991   Eldemire et al, West Indian Med J 45:82-4, 1996   National   both   47-94   47-94   1,309   1,081   Fasting glucose   497   Jamaica   1995   MacFarlane-Anderson et al, Metabolism 47:617-21, 1998   Community   urban   25+   25+   329   233   Fasting glucose   498   Jamaica   2000-2001   Jamaica Health and Lifestyle Survey   National   both   18-74   18-74   1,869   842   Fasting glucose   499   Jamaica   2007-2008   Jamaica   2010-2008   Jamaica   2010-2009   Jamaica   2010-2008   Jamaica   2010-2009   Jamaica				Yes	Lab	+	+
487   Italy   2009-2010   Factors associated with metabolic syndrome in a mediterranean population: role of calfeinated beverages   Community   both   19-88   19-88   1,119   752   Fasting glucose   489   Italy   2008-2012   Osservatorio Epidemiologico Cardiovascolare/Health Examination Survey (OEC/HES)   National   both   35-80   35-80   4,330   4,399   Fasting glucose   489   Italy   2010-2012   CArdiovascular risk MEtabolic syndrome Liver and Autoimmunity diseases (CA.ME.LLA)   Community   both   18-75   18-75   506   466   Fasting glucose   490   Italy   2011-2012   Vobamo study   Community   rural   49-62   49-62   143   107   Fasting glucose   491   Italy   2015   Bruneck Study   Community   rural   65+   65+   169   171   Fasting glucose   HbA1c   492   Italy   2016   The Tyrolean Early Vascular Ageing-study (EVA-Tyrol) - South-Tyrol   Subnational   both   18   2   Fasting glucose   HbA1c   493   Italy   2018-2019   Progetto VIP   Community   both   25-74   598   600   Fasting glucose   494   Italy   2017-2020   Moli-sani Study   Subnational   both   47-94   47-94   1,309   1,081   Fasting glucose   495   Jamaica   1991   Eldemire et al, West Indian Med J 45:82-4, 1996   National   both   60+   60+   669   649   Fasting glucose   496   Jamaica   1995   MacFarlane-Anderson et al., Metabolism 47:617-21, 1998   Community   urban   25+   25+   329   233   Fasting glucose   498   Jamaica   2000-2007   Jamaica Health and Lifestyle Survey   National   both   18-74   18-74   1,249   622   Fasting glucose   499   Jamaica   2007-2008   Jamaica Health and Lifestyle Survey   National   both   18-74   18-74   1,869   842   Fasting glucose   500   Jamaica   2016-2017   Jamaica Health and Lifestyle Survey   National   both   18-74   18-74   1,869   842   Fasting glucose   500   Jamaica   2016-2017   Jamaica Health and Lifestyle Survey   National   both   18-14   18-74   1,869   593   471   Fasting glucose   470   470   470   470   470   470   470   470   470   470   470   470   470   470   470   470   470   470   470				_	Lab	Lab	+
488         Italy         2008-2012         Osservatorio Epidemiologico Cardiovascolare/Health Examination Survey (OEC/HES)         National         both         35-80         35-80         4,330         4,369         Fasting glucose           489         Italy         2010-2012         CArdiovascular risk MEtabolic syndrome Liver and Autoimmunity diseases (CA.ME.LI.A)         Community         both         18-75         506         466         Fasting glucose           490         Italy         2011-2012         Vobarno study         Community         rural         49-62         49-62         143         107         Fasting glucose           491         Italy         2016         Bruneck Study         Community         rural         65+         65+         169         171         Fasting glucose           492         Italy         2016         The Tyrolean Early Vascular Ageing-study (EVA-Tyrol) - South-Tyrol         Subnational         both         18         2         Fasting glucose, HbA10           493         Italy         2016-2019         Progetto VIP         Community         both         25-74         598         600         Fasting glucose           494         Italy         2017-2020         Moli-san Study         Moli-san Study         Subnational         both <t< td=""><td></td><td></td><td></td><td>No</td><td>Unknown</td><td>Lab</td><td>+ -</td></t<>				No	Unknown	Lab	+ -
489   Italy   2010-2012   CArdiovascular risk MEtabolic syndrome Liver and Autoimmunity diseases (CA.ME.LI.A)   Community   both   18-75   18-75   506   466   Fasting glucose   490   Italy   2011-2012   Vobarno study   Community   rural   49-62   49-62   143   107   Fasting glucose   491   Italy   2015   Bruneck Study   Community   rural   49-62   49-62   143   107   Fasting glucose, HbA10				Yes	Lab	+	+
490   Italy   2011-2012   Vobamo study   Community   rural   49-62   49-62   143   107   Fasting glucose				Yes	Lab	+	+
491   Italy   2015   Bruneck Study   2016   The Tyrolean Early Vascular Ageing-study (EVA-Tyrol) - South-Tyrol   Subnational   both   18   2   Fasting glucose, HbA10   493   Italy   2018-2019   Progetto VIP   Community   both   25-74   25-74   598   600   Fasting glucose   494   Italy   2017-2020   Moli-sani Study   Subnational   both   47-94   1,309   1,081   Fasting glucose   495   Jamaica   1991   Eldemire et al, West Indian Med J 45:82-4, 1996   National   both   60+   60+   669   649   Fasting glucose   496   Jamaica   1995   MacFarlane-Anderson et al., Metabolism 47:617-21, 1998   Community   urban   25+   25+   329   233   Fasting glucose   498   Jamaica   2000-2001   Jamaica Health and Lifestyle Survey   Subnational   both   18-74   18-74   1,249   622   Fasting glucose   498   Jamaica   2000-2007   Jamaica Youth Risk and Resiliency Behaviour Survey 2006   National   both   18-19   18-19   192   152   Fasting glucose   499   Jamaica   2007-2008   Jamaica Health and Lifestyle Survey   National   both   18-74   18-74   1,869   842   Fasting glucose   500   Jamaica   2012   Older Persons in Jamaica 2012   National   both   50+				Yes	Lab	+	+
492   Italy   2016   The Tyrolean Early Vascular Ageing-study (EVA-Tyrol) - South-Tyrol   Subnational   both   18   2   Fasting glucose, HbA1c   493   Italy   2018-2019   Progetto VIP   Community   both   25-74   598   600   Fasting glucose   494   Italy   2017-2020   Moli-sani Study   500   Fasting glucose   495   Jamaica   1991   Eldemire et al, West Indian Med J 45:82-4, 1996   National   both   47-94   47-94   1,309   1,081   Fasting glucose   496   Jamaica   1995   MacFarlane-Anderson et al., Metabolism 47:617-21, 1998   Community   urban   25+   25+   329   233   Fasting glucose   497   Jamaica   2000-2001   Jamaica   Halith and Lifestyle Survey   National   both   18-74   18-74   1,249   622   Fasting glucose   498   Jamaica   2006-2007   Jamaica Youth Risk and Resiliency Behaviour Survey 2006   National   both   18-74   18-74   1,869   842   Fasting glucose   499   Jamaica   2007-2008   Jamaica Health and Lifestyle Survey   National   both   18-74   18-74   1,869   842   Fasting glucose   500   Jamaica   2012   Older Persons in Jamaica 2012   National   both   18-74   18-74   1,869   842   Fasting glucose   500   Jamaica   2016-2017   Jamaica   4016   Survey   National   both   18-74   18-74   1,869   842   Fasting glucose   500   Jamaica   2016-2017   Jamaica   4016   Survey   National   both   18-74   18-74   1,869   842   Fasting glucose   501   Jamaica   2016-2017   Jamaica   4016   Survey   National   both   504   504   505   471   505   505   505   Japan   1985-1986   Akabane Study   Community   Urban   40-69   40-69   593   471   Fasting glucose   1000					Lab	Lab	
494         Italy         2017-2020         Moli-sani Study         Subnational         both         47-94         47-94         1,309         1,081         Fasting glucose           495         Jamaica         1991         Eldemire et al, West Indian Med J 45:82-4, 1996         National         both         60+         60+         669         649         Fasting glucose           496         Jamaica         1995         MacFarlane-Anderson et al., Metabolism 47:617-21, 1998         Community         urban         25+         329         233         Fasting glucose           497         Jamaica         2000-2001         Jamaica Health and Lifestyle Survey         National         both         18-74         1,249         622         Fasting glucose           498         Jamaica         2006-2007         Jamaica Youth Risk and Resiliency Behaviour Survey 2006         National         both         18-19         18-19         192         152         Fasting glucose           499         Jamaica         2007-2008         Jamaica Health and Lifestyle Survey         National         both         18-74         1,869         842         Fasting glucose           500         Jamaica         2012         Older Persons in Jamaica 2012         National         both         60+ <t< td=""><td></td><td></td><td></td><td></td><td>Lab</td><td>Lab</td><td>1</td></t<>					Lab	Lab	1
495         Jamaica         1991         Eldemire et al, West Indian Med J 45:82-4, 1996         National         both         60+         60+         669         649         Fasting glucose           496         Jamaica         1995         MacFarlane-Anderson et al., Metabolism 47:617-21, 1998         Community         urban         25+         329         233         Fasting glucose           497         Jamaica         2000-2001         Jamaica Health and Lifestyle Survey         National         both         18-74         11,249         622         Fasting glucose           498         Jamaica         2006-2007         Jamaica Youth Risk and Resiliency Behaviour Survey 2006         National         both         18-19         18-19         192         152         Fasting glucose           499         Jamaica         2007-2008         Jamaica Health and Lifestyle Survey         National         both         18-74         18-74         1,869         842         Fasting glucose           500         Jamaica         2012         Older Persons in Jamaica 2012         National         both         60+         60+         208         156         HbA1c           501         Jamaica         2016-2017         Jamaica Health and Lifestyle Survey         National         both				Yes	Lab		
496         Jamaica         1995         MacFarlane-Anderson et al., Metabolism 47:617-21, 1998         Community         urban         25+         25+         329         233         Fasting glucose           497         Jamaica         2000-2001         Jamaica Health and Lifestyle Survey         National         both         18-74         18-74         1,249         622         Fasting glucose           498         Jamaica         2006-2007         Jamaica Youth Risk and Resiliency Behaviour Survey 2006         National         both         18-79         18-19         192         152         Fasting glucose           499         Jamaica         2007-2008         Jamaica Health and Lifestyle Survey         National         both         18-74         1,869         842         Fasting glucose           500         Jamaica         2012         Older Persons in Jamaica 2012         National         both         60+         60+         208         156         HbA1c           501         Jamaica         2016-2017         Jamaica Health and Lifestyle Survey         National         both         60+         208         156         HbA1c           501         Jamaica         2016-2017         Jamaica Health and Lifestyle Survey         National         both         18+	Fasting gluo	1,081	Fasting glucose	Yes	Lab		
497         Jamaica         2000-2001         Jamaica Health and Lifestyle Survey         National         both         18-74         18-74         1,249         622         Fasting glucose           498         Jamaica         2006-2007         Jamaica Youth Risk and Resiliency Behaviour Survey 2006         National         both         18-79         18-19         192         152         Fasting glucose           499         Jamaica         2007-2008         Jamaica Health and Lifestyle Survey         National         both         18-74         1,869         842         Fasting glucose           500         Jamaica         2012         Older Persons in Jamaica 2012         National         both         60+         60+         208         156         HbA1c           501         Jamaica         2016-2017         Jamaica Health and Lifestyle Survey         National         both         60+         60+         208         156         HbA1c           501         Jamaica         2016-2017         Jamaica Health and Lifestyle Survey         National         both         18+         18+         1,633         1,026         Fasting glucose, HbA1c           502         Japan         1985-1986         Akabane Study         Community         urban         40-69	Fasting gluc	649	Fasting glucose	No	Unknown		
498         Jamaica         2006-2007         Jamaica Youth Risk and Resiliency Behaviour Survey 2006         National         both         18-19         18-19         192         152         Fasting glucose           499         Jamaica         2007-2008         Jamaica Health and Lifestyle Survey         National         both         18-74         1.869         842         Fasting glucose           500         Jamaica         2012         Older Persons in Jamaica 2012         National         both         60+         208         156         HbA1c           501         Jamaica         2016-2017         Jamaica Health and Lifestyle Survey         National         both         18+         18+         1,633         1,026         Fasting glucose, HbA1c           502         Japan         1985-1986         Akabane Study         Community         urban         40-69         593         471         Fasting glucose	Fasting gluce	233	Fasting glucose	No	Unknown		
499         Jamaica         2007-2008         Jamaica Health and Lifestyle Survey         National         both         18-74         1.869         842         Fasting glucose           500         Jamaica         2012         Older Persons in Jamaica 2012         National         both         60+         60+         208         156         HbA1c           501         Jamaica         2016-2017         Jamaica Health and Lifestyle Survey         National         both         18+         18+         1,633         1,026         Fasting glucose, HbA1c           502         Japan         1985-1986         Akabane Study         Community         urban         40-69         40-69         593         471         Fasting glucose				Yes	Portable		
500         Jamaica         2012         Older Persons in Jamaica 2012         National         both         60+         60+         208         156         HbA1c           501         Jamaica         2016-2017         Jamaica Health and Lifestyle Survey         National         both         18+         18+         1,633         1,026         Fasting glucose, HbA1c           502         Japan         1985-1986         Akabane Study         Community         urban         40-69         40-69         593         471         Fasting glucose				Yes	Portable		
501         Jamaica         2016-2017         Jamaica Health and Lifestyle Survey         National         both         18+         18+         1,633         1,026         Fasting glucose, HbA1t           502         Japan         1985-1986         Akabane Study         Community         urban         40-69         40-69         593         471         Fasting glucose				Yes	Portable		$\bot$
502         Japan         1985-1986         Akabane Study         Community         urban         40-69         49-69         593         471         Fasting glucose				Yes	<b>_</b>	Lab	13
					Portable	Portable	+
503   Japan   1987   Konan i own Study   Community   rural   20-79   87   69   Fasting glucose				No	Unknown	+	+
				No.	Unknown	+	+
504 Japan 1988 Konan Town Study Community rural 20-79 20-79 85 76 Fasting glucose				No	Unknown	+	+
505         Japan         1988         The Hisayama Study         Community         rural         40+         40+         1,574         1,162         Fasting glucose           506         Japan         1989         Konan Town Study         Community         rural         20-79         20-79         63         59         Fasting glucose				Yes No	Lab Unknown	+	+
500         Japan         1989         Notinal Town Study         Community         Tural         20-79         63         99         Fasting glucose           507         Japan         1989         National Nutrition Survey         National both         30+         1,613         1,377         Fasting glucose           National Nutrition Survey         National Nutrition Survey         National Nutrition Survey         30+         30+         1,613         1,377         Fasting glucose				No	Lab	+	+
307   Japan   1989   INSTITUTION SURVEY   NATIONAL VITTOR SURVEY   1,613   1,377   Fasting glucose   1980   Konan Town Study   20-79   20-79   58   30   Fasting glucose   30-79   20-79   20-79   3				No No	Unknown	+	+-1
509 Japan 1990 National Voluntinos Survey and National Cadiovascular Survey National both 30+ 30+ 1,615 1,517 Fasting glucose				No	Lab	+	+-1
1992   Value   1992				No	Unknown	+	+
511 Japan 1991 Konan Town Study Committed United Study Committed Uni				No	Unknown	+	+
512 Japan 1991 National Nutrition Survey National both 30+ 1,523 1,444 Fasting glucose				No	Lab	1	+
513 Japan 1992 Konan Town Study Community rural 20-79 52 55 Fasting glucose				No	Unknown	1	+
514 Japan 1992 National Nutrition Survey National both 30+ 1,445 1,332 Fasting glucose				No	Lab	1	
515 Japan 1993 Konan Town Study Community rural 20-79 20-79 65 54 Fasting glucose				No	Unknown	1	
516 Japan 1993 National Nutrition Survey National both 30+ 30+ 1,360 1,251 Fasting glucose				No	Lab	1	
517         Japan         1994         Konan Town Study         Community         rural         20-79         20-79         59         42         Fasting glucose				No	Unknown		
518 Japan 1994 National Nutrition Survey National both 20-59 20-59 1,112 970 Fasting glucose				No	Lab	1	

1.00		easured in a by a point-of-car le device	laboratory or by	Information available on diabetes	Glycaemic markers available	size used I analysis	Sample for globa		Age rang for globa	Rural, urban, or	Level of representative-	Data years Survey/Study name/Citation	Country Data years	
Section   Part   Section   Section	11c	HbA1c	Glucose	treatment		Male	Female	Male	Female	both	ness			
1821   1822   1822   1823   1824   1825											,			
Section   1989		ļ										,		
1906   1906   1906   1907	$\!\!\!\!+\!\!\!\!-$	<b></b>												
Sect   Control   Control		<del> </del>										,		
Section   1985   1987														
1982   1982	+-													
200   1900   1	+-													
200   1900   1	ah.	Lah	Lab											
200   Section   201   Sectio		Lab	Lah											
1906   1906		†												
Sect	ab	Lab									,			
Section   Process										urban				
			Lab	No	Fasting glucose	968	1,088	20+	20+	both	National	2001 National Nutrition Survey	ın 2001	532
Sept	ıb	Lab		No	HbA1c	211	185	74	74	urban	Community	2002 Niigata Study	ın 2002	533
Section			Lab	No	Fasting glucose	952	1,072	20+	20+	both	National	2002 National Nutrition Survey	ın 2002	534
Section   Sect	ıb	Lab	Lab	Yes	Fasting glucose, HbA1c	1,391			40+	rural	Community	2002-2003 The Hisayama Study	n 2002-2003	535
Separa   2004   Pages Bally   2006   Pages Bally			Lab	Yes						both				
Separate   2000   Region Broady   Region Delay			<u> </u>								,	,		
Separate   1987   1989   198	b	Lab	Lab	Yes	Fasting glucose, HbA1c	1,879	2,700	20+	20+	both		2004 National Health and Nutrition Survey	ın 2004	538
Section   Depth   De			ļ									The second secon		
Section   Property														
254   Speen   2008   Noticed Seath and National Search   2014   1920   2015   1920			Lab											
			Lab											
Sept   2007   Najeun South   1907   Najeun South   1907														
			Lab											
Sept   2008   Solicons   Heach and Nation Strevy   National   Solit   2017   2029   2,149   Failing pulsone (NATC   New   Lab   Lab   1,200   2009   National Heach and Nation Strevy   National   Solit   1,200   2009   National Heach and Nation Strevy   National   Solit   1,200   2009   National Heach and Nation Strevy   National   Solit   1,200   2009   2,255   2,2018   Failing pulsone (NATC   New   Lab   Lab   1,200   2009   National Heach and Nation Strevy   National   Solit   1,200   2009   2,255   2,2018   Failing pulsone (NATC   New   Lab   Lab   1,200   2009   2,2011   National Head and National Street   National   Solit   1,200   2,2011   National Head and National Street   National   Solit   1,200   2,2011   National Head and National Street   National   Solit   1,200   2,2011   National Head and National Street   National Head and National Street   National Head and National Head and National Street   National Head and National Street   National Head and Nati	d	Lab									,	,		
	-	<del></del>										, , , ,		
			Lab									,		
			Lob											
			Lab											
			Lab											
												,		
See   Depart   2014   National Health and Allurinon Survey   National Health and N														
											,	· · ·		
	ıb	Lab		Yes	HbA1c	5,819	7,918	20+	20+	both	National	2016 National Health and Nutrition Survey	ın 2016	560
563   Japan   2017   The Tokyo Health Service Association Database   Community urban   20+   20+   14,915   27,734   Fasting glucose, HbA1c   Yes   Unknown   Unknown   565   Japan   2019   National Health and Nutrition Survey   National   both   20+   20+   1,645   1,260   Fasting glucose, HbA1c   Yes   Lab   Lab   Lab   566   Japan   2022-2023   The Tokyo Health Service Association Database   Community urban   20-79   22-79   28,852   43,488   Fasting glucose, HbA1c   Yes   Lab   Lab   Lab   Community   Urban   20-79   22-79   28,852   43,488   Fasting glucose, HbA1c   Yes   Lab   Lab	ıb	Lab	Lab	Yes	Fasting glucose, HbA1c	1,556	2,037	20+	20+	both	National	2017 National Health and Nutrition Survey	ın 2017	561
	ıb	Lab	Lab	Yes	Fasting glucose, HbA1c	2,109	4,337	75-90	75-90	both	Subnational	2015-2019 The Shizuoka KDB study	n 2015-2019	562
September   Sept											,	·		
566   Japan   2022-2023   The Tokyo Health Service Association Database   Community   urban   20-79   20-79   28.652   43.488   Fasting glucose, HbA1c   Yes   Lab   Lab   Service Association Database   Community   urban   20-79   20-79   28.652   43.488   Fasting glucose   HbA1c   Yes   Lab   Lab   Service Association Database   Community   urban   20-79   20-79   28.652   43.488   Fasting glucose   Yes   Lab   Lab														
S67   Jordan   2004   Behavioural Risk Factor Surveillence Survey   National   Tural   18+   18+   1,974   1,351   Fasting glucose   Yes   Lab   S68   Jordan   2007   Behavioural Risk Factor Surveillence Survey   National   both   18+   18+   1,075   1,949   Fasting glucose   Yes   Lab   S68   Jordan   2009   Metablic abnormatics and Vitamin   Study   St												,		
Sea   Jordan   2007   Behavioural Risk Factor Surveillence Survey   National   both   18+   18+   1,705   1,949   Fasting glucose   Yes   Lab   1,005   1,049   Metablic abnormalities and vitamin D study   National   both   18+   18+   1,705   1,949   Fasting glucose   Yes   Lab   1,007   1,0	.b	Lab												
Seg		<b>↓</b>												
S70   Jordan   2016-2017   National Cardiovascular Diseases and Diabetes Study (NCDDS)   National   both   18+   18+   2,763   1,182   Fasting glucose, HbA1c   Yes   Portable   Lab	$\longrightarrow$	<del>                                     </del>												
571         Jordan         2019         STEPS         National         both         18-69         3,324         2,202         Fasting glucose         Yes         Portable           572         Kazakhstan         2011-2012         Household Health Survey         National         both         18+69         1,384         3,77         Fasting glucose         No         Portable           573         Kazakhstan         2015         Almaty STEPS         Subnational         both         18-69         1,138         3,77         Fasting glucose         Yes         Lab           574         Kazakhstan         2015         Shymkent STEPS         Subnational         both         18-69         1,138         3,77         Fasting glucose         Yes         Lab           575         Kazakhstan         2015-2016         Aktobe STEPS         Subnational         both         18-69         18-69         1,138         347         Fasting glucose         Yes         Lab           576         Kazakhstan         2019         A health status assessment of a population of Karaganda urban region         Community         urban         18-69         1,138         347         Fasting glucose, HbAtc         No         Lab           577         Kazakhstan		<del></del>												
572         Kazakhstan         2011-2012         Household Health Survey         No         Portable           573         Kazakhstan         2015         Almay STEPS         Subnational         both         18-69         11-89	<u>a</u>	Lab												
S73   Kazakhstan   2015   Almaty STEPS   Subnational   both   18-69   18-69   1,138   377   Fasting glucose   Yes   Lab	+-	<del> </del>												
574         Kazakhstan         2015         Shymkent STEPS         Subnational         both         18-69         18-69         925         428         Fasting glucose         Yes         Lab           575         Kazakhstan         2015-2016         Aktobe STEPS         Subnational         both         18-69         18-69         1,138         347         Fasting glucose         Yes         Lab           576         Kazakhstan         2019         A health status assessment of a population of Karaganda urban region         Community         urban         18+69         11,38         347         Fasting glucose, HbA1c         No         Lab           577         Kazakhstan         2021         Prevalence of NCD Risk Factors in Kazakhstan         Subnational         both         18-69<	+-	<del> </del>												
Subnational   Doth   18-69   1,138   347   Fasting glucose   Yes   Lab   Lab	+-	<del></del>												
576         Kazakhstan         2019         A health status assessment of a population of Karaganda urban region         Community         urban         18+         664         315         Fasting glucose, HbA1c         No         Lab         Lab           577         Kazakhstan         2021         Prevalence of NCD Risk Factors in Kazakhstan         Subnational         both         18-69         18-69         813         801         Fasting glucose, HbA1c         Yes         Lab         Lab           578         Kazakhstan         2021-2022         Prevalence of risk factors for NCD in Kazakhstan         National         both         18-69         18-69         1,997         1,811         Fasting glucose, HbA1c         Yes         Lab         Lab           579         Kernya         2015         STEPS         National         both         18-69         18-69         1,997         1,811         Fasting glucose         Yes         Lab         Lab           580         Kiribati         1981         Epidemiological survey of Kiribati         Subnational         rural         20+         20+         533         474         Fasting glucose         Yes         Lab           582         Kiribati         2004         STEPS         STEPS         National         <		<del>                                     </del>												
577         Kazakhstan         2021         Prevalence of NCD Risk Factors in Kazakhstan         Subnational         both         18-69         18-69         813         801         Fasting glucose, HbA1c         Yes         Lab         Lab           578         Kazakhstan         2021-2022         Prevalence of risk factors for NCD in Kazakhstan         National         both         18-69         18-69         1,997         1,811         Fasting glucose, HbA1c         Yes         Lab         Lab           579         Kenya         2015         STEPS         National         both         18-69         18-69         2,527         1,763         Fasting glucose, HbA1c         Yes         Lab         Lab           580         Kiribati         1981         Epidemiological survey of Kiribati         Subnational         rural         20+         533         474         Fasting glucose         Yes         Lab           581         Kiribati         1981         Epidemiological survey of Kiribati         Subnational         urban         20+         20+         879         917         Fasting glucose         Yes         Lab           582         Kiribati         2004         STEPS         National         both         18-64         18-64         897 <td>-h</td> <td>l ah</td> <td></td>	-h	l ah												
578         Kazakhstan         2021-2022         Prevalence of risk factors for NCD in Kazakhstan         National         both         18-69         1,997         1,811         Fasting glucose, HbA1c         Yes         Lab         Lab           579         Kenya         2015         STEPS         National         both         18-69         18-69         2,527         1,763         Fasting glucose         Yes         Unknown           580         Kiribati         1981         Epidemiological survey of Kiribati         Subnational         rural         20+         20+         533         474         Fasting glucose         Yes         Lab           581         Kiribati         1981         Epidemiological survey of Kiribati         Subnational         urban         20+         20+         879         917         Fasting glucose         Yes         Lab           582         Kiribati         2004         STEPS         National         both         18-64         897         727         Fasting glucose         Yes         Portable           583         Kiribati         2015-2016         STEPS         National         both         18-69         1,414         963         Fasting glucose         Yes         Portable														
579         Kenya         2015         STEPS         National         both         18-69         2,527         1,763         Fasting glucose         Yes         Unknown           580         Kiribati         1981         Epidemiological survey of Kiribati         Subnational         rural         20+         20+         533         474         Fasting glucose         Yes         Lab           581         Kiribati         1981         Epidemiological survey of Kiribati         Subnational         urban         20+         879         917         Fasting glucose         Yes         Lab           582         Kiribati         2004         STEPS         National         both         18-64         897         727         Fasting glucose         Yes         Portable           583         Kiribati         2015-2016         STEPS         National         both         18-69         1,141         963         Fasting glucose         Yes         Portable														
580 Kiribati 1981 Epidemiological survey of Kiribati Subnational rural 20+ 20+ 533 474 Fasting glucose Yes Lab 581 Kiribati 1981 Epidemiological survey of Kiribati Subnational urban 20+ 20+ 879 917 Fasting glucose Yes Lab 582 Kiribati 2004 STEPS National both 18-64 18-64 897 727 Fasting glucose Yes Portable 583 Kiribati 2015-2016 STEPS National both 18-69 1,141 963 Fasting glucose Yes Portable		200												
581         Kiribati         1981         Epidemiological survey of Kiribati         Subnational         urban         20+         20+         879         917         Fasting glucose         Yes         Lab           582         Kiribati         2004         STEPS         National         both         18-64         18-64         897         727         Fasting glucose         Yes         Portable           583         Kiribati         2015-2016         STEPS         National         both         18-69         1,141         963         Fasting glucose         Yes         Portable														
582         Kiribati         2004         STEPS         National         both         18-64         18-64         897         727         Fasting glucose         Yes         Portable           583         Kiribati         2015-2016         STEPS         National         both         18-69         1,141         963         Fasting glucose         Yes         Portable												, , ,		
583 Kiribati 2015-2016 STEPS National both 18-69 1,141 963 Fasting glucose Yes Portable	-													
	-													
584 Kuwait 2006 STEPS National both 20-64 20-64 1,298 918 Fasting glucose Yes Lab	-	i e	Lab	Yes	Fasting glucose	918	1,298	20-64	20-64	both	National	2006 STEPS		

	Country	Data years	Survey/Study name/Citation	Level of representative-	Rural, urban, or		e as used Il analysis		size used Il analysis	Glycaemic markers available	Information available on diabetes		easured in a y a point-of-care e device	Notes
				ness	both	Female	Male	Female	Male		treatment	Glucose	HbA1c	
585	Kuwait	2008-2009	National Nutrition Program for the State of Kuwait	National	both	18-86	18-86	530	459	Fasting glucose, HbA1c	Yes	Lab	Lab	15
586 587	Kuwait Kuwait	2011-2014	Kuwait Diabetes Epidemiology Program	National National	both	18-82 18-69	18-82 18-69	2,144 2.406	2,781 1,458	Fasting glucose, HbA1c	Yes	Lab	Lab Lab	1
588	Kyrgyzstan	2014	STEPS STEPS	National	both both	25-64	25-64	1,605	946	Fasting glucose, HbA1c Fasting glucose	Yes Yes	Lab Portable	Lab	<del>                                     </del>
589	Lao PDR	2013	STEPS	National	both	18-64	18-64	1,471	989	Fasting glucose	Yes	Portable		
590	Latvia	2008-2009	Cardiovascular risk factor study	National	both	25-74	25-74	2,394	1,359	Fasting glucose	No	Lab		
591	Lebanon	2017	STEPS	National	both	18-69	18-69	1,086	790	Fasting glucose	Yes	Lab		ļ
592 593	Lebanon	2023-2024 2012	STEPS STEPS	National	both	18-69 25-64	18-69 25-64	1,820 1,492	1,053 774	Fasting glucose, HbA1c	Yes	Portable	Unknown	2
593	Lesotho Liberia	2012	STEPS	National Subnational	both both	25-64	25-64	1,492	1,060	Fasting glucose Fasting glucose	Yes Yes	Portable Portable		$\vdash$
595	Liberia	2022	STEPS	National	both	18-69	18-69	2,367	1,532	Fasting glucose	Yes	Portable		
596	Libya	1999	Kadiki et al., Diabetes Metab 27:647-54, 2001	Community	both	25-84	25-84	388	211	Fasting glucose	No	Unknown		
597	Libya	2009	STEPS	National	both	25-64	25-64	1,622	1,712	Fasting glucose	Yes	Portable		<u> </u>
598	Libya	2022-2023	STEPS	National	both	18-69	18-69	2,599	2,271	Fasting glucose, HbA1c	Yes	Portable	Lab	2
599 600	Lithuania Lithuania	2001-2002 2006-2008	MONICA4 Health, Alcohol and Psychosocial Factors In Eastern Europe	Community Community	urban urban	35-64 45-70	35-64 45-70	760 3,497	609 2,906	Fasting glucose Fasting glucose	Yes Yes	Portable Portable		$\vdash$
601	Luxembourg	2006-2008	Observation of cardiovascular risk factors in Luxembourg (ORISCAV-LUX)	National	both	18-69	18-69	735	697	Fasting glucose, HbA1c	Yes	Lab	Lab	$\vdash \vdash \vdash$
602	Luxembourg	2013-2015	European Health Examination Survey in Luxembourg	National	both	25-64	25-64	781	721	Fasting glucose, HbA1c	Yes	Lab	Lab	
603	Luxembourg	2016-2018	Observation of cardiovascular risk factors in Luxembourg (ORISCAV-LUX2)	Community	both	25-79	25-79	825	731	Fasting glucose, HbA1c	Yes	Lab	Lab	$ldsymbol{\Box}$
604	Malawi	2009	STEPS	National	both	25-64	25-64	3,252	1,690	Fasting glucose	Yes	Portable		$\longmapsto$
605 606	Malawi	2013-2017 2013-2017	NCD Survey Malawi Epidemiology and Intervention Research Unit	Community	rural	18+ 18+	18+ 18+	7,497 10,291	5,830 5,799	Fasting glucose	Yes	Lab	<del></del>	<del>                                     </del>
607	Malawi Malawi	2013-2017	NCD Survey Malawi Epidemiology and Intervention Research Unit STEPS	Community National	urban both	18-69	18-69	2,560	1,485	Fasting glucose Fasting glucose	Yes Yes	Lab Portable		$\vdash$
608	Malaysia	2004	Rampal et al., Public Health 122(1):11-8, 2008	National	both	18+	18+	9,840	7,362	Fasting glucose	Yes	Lab		$\vdash$
609	Malaysia	2006	National Health and Morbidity Survey (NHMS)	National	both	18+	18+	18,176	15,274	Fasting glucose	Yes	Portable		
610	Malaysia	2008	Metabolic Syndrome Study in Malaysia	National	rural	18+	18+	1,360	749	Fasting glucose, HbA1c	Yes	Lab	Lab	
611	Malaysia	2008	Metabolic Syndrome Study in Malaysia	National	urban	18+	18+	1,435	765	Fasting glucose, HbA1c	Yes	Lab	Lab	<b>↓</b>
612 613	Malaysia Malaysia	2011 2015	National Health and Morbidity Survey (NHMS)  National Health and Morbidity Survey (NHMS)	National National	both both	18+ 18+	18+ 18+	8,844 10,411	8,090 9,439	Fasting glucose Fasting glucose	Yes Yes	Portable Portable		<b>├</b>
614	Malaysia	2019	National Health and Morbidity Survey (NHMS)	National	both	18+	18+	5,555	4,783	Fasting glucose	Yes	Portable		$\vdash$
615	Maldives	2004	STEPS	Subnational	urban	25-64	25-64	839	716	Fasting glucose	No	Lab		
616	Maldives	2020-2021	STEPS	National	both	18-69	18-69	1,994	886	Fasting glucose	Yes	Portable		
617	Mali	2013	STEPS	Subnational	both	18-65	18-65	805	446	Fasting glucose	Yes	Portable		ļ
618 619	Malta Malta	1981 2014-2016	DECODE; DECODE Study Group, Diabetes Care 26:61-9, 2003	Community National	both	30-89	30-89	1,173 1,022	870	Fasting glucose	No	Unknown	<del></del>	$\vdash$
620	Marshall Islands	2014-2016	SAHHTEK - The University of Malta Health and Wellbeing Study STEPS	National	both both	18-70 18-64	18-70 18-64	1,532	834 1,081	Fasting glucose Fasting glucose	Yes Yes	Lab Portable		$\vdash$
621	Marshall Islands	2017-2018	STEPS	National	both	18+	18+	1,556	1,406	Fasting glucose	Yes	Unknown		$\vdash \lnot$
622	Mauritania	2006	STEPS	Community	urban	18-64	18-64	1,126	1,023	Fasting glucose	Yes	Lab		
623	Mauritius	1987	Mauritius Noncommunicable Disease Survey	National	both	25-74	25-74	2,662	2,355	Fasting glucose	Yes	Lab		<u> </u>
624	Mauritius	1992	Mauritius Noncommunicable Disease Survey	National	both	25-74	25-74	3,481	2,994	Fasting glucose	Yes	Lab	<del></del>	$\vdash$
625 626	Mauritius Mauritius	1992 1998	Rodrigues, Mauritius (1992)  Mauritius Noncommunicable Disease Survey	Community National	rural both	25-64 25-74	25-64 25-74	774 3,248	737 2,567	Fasting glucose Fasting glucose	Yes Yes	Lab Lab	$\overline{}$	$\vdash \vdash$
627	Mauritius	1999	Rodrigues, Mauritius (1999)	Community	rural	20+	20+	1,295	977	Fasting glucose	Yes	Lab		$\vdash \lnot$
628	Mauritius	2009	Mauritius Noncommunicable Disease Survey	National	both	19-74	19-74	3,432	2,903	Fasting glucose	Yes	Lab	i	
629	Mauritius	2015	Mauritius Noncommunicable Disease Survey	National	both	20-74	20-74	1,948	1,626	Fasting glucose, HbA1c	Yes	Lab	Lab	<u> </u>
630	Mauritius	2015	Mauritius Noncommunicable Disease Survey 1998 Follow Up	National	both	20+	20+	1,171	886	Fasting glucose, HbA1c	Yes	Lab	Lab	$\longmapsto$
631 632	Mexico Mexico	2004-2005 2006	CArdiovascular Risk factors Multiple Evaluation in Latin America (CARMELA)  Encuesta Nacional de Salud v Nutrición	Community National	urban both	25-64 20+	25-64 20+	889 24,881	833 19,673	Fasting glucose Fasting glucose	Yes Yes	Lab Unknown		$\vdash$
633	Mexico	2009-2012	Encuesta Nacional Sobre Niveles de vida de los Hogares	National	both	18+	18+	10,495	8,215	HbA1c	Yes	Olikiowii	Portable	$\vdash \vdash \vdash$
634	Mexico	2012	The Mexican Health and Aging Study	National	both	50+	50+	8,399	6,441	HbA1c	Yes		Portable	
635	Mexico	2016	Encuesta Nacional de Salud y Nutrición	National	both	20+	20+	5,477	2,865	Fasting glucose, HbA1c	Yes	Lab	Lab	
636	Mexico	2016	Cognitive Aging Linked to MHAS (Mex-Cog)	National	both	55+	55+	1,191	833	HbA1c	Yes		Portable	16
637 638	Mexico Mexico	2018-2019	Encuesta Nacional de Salud y Nutrición Encuesta Nacional de Salud y Nutrición	National National	both both	20+ 20+	20+ 20+	9,798 1,359	7,584 932	Fasting glucose, HbA1c Fasting glucose, HbA1c	Yes No	Lab Lab	Lab Lab	$\vdash$
639	Mexico	2020	Encuesta Nacional de Salud y Nutrición	National	both	20+	20+	5,243	2,947	Fasting glucose, HbA1c	Yes	Lab	Lab	$\vdash$
0.10	Mexico	2022	Encuesta Nacional de Salud y Nutrición	National	both	20+	20+	6,627	4,338	Fasting glucose, HbA1c	Yes	Lab	Lab	
	Micronesia	2002	STEPS	Subnational	both	25-64	25-64	866	580	Fasting glucose	Yes	Lab		
642	Micronesia	2006	STEPS	Subnational	both	25-64	25-64	1,246	702	Fasting glucose	Yes	Unknown		igspace
	Micronesia Micronesia	2008 2009	STEPS STEPS, Kosrae	Subnational Subnational	both both	25-64 18-64	25-64 18-64	1,255 446	857 246	Fasting glucose Fasting glucose	Yes Yes	Unknown Portable		$\vdash \!$
	Micronesia	2009	STEPS, Kostae STEPS, Yap	Subnational	both	18-64	18-64	562	435	Fasting glucose Fasting glucose	Yes	Unknown		$\vdash$
	Moldova	2013	STEPS	National	both	18-69	18-69	2,893	1,806	Fasting glucose	Yes	Portable	i	
647	Moldova	2021	STEPS	National	both	18-69	18-69	2,289	1,774	Fasting glucose	Yes	Portable		
648	Mongolia	2005	STEPS	National	both	25-64	25-64	1,374	1,323	Fasting glucose	Yes	Portable		1
	Mongolia	2009	STEPS	National	both	25-64	25-64	2,658	1,814	Fasting glucose	Yes	Unknown		$\vdash \vdash$
650	Mongolia	2013	STEPS	National	both	25-64	25-64	2,366	1,921	Fasting glucose	Yes	Portable		

	Country	Data years	Survey/Study name/Citation	Level of representative-	Rural, urban, or both		e as used Il analysis		size used Il analysis	Glycaemic markers available	Information available on diabetes		easured in a / a point-of-care e device	Notes
				ness	DOIN	Female	Male	Female	Male		treatment	Glucose	HbA1c	
	Mongolia	2019	STEPS	National	both	18-69	18-69	3,481	2,885	Fasting glucose	Yes	Portable		
652	Morocco	2000	National Survey 2000	National	both	20-89	20-89	973	689	Fasting glucose	No	Lab		<u>.</u>
653	Morocco	2017	STEPS	National	both	18+	18+	3,417	1,891	Fasting glucose	Yes	Portable		₩
654	Mozambique	2014-2015	STEPS	National	both	18-64	18-64	1,644	1,111	Fasting glucose	Yes	Portable		
655 656	Myanmar Myanmar	2003-2004	STEPS National aureau of Diabetes Malifus and risk factors for Non-communicable diseases in Museumer	Subnational National	both both	25-74 25-64	25-74 25-64	2,453 5,592	1,991 3,079	Fasting glucose	Yes Yes	Lab Portable		$\vdash$
657	Myanmar	2013-2014	National survey of Diabetes Mellitus and risk factors for Non-communicable diseases in Myanmar STEPS, Yangon	Subnational	both	25-74	25-74	740	745	Fasting glucose Fasting glucose	Yes	Lab		+
658	Namibia	2013	Demographic and Health Survey Namibia 2013	National	both	35-64	35-64	2,054	1,551	Fasting glucose	Yes	Portable		$\vdash$
659	Nauru	1982	Trends in the prevalence and incidence of non-insulin-dependent diabetes mellitus and impaired glucose tolerance	National	both	20+	20+	775	706	Fasting glucose	Yes	Unknown		
660	Nauru	1987	Trends in the prevalence and incidence of non-insulin-dependent diabetes mellitus and impaired glucose tolerance	National	both	20+	20+	662	554	Fasting glucose	Yes	Lab		
661	Nauru	1994	Trends in the prevalence and incidence of non-insulin-dependent diabetes mellitus and impaired glucose tolerance	National	both	25+	25+	735	652	Fasting glucose	Yes	Lab		
662	Nauru	2006	STEPS	National	both	18-65	18-65	231	245	Fasting glucose	Yes	Lab		<u> </u>
663	Nauru	2015	STEPS	National	both	18-69	18-69	704	649	Fasting glucose	Yes	Portable		
664	Nepal	1990	Sasaki et al., Diabetes Res Clin Pract 67:167-74, 2005	Community	rural		20+	205	85	Fasting glucose	No	Unknown		₩
665	Nepal	2000	Singh et al., Diabet Med 20:170-1, 2003	Subnational	rural	20+	20+	235	105	Fasting glucose	No	Unknown		
666	Nepal	2000	Singh et al., Diabet Med 20:170-1, 2003	Subnational	urban	20+	20+	456	442	Fasting glucose	No	Unknown		$\vdash$
667	Nepal	2006-2011	Early detection and management of Kidney disease, Hypertension, Diabetes and Cardiovascular disease (KHDC Nepal), Tarahara	Community	rural	18+	18+	2,351	1,176	Fasting glucose	Yes	Lab		
668	Nepal	2006-2011	Early detection and management of Kidney disease, Hypertension, Diabetes and Cardiovascular disease (KHDC Nepal), Damak	Community	urban	18+	18+	1,577	1,095	Fasting glucose	Yes	Lab		
669	Nepal	2006-2011	Early detection and management of Kidney disease, Hypertension, Diabetes and Cardiovascular disease (KHDC Nepal), Dharan	Community	urban	18+	18+	6,130	4,130	Fasting glucose	Yes	Lab		
670	Nepal	2012-2013	STEPS	National	both	18-69	18-69	2,702	1,276	Fasting glucose	Yes	Lab		igsquare
671	Nepal	2015	Community based intervention for prevention and control of non-communicable diseases risk factors (CIPCON) baseline survey, Dhankuta	Subnational	rural	18-69	18-69	779	555	Fasting glucose	Yes	Lab		
672	Nepal	2015	Community based intervention for prevention and control of non-communicable diseases risk factors (CIPCON) baseline survey, llam	Subnational	rural	18-69	18-69	717	553	Fasting glucose	Yes	Lab		
673	Nepal	2016-2018	The Population Based Prevalence of Selected Non-Communicable Diseases In Nepal	National	both	20+	20+	7,423	4,834	Fasting glucose	Yes	Lab		
674	Nepal	2019	STEPS	National	both	18-69	18-69	3,400	1,910	Fasting glucose	Yes	Portable		
675	Netherlands	1990	DECODE; DECODE Study Group, Diabetes Care 26:61-9, 2003	Community	urban	50-79	50-79	1,322	1,131	Fasting glucose	No	Unknown		
676	Netherlands	1990	Zutphen Elderly Study	Community	urban		69-90		555	Fasting glucose	Yes	Lab		<u>.</u>
677	Netherlands	1995-1996	The Longitudinal Aging Study Amsterdam (LASA)	Subnational	both	65-88	65-88	780	725	Fasting glucose	Yes	Unknown		17
678	Netherlands	1998-2001	Regenboog Project	National	both	18-89	18-89 35-60	2,305	2,323	Fasting glucose	Yes	Lab		
679 680	Netherlands Netherlands	2001-2003 2008-2009	Surinamese in the Netherlands: Study on Ethnicity and Health (SUNSET) The Longitudinal Aging Study Amsterdam (LASA)	Community Subnational	urban both	35-60 60-100	60-100	257 827	251 665	Fasting glucose Fasting glucose	Yes Yes	Lab Unknown		17
681	Netherlands	2012-2013	The Longitudinal Aging Study Amsterdam (LASA)	Subnational	both	55-65	55-65	454	433	Fasting glucose	Yes	Unknown		17
682	New Zealand	2008-2009	New Zealand Adult Nutrition Survey	National	both	18+	18+	2,284	1,806	HbA1c	Yes		Lab	
683	Nicaragua	2003-2004	CAMDI	Community	urban	20+	20+	919	781	Fasting glucose	Yes	Lab		
684	Niger	2021	STEPS	National	both	18-69	18-69	3,076	2,283	Fasting glucose	Yes	Portable		
685	Nigeria	1998	Okesina et al., East Afr Med J 76:212-6, 1999	Community	rural	21+	21+	120	222	Fasting glucose	No	Unknown		
686	Nigeria	1999-2009	Prostate cancer dietary risk factors study	Subnational	both		35+		447	Fasting glucose	No	Portable		<u> </u>
687	Nigeria	2006	Clustering of cardiovascular disease risk-factors in semiurban population in Northern Nigeria	Community	urban	18+	18+	106	87	Fasting glucose	No	Lab		<b>↓</b>
688	Nigeria	2007	Southeast Nigeria kidney disease study	Community	rural	25-64	25-64	442	169	Fasting glucose	Yes	Portable		
689 690	Nigeria Norway	2009-2011	Anthropometric indices in Calabar The Tromsø Study: Tromsø 5, Tromsø Study Panel	Community Community	urban both	18-79 30-89	18-79 30-89	331 3,576	381 2,536	Fasting glucose HbA1c	No Yes	Lab	Lab	$\vdash$
691	Norway	2007-2008	The Tromsø Study: Tromsø 6	Community	both	30-89	30-89	6,709	5,927	HbA1c	Yes	1	Lab	$\vdash$
692	Norway	2017-2019	HUNT4 study	Community	rural	19+	19+	29,556	24,585	HbA1c	Yes	1	Lab	$\vdash$
693	Oman	1991	The 1991 National Diabetes Survey of Oman	National	both	20+	20+	2,809	1,989	Fasting glucose	Yes	Lab		
694	Oman	2000	Oman National Health Survey	National	both	20+	20+	2,933	2,905	Fasting glucose	No	Lab		
695	Oman	2001	Nizwa Healthy Lifestyle Project	Community	urban	20+	20+	692	600	Fasting glucose	Yes	Lab		┷
696	Oman	2008	Gulf Cooperation Council World Health Survey	National	both	18+	18+	2,264	2,446	Fasting glucose	Yes	Unknown		$\vdash$
697 698	Oman Pakistan	2017 1994	STEPS  Pacifiet at J. LiPak Med Acces 52:257.60, 2002	National Subnational	both rural	18+ 25+	18+ 25+	2,997 1,362	3,365	Fasting glucose	Yes	Portable Unknown		$\vdash$
699	Pakistan	2002	Basit et al., J Pak Med Assoc 52:357-60, 2002 Basit et al., Diabetes Res Clin Pract 94:456-62, 2011; Study 1	Subnational	rural	25+	25+	1,362	761 670	Fasting glucose Fasting glucose	No No	Lab		$\vdash$
700	Pakistan	2002	COBRA-1	Community	urban	40+	40+	1,502	1,376	Fasting glucose	No	Unknown		$\vdash \vdash \vdash$
701	Pakistan	2009-2010	Basit et al., Diabetes Res Clin Pract 94:456-62, 2011; Study 2	Subnational	rural	25+	25+	840	272	Fasting glucose	No	Lab		
702	Pakistan		National Diabetes Survey of Pakistan	National	both	20+	20+	4,261	3,273	Fasting glucose, HbA1c	No	Lab	Lab	18
	Palau	2011-2013	STEPS	National	both	25-64	25-64	1,124	1,038	Fasting glucose	Yes	Portable		
	Palau	2016	STEPS	National	both	18+	18+	854	872	Fasting glucose	Yes	Portable		igsquare
	Panama	2010-2011	Prevalencia de factores de riesgo asociados a enfermedad cardiovascular 2010-2011	Subnational	both	18+	18+	2,472	1,072	Fasting glucose, HbA1c	Yes	Lab	Lab	igwdapprox
	Panama Papua New	2019	Encuesta Nacional de Salud de Panama (ENSPA)	National	both	18+	18+	12,013	4,846	Fasting glucose, HbA1c	Yes	Lab	Lab	<del>                                     </del>
707	Guinea	1991	Dowse et al., Med J Aust 160:767-74, 1994	Subnational	both	25-88	25-88	1,009	837	Fasting glucose	Yes	Lab		<u> </u>
	Peru Peru	2004 2005	Factores de Riesgo de Enfermedades No Transmisibles  PREVENCION Study; Medina-Lezama et al., J Am Soc Hypertens 1:216-25, 2007	Community	urban urban	18+ 20-80	18+ 20-80	430 1,011	209 867	Fasting glucose Fasting glucose	Yes No	Lab Unknown		$\vdash \vdash$
	Peru	2005	CArdiovascular Risk factors Multiple Evaluation in Latin America (CARMELA)	Community	urban urban	25-64	25-64	1,011	769	Fasting glucose Fasting glucose	Yes	Lab		$\vdash \vdash$
110	ı cıu	2004-2003	Ontoropoular trait factors intultible Evaluation in Eath America (OARMELA)	Community	uivall	2J-04	23-04	003	109	i asiiiy yiucuse	162	Lan		

	Country	Data years	Survey/Study name/Citation	Level of representative-	Rural, urban, or		e as used Il analysis		size used I analysis	Glycaemic markers available	Information available on diabetes		easured in a / a point-of-care e device	Notes
				ness	both	Female	Male	Female	Male		treatment	Glucose	HbA1c	
711	Peru	2004-2005	Encuesta Nacional de Indicadores Nutricionales, Bioquímicos, Socioeconómicos y Culturales Relacionados con las Enfermedades Crónicas Degenerativas	National	both	20+	20+	2,095	2,095	Fasting glucose	Yes	Lab		
712	Peru	2005	Factores de Riesgo de Enfermedades No Transmisibles	Community	urban	18+	18+	532	199	Fasting glucose	Yes	Lab		<u> </u>
713	Peru	2006	Factores de Riesgo de Enfermedades No Transmisibles	Community	urban	18+	18+	1,030	608	Fasting glucose	Yes	Lab		<u> </u>
714	Peru	2007-2008	PERU MIGRANT Study	Community	both	30+	30+	521	466	Fasting glucose, HbA1c	Yes	Lab	Lab	$\vdash$
715 716	Peru Peru	2009-2012 2013	CRONICAS Cohort Study  Clinical functional and sociofamilial profiles of the elderly from a community in a district of Lima, Peru	Subnational Community	both urban	35+ 60+	35+ 60+	1,849 200	1,737 114	Fasting glucose, HbA1c Fasting glucose	Yes No	Lab Lab	Lab	$\vdash$
717	Peru	2013-2014	CRONICAS Cohort Study	Subnational	both	36+	36+	1,387	1,309	Fasting glucose, HbA1c	Yes	Lab	Lab	$\vdash$
718	Peru	2015-2016	PERU MIGRANT Study	Community	both	38+	38+	437	344	Fasting glucose	Yes	Lab		
719	Peru	2016-2017	Screening of T2DM	Community	urban	30-70	30-70	804	792	Fasting glucose, HbA1c	Yes	Lab	Lab	
720	Peru	2017-2018	Vigilancia Alimentario Nutricional por Etapas de Vida (VIANEV) 2017-2018	National	both	18-59	18-59	618	463	Fasting glucose	Yes	Portable		<u> </u>
721	Philippines	1998	National Nutrition Survey; Tanchoco et al., Asia Pac J Clin Nutr 12:271-6, 2003	National	both	20+	20+	927	1,030	Fasting glucose	No	Unknown		$\vdash$
722 723	Philippines Philippines	2003 2005	6th National Nutrition Survey  Cebu Longitudinal Health and Nutrition Survey 2005 Child Follow-up	National Community	both both	20+ 20-22	20+ 20-22	2,497 764	2,255 927	Fasting glucose Fasting glucose	Yes No	Lab Unknown		₩
724	Philippines	2005	Cebu Longitudinal Health and Nutrition Survey 2005 Child Follow-up  Cebu Longitudinal Health and Nutrition Survey 2005 Mother Follow-up	Community	both	35-69	20-22	1,872	927	Fasting glucose	No	Unknown		$\vdash \vdash \vdash$
725	Philippines	2008	7th National Nutrition Survey	National	both	20+	20+	3,719	3,318	Fasting glucose	Yes	Lab		
726	Philippines	2008	Philippines LIFECARE Cohort	National	both	20-50	20-50	1,743	1,329	Fasting glucose	No	Lab		
727	Philippines	2013-2014	8th National Nutrition Survey	National	both	18+	18+	10,427	9,797	Fasting glucose	Yes	Lab		
728	Philippines	2018-2021	Philippine Expanded National Nutrition Survey	National	both	18+	18+	42,650	34,265	Fasting glucose	No	Lab		<u> </u>
729	Poland	1989-1990	Polish Program CINDI (CINDI Lodz 1989-1990)	Community	urban	25-64	25-64	945	812	Fasting glucose	Yes	Lab		
730 731	Poland Poland	1993 2000	DECODE; DECODE Study Group, Diabetes Care 26:61-9, 2003  The health status, risk factors of exercis diseases and health behaviors of residents of Taxus (CIND) Taxus 2000)	Community	urban	40-79 18-83	40-79 18-83	192 1,010	172 929	Fasting glucose	No Yes	Unknown		┾┷┩
732	Poland	2001-2002	The health status, risk factors of chronic diseases and health behaviors of residents of Torun (CINDI Torun 2000)  The health status, risk factors of chronic diseases and health behaviors of residents of Lodz (CINDI Lodz 2001)	Community Community	urban urban	18-64	18-64	841	1,003	Fasting glucose Fasting glucose	Yes Yes	Lab Lab		$\vdash$
733	Poland	2002	The health status, risk factors of chronic diseases and health behaviors of residents of Lodz - seniors (CINDI Lodz 2002)	Community	urban	65+	65+	539	291	Fasting glucose	Yes	Lab		$\vdash$
734	Poland	2002	NATPOL	National	both	18+	18+	1,294	1,014	Fasting glucose	Yes	Lab		
735	Poland	2001-2002	Young Men Cardiovascular Association Study	Community	urban		18+		944	Fasting glucose	Yes	Lab		19
736	Poland	2003	The European Male Ageing Study	Community	both		40+		394	Fasting glucose	Yes	Lab		<u> </u>
737	Poland	2002-2005	Health, Alcohol and Psychosocial Factors In Eastern Europe	Community	urban	45-70	45-70	5,484	5,214	Fasting glucose, HbA1c	Yes	Lab	Lab	<u> </u>
738 739	Poland Poland	2003-2005 2006	National Multicenter Health Survey in Poland. Project WOBASZ  The health right factors for changing dispages, gifting and helpolices of health residents of Terra (CINDI Terra 2006).	National	both	20-74 18-65	20-74 18-65	6,960	6,310 750	Fasting glucose	Yes	Lab		<b>├</b>
740	Poland	2008	The health, risk factors for chronic diseases, attitudes and behaviors of health residents of Torun (CINDI Torun 2006)  The European Male Ageing Study	Community	urban both	10-05	45+	1,115	308	Fasting glucose Fasting glucose	Yes Yes	Lab Lab		$\vdash$
741	Poland	2007-2011	Medical, psychological and socioeconomic aspects of aging in Poland	National	both	55+	55+	2,671	2,775	Fasting glucose	Yes	Lab		$\vdash$
742	Poland	2010	Zatonska et al., Ann Agric Environ Med 18:265-9, 2011	Community	both	45-64	45-64	2,570	1,289	Fasting glucose	No	Lab		
743	Poland	2011	NATPOL	National	both	18-79	18-79	1,210	1,145	Fasting glucose, HbA1c	Yes	Lab	Lab	
744	Poland	2011-2014	Mogielica Human Ecology Study Site	Community	rural	45+	45+	405	142	Fasting glucose	Yes	Lab		<u> </u>
745	Poland	2013-2014	National Multicenter Health Survey in Poland. Project WOBASZ II	National	both	20+	20+	3,360	2,747	Fasting glucose	Yes	Lab		₩
746	Poland	2015-2016	LIPIDOGRAM2015 & LIPIDOGEN2015 Study - National epidemiological study of lipid disorders and selected risk factors of cardiovascular disease in primary health care in Poland	National	both	18+	18+	8,690	5,034	Fasting glucose, HbA1c	Yes	Portable	Lab	
747	Portugal	1999-2003	EPIPorto study	Community	urban	18+	18+	1,539	946	Fasting glucose	Yes	Lab		
748 749	Portugal	2011-2013 2015	EPITeen - Epidemiological Health Investigation of Teenagers in Porto Inquérito Nacional de Saúde com Exame Físico (INSEF)	Community National	urban both	20-23 25-74	20-23 25-74	868 2,615	810 2,265	Fasting glucose, HbA1c HbA1c	No Voc	Lab	Unknown Lab	<b>├</b>
750	Portugal Puerto Rico	2015	Pérez et al., Ethn Dis 18:434-41, 2008	Community	urban	25-74	25-74	532	2,205	Fasting glucose	Yes No	Unknown	Lab	$\vdash$
751	Qatar	2012	STEPS	National	both	18-64	18-64	1,379	1,053	Fasting glucose	Yes	Portable		
752	Romania	1997	Valorile medii si limitele normalitatii unor constante biologice	National	both	30-85	30-85	5,050	3,964	Fasting glucose	No	Unknown		
753	Romania	2011-2012	SEPHAR II (Study for the Evaluation of Prevalence of Hypertension and Cardiovascular Risk in Romania - 2nd edition)	National	both	18-80	18-80	1,038	936	Fasting glucose, HbA1c	Yes	Lab	Lab	
754	Romania	2012-2014	PREDATORR	National	both	20-79	20-79	1,431	1,285	Fasting glucose, HbA1c	Yes	Lab	Lab	<u> </u>
755	Romania	2021	SEPHAR IV (Study for the Evaluation of Prevalence of Hypertension and Cardiovascular Risk in Romania - 4th edition)	National	both	18-80	18-80	884	588	Fasting glucose, HbA1c	Yes	Lab	Lab	$\vdash$
756	Russian Federation	2002-2005	Health, Alcohol and Psychosocial Factors In Eastern Europe	Community	urban	45-70	45-70	5,079	4,244	Fasting glucose, HbA1c	Yes	Lab	Lab	
757	Russian Federation	2012-2014	Epidemiology of Cardiovascular dieseases in different regions of Russia (ESSE-RF)	National	both	25-64	25-64	11,639	6,979	Fasting glucose	Yes	Lab		
758	Russian Federation	2015-2017	Ural Eye and Medical Study (UEMS)	Subnational	rural	40+	40+	1,869	1,524	Fasting glucose	Yes	Lab		
759	Russian Federation	2015-2017	Ural Eye and Medical Study (UEMS)	Community	urban	40+	40+	1,437	1,035	Fasting glucose	Yes	Lab		
760	Russian Federation	2017	Epidemiology of Cardiovascular Diseases in Different Regions of Russia - 2 (ESSE-RF-2)	Subnational	both	25-64	25-64	3,699	2,980	Fasting glucose	Yes	Lab		
761	Russian Federation	2017-2020	Ural Very Old Study	Community	both	85+	85+	635	231	Fasting glucose	Yes	Lab		
	Rwanda	2012	STEPS	National	both	18-64	18-64	4,066	2,524	Fasting glucose	Yes	Portable		
	Rwanda	2021-2022	STEPS	National	both	18-69	18-69	3,389	2,130	Fasting glucose	Yes	Portable		<b></b> '
764	Saint Lucia	2012	STEPS	National	both	25-64	25-64	1,126	693	Fasting glucose	Yes	Portable		₩
	Saint Lucia	2019-2020	STEPS	National	both	18-69	18-69	1,651	1,296	Fasting glucose	Yes	Portable		$\vdash \vdash \vdash$
766	Saint Vincent and the Grenadines	2013-2014	STEPS  Non-Communicable Disease Birls Forter (NCDRF)	National	both	18-69	18-69	1,923	1,544	Fasting glucose	Yes	Portable		<u> </u>
	Samoa	1991 1991	Non-Communicable Disease Risk Factor (NCDRF)  Non-Communicable Disease Risk Factor (NCDRF)	Subnational	rural	25+	25+ 25+	495 443	466	Fasting glucose	Yes	Lab		<del></del>
768	Samoa	1991	Indirecommunicable disease Kisk Factor (NCDKF)	Subnational	urban	25+	Z0+	443	328	Fasting glucose	Yes	Lab		

	Country	Data years	a years Survey/Study name/Citation	Level of representative-	Rural, urban, or		e as used Il analysis		size used Il analysis	Glycaemic markers available	Information available on diabetes	Whether me laboratory or by portable	a point-of-care	Notes
				ness	both	Female	Male	Female	Male		treatment	Glucose	HbA1c	
	Samoa	1995	McGarvey, Pac Health Dialog 8(1):157-62, 2001	National	both	29+	29+	153	145	Fasting glucose	Yes	Lab		1
770	Samoa	2002	STEPS	National	both	25-64	25-64	1,339	1,188	Fasting glucose	Yes	Portable		20
771	Samoa Sao Tome and	2010	Samoan Genome-Wide Association Study	National	both	24-65	24-65	2,056	1,404	Fasting glucose	Yes	Lab		20
772	Principe	2019	STEPS	National	both	18-69	18-69	1,367	979	Fasting glucose	Yes	Portable		1 ,
773	Saudi Arabia	1998	National Epidemiological Health Survey; Al-Nozha et al., Saudi Med J 25:1603-10, 2004	National	both	30-70	30-70	8,804	8,002	Fasting glucose	No	Unknown		
774	Saudi Arabia	2004-2005	STEPS	National	both	15-64	18-64	2,231	2,155	Fasting glucose	No	Lab		3
775	Saudi Arabia	2009	RIYADH Cohort 2; Al-Daghri et al., BMC Med 9:76, 2011	Community	urban	18-80	18-80	1,267	1,409	Fasting glucose	No	Lab		$\longmapsto$
776	Saudi Arabia	2011-2012	Jeeluna Study- National Assessment of the Health Needs of Adolescents in Saudi Arabia	National	both	18-19	18-19	256	430	Fasting glucose	No	Lab		1
777	Senegal	2010-2012	Biocultural determinants of overweight and obesity in the context of nutrition transition in Senegal: a holistic anthropological approach	Subnational	both	18+	18+	242	216	Fasting glucose	Yes	Portable		
778	Senegal	2015	Les maladies chroniques au Sénégal: Une écologie de la santé comparative entre Dakar et Widou Thiengoly	Community	both	20+	20+	765	732	Fasting glucose	Yes	Portable		
779	Senegal	2015	Enquête Nationale sur les Facteurs de Risque des Maladies Non Transmissibles (STEPS)	National	both	18-70	18-70	3,336	1,923	Fasting glucose	Yes	Portable		<u> </u>
780	Seychelles	1989	Seychelles Heart Survey I	National	both	25-64	25-64	568	513	Fasting glucose	Yes	Portable		1
781 782	Seychelles Seychelles	2004 2013-2014	Seychelles Heart Survey IV Seychelles Heart Survey IV	National National	both both	25-64 25-64	25-64 25-64	687 699	568 531	Fasting glucose Fasting glucose, HbA1c	Yes Yes	Portable Portable	Lab	-
783	Seychelles	2023	Seychelles Heart Survey V	National	both	18-74	18-74	654	531	Fasting glucose, HbA1c	Yes	Lab	Lab	2
784	Singapore	1982-1985	Thyroid Heart Study	National	both	18+	18+	900	959	Fasting glucose	Yes	Lab	Lab	
785	Singapore	1992	National Health Survey	National	both	18-64	18-64	1,703	1,744	Fasting glucose	No	Lab		
786	Singapore	1993-1995	NUH Heart Study	National	both	26+	26+	484	495	Fasting glucose	Yes	Lab		
787	Singapore	1998	National Health Survey	National	both	18-69	18-69	2,262	2,280	Fasting glucose	No	Lab		ļ
788	Singapore	2004	National Health Survey	National	both	18-74	18-74	1,842	1,796	Fasting glucose	No	Unknown		<b>↓</b>
789 790	Singapore Singapore	2003-2005 2004-2007	Singapore Longitudinal Ageing Study - Cohort 1; SLAS-1 Singapore Cardiovascular Cohort Study and Singapore Prospective Study Program	Community National	both both	55+ 21+	55+ 21+	1,766 4,489	1,031 3,868	Fasting glucose Fasting glucose, HbA1c	Yes Yes	Lab Lab	Lab	21
790	Singapore	2004-2007	The Singapore Chinese Eye Study	Community	both	40-80	40-80	1,597	1,590	HbA1c	Yes	Lau	Lab	21
792	Singapore	2008-2013	Singapore Longitudinal Ageing Study - Cohort 2; SLAS-2	Community	both	55+	55+	2,000	1,216	Fasting glucose	Yes	Lab	Lab	$\vdash$
793	Singapore	2012-2013	Singapore Health Study	National	both	18-79	18-79	1,192	1,128	Fasting glucose, HbA1c	Yes	Lab	Lab	22
794	Singapore	2014-2015	Singapore Health 2	National	both	18-79	18-79	1,476	1,203	Fasting glucose, HbA1c	Yes	Lab	Lab	22
795	Singapore	2015-2017	The Singapore Chinese Eye Study	Community	both	50+	50+	1,193	1,169	HbA1c	Yes		Lab	1
796	Slovakia	1993	Countrywide Integrated Noncommunicable Diseases Intervention Programme	National	both	18-64	18-64	1,177	719	Fasting glucose	No	Lab		$\longmapsto$
797 798	Slovakia Slovakia	1998 2003	Countrywide Integrated Noncommunicable Diseases Intervention Programme  Countrywide Integrated Noncommunicable Diseases Intervention Programme	National National	both both	18-64 18-64	18-64 18-64	1,045 866	855 620	Fasting glucose Fasting glucose	No No	Lab Lab		$\vdash$
799	Slovakia	2003	Countrywide Integrated Noncommunicable Diseases Intervention Programme	National	both	18-64	18-64	561	390	Fasting glucose	No	Lab		$\vdash$
800	Slovakia	2011-2012	European Health Examination Survey	National	both	18-64	18-64	1,091	893	Fasting glucose	Yes	Lab		
801	Solomon Islands	2004	A genetic-ecological study of the risk factors for lifestyle-related diseases in Oceanian populations	Community	rural	18+	18+	107	106	Fasting glucose	Yes	Lab		
802	Solomon Islands	2004	A genetic-ecological study of the risk factors for lifestyle-related diseases in Oceanian populations	Community	urban	18+	18+	95	91	Fasting glucose	Yes	Lab		<u> </u>
803	Solomon Islands	2006	STEPS	Subnational	both	18-64	18-64	1,368	1,159	Fasting glucose	Yes	Portable	5	$\longmapsto$
804 805	Solomon Islands Solomon Islands	2009-2010 2009-2010	Furusawa et al., N Z Med J 124(1333):17-28, 2011  Furusawa et al., N Z Med J 124(1333):17-28, 2011	Community	rural urban	18+ 18-70	18+ 18-70	215 70	153 38	HbA1c HbA1c	Yes Yes		Portable Portable	$\vdash$
806	Solomon Islands	2015	STEPS	National	both	18-69	18-69	1.318	1,087	Fasting glucose	Yes	Portable	Tortable	$\vdash$
807	Solomon Islands	2017-2018	Impact of sea-level rise and relocation projects on health, ecology, and society in Oceania	Community	both	18+	18+	261	117	HbA1c	Yes		Portable	
808	Somalia	2016	The prevalence of selected risk factors for non-communicable diseases in Hargeisa, Somaliland: a cross-sectional study	Community	urban	20-69	20-69	955	145	Fasting glucose, HbA1c	Yes	Lab	Lab	
809	South Africa	2008-2009	Cape Town Bellville South Cohort Study - Baseline evaluation I	Community	urban	35-65	35-65	494	140	Fasting glucose, HbA1c	Yes	Lab	Lab	<u> </u>
810	South Africa	2011-2012	South Africa National Health and Nutrition Examination Survey	National	both	18+	18+	8,081	5,576	HbA1c	Yes		Lab	$\longmapsto$
811 812	South Africa South Korea	2018-2020 2001	Vukuzazi Study Kim et al., Br J Psychiatry 185:102-7, 2004	Community	both both	18+ 65+	18+ 65+	10,976 432	4,874 300	HbA1c Fasting glucose	Yes No	Unknown	Lab	<del></del>
813	South Korea	2001	Korea National Health and Nutrition Examination Survey	National	both	18+	18+	3,366	2,647	Fasting glucose	Yes	Lab		
814	South Korea	2005	Korea National Health and Nutrition Examination Survey	National	both	18+	18+	3,101	2,313	Fasting glucose	Yes	Lab		
815	South Korea	2007	Korea National Health and Nutrition Examination Survey	National	both	18+	18+	1,734	1,257	Fasting glucose	Yes	Lab		
816	South Korea	2008	Korea National Health and Nutrition Examination Survey	National	both	18+	18+	3,967	2,895	Fasting glucose	Yes	Lab		
817	South Korea	2009	Korea National Health and Nutrition Examination Survey	National	both	18+	18+	4,264	3,314	Fasting glucose	Yes	Lab		igwdapprox
818	South Korea	2008-2009 2010	Korean National Health Insurance	National	both	40+ 18+	40+ 18+	6,087,857 3,543	5,763,109 2,740	Fasting glucose	Yes	Lab		-
819 820	South Korea South Korea	2010	Korea National Health and Nutrition Examination Survey  Korea National Health and Nutrition Examination Survey	National National	both both	18+	18+	3,543	2,740	Fasting glucose Fasting glucose, HbA1c	Yes Yes	Lab Lab	Lab	$\vdash$
821	South Korea	2010-2011	Korean National Health Insurance	National	both	40+	40+	7,126,632	6,670,305	Fasting glucose	Yes	Lab		
	South Korea	2012	Korea National Health and Nutrition Examination Survey	National	both	18+	18+	3,300	2,344	Fasting glucose, HbA1c	Yes	Lab	Lab	
	South Korea	2013	Korea National Health and Nutrition Examination Survey	National	both	18+	18+	3,115	2,303	Fasting glucose, HbA1c	Yes	Lab	Lab	
824	South Korea	2012-2013	Korean National Health Insurance	National	both	40+	40+	7,782,567	7,256,160	Fasting glucose	Yes	Lab		<b></b>
825 826	South Korea	2014 2015	Korea National Health and Nutrition Examination Survey	National National	both both	18+ 18+	18+ 18+	2,943 2,963	2,115 2,308	Fasting glucose, HbA1c Fasting glucose, HbA1c	Yes	Lab Lab	Lab Lab	$\vdash$
826	South Korea South Korea	2015-2015	Korea National Health and Nutrition Examination Survey  Korean National Health Insurance	National	both	40+	40+	2,963 8,356,468	7,869,529	Fasting glucose, HBA1c	Yes Yes	Lab	Lau	$\vdash$
828	South Korea	2014-2013	Korea National Health and Nutrition Examination Survey	National	both	18+	18+	3,474	2,682	Fasting glucose, HbA1c	Yes	Lab	Lab	
829	South Korea	2017	Korea National Health and Nutrition Examination Survey	National	both	18+	18+	3,438	2,780	Fasting glucose, HbA1c	Yes	Lab	Lab	
830	South Korea	2017	Korean National Health Insurance	National	both	40+	40+	9,074,778	8,535,547	Fasting glucose	Yes	Lab		
	South Korea	2018	Korea National Health and Nutrition Examination Survey	National	both	18+	18+	3,522	2,763	Fasting glucose, HbA1c	Yes	Lab	Lab	1
832	South Korea South Korea	2019	Korea National Health and Nutrition Examination Survey	National	both	18+	18+	3,504	2,835	Fasting glucose, HbA1c	Yes	Lab	Lab	$\vdash$
833	South Kofea	2019	Korean National Health Insurance	National	both	40+	40+	9,809,734	9,238,805	Fasting glucose	Yes	Lab		

	Country	Data years	ness both			size used Il analysis	Glycaemic markers available	Information available on diabetes		easured in a y a point-of-care e device	Notes			
				ness	both	Female	Male	Female	Male		treatment	Glucose	HbA1c	
834	South Korea	2020	Korea National Health and Nutrition Examination Survey	National	both	18+	18+	3,283	2,685	Fasting glucose, HbA1c	Yes	Lab	Lab	
835 836	South Korea	2021	Korea National Health and Nutrition Examination Survey	National National	both	18+	18+ 40+	3,203 9,735,163	2,510 9,234,806	Fasting glucose, HbA1c	Yes	Lab	Lab	igwdot
837	South Korea South Korea	2021	Korean National Health Insurance Korean National Health Insurance	National	both both	40+ 40+	40+	10,590,978	10,102,790	Fasting glucose Fasting glucose	Yes Yes	Lab Lab	$\overline{}$	2
838	Spain	1989	Cardiovascular Risk Factors Study in Catalonia	Subnational	both	18+	18+	115	109	Fasting glucose	No	Lab	i	
839	Spain	1994	DECODE; DECODE Study Group, Diabetes Care 26:61-9, 2003	Community	urban	30-89	30-89	2,699	2,108	Fasting glucose	No	Unknown	i	
840	Spain	1996	REGICOR Study; Masia et al., Rev Esp Cardiol 57:261-4, 2004	Subnational	both	25-74	25-74	874	874	Fasting glucose	No	Unknown		
841	Spain	1999	The Asturias Study; Botas, et al., Diabet Med 20:904-8, 2003	Subnational	both	30-79	30-79	542	445	Fasting glucose	No	Unknown		igwdown
842 843	Spain Spain	1999-2000 2001-2002	Factores de riesgo en las islas Baleares: Estudio CORSAIB  Catalan Health Interview Survey	Subnational Subnational	both both	35-74 18-74	35-74 18-74	865 755	804 599	Fasting glucose	Yes Yes	Lab Portable		$\vdash \vdash$
844	Spain	2001-2002	Dlabetes, Nutrición y Obesidad en la población adulta de la Región de Murcia (DINO)	Subnational	both	20+	20+	837	719	Fasting glucose Fasting glucose, HbA1c	Yes	Lab	Lab	$\vdash$
845	Spain	2000-2005	CDC of the Canary Islands	Subnational	both	18-75	18-75	3,763	2,908	Fasting glucose	Yes	Lab		$\vdash$
846	Spain	2003	The European Male Ageing Study	Community	both		40+		402	Fasting glucose	Yes	Lab	i	
847	Spain	2004	Vioque J et al., Obesity 16(3):664-70, 2008	Community	urban	24+	24+	117	87	Fasting glucose, HbA1c	Yes	Portable	Portable	
848	Spain	2004	Cardiovascular Risk Study in Castilla y León (RECCyL)	Subnational	both	18+	18+	2,003	1,816	Fasting glucose	Yes	Lab		igwdown
849 850	Spain Spain	2003-2005 2004-2006	Registre Gironi del Cor (REGICOR) PREVICTUS	Subnational National	both both	35-79 60+	35-79 60+	3,200 3,905	2,883 3,405	Fasting glucose Fasting glucose	Yes Yes	Lab Lab		$\vdash \!$
851	Spain	2004-2000	Lopez Suarez et al., Rev Esp Cardiol 61:1150-8, 2008	Community	urban	50-75	50-75	460	398	Fasting glucose	No	Unknown	(	$\vdash \vdash \vdash$
852	Spain	2008	The European Male Ageing Study	Community	both		45+		289	Fasting glucose	Yes	Lab	1	$\overline{}$
853	Spain	2007-2009	Harmonizing Equation of Risk in Mediterraneon countries EXtremadura (HERMEX)	Subnational	both	25-79	25-79	1,498	1,298	Fasting glucose, HbA1c	Yes	Lab	Lab	
854	Spain	2008-2010	Study on Nutrition and Cardiovascular Risk in Spain	National	both	18+	18+	6,803	6,123	Fasting glucose, HbA1c	Yes	Lab	Lab	igsquare
855	Spain	2009	Cardiovascular Risk Study in Castilla y León (RECCyL)	Subnational	both	20+	20+	1,579	1,299	Fasting glucose	Yes	Lab		igwdown
856 857	Spain Spain	2014	Cardiovascular Risk Study in Castilla y León (RECCyL) Study on Nutrition and Cardiovascular Risk in Spain (ENRICA)	Subnational National	both both	20+ 65+	20+ 65+	1,495 952	1,226 871	Fasting glucose Fasting glucose	Yes Yes	Lab Lab		$\vdash$
858	Spain	2016-2017	Estudio de Nutrición y Riesgo Cardiovascular en España (ENRICA)-Seniors cohort	Subnational	urban	65-94	65-94	1,739	1,532	Fasting glucose, HbA1c	Yes	Lab	Lab	$\vdash$
859	Spain	2019	Estudio de Nutrición y Riesgo Cardiovascular en España (ENRICA)-Seniors cohort	Subnational	urban	65-95	65-95	968	926	Fasting glucose, HbA1c	Yes	Lab	Lab	$\vdash$
860	Sri Lanka	2000	Malavige et al., Diabetes Res Clin Pract 57:143-5, 2002	Community	urban	30-64	30-64	621	421	Fasting glucose	No	Unknown	i	
861	Sri Lanka	2005-2006	Sri Lanka Diabetes, Cardiovascular study (SLDCS)	National	both	18+	18+	2,717	1,777	Fasting glucose	Yes	Lab		
862	Sri Lanka	2007	Pinidiyapathirage et al., Diabet Med 30:326-32, 2013	Community	urban	35-64	35-64	1,636	1,349	Fasting glucose, HbA1c	No	Lab	Lab	igspace
863 864	Sri Lanka Sri Lanka	2010	Pubudu De Silva et al., Int J Equity Health 11:76, 2012 STEPS	Subnational National	both both	35-64 18-69	35-64 18-69	606 3,082	628 2,017	Fasting glucose	No Yes	Lab Portable		$\vdash$
865	Sri Lanka	2014-2019	The Sri Lanka Health and Ageing Study (SLHAS)	National	both	18+	18+	3,239	3,109	Fasting glucose Fasting glucose, HbA1c	Yes	Lab	Lab	$\vdash$
866	Sri Lanka	2021	STEPS	National	both	18-69	18-69	3,767	2,447	Fasting glucose	Yes	Portable	1	
867	State of Palestine	1996-1998	Ramallah study	Community	rural	18-64	18-64	443	206	Fasting glucose	Yes	Portable	ı	
868	State of Palestine	1996-1998	Ramallah study	Community	urban	18-64	18-64	458	182	Fasting glucose	Yes	Portable	ı——	ш
869	State of Palestine	2010	STEPS	National	both	18-64	18-64	3,834	2,330	Fasting glucose	Yes	Unknown		igwdot
870 871	State of Palestine Sudan	2022 2005-2006	STEPS STEPS	National Subnational	both both	18-69 25-64	18-69 25-64	3,661 321	1,701 145	Fasting glucose Fasting glucose	Yes No	Portable Unknown		$\vdash \vdash$
872	Sudan	2005-2006	STEPS	National	both	18-69	18-69	4,594	2,707	Fasting glucose	Yes	Portable	i	$\vdash \vdash \vdash$
873	Suriname	2013-2015	The Healthy Life in Suriname Study (HELISUR)	Subnational	urban	18-70	18-70	722	422	Fasting glucose, HbA1c	Yes	Lab	Lab	
874	Sweden	1980-1981	Population Study of Women in Gothenburg	Community	urban	50-72		1,153		Fasting glucose	Yes	Lab	ı	
875	Sweden	1980-1985	Uppsala Longitudinal Study of Adult Men	Community	both		55-64		1,814	Fasting glucose	No	Lab	ı——	23
876	Sweden	1991	Asplund-Carlson et al., J Intern Me 236:57-64, 1994	Subnational	both	20.04	40-50	20.4	1,564	Fasting glucose	No	Unknown		igwdot
877 878	Sweden Sweden	1992-1993 1991-1995	Population Study of Women in Gothenburg Uppsala Longitudinal Study of Adult Men	Community Community	urban both	62-84	69-74	834	1,151	Fasting glucose Fasting glucose	Yes Yes	Lab Lab		23
879	Sweden	1994	DECODE; DECODE Study Group, Diabetes Care 26:61-9, 2003	Community	urban	30-79	30-79	1,120	1,058	Fasting glucose	No	Unknown	,	
880	Sweden	1994	Nilson et al., Scand J Prim Health Care 18(2):111-112, 2000	Community	urban	56-65	56-65	217	170	Fasting glucose	No	Lab	<u></u> 1	
881	Sweden	1995	MONICA Gothenburg	Community	urban	25-64	25-64	865	745	Fasting glucose	Yes	Unknown		$\Box$
882	Sweden	1997	SWESTONIA; Johansson et al., J Intern Med 252:551-60, 2002	Community	urban	35-55	35-55	135	137	Fasting glucose	No	Unknown		
883 884	Sweden Sweden	1997-2001 2003	Uppsala Longitudinal Study of Adult Men The European Male Ageing Study	Community Community	both both		73-80 40+		781 404	Fasting glucose Fasting glucose	Yes Yes	Lab Lab		23
885	Sweden	2003	Swedish INTERGENE Cohort Study	Subnational	both	24-76	24-76	1,907	1,694	Fasting glucose Fasting glucose	Yes	Lab	,	$\vdash$
886	Sweden	2001-2004	PIVUS Study	Community	both	70	70	1,527	1,512	Fasting glucose	Yes	Portable		
887	Sweden	2004	Welin et al., BMC Public Health 8:403, 2008	Community	urban	50-61	50-61	667	1,250	Fasting glucose	No	Unknown		
888	Sweden	2002-2006	Malmö Preventive Project Re-examination Study Cohort; Leosdottir et al., Cardiovasc Diabetol 10:118, 2011	Community	both	57-86	57-86	6,680	11,546	Fasting glucose	No	Lab	<u>_</u>	╙
	Sweden		Uppsala Longitudinal Study of Adult Men	Community	both	40.01	80-83	400	476	Fasting glucose, HbA1c	Yes	Lab	Lab	
890 891	Sweden Sweden	2004-2005 2004-2005	European Youth Heart Study (EYHS) II Population Study of Women in Gothenburg	Subnational Community	urban urban	18-21 38-50	18-21	109 500	68	Fasting glucose Fasting glucose	No Yes	Lab Lab		$\vdash \vdash \vdash$
	Sweden	2004-2005	The European Male Ageing Study	Community	both	50-50	45+	300	382	Fasting glucose	Yes	Lab	,	$\vdash$
	Sweden	2007-2009	PIVUS Study	Community	both	75	75	1,440	1,389	Fasting glucose	Yes	Portable	,	
894	Sweden	2008-2009	Uppsala Longitudinal Study of Adult Men	Community	both		84-88		275	Fasting glucose, HbA1c	Yes	Lab	Unknown	
	Sweden	2011-2014	PIVUS Study	Community	both	80	80	1,275	1,161	Fasting glucose	Yes	Portable	<u>_</u>	╙
	Sweden	2014-2016	Swedish INTERGENE Cohort Study	Subnational	urban	37-88	37-88	659	606	Fasting glucose	Yes	Lab		igwdot
897 898	Sweden Switzerland	2016-2017 2005	Population Study of Women in Gothenburg Bus Santé Study	Community Subnational	urban urban	38-50 20-80	20-80	573 74	53	Fasting glucose Fasting glucose	Yes Yes	Lab Lab		$\vdash \vdash \vdash$
898	Switzerland	2003-2006	Cohorte Lausannoise	Community	urban	35-75	35-75	3,454	3,104	Fasting glucose Fasting glucose	Yes	Lab	<del>,                                    </del>	$\vdash$
000	LOTTOFIU	2000 2000	and the second s	Community	G.Dan	55 15	5575	5, 757	5,104	r doing gluooso		Lab		

	Country	Data years	Survey/Study name/Citation	Level of representative-	Rural, urban, or both		e as used Il analysis		size used Il analysis	Glycaemic markers available	Information available on diabetes	Whether me laboratory or by portable	a point-of-care	Notes
				Hess	both	Female	Male	Female	Male		treatment	Glucose	HbA1c	
900	Switzerland	2006	Bus Santé Study	Subnational	urban	20-80	20-80	53	67	Fasting glucose	Yes	Lab		
901	Switzerland	2007	Bus Santé Study	Subnational	urban	20-80	20-80	46	48	Fasting glucose	Yes	Lab		
902	Switzerland	2008	Bus Santé Study	Subnational	urban	20-80	20-80	209	215	Fasting glucose	Yes	Lab		—
903 904	Switzerland Switzerland	2009 2010	Bus Santé Study Bus Santé Study	Subnational Subnational	urban urban	20-80 20-80	20-80 20-80	505 483	455 467	Fasting glucose Fasting glucose	Yes Yes	Lab Lab		<b>├</b> ──
904	Switzerland	2010	Bus Sante Study Bus Santé Study	Subnational	urban	20-80	20-80	483	434	Fasting glucose	Yes	Lab		
906	Switzerland	2009-2012	Cohorte Lausannoise	Community	urban	40-75	40-75	29	2,337	Fasting glucose	Yes	Lab		†
907	Switzerland	2012	Bus Santé Study	Subnational	urban	20-80	20-80	451	473	Fasting glucose	Yes	Lab		
908	Switzerland	2013	Bus Santé Study	Subnational	urban	20-80	20-80	514	461	Fasting glucose	Yes	Lab		1
909	Switzerland	2014	Bus Santé Study	Subnational	urban	20-80	20-80	512	464	Fasting glucose	Yes	Lab		
910	Switzerland	2015	Bus Santé Study	Subnational	urban	20-80	20-80	499	501	Fasting glucose	Yes	Lab		
911	Switzerland	2016	Bus Santé Study	Subnational	urban	20-80	20-80	496	460	Fasting glucose	Yes	Lab		
912	Switzerland	2014-2017	Cohorte Lausannoise	Community	urban	45-87	45-87	2,398	1,938	Fasting glucose, HbA1c	Yes	Lab	Lab	
913	Switzerland	2017	Bus Santé Study	Subnational	urban	20-80	20-80	577	543	Fasting glucose	Yes	Lab		
914	Switzerland	2018	Bus Santé Study	Subnational	urban	20-80	20-80	545	513	Fasting glucose	Yes	Lab		
915	Switzerland	2019	Bus Santé Study	Subnational	urban	20-80	20-80	418	350	Fasting glucose	Yes	Lab		<del></del>
916	Switzerland	2018-2021	Cohorte Lausannoise	Community	urban	49-90	49-90	1,866	1,510	Fasting glucose, HbA1c	Yes	Lab	Lab	—
917	Syrian Arab Republic	2002	National Survey on non-communicable diseases and factors affecting their development	National	both	15-65	15-65	2,958	1,784	Fasting glucose	No	Unknown		3
918	Taiwan	1993-1996	Nutrition and Health Survey in Taiwan	National	both	18+	18+	2,608	2,598	Fasting glucose	Yes	Lab		₩
919 920	Taiwan Taiwan	1996 1997	Chen et al., Diabetes Res Clin Pract 44:59-69, 1999 Chen et al., Diabetes Res Clin Pract 51:59-66, 2001	Community	both both	40-79 50-79	40-79 50-79	822 1,053	779 540	Fasting glucose	No No	Unknown Unknown		$\vdash$
920	Taiwan	1997	Lai et al., J Gerontol A Biol Sci Med Sci 55:M257-9, 2000	Community	both	65-80	65-80	1,053	387	Fasting glucose Fasting glucose	No	Unknown		+
921	Taiwan	1999-2000	Nutrition and Health Survey in Taiwan	National	both	65+	65+	754	796	Fasting glucose	Yes	Lab		+
923	Taiwan	2000	Social Environment and Biomarkers of Aging Study	National	both	50+	50+	432	590	Fasting glucose, HbA1c	Yes	Lab	Lab	+
924	Taiwan	2002	Taiwanese Survey on Hypertension, Hyperglycemia and Hyperlipidemia	National	both	18+	18+	3,215	2,944	Fasting glucose, HbA1c	No	Lab	Lab	1
925	Taiwan	2005	TCHS; Lin et al., Eur J Clin Invest 37:783-90, 2007	Community	urban	40+	40+	1,212	1,147	Fasting glucose	No	Unknown		1
926	Taiwan	2006	Social Environment and Biomarkers of Aging Study	National	both	53+	53+	604	679	Fasting glucose, HbA1c	Yes	Lab	Lab	1
927	Taiwan	2005-2008	Nutrition and Health Survey in Taiwan	National	both	19+	19+	1,363	1,327	Fasting glucose	Yes	Lab		
928	Taiwan	2007	Taiwanese Survey on Hypertension, Hyperglycemia and Hyperlipidemia	National	both	20+	20+	2,508	2,174	Fasting glucose, HbA1c	Yes	Lab	Lab	
929	Taiwan	2013-2016	Nutrition and Health Survey in Taiwan	National	both	18+	18+	2,861	2,863	Fasting glucose, HbA1c	Yes	Lab	Lab	
930	Taiwan	2017-2020	Nutrition and Health Survey in Taiwan	National	both	18+	18+	3,399	3,379	Fasting glucose, HbA1c	Yes	Lab	Lab	
931	Tajikistan	2016	STEPS	National	both	18-69	18-69	1,567	1,098	Fasting glucose	Yes	Portable		
932	Tanzania	1987	Swai et al., BMJ 305:1057-62, 1992	Community	rural	15+	18+	4,283	3,301	Fasting glucose	No	Lab		3
933	Tanzania	1997 1997	Aspray et al., Trans R Soc Trop Med Hyg 94:637-44, 2000	Community	rural	18-64	18-64	527 438	401	Fasting glucose	No No	Unknown		
934 935	Tanzania Tanzania	2011	Aspray et al., Trans R Soc Trop Med Hyg 94:637-44, 2000 STEPS	Community Subnational	urban both	18-64 25-64	18-64 25-64	1,524	332 1,011	Fasting glucose Fasting glucose	No Yes	Unknown Portable		+
936	Tanzania	2012	STEPS	National	both	25-64	25-64	2,849	2,601	Fasting glucose	Yes	Portable		+
937	Tanzania	2023	STEPS	National	both	18-69	18-69	1,945	1,459	Fasting glucose	Yes	Portable		2
938	Thailand	1983	Vannasaeng et al., J Med Assoc Thai 70 Suppl 2:126-30, 1987	Subnational	urban	20-79	20-79	681	442	Fasting glucose	No	Unknown		<del></del>
939	Thailand	1991	Thailand National Health Examination Survey I	National	both	18+	18+	7,255	5,363	Fasting glucose	Yes	Lab		
940	Thailand	1997	Thailand National Health Examination Survey II	National	both	18-59	18-59	1,682	1,022	Fasting glucose	No	Lab		1
941	Thailand	2000	InterASIA	National	both	35+	35+	3,212	2,093	Fasting glucose	Yes	Lab		
942	Thailand	2004	Thailand National Health Examination Survey III	National	both	18+	18+	19,942	18,500	Fasting glucose	Yes	Lab		
943	Thailand	2009	Thailand National Health Examination Survey IV	National	both	18+	18+	10,225	9,271	Fasting glucose	Yes	Lab		Ь
944	Thailand	2014	Thailand National Health Examination Survey V	National	both	18+	18+	10,566	7,714	Fasting glucose	Yes	Lab		<b>↓</b>
945	Thailand	2019-2020	Thailand National Health Examination Survey VI	National	both	18+	18+	12,463	8,976	Fasting glucose, HbA1c	Yes	Lab	Lab	₩
946	Timor-Leste	2009-2010	Timor-Leste Eye Health Survey	Subnational	both	40+	40+	248	246	HbA1c	Yes	Dodoblo	Lab	—
947 948	Timor-Leste Togo	2014	STEPS STEPS	National National	both both	18-69 18-64	18-69 18-64	1,465 1,967	1,080 1,897	Fasting glucose Fasting glucose	Yes Yes	Portable Portable		+-
948	Togo	2010	STEPS	National	both	18-69	18-69	2,172	1,586	Fasting glucose Fasting glucose	Yes	Portable		<del>                                     </del>
950	Tokelau	2021-2022	STEPS	National	both	18-64	18-64	267	241	Fasting glucose	Yes	Portable		$\vdash$
951	Tokelau	2014	STEPS	National	both	18-64	18-64	282	262	Fasting glucose	Yes	Portable		<b>†</b>
952	Tonga	2004	STEPS	National	both	18-64	18-64	301	237	Fasting glucose	No	Portable		
953	Tonga	2011	STEPS	National	both	18-64	18-64	1,464	928	Fasting glucose	Yes	Portable		
954	Trinidad and Tobago	2011	STEPS	National	both	18-64	18-64	1,538	1,050	Fasting glucose	Yes	Portable		
	Tunisia	1981	Papoz et al., Int J Epidemiol 17:419-22, 1988	Community	rural	20+	20+	893	618	Fasting glucose	No	Unknown		<b>†</b>
956	Tunisia	1990	Gharbi et al., Rev Epidemiol Sante Publique 50:349-55, 2002	Community	both	35-50	35-50	345	345	Fasting glucose	No	Unknown		
957	Tunisia	1996-1997	Ariana Healthy Project 1997	Community	both	35-65	35-65	2,724	2,650	Fasting glucose	Yes	Lab		
958	Tunisia	1996-1997	Tunisian National Nutrition Survey 1996-1997	National	both	18+	18+	2,674	1,397	Fasting glucose	Yes	Lab		
959	Tunisia	2001	Romdhane et al., Tunis Med 83 Suppl 5:41-6, 2005	Community	both	40-69	40-69	1,092	744	Fasting glucose	No	Unknown	_	
960	Tunisia	2005	Tunisian National Survey 2005 (TAHINA)	National	both	35-71	35-71	4,436	3,314	Fasting glucose	Yes	Lab		
961	Tunisia	2009-2010	ObeMaghreb	Subnational	urban	18-49	18-49	696	998	Fasting glucose	Yes	Lab		
962	Tunisia	2016	Tunisian Health Examination Survey	Community	both	18+	18+	4,702	4,196	HbA1c	Yes		Portable	<u> </u>
963	Türkiye	1990	Turkish Adult Risk Factor Study	National	both	20+	20+	1,130	1,028	Fasting glucose	No	Lab		<del></del>
964	Türkiye	1995	Turkish Adult Risk Factor Study	National	both	25+	25+	606	584	Fasting glucose	No	Lab		1

966 Tür 967 Tür			Data years Survey/Study name/Citation	Level of representative-ness	Rural, urban, or	Age range as used for global analysis		Sample size used for global analysis  Female Male		Glycaemic markers available	available on diabetes	etes portable device		Notes
966 Tür 967 Tür 968 Tür				ness	both	Female	Male	Female	Male		treatment	Glucose	HbA1c	
967 Tür 968 Tür	irkiye	1998	Turkish Adult Risk Factor Study	National	both	28+	28+	781	718	Fasting glucose	Yes	Lab		
968 Tür	irkiye	2000	Turkish Adult Risk Factor Study	National	both	30+	30+	952	904	Fasting glucose	Yes	Lab		
	irkiye	2001	Yumuk et al., Diabetes Res Clin Pract 70:151-8, 2005	Community	urban	20+	20+	1,789	688	Fasting glucose	No	Unknown		
	irkiye	2001	Sekuri et al., Jpn Heart J 45:119-31, 2004 The Healthy Nutrition for Healthy Heart Study	Community National	rural both	45+ 30+	30+	205 10,657	4,778	Fasting glucose Fasting glucose	No No	Unknown Lab		
	irkiye	2000-2002	Gokcel et al., Diabetes Care 26:3031-4, 2003	Subnational	both	20-79	20-79	1,030	607	Fasting glucose Fasting glucose	No	Unknown		
	irkiye	2002	Soysal et al., Anadolu Kardiyol Derg 5:196-201, 2005	Subnational	urban	25-39	25-39	469	289	Fasting glucose	No	Unknown		_
	irkiye	2001-2002	Turkish Adult Risk Factor Study	National	both	32+	32+	1,247	1,137	Fasting glucose	Yes	Lab		
	irkiye	2003-2005	Prevalence of prehypertension and associated risk factors among Turkish adults: Trabzon Hypertension Study	Subnational	both	20+	20+	2,601	2,208	Fasting glucose	No	Lab		1
974 Tür	irkiye	2005-2006	Turkish Adult Risk Factor Study	National	both	35+	35+	1,473	1,401	Fasting glucose	Yes	Lab		
	irkiye	2007-2009	Balcova Heart Study	Community	urban	30+	30+	8,441	4,187	Fasting glucose	Yes	Lab		
	irkiye	2007-2008	Turkish Adult Risk Factor Study	National	both	37+	37+	1,473	1,401	Fasting glucose	Yes	Lab		
	irkiye	2009-2010	Turkish Adult Risk Factor Study	National	both	39+	39+	1,474	1,403	Fasting glucose	Yes	Lab		
	irkiye irkiye	2010 2010	TURDEP-II; Satman et al., Eur J Epidemiol 28:169-80, 2013 TURDEP-II; Satman et al., Eur J Epidemiol 28:169-80, 2013	National National	urban rural	20+	20+ 20+	9,943 6,578	5,840 3,863	Fasting glucose Fasting glucose	No No	Lab Lab		
	irkiye	2009-2012	Prevalence of diabetes and associated risk factors among adult population in Trabzon city	Subnational	both	20+	20+	2,125	1,574	Fasting glucose	No	Lab		_
	irkiye	2012-2013	Turkish Adult Risk Factor Study	National	both	37+	37+	1,115	1,028	Fasting glucose, HbA1c	Yes	Lab	Lab	
	irkiye	2014-2015	Turkish Adult Risk Factor Study	National	both	44+	44+	756	712	Fasting glucose	No	Lab	-	
	irkiye	2017	STEPS	National	both	18+	18+	3,493	2,369	Fasting glucose	Yes	Portable		
	urkmenistan	2013	STEPS	National	both	18-64	18-64	2,874	2,033	Fasting glucose	Yes	Portable		$oxed{oxed}$
	urkmenistan	2018	STEPS	National	both	18-69	18-69	2,244	1,716	Fasting glucose	Yes	Portable		
	ıvalu	2015	STEPS	National	both	18-69	18-69	616	525	Fasting glucose	Yes	Portable		
	ganda	2014	STEPS	National	both	18-69	18-69	2,132	1,570	Fasting glucose	Yes	Portable		
	ganda kraine	2023	STEPS STEPS	National National	both both	18-69 18-69	18-69 18-69	2,069 2,662	1,426 1,610	Fasting glucose	Yes	Portable Portable		2
gan Uni	nited Arab mirates	1999-2000	Emirates National Diabetes and Coronary Artery Disease Risk Factor Study	National	both	20-80	20-80	3,773	2,839	Fasting glucose Fasting glucose	Yes No	Lab		
	nited Arab mirates	2017-2018	STEPS	National	both	18+	18+	2,356	2,172	Fasting glucose	Yes	Portable		
992 Uni	nited Kingdom	1984-1986	Scottish Heart Health Survey	Subnational	both	40-59	40-59	4,447	4,381	Fasting glucose	Yes	Lab		
	nited Kingdom	1987-1988	Edinburgh Artery Study	Community	urban	54-75	54-75	769	797	Fasting glucose	No	Lab		
	nited Kingdom	1993	DECODE; DECODE Study Group, Diabetes Care 26:61-9, 2003	Community	urban	30-79	30-79	384	415	Fasting glucose	No	Unknown		
	nited Kingdom nited Kinadom	1993 1998-2000	Whickham Survey; Vanderpump et al., Diabet Med 13:741-7, 1996 The British Regional Heart Study	Community	urban	35+	35+ 60-79	938	761 4,105	Fasting glucose	No	Unknown		
	nited Kingdom nited Kingdom	1998-2000	British Women's Heart and Health Study	National National	urban both	60-79	60-79	3,909	4,105	Fasting glucose Fasting glucose	Yes Yes	Lab Lab		24
	nited Kingdom	1999-2004	Hertfordshire Cohort Study	Subnational	both	59-73	59-73	1,418	1,579	Fasting glucose	Yes	Lab		2.7
	nited Kingdom	2003	The European Male Ageing Study	Community	both	00.10	40+	1,110	389	Fasting glucose	Yes	Lab		
	nited Kingdom	2003	Health Survey for England	National	both	18+	18+	7,823	6,408	Fasting glucose, HbA1c	Yes	Lab	Lab	
1001 Uni	nited Kingdom	2003	Scottish Health Survey (SHeS)	Subnational	both	18+	18+	4,370	3,497	HbA1c	Yes		Lab	
	nited Kingdom	2003-2005	Hertfordshire Ageing Study	Subnational	both	72-82	72-82	151	208	Fasting glucose, HbA1c	Yes	Lab	Lab	
	nited Kingdom	2004-2005	English Longitudinal Study of Ageing Wave 2	National	both	52-80	52-80	3,715	3,153	Fasting glucose, HbA1c	Yes	Lab	Lab	
	nited Kingdom	2005	Health Survey for England	National	both	65+	65+	2,372	1,897	HbA1c	Yes		Lab	
	nited Kingdom nited Kingdom	2006 2006-2007	Health Survey for England Newcastle 85+ Study	National Community	both urban	18+ 84+	18+ 84+	6,561 431	5,444 283	HbA1c Fasting glucose, HbA1c	Yes No	Lab	Lab Lab	25
	nited Kingdom	2008	The European Male Ageing Study	Community	both	04+	45+	431	335	Fasting glucose, HDATC	Yes	Lab	Lau	25
	nited Kingdom	2008	Health Survey for England	National	both	18+	18+	7,960	6,550	HbA1c	Yes		Lab	
	nited Kingdom	2007-2009	Newcastle 85+ Study	Community	urban	85+	85+	311	187	HbA1c	No		Lab	25
	nited Kingdom	2008	Scottish Health Survey (SHeS)	Subnational	both	18+	18+	3,475	2,772	HbA1c	Yes		Lab	
	nited Kingdom	2008-2009	English Longitudinal Study of Ageing Wave 4	National	both	50+	50+	4,499	3,674	Fasting glucose, HbA1c	Yes	Lab	Lab	ldot
	nited Kingdom	2009	Health Survey for England	National	both	18+	18+	2,420	2,050	HbA1c	Yes		Lab	$\longmapsto$
	nited Kingdom	2006-2010	MRC National Survey of Health and Development	National	both	60-65	60-65	1,068	969	Fasting glucose, HbA1c	Yes	Lab	Lab	$\vdash$
	nited Kingdom	2009	Scottish Health Survey (SHeS) Health Survey for England	Subnational National	both	18+ 18+	18+ 18+	4,089 4,486	3,226 3,598	HbA1c HbA1c	Yes		Lab Lab	$\vdash$
	nited Kingdom nited Kingdom	2010	National Diet and Nutrition Survey (NDNS)	National National	both both	18+	18+ 18+	4,486 914	3,598 674	Fasting glucose, HbA1c	Yes No	Lab	Lab	$\vdash$
	nited Kingdom	2009-2010	Newcastle 85+ Study	Community	urban	85+	85+	258	160	Fasting glucose, HbA1c	No	Lab	Lab	25
	nited Kingdom	2010	Scottish Health Survey (SHeS)	Subnational	both	18+	18+	3,981	3,047	HbA1c	Yes		Lab	
1019 Uni	nited Kingdom	2010	Understanding Society: the UK Household Longitudinal Study	National	both	18+	18+	4,709	3,716	HbA1c	No		Lab	26
	nited Kingdom	2011	British Household Panel Survey	National	both	18+	18+	1,385	1,157	HbA1c	No		Lab	26
	nited Kingdom	2010-2012	The British Regional Heart Study	National	urban		72-91		1,671	Fasting glucose, HbA1c	Yes	Lab	Unknown	igspace
	nited Kingdom	2011	Health Survey for England	National	both	18+	18+	4,584	3,725	HbA1c	Yes		Lab	$\longmapsto$
	nited Kingdom	2011	Scottish Health Survey (SHeS)	Subnational	both	18+	18+	4,093	3,219	HbA1c	Yes		Lab	$\vdash$
	nited Kingdom	2012	Health Survey for England  English Longitudinal Study of Agains Ways 6	National	both	18+	18+	4,447 4,874	3,580	HbA1c	Yes	Loh	Lab Lab	$\vdash$
	nited Kingdom nited Kingdom	2012-2013	English Longitudinal Study of Ageing Wave 6 Health Survey for England	National National	both both	50+ 18+	50+ 18+	4,874	3,945 3,815	Fasting glucose, HbA1c HbA1c	Yes Yes	Lab	Lab	$\vdash$
	nited Kingdom nited Kingdom	2013	Health Survey for England Health Survey for England	National	both	18+	18+	4,853	3,815	HbA1c	Yes		Lab	$\vdash$
	nited Kingdom	2013-2014	National Diet and Nutrition Survey (NDNS)	National	both	18+	18+	407	281	Fasting glucose, HbA1c	No	Lab	Lab	$\vdash$
	nited Kingdom	2015	Health Survey for England	National	both	18+	18+	4,290	3,503	HbA1c	Yes		Lab	

	Country	Data years	Survey/Study name/Citation	Level of representative-	Rural, urban, or		e as used Il analysis		size used I analysis	Glycaemic markers available	Information available on diabetes	laboratory or b	easured in a y a point-of-care e device	Notes
				ness	both	Female	Male	Female	Male		treatment	Glucose	HbA1c	
1030	United Kingdom	2015	MRC National Survey of Health and Development	National	both	69-70	69-70	1,016	949	HbA1c	Yes		Lab	
1031	United Kingdom	2016	Health Survey for England	National	both	18+	18+	4,274	3,461	HbA1c	Yes		Lab	1
1032	United Kingdom United Kingdom	2015-2016 2015-2018	National Diet and Nutrition Survey (NDNS) British Cohort Study 1970	National National	both both	18+ 45-48	18+ 45-48	370 4,426	273 4,154	Fasting glucose, HbA1c HbA1c	No Yes	Lab	Lab Lab	
1034	United Kingdom	2016-2017	English Longitudinal Study of Ageing Wave 8	National	both	50+	50+	3,650	2,891	Fasting glucose, HbA1c	Yes	Lab	Lab	
1035	United Kingdom	2017	Health Survey for England	National	both	18+	18+	4,318	3,454	HbA1c	Yes		Lab	
1036	United Kingdom	2018	Health Survey for England	National	both	18+	18+	4,347	3,576	HbA1c	Yes		Lab	
1037	United Kingdom	2016-2019	National Diet and Nutrition Survey (NDNS)	National	both	18+	18+	318	234	Fasting glucose, HbA1c	No	Lab	Lab	
1038	United Kingdom	2018-2019	English Longitudinal Study of Ageing Wave 9	National	both	50+	50+	3,981	3,053	Fasting glucose, HbA1c	Yes	Lab	Lab	$\vdash$
1039 1040	United Kingdom United Kingdom	2019 2019	Health Survey for England Understanding Society: Innovation Panel	National National	both both	18+ 18+	18+ 18+	4,363 214	3,588 181	HbA1c HbA1c	Yes No		Lab Lab	$\vdash$
1041	United States of America	1976-1980	US NHANES II	National	both	18-74	18-74	6,236	5,594	Fasting glucose	Yes	Lab	Edb	27
1042	United States of America	1985-1986	Coronary Artery Risk Development in Young Adults (CARDIA)	Subnational	urban	18-30	18-30	2,633	2,198	Fasting glucose	No	Lab		
1043	United States of America	1987-1989	Atherosclerosis Risk in Communities Study	Subnational	both	44-66	44-66	6,222	5,051	Fasting glucose	Yes	Lab		
1044	United States of America	1989-1990	Cardiovascular Health Study	Subnational	both	65+	65+	3,321	2,462	Fasting glucose	Yes	Lab		$\sqcup \sqcup$
1045	United States of America United States of	1990-1992	Atherosclerosis Risk in Communities Study	Subnational	both	46-70	46-70	5,631	4,546	Fasting glucose	Yes	Lab		$\vdash \vdash \vdash$
1046	America United States of	1988-1994	US NHANES III	National	both	20+	20+	9,702	8,806	Fasting glucose	Yes	Lab		
1047	America United States of	1992-1993	Coronary Artery Risk Development in Young Adults (CARDIA)	Subnational	urban	25-37	25-37	1,979	1,670	Fasting glucose	No	Lab		
1048	America United States of	1992-1993	Cardiovascular Health Study	Subnational	both	65+	65+	2,971	2,105	Fasting glucose	Yes	Lab		
1049	America United States of	1993-1995	Atherosclerosis Risk in Communities Study	Subnational	both	48-73	48-73	5,019	4,004	Fasting glucose	Yes	Lab		$\vdash$
1050	America United States of	1995-1996 1995-1996	The Bogalusa Heart Study  Coronary Artery Risk Development in Young Adults (CARDIA)	Community	rural	20-37	20-37	790 1,923	515 1,595	Fasting glucose Fasting glucose	No No	Lab Lab		$\vdash$
1051	America United States of	1996-1998	Atherosclerosis Risk in Communities Study	Subnational	both	50-75	50-75	4,497	3,561	Fasting glucose	Yes	Lab		
1053	America United States of	1996-1997	Cardiovascular Health Study	Subnational	both	67+	67+	2,504	1,567	Fasting glucose	Yes	Lab		
1054	America United States of America	1996-1997	Study of Women's Health Across the Nation	Subnational	both	40-55		3,240		Fasting glucose	Yes	Lab		28
1055	United States of America	1997-1999	Study of Women's Health Across the Nation	Subnational	both	40-55		2,862		Fasting glucose	Yes	Lab		28
1056	United States of America	1999-2000	US NHANES 1999-2000	National	both	18+	18+	2,586	2,546	Fasting glucose, HbA1c	Yes	Lab	Lab	
1057	United States of America	1999-2001	Study of Women's Health Across the Nation	Subnational	both	40-56		2,694		Fasting glucose	Yes	Lab		28
1058	United States of America	2000-2001	Coronary Artery Risk Development in Young Adults (CARDIA)	Subnational	urban	33-45	33-45	1,823	1,474	Fasting glucose	No	Lab		
1059	United States of America	2000-2002	Study of Women's Health Across the Nation	Subnational	both	40-57		2,658		Fasting glucose	Yes	Lab		28
1060	United States of America	2001-2002	US NHANES 2001-2002	National	both	18+	18+	2,800	2,844	Fasting glucose, HbA1c	Yes	Lab	Lab	igsquare
1061	United States of America	2003-2004	US NHANES 2003-2004	National	both	18+	18+	2,654	2,695	Fasting glucose, HbA1c	Yes	Lab	Lab	
1062	United States of America	2004	2004 New York City HANES	Community	urban	20+	20+	1,129	824	Fasting glucose, HbA1c	Yes	Lab	Lab	$\vdash \vdash \vdash$
1063	United States of America United States of	2005-2006	Coronary Artery Risk Development in Young Adults (CARDIA)	Subnational	urban	38-50	38-50	2,010	1,531	Fasting glucose	Yes	Lab		<del>                                     </del>
1064	America United States of		Cardiovascular Health Study	Subnational	both	70+	70+	1,018	536	Fasting glucose	Yes	Lab		<b> </b>
1065	America United States of	2005-2006	US NHANES 2005-2006	National	both	18+	18+	2,519	2,671	Fasting glucose, HbA1c	Yes	Lab	Lab	$\vdash$
1066	America United States of	2007-2008	US NHANES 2007-2008	National	both	18+	18+	3,100	3,063	Fasting glucose, HbA1c	Yes	Lab	Lab	20
1067	America United States of	2008-2009	National Longitudinal Study of Adolescent Health Wave IV  US NHANES 2009-2010	National National	both	24-34	24-34	2,761 3,281	2,353 3,174	HbA1c	Yes	Lab	Lab Lab	29
1068	America United States of	2009-2010	US NHANES 2009-2010  Coronary Artery Risk Development in Young Adults (CARDIA)	Subnational	urban	43-55	43-55	1,971	1,510	Fasting glucose, HbA1c Fasting glucose	Yes Yes	Lab	Lab	
1009	America	2010-2011	Contribing Artery reson Development in Touring Adults (CARDIA)	Subhational	uibaii	43-33	43-33	1,971	1,510	rasilily glucose	162	Lau		

	Country	Data years	Survey/Study name/Citation	Level of representative-	Rural, urban, or		e as used Il analysis		size used Il analysis	Glycaemic markers available	Information available on diabetes	n portable device		Notes
					both	Female	Male	Female	Male	]	treatment	Glucose	HbA1c	Ī
1070	United States of America	2011-2013	Atherosclerosis Risk in Communities Study		both	67-90	67-90	2,210	1,605	Fasting glucose, HbA1c	Yes	Lab	Lab	
1071	United States of America	2011-2012	US NHANES 2011-2012	National	both	18+	18+	2,907	2,894	Fasting glucose, HbA1c	Yes	Lab	Lab	
1072	United States of America	2013-2014	US NHANES 2013-2014	National	both	18+	18+	3,128	2,914	Fasting glucose, HbA1c	Yes	Lab	Lab	
1073	United States of America	2015-2016	US NHANES 2015-2016		both	18+	18+	3,030	2,886	Fasting glucose, HbA1c	Yes	Lab	Lab	
1074	United States of America	2017-2018	US NHANES 2017-2018	National	both	18+ 18+		2,959	2,839	Fasting glucose, HbA1c	Yes	Lab	Lab	
1075	United States of America	2019-2020	US NHANES 2019-2020	Subnational	both	18+	18+ 18+ 1,92		1,878	Fasting glucose, HbA1c	Yes	Lab	Lab	30
1076	Uruguay	1992	Enfermedades Cardiovasculares	National	both	19+ 19+		574	449	Fasting glucose	No	Unknown		
1077	Uruguay	2011-2012	CESCAS Study	Community	urban	35-74	35-74	903	639	Fasting glucose	Yes	Lab		
1078	Uruguay	2013	STEPS	National	urban	18-64	18-64	1,468	872	Fasting glucose	Yes	Lab		
1079	Uruguay	2012-2016	Genotype, Phenotype and Environment of Hypertension in Uruguay (GEFA-HT-UY)	Community	urban	19+	19+	193	130	Fasting glucose	Yes	Lab		
1080	Uzbekistan	2014	STEPS	National	both	18-64	18-64	2,176	1,543	Fasting glucose	Yes	Portable		
1081	Uzbekistan	2015-2016	Epidemiology of Diabetes and Prediabetes in Uzbekistan Screening Results	Subnational	both	35+	35+	1,511	714	Fasting glucose, HbA1c	Yes	Portable	Lab	
1082	Uzbekistan	2019	STEPS	National	both	18-69	18-69 18-69 2,260		1,488	Fasting glucose	Yes	Portable		
1083	Vanuatu	1998	Vanuatu Non-communicable Disease Survey	National	both	20-59	20-59	179	185	Fasting glucose	No	Unknown		
1084	Vanuatu	2011	STEPS	National	both	25-64	25-64	2,236	2,312	Fasting glucose	Yes	Portable		
1085	Venezuela	2000	Zulia Coronary Heart Disease Risk Factor Study; Florez et al., Diabetes Res Clin Pract 69:63-77, 2005	Subnational	both	25+	25+	2,552	1,120	Fasting glucose	No	Unknown		
1086	Venezuela	1998-2001	Maracaibo aging study Santa lucia cohort	Community	urban	55+	55+	1,609	787	Fasting glucose	Yes	Lab		
1087	Venezuela	2004-2005	CArdiovascular Risk factors Multiple Evaluation in Latin America (CARMELA)	Community	urban	25-64	25-64	1,135	713	Fasting glucose	Yes	Lab		1
1088	Venezuela	2005-2006	Brajkovich et al., Rev Ven Endoc Metab 4(3):31-32, 2006	Community	urban	20-65	20-65	331	114	Fasting glucose	No	Lab		
1089	Venezuela	2007-2008	Venezuelan Study of Metabolic Syndrome, Obesity and Lifestyle (VEMSOLS)	Community	urban	20+	20+	229	107	Fasting glucose	Yes	Lab		1
1090	Venezuela	2008-2009	Venezuelan Study of Metabolic Syndrome, Obesity and Lifestyle (VEMSOLS)	Community	rural	20+	20+	89	51	Fasting glucose	Yes	Lab		1
1091	Venezuela	2010-2011	Cardiometabolic risk factors in schoolchildren and adolescents of Mérida, Venezuela (CREDEFAR)	Community	urban	18	18	13	12	Fasting glucose	Yes	Lab		1
1092	Venezuela	2010-2011	Venezuelan Study of Metabolic Syndrome, Obesity and Lifestyle (VEMSOLS)	Community	urban	20+	20+	154	51	Fasting glucose	No	Lab		
1093	Venezuela	2015-2017	Cardio-Metabolic Health Venezuelan Study (EVESCAM)	National	both	20+	20+	2,347	1,059	Fasting glucose	Yes	Lab		1
1094	Venezuela	2014-2017	Maracaibo aging study Santa Rosa cohort	Community	urban	37+	37+	292	116	Fasting glucose, HbA1c	Yes	Lab	Lab	
1095	Venezuela	2018-2020	Cardio-metabolic Health Venezuelan Study (EVESCAM) follow-up	National	both	22+	22+	890	354	Fasting glucose	Yes	Lab		
1096	Viet Nam	1990	Quoc et al., Am J Epidemiol 139:713-22, 1994	Community	both	25-69	25-69	2,061	1,712	Fasting glucose	No	Unknown		
1097	Viet Nam	2001	Duc Son et al., Diabet Med 21:371-6, 2004	Community	both	25+	25+	2,001	654	Fasting glucose	No	Unknown		1
1098	Viet Nam	2005	Non-communicable disease risk factors in Ho Chi Minh City	Community	urban	25-64	25-64	1,029	887	Fasting glucose	No	Portable		
1099	Viet Nam	2008-2009	The Survey on Diabetes and Its Risk Factors	Subnational	both	25+ 25+		1,357	795	Fasting glucose, HbA1c	Yes	Lab	Lab	1
1100	Viet Nam	2009	STEPS	National	both	25-64 25-64		7,754	6,677	Fasting glucose	Yes	Portable		
1101	Viet Nam	2012	National Survey of Diabetes in Vietnam	National	both	30-69	30-69	3,869	3,495	Fasting glucose	No	Portable		
1102	Viet Nam	2015	STEPS	National	both	18-69 18-69		2,043	1,669	Fasting glucose	Yes	Portable		
1103	Viet Nam	2021	STEPS	National	both	18+ 18+		2,172	2,200	Fasting glucose	Yes	Portable		
1104	Yemen	2007-2009	Hypertension and Diabetes in Yemen (HYDY)	National	rural	18-70 18-70		2,408	2,358	Fasting glucose	Yes	Portable		
1105	Yemen	2007-2009	Hypertension and Diabetes in Yemen (HYDY)	National	urban	18-70 18-70		2,432	2,351	Fasting glucose	Yes	Portable		
1106	Zambia	2008	STEPS	Subnational	urban	25+ 25+		1,219	631	Fasting glucose	Yes	Unknown		
1107	Zambia	2017	STEPS	National	both	18-69	18-69	2,513	1,614	Fasting glucose	Yes	Portable		
1108	Zimbabwe	2005	Zimbabwe Non-Communicable Disease Risk Factors (ZiNCoDs/STEPS)	National	both	25+	25+	1,393	444	Fasting glucose	No	Lab		

- 1. Sciensano, OD Public health and surveillance (2020), Health Interview Survey 2018 [Data file and code book]. Conditionally obtainable from the Sciensano website: https://www.sciensano.be/en/node/55737/health-interview-survey-microdata-request-procedure.
- $2.\ National\ studies\ from\ 2023\ were\ assigned\ to\ 2022\ so\ that\ they\ inform\ the\ estimates\ in\ countries\ with\ slightly\ later\ national\ data.$
- The first age group started from 15 years old, but had mean age ≥18 years.
- 4. The bibliographic citation for this data source is: Zeng, Yi, and Vaupel, James W. Chinese Longitudinal Healthy Longevity Survey (CLHLS), Biomarkers Datasets, 2009, 2012, 2014. Inter-university Consortium for Political and Social Research (distributor), 2019-01-15. https://doi.org/10.3886/ICPSR37226.v1.
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- 6. The bibliographic citation for this data source is: Center for Healthy Aging and Development Studies, 2020, "The Chinese Longitudinal Healthy Longevity Survey (CLHLS)-Longitudinal Data (1998-2018) ", https://doi.org/10.18170/DVN/WBO7LK, Peking University Open Research Data Platform, V2.
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- 12. Data on diabetes treatment were collected in the questionnaire from all participants, and at the time of examination which was done in randomly selected subset of 2,500 of the 30,000 PSUs. At the time of this analysis, only the data on diabetes treatment collected at the time of physical examination was available to us and was used in the analysis.
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- 15. The National Nutrition Survey of Kuwait (2008-2009) which was supported by The Kuwait Foundation for Advancement Sciences (KFAS) Grant # 2003-1202-02.
- 16. The MHAS Cognitive Aging Ancillary Study (Mex-Cog) is sponsored by the National Institutes of Health/National Institute on Aging (NIH R01AG051158). Data files and documentation are public use and available at www.MHASweb.org.

- 17. The Longitudinal Aging Study Amsterdam is supported by a grant from the Netherlands Ministry of Health Welfare and Sports, Directorate of Long-Term Care.
- 18. Data on 1,216 participants from one specific area in this study, which had high prevalence of the high thalassemia, were excluded. Glycaemic measurements in these participants were systematically different from the rest from the same study, possibly because of the high thalassemia prevalence.
- 19. The bibliographic citation for this data source is: Am J Hypertens 2009 Jan;22(1):100-5 and Atherosclerosis. 2009 Mar;203(1):257-62.
- 20. Dr Take Naseri (Ministry of Health, Samoa), and Muagututia Sefuiva Reupena (Lutia I Puava Ae Mapu I Fagalele) contributed to the GWAS studies in Samoa.
- 21. The SP2 and SCCS2 studies are supported by individual research and clinical scientist award schemes from National Medical Research Council (NMRC) and the Biomedical Research Council (BMRC) of Singapore, the Singapore Ministry of Health, National University of Singapore and University Health System, Singapore
- 22. The SH2012 and SH2 studies are supported by infrastructure funding from the Singapore Ministry of Health (Population Health Metrics Population Health Metrics and Analytics PHMA), National University of Singapore and National University Health System, Singapore.
- 23. The ULSAM study was supported by Uppsala University and Uppsala University Hospital.
- 24. The British Women's Heart and Health Study is supported by the British Heart Foundation (PG/13/66/30442). British Women's Heart and Health Study data are available to bona fide researchers for research purposes. Please refer to the BWHHS data sharing policy at http://www.ucl.ac.uk/british-womens-heart-health-study.
- 25. The Newcastle 85+ Study has been funded by the Medical Research Council, Biotechnology and Biological Sciences Research Council, the Dunhill Medical Trust and the National Institute for Health Research School for Primary Care. Parts of the work have also been funded by the British Heart Foundation, Unilever Corporate Research, Newcastle University, NHS North of Tyne (Newcastle Primary Care Trust). Morthof Length Research Clinical Research Network North East and North Cumbria, local general practitioners and their staff. We thank the research nurses, laboratory technicians, data management and clerical team for their work throughout, as well as many colleagues for their expert advice. Thanks are due especially to the study participants and, where appropriate, their families and carers.
- 26. University of Essex. Institute for Social and Economic Research and National Centre for Social Research, Understanding Society: Waves 2 and 3 Nurse Health Assessment, 2010-2012 [data collection]. 5th Edition. UK Data Service. SN:7251. http://doi.org/10.5255/UKDA-SN-7251-5.
- 27. National studies for the 3 years prior to 1980 were assigned to 1980 so that they inform the estimates in countries with slightly earlier national data.
- 28. The bibliographic citation for this data source is: Sutton-Tyrrell, Kim, Faith Selzer, MaryFran Sowers, Robert Neer, Lynda Powell, Ellen Gold, Gail Greendale, Gerson Weiss, Karen Matthews, and Sonja McKinlay. Study of Women's Health Across the Nation (SWAN), 1996-1997: Baseline Dataset. ICPSR28762-v2. Ann Arbor, Ml: Inter-university Consortium for Political and Social Research[distributor], 2014-02-04. http://doi.org/10.3886/ICPSR28762-v2.
- 29. This research uses data from Add Health, a program project designed by J. Richard Udry, Peter S. Bearman, and Kathleen Mullan Harris, and funded by a grant P01-HD31921 from the Eunice Kennedy Shriver National Institute of Child Health and Human Development, with cooperative funding from 17 other agencies. Special acknowledgment is due Ronald R. Rindfuss and Barbara Entwisle for assistance in the original design. Persons interested in obtaining data files from Add Health should contact Add Health Should contact Add Health (and Institute of Child Health (and Institute of Child Health (and Institute of Child Health and Human Development, with cooperative funding from 17 other agencies. Special acknowledgment is due Ronald R. Rindfuss and Barbara Entwisle for assistance in the original design. Persons interested in obtaining data files from Add Health (and Institute of Child Health (and Institute of Child Health and Human Development, with cooperative funding from 17 other agencies. Special acknowledgment is due Ronald R. Rindfuss and Barbara Entwisle for assistance in the original design. Persons interested in obtaining data files from Add Health (and Institute of Child Health (and Institute of Child Health and Human Development, with cooperative funding from 17 other agencies. Special acknowledgment is due Ronald R. Rindfuss and Barbara Entwisle for assistance in the original design. Persons interested in obtaining data files from Add Health (and Institute of Child Health (and Institute of Ch
- 30. Due to the COVID-19 pandemic the NHANES 2019-2020 cycle was not completed. As a result the data are not nationally representative and were considered subnational.

Appendix Table 2. List of analysis regions and super-regions, and countries in each region.

Super-region	Region
Central and eastern	<b>Central Europe</b> : Albania, Bosnia and Herzegovina, Bulgaria, Croatia, Czechia, Hungary, Montenegro, North Macedonia, Poland, Romania, Serbia, Slovakia, Slovenia
Europe	<b>Eastern Europe</b> : Belarus, Estonia, Latvia, Lithuania, Moldova, Russian Federation, Ukraine
Control Asia Middle	<b>Central Asia</b> : Armenia, Azerbaijan, Georgia, Kazakhstan, Kyrgyzstan, Mongolia, Tajikistan, Turkmenistan, Uzbekistan
Central Asia, Middle East and north Africa	<b>Middle East and north Africa</b> : Algeria, Bahrain, Egypt, Iran, Iraq, Jordan, Kuwait, Lebanon, Libya, Morocco, Oman, Qatar, Saudi Arabia, State of Palestine, Syrian Arab Republic, Tunisia, Türkiye, United Arab Emirates, Yemen
	<b>High-income English-speaking countries*</b> : Australia, Canada, Ireland, New Zealand, United Kingdom, United States of America
High-income western	Northwestern Europe: Austria, Belgium, Denmark, Finland, Germany, Greenland, Iceland, Luxembourg, Netherlands, Norway, Sweden, Switzerland
	<b>Southwestern Europe</b> : Andorra, Cyprus, France, Greece, Israel, Italy, Malta, Portugal, Spain
	Andean Latin America: Bolivia, Ecuador, Peru
Latin America and the Caribbean	<b>The Caribbean</b> : Antigua and Barbuda, Bahamas, Barbados, Belize, Bermuda, Cuba, Dominica, Dominican Republic, Grenada, Guyana, Haiti, Jamaica, Puerto Rico, Saint Kitts and Nevis, Saint Lucia, Saint Vincent and the Grenadines, Suriname, Trinidad and Tobago
	Central Latin America: Colombia, Costa Rica, El Salvador, Guatemala, Honduras, Mexico, Nicaragua, Panama, Venezuela
	Southern Latin America: Argentina, Brazil, Chile, Paraguay, Uruguay
	Melanesia: Fiji, Papua New Guinea, Solomon Islands, Vanuatu
Pacific island nations	Polynesia and Micronesia: American Samoa, Cook Islands, French Polynesia, Kiribati, Marshall Islands, Federated States of Micronesia, Nauru, Niue, Palau, Samoa, Tokelau, Tonga, Tuvalu
South Asia	South Asia: Afghanistan, Bangladesh, Bhutan, India, Nepal, Pakistan, Sri Lanka
Fact and southeest	East Asia and the Pacific: China, Japan, Singapore, South Korea, Taiwan
East and southeast Asia and the Pacific	Southeast Asia: Brunei Darussalam, Cambodia, Indonesia, Lao PDR, Malaysia, Maldives, Myanmar, North Korea, Philippines, Thailand, Timor-Leste, Viet Nam
	<b>Central and southern Africa</b> : Angola, Botswana, Central African Republic, Congo, DR Congo, Equatorial Guinea, Gabon, Namibia
Sub-Saharan Africa	<b>East Africa</b> : Burundi, Comoros, Djibouti, Eritrea, Eswatini, Ethiopia, Kenya, Lesotho, Madagascar, Malawi, Mozambique, Rwanda, Somalia, South Sudan, Sudan, Tanzania, Uganda, Zambia, Zimbabwe
	<b>West Africa</b> : Benin, Burkina Faso, Cabo Verde, Cameroon, Chad, Côte d'Ivoire, Gambia, Ghana, Guinea, Guinea Bissau, Liberia, Mali, Mauritania, Niger, Nigeria, Sao Tome and Principe, Senegal, Sierra Leone, Togo
	Other sub-Saharan Africa: Mauritius, Seychelles, South Africa

<sup>\*</sup> Although high-income English-speaking countries are geographically separated, they experienced similar trends in cardiometabolic risk factors and outcomes. 1,2,16,32-38 They were therefore grouped together so that the statistical model shares information amongst them more than it does with other countries that are geographically closer but epidemiologically more distinct.

**Appendix Table 3.** Global age-standardised diabetes prevalence and number of people with diabetes in 1990 and 2022 for women and men aged 18+ years and 30+ years. Numbers in brackets are 95% credible intervals.

			18+ years	30+ years					
	Women	1990	6.9 (5.7-8.1)	9.1 (7.6-10.7)					
Prevalence	vvoilleii	2022	2022 13.9 (12.3-15.8) 17.3 (						
(%)	Men	1990	6.8 (5.7-8.0)	8.9 (7.6-10.4)					
	Wen	2022	14.3 (12.5-16.4)	17.9 (15.7-20.4)					
	Women	1990	103.7 (86.1-123.0)	97.0 (81.3-114.3)					
Number	vvoillen	2022	419.8 (372.2-473.5)	381.3 (340.0-428.4)					
(millions)	Men	1990	93.8 (78.4-110.9)	86.1 (72.5-100.8)					
	MEH	2022	408.0 (356.2-466.8)	367.4 (322.2-418.1)					

**Appendix Table 4.** Comparison of age-standardised diabetes prevalence estimates in this work with recent global studies.

		Ong et al., 2023 <sup>39</sup>	Sun et al., 2022 <sup>40</sup>	NCD-RisC (this study)
Women	0+ years	5.8 (5.4-6.1)	n.a.	n.a.
	20-79 years	n.a.	10.2	14.1 (12.4-16.0)
	18+ years	n.a.	n.a.	13.9 (12.3-15.8)
Men	0+ years	6.5 (6.2-7.0)	n.a.	n.a.
	20-79 years	n.a.	10.8	14.6 (12.7-16.7)
	18+ years	n.a.	n.a.	14.3 (12.5-16.4)

Estimates from Ong et al. (2023) and Sun et al. (2022) are for 2021. Estimates from this study are for 2022. n.a. = not available.

**Appendix Table 5.** Comparison of data sources and estimates of diabetes prevalence in this work with recent global studies, for the 30 countries with the largest adult population in 2022. For each country, the table shows the year of the most recent data source used and diabetes prevalence estimates.

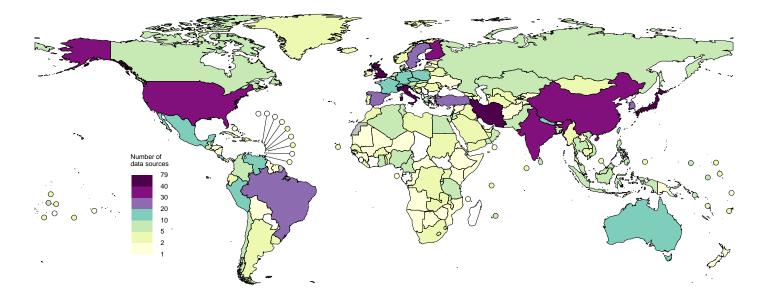
Country	Years of the most recent data sources †			Estimates of age-standardised diabetes prevalence (%) $^{\ddagger}$			
	Ong et al., 2023 <sup>39</sup>	Sun et al., 2022 <sup>40</sup>	NCD-RisC (this study)	Ong et al., 2023 <sup>39</sup> (0+ years)	Sun et al., 2022 <sup>40</sup> (20-79 years)	NCD-RisC (18+ years)	
						Women	Men
China	2016 (2013)	2017 (2017)	2018 (2015)	6.2 (5.7-6.6)	10.6	9.4 (5.4-14.7)	12.9 (7.4-19.7)
India	2017 (2014)	2015 (2015)	2021 (2018)	5.8 (5.4-6.4)	9.6	23.7 (18.0-29.9)	21.4 (16.4-27.1)
USA	2018 (2018)	2016 (2016)	2020 (2018)	9.0 (8.6-9.5)	10.7	11.4 (7.5-16.3)	13.6 (9.4-18.6)
Indonesia	2008 (2008)	2018 (2018)	2018 (2018)	4.6 (4.2-5.0)	10.6	14.5 (8.4-22.7)	10.9 (5.9-17.6)
Brazil	2008 (n.a.)	2019* (2019*)	2016 (2013)	5.4 (5.0-5.9)	8.8	14.0 (7.4-23.2)	11.7 (5.7-20.4)
Pakistan	2011 (n.a.)	2017 (2017)	2017 (2017)	7.1 (6.6-7.8)	30.8	30.9 (21.0-42.1)	30.8 (21.3-42.1)
Bangladesh	2018 (2018)	2018 (2018)	2018 (2018)	7.1 (6.7-7.6)	14.2	19.7 (12.9-27.7)	16.1 (9.9-23.7)
Russian Federation	2020 (n.a.)	2020 (2020)	2020 (2014)	4.0 (3.7-4.3)	5.6	8.3 (4.5-13.6)	7.6 (4.0-12.7)
Nigeria	2005 (n.a.)	2007 (2007)	2011 (n.a.)	3.7 (3.4-4.0)	3.6	11.2 (3.2-24.2)	10.7 (3.4-22.6)
Japan	2017 (2017)	2011* (2011*)	2023 (2019)	5.9 (5.4-6.4)	6.6	4.3 (2.8-6.2)	8.5 (6.2-11.4)
Mexico	2019 (2019)	2012 (2012)	2022 (2022)	9.1 (8.4-9.8)	16.9	15.2 (12.4-18.2)	13.2 (10.5-16.1)
Philippines	2014 (2014)	2008 (2008)	2021 (2021)	4.4 (4.1-4.8)	7.1	13.8 (8.6-19.9)	12.5 (7.8-18.8)
Viet Nam	2015 (2015)	2013 (2009)	2021 (2021)	4.5 (4.3-4.8)	6.1	11.1 (7.0-16.5)	10.4 (6.4-15.7)
Germany	2008 (n.a.)	2015* (2015*)	2019 (2012)	4.8 (4.4-5.2)	6.9	5.2 (2.5-9.2)	8.0 (4.4-12.8)
Egypt	2012 (2012)	2017 (2017)	2017 (2017)	8.4 (7.7-9.2)	20.9	28.8 (18.7-40.2)	26.3 (16.4-37.6)
Ethiopia	2015 (2015)	2015 (2015)	2015 (2015)	3.4 (3.1-3.6)	5.0	7.4 (2.6-14.7)	6.2 (2.3-12.6)
ran	2011 (2011)	2007 (2007)	2021 (2021)	6.9 (6.3-7.5)	9.1	15.2 (12.3-18.6)	14.2 (11.2-17.5)
Гürkiye	2011 (2011)	2010 (2010)	2017 (2017)	6.6 (6.1-7.2)	14.5	17.1 (10.6-24.7)	16.0 (10.0-23.3)
Thailand	2009 (2009)	2014 (2014)	2020 (2020)	5.4 (4.9-5.9)	9.7	12.8 (8.6-18.0)	11.7 (7.7-16.6)
UK	2013 (2013)	2021* (2021*)	2019 (2019)	7.8 (7.2-8.3)	6.3	7.9 (5.5-10.8)	9.7 (7.0-12.7)

France	2007 (2007)	2016 (2016)	2023 (2019)	3.6 (3.4-4.0)	5.3	1.8 (1.1-2.7)	3.7 (2.5-5.2)
Italy	2012 (2012)	2005* (2005*)	2020 (2012)	4.7 (4.3-5.2)	6.4	5.1 (2.7-8.4)	9.5 (5.7-14.1)
DR Congo	2007 (2005)	n.a. (n.a.)	2017 (n.a.)	4.8 (4.3-5.1)	5.8	10.0 (3.8-19.5)	8.7 (3.4-17.1)
South Korea	2019 (2019)	2015 (2015)	2023 (2023)	10.3 (9.8-10.8)	6.8	8.1 (6.7-9.6)	12.6 (10.6-14.7)
South Africa	2005 (n.a.)	2012* (2012*)	2020 (2012)	5.9 (5.5-6.3)	10.8	15.5 (7.8-25.6)	11.9 (5.8-20.1)
Spain	2011 (2010)	2010 (2010)	2019 (2015)	6.9 (6.4-7.3)	10.3	2.6 (1.3-4.4)	4.6 (2.6-7.2)
Colombia	2015 (2015)	2010 (n.a.)	2016 (2016)	6.1 (5.7-6.5)	8.3	12.2 (6.0-20.6)	12.3 (6.0-21.6)
Myanmar	2014 (2014)	2014 (2014)	2014 (2014)	8.0 (7.5-8.5)	7.1	14.9 (7.2-25.1)	12.5 (5.9-21.6)
Argentina	1995 (n.a.)	1995 (n.a.)	2018 (2018)	5.5 (5.0-6.0)	5.4	10.4 (5.4-17.0)	10.7 (5.8-17.1)
Tanzania	2012 (2012)	2012 (2012)	2023 (2023)	2.6 (2.4-2.8)	12.3	7.3 (4.1-11.6)	5.8 (3.3-9.3)

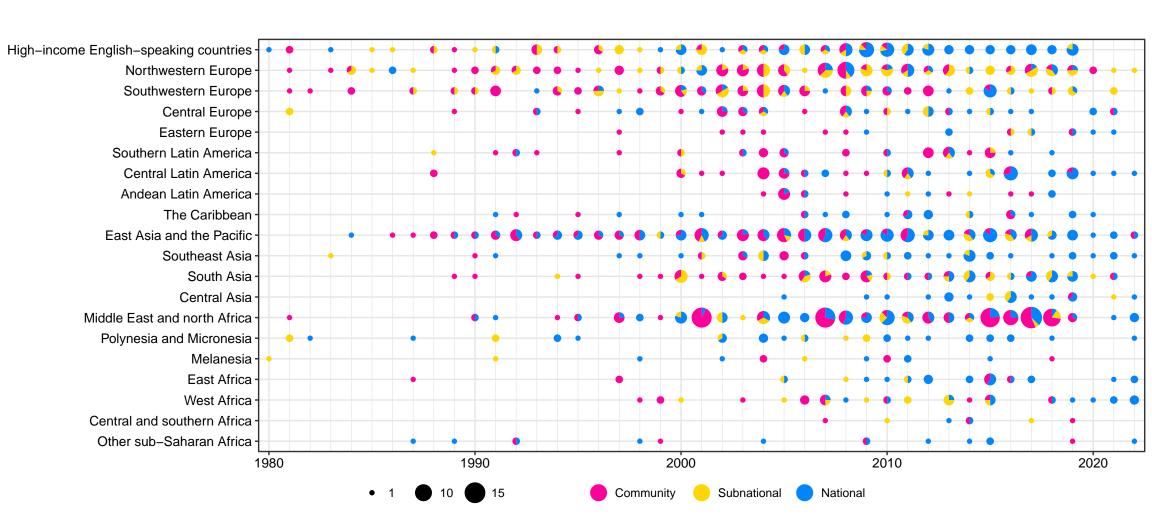
<sup>†</sup> For each country, the first number shows the year of the most recent data source, and the number in brackets shows the year of the most recent national data source. Data sources that only reported type 1 diabetes are not included in this table because type 1 diabetes is an outcome only in Ong et al. (2023) but not in Sun et al. (2022) or this study. ‡ Estimates from Ong et al. (2023) and Sun et al. (2022) are for both sexes in 2021. Estimates from this study are for 2022. Numbers in brackets are 95% uncertainty intervals, when reported.

<sup>\*</sup> Data were based on self-reported diabetes diagnosis, medical records, or a registry of people with diagnosed diabetes, handled as described in Research in Context panel. n.a. = not available.

**Appendix Figure 1.** Number of data sources used in this analysis, by country.



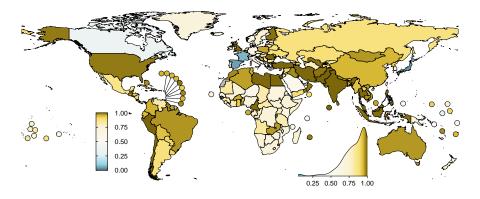
**Appendix Figure 2.** Number of data sources used in this paper, by region and year. The size of each circle shows the number of data sources for each region and year, and the colours indicate the relative count of national, subnational and community data sources.



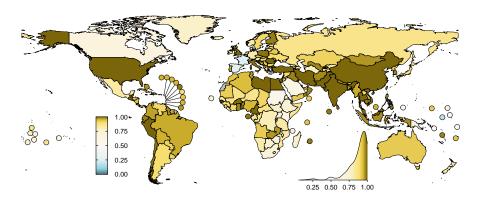
**Appendix Figure 3.** Posterior probability that age-standardised diabetes prevalence for people aged 18+ years increased from 1990 to 2022.

The maps show the PP of an increase from 1990 to 2022. The PP of a decrease is one minus that of an increase. The density plot alongside each map shows the smoothed distribution of estimates across countries.

#### Women



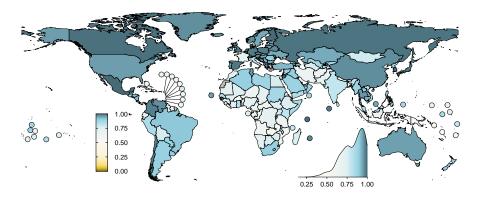
### Men



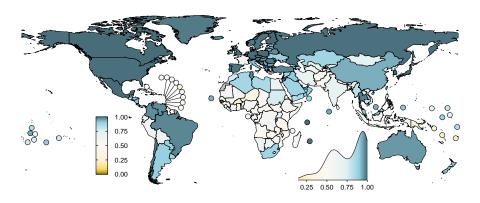
**Appendix Figure 4.** Posterior probability that age-standardised diabetes treatment coverage for people aged 30+ years increased from 1990 to 2022.

The maps show the PP of an increase from 1990 to 2022. The PP of a decrease is one minus that of an increase. The density plot alongside each map shows the smoothed distribution of estimates across countries.

### Women

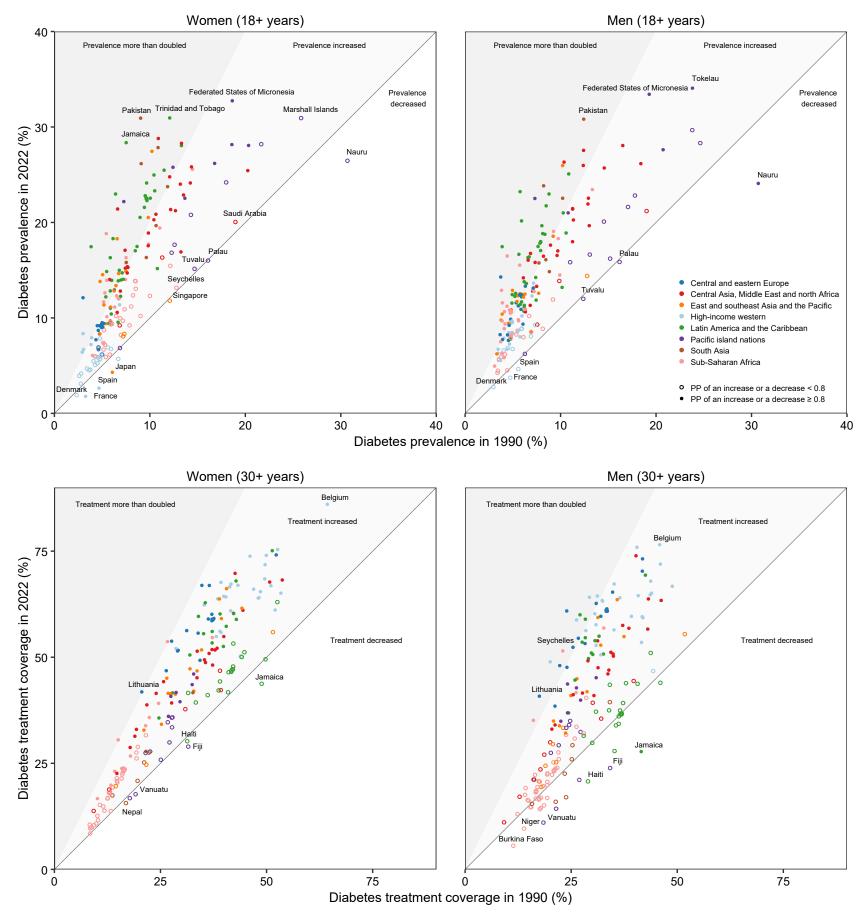


### Men

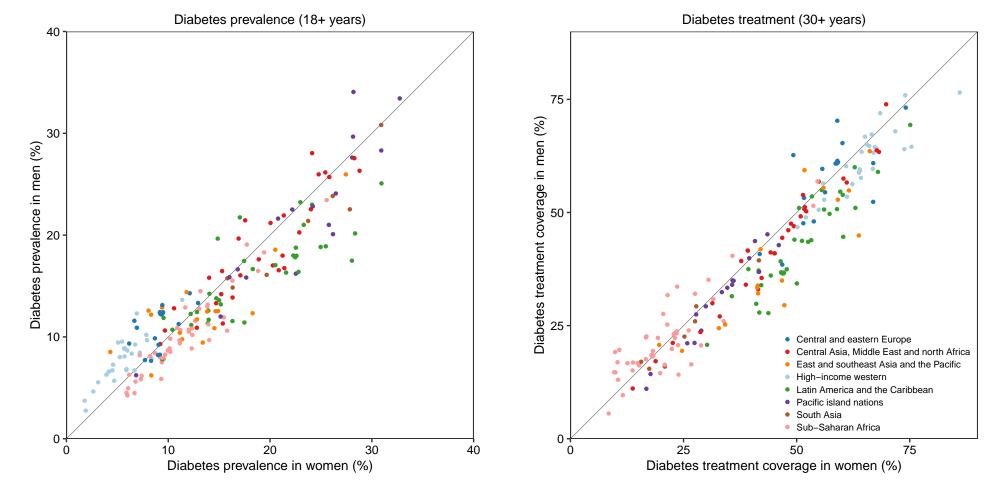


**Appendix Figure 5.** Relationship between levels in 1990 and 2022 for age-standardised diabetes prevalence for people aged 18+ years and age-standardised treatment coverage for people aged 30+ years.

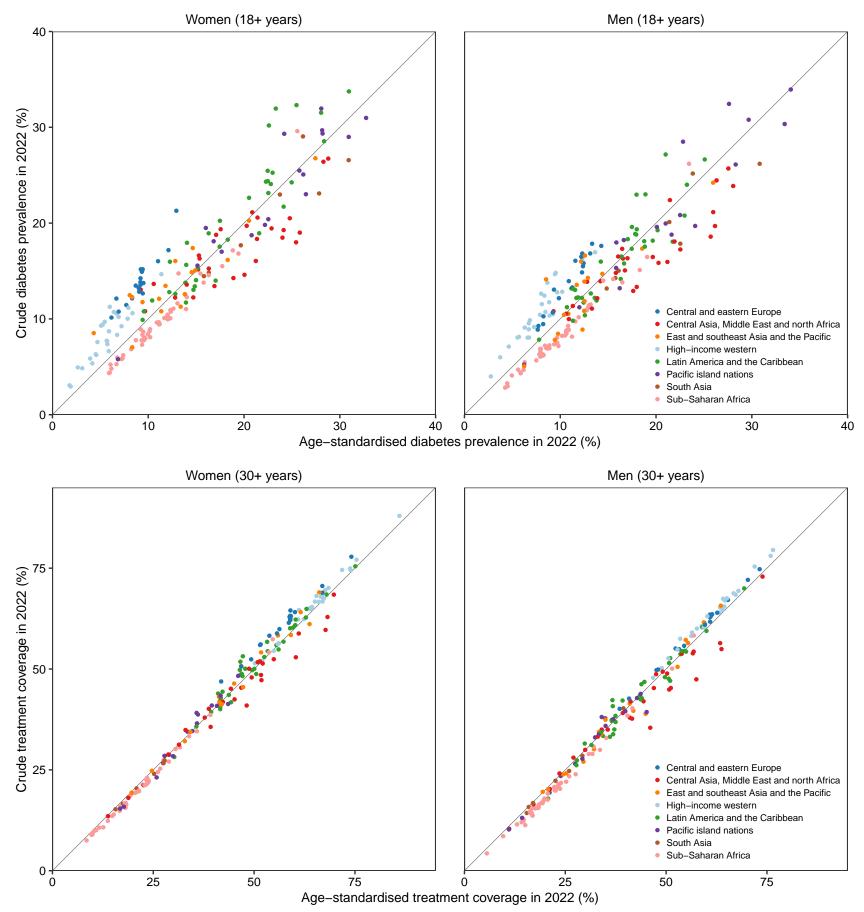
Each point shows one country. Points that are filled have a posterior probability (PP) >0.80 of the observed change being a true decrease or increase. If neither an increase nor a decrease was detected at PP = 0.80 the point is hollow.



**Appendix Figure 6.** Relationship of age-standardised diabetes prevalence and treatment coverage between women and men in 2022. Each point shows one country.

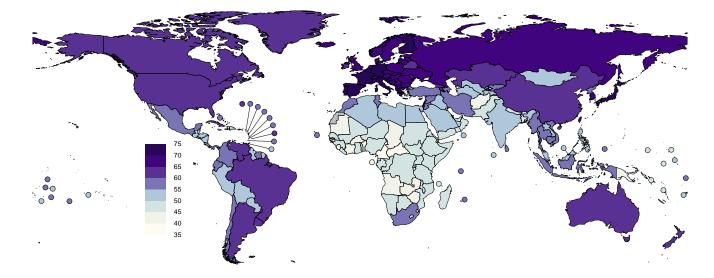


**Appendix Figure 7.** Relationship between age-standardised and crude prevalence of diabetes among women and men aged 18+ years in 2022. Each point shows one country.

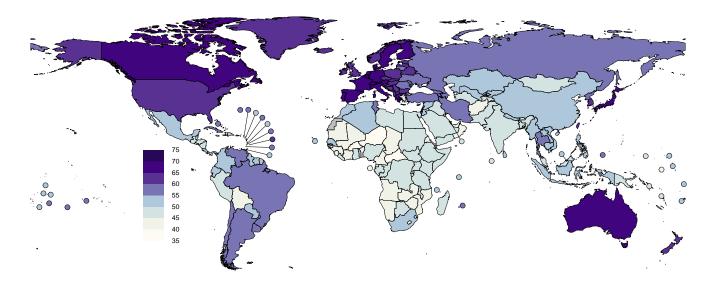


Appendix Figure 8. Median age of women and men with diabetes in 2022 by country.

### Women

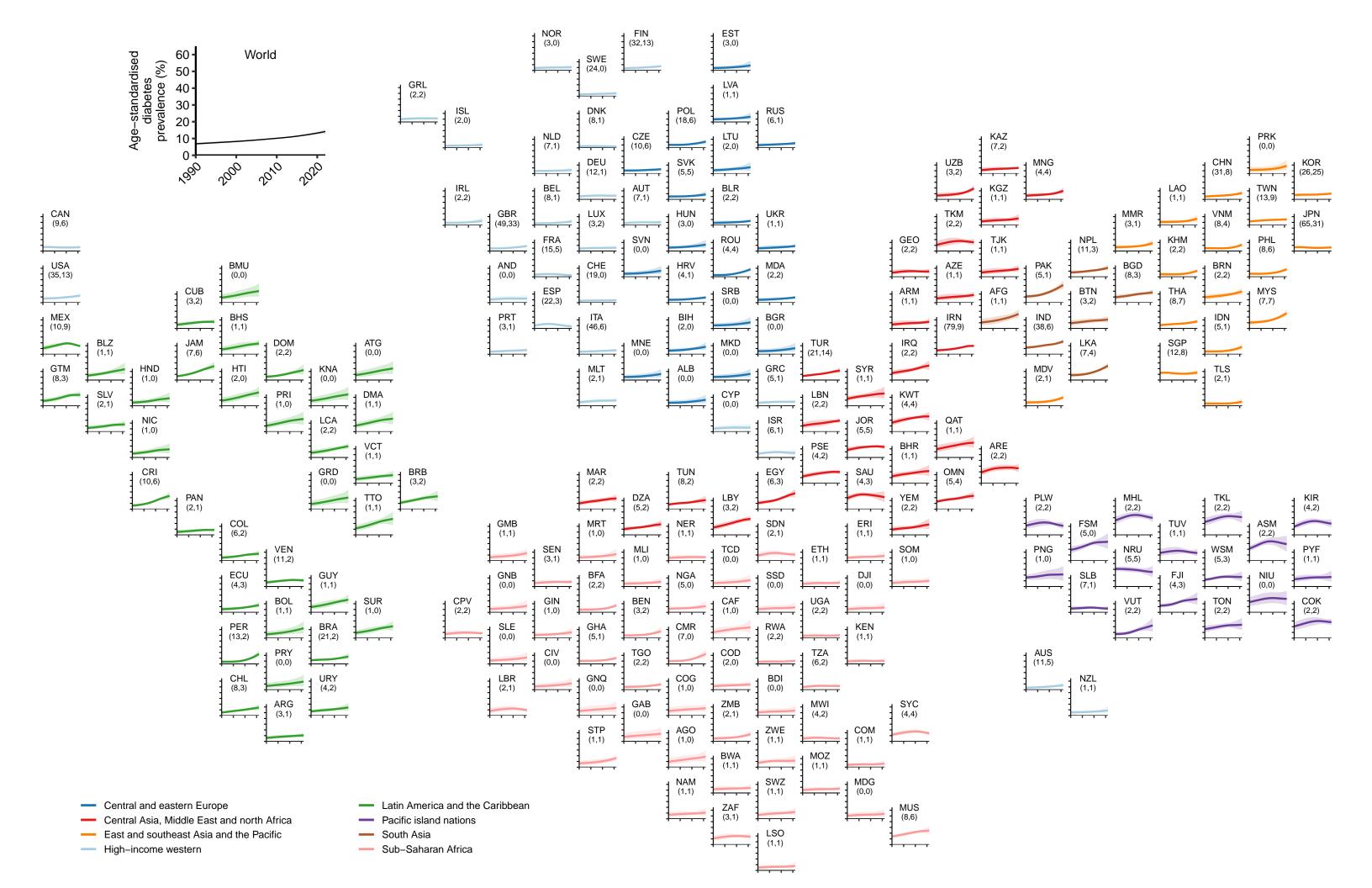


### Men



**Appendix Figure 9.** Age-standardised diabetes prevalence for people aged 18+ years from 1990 to 2022 by country, for both sexes combined.

See main paper Figure 1 caption for descriptions of the contents of the figure and for definitions.

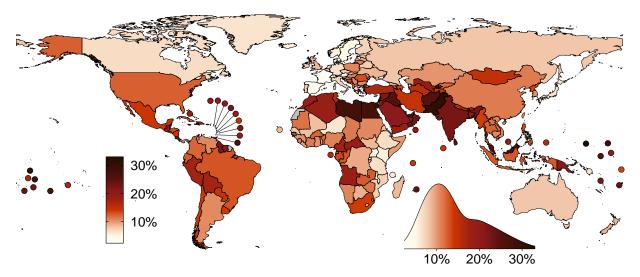


**Appendix Figure 10.** Levels in 2022 and change from 1990 to 2022 by country of (A) agestandardised diabetes prevalence for people aged 18+ years, and (B) age-standardised treatment coverage for people aged 30+ years, for both sexes combined.

The density plot alongside each map shows the smoothed distribution of estimates across countries.

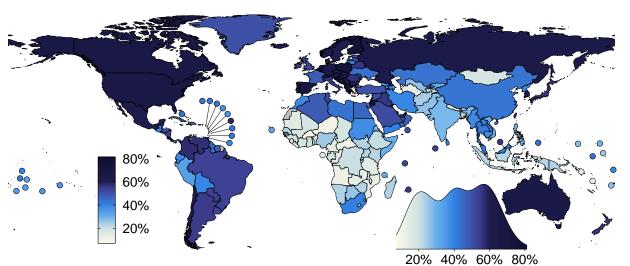
## A Both sexes (18+ years)

Prevalence in 2022

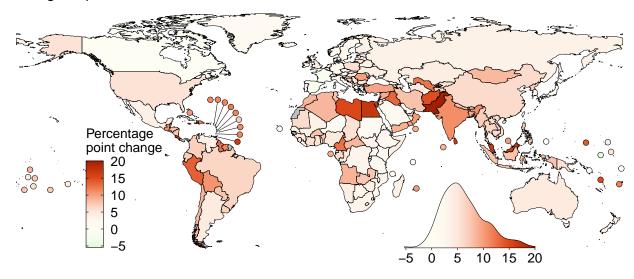


# B Both sexes (30+ years)

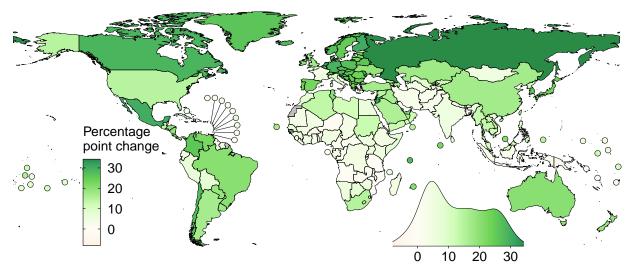
Treatment in 2022



## Change in prevalence from 1990 to 2022



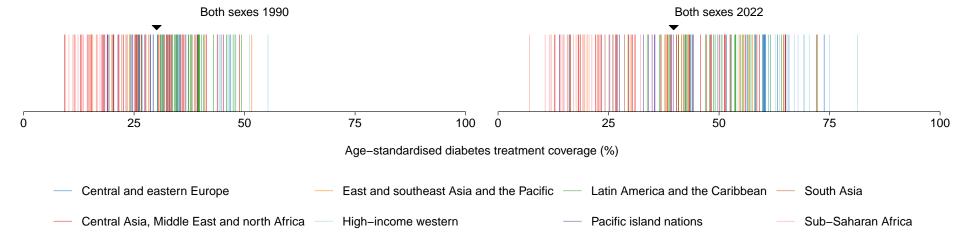
# Change in treatment from 1990 to 2022



**Appendix Figure 11.** Age-standardised treatment coverage for people aged 30+ years in 1990 and 2022, for both sexes combined.

Each line represents a country, with countries coloured by the super-region in which they fall.

The black triangle shows the age-standardised treatment coverage for the world.



#### References

- 1. NCD Risk Factor Collaboration (NCD-RisC). Worldwide trends in diabetes since 1980: a pooled analysis of 751 population-based studies with 4.4 million participants. *Lancet* 2016; **387**(10027): 1513-30.
- 2. NCD Risk Factor Collaboration (NCD-RisC). Worldwide trends in underweight and obesity from 1990 to 2022: a pooled analysis of 3663 population-representative studies with 222 million children, adolescents, and adults. *Lancet* 2024; **403**(10431): 1027-50.
- 3. Danaei G, Finucane MM, Lin JK, et al. National, regional, and global trends in systolic blood pressure since 1980: systematic analysis of health examination surveys and epidemiological studies with 786 country-years and 5·4 million participants. *Lancet* 2011; **377**(9765): 568-77.
- 4. Danaei G, Finucane MM, Lu Y, et al. National, regional, and global trends in fasting plasma glucose and diabetes prevalence since 1980: systematic analysis of health examination surveys and epidemiological studies with 370 country-years and 2·7 million participants. *Lancet* 2011; **378**(9785): 31-40.
- 5. Farzadfar F, Finucane MM, Danaei G, et al. National, regional, and global trends in serum total cholesterol since 1980: systematic analysis of health examination surveys and epidemiological studies with 321 country-years and 3·0 million participants. *Lancet* 2011; **377**(9765): 578-86.
- 6. Finucane MM, Stevens GA, Cowan MJ, et al. National, regional, and global trends in body-mass index since 1980: systematic analysis of health examination surveys and epidemiological studies with 960 country-years and 9·1 million participants. *Lancet* 2011; **377**(9765): 557-67.
- 7. Ogurtsova K, Guariguata L, Barengo NC, et al. IDF Diabetes Atlas: global estimates of undiagnosed diabetes in adults for 2021. *Diabetes Res Clin Pract* 2022; **183**: 109118.
- 8. NCD Risk Factor Collaboration (NCD-RisC). Global variation in diabetes diagnosis and prevalence based on fasting glucose and hemoglobin A1c. *Nat Med* 2023; **29**(11): 2885-901.
- 9. John WG, Mosca A, Weykamp C, Goodall I. HbA1c standardisation: history, science and politics. *Clin Biochem Rev* 2007; **28**(4): 163-8.
- 10. Carstensen B, Lindstrom J, Sundvall J, Borch-Johnsen K, Tuomilehto J, DPS Study Group. Measurement of blood glucose: comparison between different types of specimens. *Ann Clin Biochem* 2008; **45**(Pt 2): 140-8.
- 11. Cowie CC, Rust KF, Ford ES, et al. Full accounting of diabetes and pre-diabetes in the U.S. population in 1988-1994 and 2005-2006. *Diabetes Care* 2009; **32**(2): 287-94.
- 12. Anjana RM, Unnikrishnan R, Deepa M, et al. Metabolic non-communicable disease health report of India: the ICMR-INDIAB national cross-sectional study (ICMR-INDIAB-17). *Lancet Diabetes Endocrinol* 2023; **11**(7): 474-89.
- 13. Diabetes Control Complications Trial Research Group, Nathan DM, Genuth S, et al. The effect of intensive treatment of diabetes on the development and progression of long-term complications in insulin-dependent diabetes mellitus. *N Engl J Med* 1993; **329**(14): 977-86.
- 14. UK Prospective Diabetes Study (UKPDS) Group. Intensive blood-glucose control with sulphonylureas or insulin compared with conventional treatment and risk of complications in patients with type 2 diabetes (UKPDS 33). *Lancet* 1998; **352**(9131): 837-53.
- 15. Finucane MM, Paciorek CJ, Danaei G, Ezzati M. Bayesian estimation of population-level trends in measures of health status. *Stat Sci* 2014; **29**(1): 18-25.
- 16. NCD Risk Factor Collaboration (NCD-RisC). Worldwide trends in hypertension prevalence and progress in treatment and control from 1990 to 2019: a pooled analysis of 1201 population-representative studies with 104 million participants. *Lancet* 2021; **398**(10304): 957-80.
- 17. Finucane MM, Paciorek CJ, Stevens GA, Ezzati M. Semiparametric Bayesian density estimation with disparate data sources: a meta-analysis of global childhood undernutrition. *J Am Stat Assoc* 2015; **110**(511): 889-901.
- 18. Gelman A. Prior distributions for variance parameters in hierarchical models (comment on article by Browne and Draper). *Bayesian Anal* 2006; **1**(3): 515-34.

- 19. Breslow NE, Clayton DG. Approximate inference in generalized linear mixed models. *J Am Stat Assoc* 1993; **88**(421): 9-25.
- 20. Wood SN. Generalized Additive Models: An Introduction with R. Boca Raton, FL: Chapman & Hall/CRC; 2006.
- 21. Rue H, Held L. Gaussian Markov Random Fields: Theory and Applications. New York: CRC Press; 2005.
- 22. Harville DA. Matrix Algebra from a Statistician's Perspective: Springer-Verlag; 2008.
- 23. Danaei G, Finucane MM, Lin JK, et al. National, regional, and global trends in systolic blood pressure since 1980: systematic analysis of health examination surveys and epidemiological studies with 786 country-years and 5.4 million participants. *Lancet* 2011; **377**(9765): 568-77.
- 24. Singh GM, Danaei G, Pelizzari PM, et al. The age associations of blood pressure, cholesterol, and glucose: analysis of health examination surveys from international populations. *Circulation* 2012; **125**(18): 2204-11.
- 25. United Nations Department of Economic and Social Affairs, Population Division. World Urbanization Prospects: The 2018 Revision. New York: United Nations, 2019.
- 26. Smith AF, Roberts GO. Bayesian computation via the Gibbs sampler and related Markov chain Monte Carlo methods. *J R Stat Soc Series B Stat Methodol* 1993; **55**(1): 3-23.
- 27. Vehtari A, Gelman A, Simpson D, Carpenter B, Bürkner P-C. Rank-normalization, folding, and localization: an improved R for assessing convergence of MCMC (with discussion). *Bayesian Anal* 2021; **16**(2): 667-718.
- 28. Gelman A, Hwang J, Vehtari A. Understanding predictive information criteria for Bayesian models. *Stat Comput* 2014; **24**(6): 997-1016.
- 29. Stan Development Team. RStan: the R interface to Stan. 2022. https://mc-stan.org/. .
- 30. Ahmad OB, Boschi-Pinto C, Lopez AD, Murray CJ, Lozano R, Inoue M. Age standardization of rates: a new WHO standard. GPE Discussion Paper Series 31. Geneva: World Health Organization, 2001.
- 31. United Nations, Department of Economic and Social Affairs, Population Division. World Population Prospects 2024, Online Edition., 2024.
- 32. NCD Risk Factor Collaboration (NCD-RisC). Repositioning of the global epicentre of non-optimal cholesterol. *Nature* 2020; **582**(7810): 73-7.
- 33. NCD Risk Factor Collaboration (NCD-RisC). Trends in adult body-mass index in 200 countries from 1975 to 2014: a pooled analysis of 1698 population-based measurement studies with 19.2 million participants. *Lancet* 2016; **387**(10026): 1377-96.
- 34. NCD Risk Factor Collaboration (NCD-RisC). Rising rural body-mass index is the main driver of the global obesity epidemic in adults. *Nature* 2019; **569**(7755): 260-4.
- 35. NCD Risk Factor Collaboration (NCD-RisC). Height and body-mass index trajectories of school-aged children and adolescents from 1985 to 2019 in 200 countries and territories: a pooled analysis of 2181 population-based studies with 65 million participants. *Lancet* 2020; **396**(10261): 1511-24.
- 36. NCD Risk Factor Collaboration (NCD-RisC). Diminishing benefits of urban living for children and adolescents' growth and development. *Nature* 2023; **615**(7954): 874-83.
- 37. NCD Risk Factor Collaboration (NCD-RisC). Worldwide trends in blood pressure from 1975 to 2015: a pooled analysis of 1479 population-based measurement studies with 19.1 million participants. *Lancet* 2017; **389**(10064): 37-55.
- 38. NCD Risk Factor Collaboration (NCD-RisC). Worldwide trends in body-mass index, underweight, overweight, and obesity from 1975 to 2016: a pooled analysis of 2416 population-based measurement studies in 128.9 million children, adolescents, and adults. *Lancet* 2017; **390**(10113): 2627-42.
- 39. GBD Diabetes Collaborators. Global, regional, and national burden of diabetes from 1990 to 2021, with projections of prevalence to 2050: a systematic analysis for the Global Burden of Disease Study 2021. *Lancet* 2023; **402**(10397): 203-34.
- 40. Sun H, Saeedi P, Karuranga S, et al. IDF Diabetes Atlas: global, regional and country-level diabetes prevalence estimates for 2021 and projections for 2045. *Diabetes Res Clin Pract* 2022; **183**: 109119.

41. Shabbir S, Nadeem M, Sattar A, et al. Type and frequency of hemoglobinopathies, diagnosed in the area of Karachi, in Pakistan. *Cogent Med* 2016; **3**: 1188875.