Child mortality

Data sources
Health indicator and dimension of inequality data were sourced from publicly available Demographic and Health Surveys (DHS). DHS is a large-scale, nationally representative household survey that collects data through standardized, face-to-face interviews with women aged 15–49 years in low- and middle-income countries. Country income group was determined using the World Bank classification as of July 2017.

The disaggregated data are the product of a reanalysis of DHS micro-data by the WHO Collaborating Center for Health Equity Monitoring (International Center for Equity in Health based in the Federal University of Pelotas, Brazil), and can be freely accessed from the WHO Health Equity Monitor database (www.who.int/gho/health_equity/en/).

The survey tools used by DHS permit direct comparisons across countries and over time. The data were taken from rounds of DHS that were not conducted in the same year in all countries. In a few cases there may be minor differences between the data reported here and in previous DHS country reports due to small discrepancies in the time span, definition and/or calculation of some indicators.

Health indicators
Child mortality indicators include neonatal mortality rate, infant mortality rate and under-five mortality rate.

- **Neonatal mortality rate** is defined as the probability (expressed as a rate per 1000 live births) of a child born in a specific year or period dying in the first 30 days of life, if subject to age-specific mortality rates of that period. Note that standard DHS calculations (as applied in this analysis) specify the first 30 days of life; however, other accepted definitions may use the first 28 days.
- **Infant mortality rate** is defined as the probability (expressed as a rate per 1000 live births) of a child born in a specific year or period dying before reaching the age of one year, if subject to age-specific mortality rates of that period.
- **Under-five mortality rate** is defined as the probability (expressed as a rate per 1000 live births) of a child born in a specific year or period dying before reaching the age of five years, if subject to age-specific mortality rates of that period.

Detailed information about the criteria used to calculate the numerator and denominator values for each indicator is available in the WHO Indicator and Measurement Registry, under the topic “Health Equity Monitor” (www.who.int/gho/indicator_registry/en/).

Dimensions of inequality
Health data were disaggregated by four dimensions of inequality: economic status, education, place of residence and sex. Economic status was determined at the household level, using a wealth index. Country-specific indices were based on owning selected assets and having access to certain services, and constructed using principal component analysis. Within each country the index was used to
create quintiles, thereby identifying five equal subgroups that each account for 20% of the population. Note that certain indicators have denominator criteria that do not include all households and/or are more likely to include households from a specific quintile; thus the share of the population for a given indicator may not equal 20%. Education refers to the highest level of schooling attained by the mother. Three subgroups were defined: no education, primary school, and secondary school or higher. For place-of-residence classifications (that is, urban or rural), country-specific criteria were applied.

**Country selection**

Countries were selected on the basis of data availability and survey year. When a survey was conducted over more than one calendar year, the year of survey was assigned based on the initial year of data collection. Countries with surveys dating from the past 10 years were selected to illustrate the “latest situation” of inequality. Latest-situation analyses were possible for 54 countries representing all WHO regions with the year of their most recent survey falling between 2006 and 2015. “Change over time” was analysed for each study country that had surveys from two time points (a recent survey falling between 2006 and 2015 and an older survey falling between 1996 and 2005), and reflects the change in national averages and inequalities within countries over a period of about 10 years. Change-over-time analyses were possible for 36 countries, representing all WHO regions. The number of years between surveys within countries ranged from 5 to 15 years, as per data availability. When more than one older survey was available, the survey closest to 10 years prior to the most recent survey was selected.

Study countries were excluded on a case-by-case basis if data about the relevant health indicator and/or the dimension of inequality were not available or if the sample size was too low to report a valid estimate for one or more of the relevant subgroups (that is, less than 250 unweighted person-years of exposure to the risk of death). Situations of low sample size (that is, 250–499 unweighted person-years of exposure to the risk of death) were noted.

**Analysis**

Micro-level DHS data were used to generate national average and disaggregated estimates for each indicator. Survey design specifications were taken into consideration in the estimation. The same methods of calculation were applied across all surveys to generate comparable estimates across countries and over time.

Point estimates of disaggregated data are presented alongside 95% confidence intervals (CIs), and the population share of the subgroup. The population share for each indicator is the percentage of the affected population — the indicator denominator — represented by the subgroup in a given country.