scientific reports

Check for updates

OPEN Global, regional and national epidemiology and prevalence of child stunting, wasting and underweight in lowand middle-income countries, 2006-2018

Paddy Ssentongo^{1,2,3^{IZ}}, Anna E. Ssentongo^{3,4}, Djibril M. Ba^{3,5}, Jessica E. Ericson⁶, Muzi Na⁷, Xiang Gao⁷, Claudio Fronterre⁸, Vernon M. Chinchilli³ & Steven J. Schiff^{1,2,9,10,11}

In 2016, undernutrition, as manifested in childhood stunting, wasting, and underweight were estimated to cause over 1.0 million deaths, 3.9% of years of life lost, and 3.8% of disability-adjusted life years globally. The objective of this study is to estimate the prevalence of undernutrition in lowand middle-income countries (LMICs) using the 2006–2018 cross-sectional nationally representative demographic and health surveys (DHS) data and to explore the sources of regional variations. Anthropometric measurements of children 0–59 months of age from DHS in 62 LMICs worldwide were used. Complete information was available for height-for-age (n = 624,734), weight-for-height (n = 625,230) and weight-for-age (n = 626,130). Random-effects models were fit to estimate the pooled prevalence of stunting, wasting, and underweight. Sources of heterogeneity in the prevalence estimates were explored through subgroup meta-analyses and meta-regression using generalized linear mixed-effects models. Human development index (a country-specific composite index based on life expectancy, literacy, access to education and per capita gross domestic product) and the United Nations region were explored as potential sources of variation in undernutrition. The overall prevalence was 29.1% (95% CI 26.7%, 31.6%) for stunting, 6.3% (95% CI 4.6%, 8.2%) for wasting, and 13.7% (95% CI 10.9%, 16.9%) for underweight. Subgroup analyses suggested that Western Africa, Southern Asia, and Southeastern Asia had a substantially higher estimated prevalence of undernutrition than global average estimates. In multivariable meta-regression, a combination of human development index and United Nations region (a proxy for geographical variation) explained 54%, 56%, and 66% of the variation in stunting, wasting, and underweight prevalence, respectively. Our findings demonstrate that regional, subregional, and country disparities in undernutrition remain, and the residual gaps to close towards achieving the second sustainable development goal-ending undernutrition by 2030.

¹Center for Neural Engineering, The Pennsylvania State University, University Park, PA 16802, USA. ²Department of Engineering Science and Mechanics, The Pennsylvania State University, University Park, PA 16802, USA. ³Department of Public Health Sciences, The Pennsylvania State University College of Medicine, 500 University Drive, Hershey, PA 17033, USA. ⁴Department of Surgery, The Pennsylvania State University College of Medicine, Hershey, PA 17033, USA. ⁵Center for Applied Studies in Health Economics, The Pennsylvania State University College of Medicine, Hershey, PA 17033, USA. ⁶Department of Pediatrics, The Pennsylvania State University College of Medicine, Hershey, PA 17033, USA. ⁷Department of Nutritional Sciences, College of Health and Human Development, The Pennsylvania State University, University Park, PA 16802, USA. ⁸Centre for Health Informatics, Computing, and Statistics, Lancaster University, Lancaster, UK. ⁹The Center for Infectious Disease Dynamics, The Pennsylvania State University, University Park, PA 16802, USA. ¹⁰Department of Neurosurgery, The Pennsylvania State University College of Medicine, Hershey, PA 17033, USA. ¹¹Department of Physics, The Pennsylvania State University, University Park, PA 16802, USA. Eemail: pssentongo@pennstatehealth.psu.edu

Abbreviations

- LMICs Low- and middle-income countries
- WHO World Health Organization
- SSA Sub-Saharan Africa
- DHS Demographic and health surveys
- UN United Nations CI Confidence Inte
- CI Confidence Interval GDP Gross domestic product
- HDI Human development index

In 2016, undernutrition (stunting, wasting and underweight) was estimated to cause 1.0 million deaths, 3.9% of years of life lost, and 3.8% of disability-adjusted life years (DALYs) globally¹. Since then, undernutrition has decreased globally but remains endemic in southeastern Asia (SA) and sub-Saharan Africa (SSA)^{2,3}. Importantly, heterogeneity still exists in the trends of undernutrition, with Africa being the only region where the number of stunted children continues to rise, from 50 million in 2000 to 59 million in 2018⁴.

Early childhood is a critical window during which significant growth and development occur. The nutrition of the mother and the early life nutrition of the child have a substantial impact on the child's future physical and mental health. Undernutrition during this period is related to poor outcomes in overall health, neurobehavioral and cognitive development, and educational and economic attainment later in life^{5,6}. Therefore, exploring country-level heterogeneity within 'hot spots' of child undernutrition is crucial to guide efforts to develop informed and focused control and prevention strategies.

Nutrition status is primarily assessed through the measurement of a child's height (or length in the youngest children) and weight and comparing the child to the standard metrics. Stunting (height-for-age z-score below – 2 standard deviations (SD) from the global median, as defined by the 2006 World Health Organization Child Growth Standards), wasting (weight-for-height z-score below – 2 SD from the global median) and underweight (weight-for-age z-score below – 2 SD from the global median) are indicators of a child's undernutrition⁷. These anthropometric measures on a country level are updated regularly through the demographic and health surveys (DHS) program, which collects nationally representative health data to monitor and evaluate population health and nutrition programs in low- and middle-income countries (LMICs)⁸. Stunting, wasting and underweight have been assessed in SSA³, and stunting and wasting estimates are regularly updated by the United Nations' Food and Agriculture Organization reports. However, we are unaware of prior studies of the combined influence of human development index (a country-specific composite index based on life expectancy, literacy, access to education and per capita gross domestic product) on all three undernutrition forms, at the global, regional and country-level.

To address this gap, we conducted a pooled analysis of stunting, wasting and underweight prevalence and explored the sources of undernutrition heterogeneity. We used country-level data on the prevalence of undernutrition from the DHS for the past twelve years (2006–2018). We limited our estimates to 2006–2018 due to the introduction of the World Health Organization (WHO) child growth standards in 2006, which estimated new values for assessing child nutritional status. These standards replaced the National Center for Health Statistics/ World Health Organization (NCHS/WHO) growth reference, which had been in international use since the late 1970s but underestimated the prevalence of undernutrition, especially for infants.

The present study focuses on the spatial distribution of childhood stunting, wasting and underweight prevalence in 62 LMICs. Global, regional and country-specific information on the prevalence of the three forms of undernutrition will help guide policymakers, national and international efforts to control and prevent factors that drive undernutrition.

Materials and methods

Sampling process of demographic and health surveys. In this study, nationally representative DHS data between 2006 and 2018, including anthropometric indices for each country, were extracted⁹. Surveys without anthropometric data were excluded from the analysis (Supplementary Figure S1). The DHS Program collects nationally representative health data in LMICs every 5 years to monitor and evaluate population health and nutrition programs. The survey designs are based on stratified multistage sampling designs where each country is divided into administrative regions¹⁰. Populations within these regions are then stratified further by urban and rural areas of residence. The definition of urban or rural varies across countries. A random selection of enumeration areas or primary sampling units (PSUs) are drawn within rural or urban regions. PSUs are selected based on a probability proportional to the population size using the latest census. In most countries, a PSU is equivalent to the lowest administrative geographical unit such as a village. In the second sampling stage, all households within a PSU are listed from the most recent population census, and ~ 30 households per PSU are randomly selected for an interview. For each sampled household, all members are listed and have an equal chance of being sampled. Children between the ages of 0 and 59 months are eligible for anthropometric measurements, and had their heights (or lengths) and weights measured by trained field-workers.

Anthropometry methods and assessment of undernutrition status. Children of age 24 months and younger had their length measured in recumbent position but for ages above two years had their height measured while standing. Length was measured with the portable Harpenden Infantometer (range 30–110 cm, with digit counter readings precise to 1 mm), and the height with the Harpenden Portable Stadiometer (range 65–206 cm, digit counter reading). Portable electronic scales with a taring capability, calibrated to 0.1 kg, were used to measure weight^{11,12}. The z-scores for weight-for-age, weight-for-height, and height-for-age were provided in the DHS data and were calculated using the 2006 WHO Child Growth Standards. The 2006 WHO Child

Growth Standards replaced the NCHS/WHO growth reference curves, which had been in used as an international growth reference since $1977^{13,14}$. Unlike the NCHS/WHO growth reference, which is based on children from a single country, the WHO standards depict normal early childhood growth under optimal environmental conditions and can be used to assess children everywhere, regardless of ethnicity, socioeconomic status and type of feeding¹⁵. Another difference lies in the methodology applied to construct the growth curves. The computation of growth curves and the z-scores for the new WHO standards uses formulae based on the LMS method¹⁶. For these reasons, the DHS conducted after 2006 that used the new growth standards were analyzed. A child was stunted, wasted, or underweight if he or she exhibited a *z* score below – 2. The present study is a secondary data analysis. Country-specific number of children with anthropometric measurements are reported in Supplementary Table S1.

Human development index (HDI). To assess sources of variability in the prevalence of undernutrition in LMICs, we fitted our generalized linear mixed-effects models with the 2018 HDI (Supplementary Figure S2)¹⁷. The HDI is the geometric mean of normalized indices for each of the three human development measures: education, life expectancy and economy. The education dimension is measured by average years of schooling for adults 25 years and older and expected years of schooling for children entering school age. The economic dimension is measured by gross national income per capita, and the health dimension is assessed by life expectancy at birth. The HDI ranges from 0 to 1, with a higher score indicative of higher HDI. The official categorization by the United Nations is low, medium, high and very high. Educational attainment for women of reproductive age is one of the leading social determinants of health, with higher attainment associated with improved child nutritional outcomes in LMICs^{18,19}.

Study selection and inclusion criteria. All countries with complete DHS anthropometric information between 2006 and 2018 were included in the analysis. The following variables were extracted: year of the survey, continent, United Nations (UN) region, UN subregions (Supplementary Figure S3), number of children stunted, number of children wasted, number of children underweight, and the total population of children whose weight and lengths/heights were measured. We excluded surveys that were not within the time frame of (2006–2018) or did not report anthropometric indicators of children under 5 years.

Statistical analysis. The primary outcome was an estimation of the prevalence of stunting, wasting and underweight. The DHS program reported undernutrition prevalence based on the total number of children with completed anthropometric measurements; therefore, we reported the prevalence as a percent of the children under 5 years with anthropometric measurements.

We applied random-effects models to estimate the overall global prevalence of undernutrition and their respective 95% confidence intervals (CIs) using DerSimonian and Laird random-effects meta-analytic method (See equations in supplementary material)²⁰. A random-effects model assumes the observed prevalence estimates can vary across surveys because of real differences in the measured effect that are independent of time and represent unmodeled variance. To pool the estimates, we built random-effects models using the inverse variance method with logit transformed proportions^{21,22}. Individual and pooled estimates were graphically displayed via forest plots. Between-study variation (heterogeneity) was assessed using I^2 , which describes the percentage of total variation across surveys that is due to heterogeneity rather than chance, expressed as percent (low (25%), moderate (50%), and high (75%)²³.

A generalized linear mixed-effects meta-regression model with a logit link function was fit to investigate the sources of heterogeneity and the results were reported as odds ratios (OR) and their corresponding 95% confidence intervals (95% CI). We examined the associations of each of the explanatory variables included in the model in relation to undernutrition prevalence. These included country-level HDI and the subregions classified by the United Nations²⁴. The *metaprop*, *escalc*, *rma* functions from the R packages *meta* and *metafor* were used for the analysis²⁵.

Ethics approval and consent to participate. The DHS program takes strict measures for protecting the privacy of all survey respondents. Procedures and questionnaires for standard DHS surveys have been reviewed and approved by the ICF International Institutional Review Board (IRB) and the IRB of the host country. ICF International provides both writing and oral informed consent to each survey respondent before the beginning of each survey question and biomarker tests. Each participant's participation was voluntary. This study protocol was submitted to the Pennsylvania State University institutional review board and was not considered to be human subject research, as defined by the US Department of Health and Human Services.

Consent for publication. No consent to publish was needed for this study as we did not use any details, images or videos related to individual participants.

Results

We identified 93 potentially relevant DHS from LMICs. Thirty-one surveys were excluded because they were conducted before 2006 or had no data on anthropometric measures. The remaining 62 surveys, each representing one country, provided data for this pooled analysis. Thirty-eight countries were from Africa, 14 from Asia and 8 from Latin America and the Caribbean, and 1 each from Oceania and Europe. Supplemental Table S1 shows the characteristics of the 62 surveys that we included in the analyses.



Figure 1. Prevalence of undernutrition. Countries are shaded according to prevalence (%) of stunting (top row), wasting (middle row) and underweight (bottom row).

The overall prevalence was 29.1% (95% CI 26.7%, 31.6%) for stunting, 6.3% (95% CI 4.6%, 8.2%) for wasting, and 13.7% (95% CI 10.9%, 16.9%) for underweight. Substantial heterogeneity was evident ($I^2 = 100\%$; p for heterogeneity <0.0001). Figure 1 is a world map displaying country-level prevalence of stunting, wasting and underweight.

Stunting. Figure 2 shows the prevalence of stunting among children under age five years on a subregional and country level. Overall, Africa (Supplemental Figure S4) had the highest prevalence of stunting at 32% (95% CI 29.5–35.9%), followed by Oceania 27% (95% CI 26.2–28.1%) and Asia at 27.4% (95% CI 22.1–32.0%). The Americas and Europe had the lowest prevalence of stunting: 20% (95% CI 13.1–29.1%) and 11.3% (95% CI 10.0–12.6%), respectively. The prevalence of stunting in Africa was significantly different from America, Europe and Oceania but not Asia.

Within the Africa region, the prevalence of stunting in Northern Africa was on average 10 percentage points less than that of Middle, Eastern, Western and Southern Africa. Within the SSA region, the prevalence of stunting in Southern Africa was less than half that of Middle, Eastern, and Western Africa. Similarly, the prevalence

			Events per 100			
Country, Year of survey	Stunted	Total	observations E	vents	95%-CI	Weight
Southern Europe						
Albania, 2018	262	2322	•	11.28	[10.03; 12.60]	1.6%
Random effects model		2322	•	11.28	[10.03; 12.60]	1.6%
neterogeneity. not applicable						
Western Africa						
Angola, 2016	2778	7388	_ =	37.60	[36.50; 38.71]	1.6%
Benin, 2018 Burkina Faso, 2010	24114	6994		32.20	[31.39; 33.01]	1.6%
Cote d'Ivoire, 2012	1067	3581	÷	29.80	[28.31; 31.31]	1.6%
Gambia, 2013	826	3372	-	24.50	[23.06; 25.96]	1.6%
Ghana, 2014 Guipea, 2012	544	2895	•	18.79	[17.39; 20.24]	1.6%
Liberia, 2012	1112	3520	-	31.59	[29.69, 32.75]	1.6%
Mali, 2013	1860	4857	-	38.30	[36.93; 39.67]	1.6%
Niger, 2012	2406	5481	_ =	43.90	[42.59; 45.21]	1.6%
Nigeria, 2013 Sepagal 2017	9638	26190		36.80	[36.22; 37.39]	1.6%
Seriegal, 2017 Sierra Leone. 2013	1/93	5094		37.91	[36.58: 39.24]	1.6%
Togo, 2014	903	3282	-	27.51	[26.00; 29.05]	1.6%
Random effects model		99826		31.24	[27.03; 35.62]	22.6%
Heterogeneity: $I^{2} = 100\%$, $\tau^{2} = 0.0078$, $p = 0$						
Western Asia						
Armenia, 2016	148	1573	•	9.41	[8.01; 10.90]	1.6%
Azerbaijan, 2006	497	1979		25.11	[23.23; 27.05]	1.6%
Turkey, 2013 Yaman, 2012	239	2519	•	9.49	[8.37; 10.66]	1.6%
Random effects model	0420	19894		20.93	[4.94; 44.10]	6.4%
Heterogeneity: $I^2 = 100\%$, $\tau^2 = 0.0654$, $p = 0$					• • •	
Cauthann Aaia						
Bangladesh, 2014	2642	7318		36.10	[35.01: 37 21]	1.6%
India. 2016	84402	219796		38.40	[38.20: 38.60]	1.6%
Maldives, 2017	344	2246		15.32	[13.86; 16.84]	1.6%
Nepal, 2016	867	2421	•	35.81	[33.91; 37.73]	1.6%
Pakistan, 2018	1324	3522		37.59	[36.00; 39.20]	1.6%
Heterogeneity: $I^2 = 99\%$, $\tau^2 = 0.0054$, $\rho < 0.0$	1	235303		32.24	[26.33; 38.44]	8.1%
Latin America and the Caribbean			_			
Bolivia, 2008	2282	8422	_ =	27.10	[26.15; 28.05]	1.6%
Colombia, 2010 Dominican Bonublia, 2012	2073	15/02	_ D	13.20	[12.68; 13.74]	1.6%
Guatemala, 2015	5844	12567		46.50	[45.63: 47.38]	1.6%
Guyana, 2009	277	1522	-	18.20	[16.30; 20.18]	1.6%
Haiti, 2017	1449	6618	•	21.89	[20.91; 22.90]	1.6%
Honduras, 2012	2298	10167	_ 0	22.60	[21.79; 23.42]	1.6%
Random effects model	1009	67785		20.91	[17.31, 18.69]	12.9%
Heterogeneity: $l^2 = 100\%$, $\tau^2 = 0.0228$, $p = 0$		01100		20.01	[10:00, 20:00]	12.0 /0
Eastern Africa Burundi 2017	3602	6444		55 00	154 68· 57 111	1.6%
Comoros 2012	3267	10854		30.10	[29.24: 30.97]	1.6%
Ethiopia, 2016	3984	10376		38.40	[37.46; 39.33]	1.6%
Kenya, 2014	4936	18986		26.00	[25.38; 26.62]	1.6%
Madagascar, 2009	2723	5436		50.09	[48.76; 51.42]	1.6%
Malawi, 2016 Mazambigua, 2011	2117	5/0/	"_	37.09	[35.85; 38.35]	1.6%
Rwanda, 2015	1445	3813		37.90	[36.36: 39.44]	1.6%
Tanzania, 2016	3387	9846		34.40	[33.46; 35.34]	1.6%
Uganda, 2016	1479	5117	ė.	28.90	[27.67; 30.15]	1.6%
Zambia, 2014	4944	12328		40.10	[39.24; 40.97]	1.6%
ZIMDADWE, 2015 Random effects model	1690	0305 105525		20.80	[25.72; 27.90]	1.6%
Heterogeneity: $l^2 = 100\%$, $\tau^2 = 0.0085$, $p = 0$		103323	_	57.17	[52.20, 42.20]	13.470
Southeastern Asia	1585	1803	_	32 30	[31.00:33.71]	1.6%
Mvanmar. 2016	1194	4093		29.20	[27.82: 30.60]	1.6%
Timor-Leste, 2016	3062	6714		45.61	[44.42; 46.80]	1.6%
Random effects model		15696		35.60	[25.78; 46.07]	4.8%
Heterogeneity: $I^2 = 99\%$, $\tau^2 = 0.0088$, $p < 0.0$	11					
Middle Africa						
Cameroon, 2011	1905	5860	•	32.51	[31.31; 33.71]	1.6%
Chad, 2015	4331	10854	-	39.90	[38.98; 40.83]	1.6%
Equatorial Guinea. 2011	3656 287	5030 1094		+2.70	[23.67: 28.881	1.6%
Gabon, 2012	636	3856	• -	16.49	[15.34; 17.68]	1.6%
Republic of Congo, 2012	1120	4591	•	24.40	[23.16; 25.65]	1.6%
Sao Tome and Principe, 2009	452	1544	.	29.27	[27.03; 31.57]	1.6%
Random effects model Heterogeneity: $l^2 = 100\%$, $r^2 = 0.0115$, p.c.0	01	36829		29.88	[22.88; 37.39]	11.3%
p < 0.	~ .					
Northern Africa			_			
Egypt, 2014 Random effects model	2911	13601 13601		21.40	[20.72; 22.10]	1.6%
Heterogeneity: not applicable		13001	· · · · ·	21.40	[20.72, 22.10]	1.0 /6
Southern Africa	050	2040	1	20.04	127 20- 20 503	1 60/
⊑swauni, 2007 Lesotho, 2014	850	2940 1860		∠8.91 33.22	[27.29; 30.56] [31.11:35.39]	1.6%
Namibia, 2013	542	2287	• ⁻	23.70	[21.98; 25.46]	1.6%
South Africa, 2016	385	1404	#	27.42	[25.12; 29.79]	1.6%
Random effects model		8500	+	28.25	[24.47; 32.18]	6.4%
rreterogeneity: I ⁻ = 94%, τ ⁻ = 0.0018, p < 0.0						
Central Asia						
Kyrgyz Republic, 2012	768	4337		17.71	[16.59; 18.86]	1.6%
Tajikistan, 2017	1171	6694		17.49	[16.59; 18.41]	1.6%
Kandom effects model Heterogeneity: $y^2 = 0.9$, $z^2 = 0.9$, $z = 0.77$		11031	•	17.58	[16.87; 18.29]	3.2%
$r_{0000} = 0.000 = 0.0000 = 0.0000000000000$						
Polynesia						
Samoa, 2008	2282	8422	•	27.10	[26.15; 28.05]	1.6%
Random effects model		8422	•	21.10	[∠0.15; 28.05]	1.0%
посотоденных, посаррисацие						
Random effects model		624734 _	<u> </u>	29.12	[26.67; 31.63]	100.0%
Heterogeneity: $l^2 = 100\%$, $\tau^2 = 0.0120$, $p = 0$		Г С	20 40			
rvesidual neterogeneity: /* = 100%, p = 0		0	20 40 60 Prevalence of stunting (%)			
			diorioo or ordining (/u)			

Figure 2. Forest plot of stunting prevalence by UN subregions of LMICs: event values represent the number of cases of stunting expressed as a percentage. Blue squares and their corresponding lines are the point estimates and 95% confidence intervals (95% CI). Maroon diamonds represent the pooled estimate of the prevalence for each subgroup (width denotes 95% CI). Weights are from the random-effects meta-analysis model described by DerSimonian and Laird²⁰ (*p* for interaction comparing the different subgroups < 0.0001).

of stunting for Southern and Southeastern Asia was almost two times the burden of stunting in Western and Central Asia, and 10 percentage points greater than in Eastern Asia.

At the country level, nations with high stunting prevalence (all with prevalence greater than 40%) were mostly SSA countries—Burundi, Democratic Republic of Congo, Madagascar, Mozambique, Niger and Zambia. Yemen in Western Asia, Timor-Leste in Southeastern Asia and Guatemala in Southern America also displayed a stunting prevalence of 40% and higher. Several countries had high prevalence of both stunting and wasting. For example, Timor-Leste in Southeastern Asia and India in Southern Asia had high prevalence of both stunting and wasting.

Wasting. Figure 3 shows the prevalence of wasting on a subregional and country level. On a regional level (Supplemental Figure S5), the prevalence of wasting in Asia was 3 percentage points greater than that of Africa but 7 to 8 percentage points greater than that of Europe, the Americas and Oceania. On a subregional level, substantial variations in the burden of wasting existed. For example, in Africa, the prevalence in Western Africa was almost twice as high as that of Eastern and Middle Africa and three times that of Southern Africa. At the country level, 13 nations had > 10% prevalence of wasting. These included five countries from Asia—Timor-Leste, India, Bangladesh, Yemen and Turkey, and eight countries from SSA—Nigeria, Niger, Burkina Faso, Mali, Chad, Sao Tome Principe, The Gambia, and Comoros.

Underweight. Figure 4 shows the prevalence of underweight on the subregional and country level. The prevalence of underweight in Africa and Asia was twice that of the Americas, four times that of Oceania and more than ten times that of Europe (Supplemental Figure S6). Within SSA, Western Africa had the highest prevalence of underweight, with its pooled estimates twice that of Southern Africa. However, Western African underweight prevalence was not substantially different compared to Eastern and Middle Africa. Similarly, the prevalence of underweight in Central Asia was, on average, five times less than that of Eastern, Southeastern and Southern Asia. Countries that displayed a very high prevalence of underweight (all with a prevalence $\geq 30\%$) were SSA countries, Niger and Burundi, and in Asia, India, Timor-Leste, Bangladesh and Yemen. On the contrary, the prevalence of underweight in Latin America and the Caribbean were, on average, substantially lower than the pooled overall prevalence.

Association of human development index and UN region with stunting, wasting and underweight. A moderate negative correlation was observed between the country's HDI and all undernutrition forms (Fig. 5). Stunting (Spearman's rho; -0.65, p value < 0.0001), wasting (Spearman's rho; -0.43, p value = 0.0006) and underweight (Spearman's rho; -0.67, p value < 0.0001). The generalized linear mixed-effects meta-regression model suggested higher HDI was associated with lower odds of all three forms of undernutrition. For each increase in the level of HDI, the odds were 40%, 37%, and 51% lower for stunting, wasting, and underweight, respectively.

The United Nations region was associated with the prevalence of undernutrition. Eastern and Middle Africa, Southern and Southeastern Asia, and Polynesia displayed higher odds of stunting compared to Central Asia (Table 1). Wasting prevalence odds were significantly higher in Western, Southern, and Southeastern Asia, and Northern Africa than in Central Asia. However, Latin America and the Caribbean had lower odds of wasting compared to Central Asia. In the model, combining HDI and UN subregions (as a proxy for spatial variation) explained 54%, 56%, and 66% for stunting, wasting and underweight prevalence, respectively.

Discussion

We estimated the prevalence of childhood (0–59 months) undernutrition (stunting, wasting and underweight) using data from the most recent DHS from 62 LMICs. Our results suggest that exploring undernutrition on a global or regional level could mask the unique differences of the disease burden within a sub-region level. Western Africa, Southern, and Southeastern Asia consistently displayed a substantial burden of stunting, wasting and underweight. Six of the nine countries with the highest burden of stunting were from SSA and two were from Asia. A combination of human development index and United Nations region (a proxy for geographical variation) explained 54%, 56%, and 66% of the variation in stunting, wasting, and underweight prevalence, respectively. The residual unexplained variance implied by these figures after optimizing for random effects suggest there are additional factors involved in these disparities.

The results of this study demonstrate a moderate association between a country's HDI with undernutrition. One of the components of HDI is the gross national income (GNI). Thus, countries with lower GNI had higher prevalence of undernutrition. Eleven percent of the world's population is living in poverty, defined by The World Bank as living on less than US\$1.90 per day. Poverty disproportionally impacts children particularly those living in SSA and Southern Asia²⁶. As a result, low- and middle-income countries have the highest burden of stunting, wasting and underweight and children in SSA and Southern Asia are disproportionately affected. The United Nation's Millennium Development Goals include eradicating extreme poverty and hunger as the priority goal²⁷, and continues to be a key global development agenda under the Sustainable Development Goals (SDGs)²⁸. Like poverty, undernutrition often occurs in an intergenerational cycle. In addition to being a risk factor for infections and food insecurity—both of which are drivers of undernutrition—poverty is a pivotal contributor to allostatic load (the cumulative wear and tear on the body due to adapting to adverse physical or psychosocial stress), which modulates the biological mechanisms that influence growth²⁶. Consequently, early childhood growth failure in LMICs has persisted despite decades of nutritional interventions. Children born into low-income families have, on average, poorer growth, poorer neurocognitive outcomes, and poorer educational attainment than wealthier peers. Such setbacks are, in turn, associated with lower economic productivity, thus contributing to the intergenerational transmission of poverty and undernutrition.

				Events per 100			
Country, Year of survey	Wasted	Total		observations	Events	95%-CI	Weight
Southern Europe							
Albania, 2018	34	2299			1.48	[1.02; 2.02]	1.6%
Random effects model		2299	•		1.48	[1.02; 2.02]	1.6%
Heterogeneity: not applicable							
Nestern Africa							
Burkina Faso. 2010	1084	6994		+	15.50	[14.66: 16.36]	1.6%
Cote d'Ivoire, 2012	269	3581		+	7.51	[6.67; 8.40]	1.6%
Guinea, 2012	339	3531		+	9.60	[8.65; 10.59]	1.6%
Niger, 2012	987	5481		•	18.01	[17.00; 19.04]	1.6%
Jambia, 2013	388	3372			11.51	[10.45; 12.61]	1.6%
Joena, 2013 Mali 2013	617	4857			12 70	[5.23, 6.60]	1.6%
Nigeria, 2013	4714	26190			18.00	[17.54: 18.47]	1.6%
Sierra Leone, 2013	474	5094			9.31	[8.52; 10.12]	1.6%
Ghana, 2014	136	2895	-		4.70	[3.96; 5.50]	1.6%
Togo, 2014	213	3282	_	-	6.49	[5.67; 7.36]	1.6%
Angola, 2016 Sepagal 2017	360	10090	-	_	4.79	[4.32; 5.29]	1.6%
Senin 2018	650	13260		-	4 90	[4.54: 5.28]	1.6%
Random effects model		100547	-	-	9.41	[6.78; 12.43]	23.0%
Heterogeneity: I ² = 100%, τ ² = 0.0085, ρ = 0							
Western Asia	105	1070		_	6 00	1 5 75, 7 001	4 69/
Azerbaijan, 2006 Furkov, 2012	135	1979		.	10.02	[5.75; 7.98]	1.6%
Vemen 2013	2253	13823		· · ·	16.30	[15.69:16.92]	1.6%
Armenia, 2016	62	1555	-	-	3.99	[3.07; 5.02]	1.6%
Random effects model		19876	_		9.01	[4.19; 15.42]	6.5%
Heterogeneity: $I^2 = 99\%$, $\tau^2 = 0.0100$, $p < 0.0$	01						
Southorn Asia							
Southern Asia	1046	7318			1/ 20	[13 50: 15 10]	1.6%
ndia. 2016	46157	219796			21.00	[20.83: 21.17]	1.6%
Nepal, 2016	232	2417		+	9.60	[8.46; 10.81]	1.6%
Maldives, 2017	203	2260		. +	8.98	[7.84; 10.20]	1.6%
Pakistan, 2018	245	3547		=	6.91	[6.10; 7.77]	1.6%
Random effects model	01	235338			11.76	[6.54; 18.26]	8.2%
Heterogeneity: $T = 100\%$, $\tau = 0.0108$, $p < 0$.01						
Latin America and the Caribbean							
Bolivia, 2008	118	8422	0		1.40	[1.16; 1.66]	1.6%
Guyana, 2009	81	1522	-	÷	5.32	[4.25; 6.51]	1.6%
Colombia, 2010	141	15702			0.90	[0.76; 1.05]	1.6%
Honduras, 2012	142	10167			1.40	[1.18; 1.63]	1.6%
Cominican Republic 2013	55	3610	*		1 00	[0.45, 0.77]	1.0%
Guatemala, 2015	88	12567	a 🗌		0.70	[0.56; 0.85]	1.6%
Haiti, 2017	237	6589			3.60	[3.16; 4.06]	1.6%
Random effects model		67756	+		1.71	[1.06; 2.51]	13.1%
Heterogeneity: $I^2 = 98\%$, $\tau^2 = 0.0016$, $p < 0.0$	01						
Eastern Africa							
Madagascar, 2009		5436					0.0%
Mozambique, 2011	608	10313			5.90	[5.45; 6.36]	1.6%
Comoros, 2012	1205	10854			11.10	[10.52; 11.70]	1.6%
Kenya, 2014	759	18986			4.00	[3.72; 4.28]	1.6%
Zambia, 2014	740	12328		-	6.00	[5.59; 6.43]	1.6%
Zimbahwa 2015	10/	6255			2.20	[1.76, 2.69]	1.0%
Ethiopia, 2016	1020	10412	-		9.80	[9.23: 10.37]	1.6%
Malawi, 2016	156	5764			2.71	[2.30; 3.14]	1.6%
Tanzania, 2016	432	9811			4.40	[4.01; 4.82]	1.6%
Jganda, 2016	176	5191	•		3.39	[2.91; 3.90]	1.6%
Burundi, 2017	323	6453			5.01	[4.49; 5.55]	1.6%
Heterogeneity: $l^2 = 99\%$, $r^2 = 0.0035$, $p < 0.0000000000000000000000000000000000$	n 1	103010	-		4.95	[3.54; 0.57]	10.1%
leterogeneity. 1 = 33 %, t = 0.0033, p < 0.0	51						
Southeastern Asia							
Cambodia, 2014	470	4893		+	9.61	[8.80; 10.45]	1.6%
Myanmar, 2016	281	4076		*	6.89	[6.14; 7.69]	1.6%
Timor-Leste, 2016	1464	6476		+	22.61	[21.60; 23.63]	1.6%
Heterogeneity: $l^2 = 100\%$ $\tau^2 = 0.0158$ $p < 0$	01	15445			12.33	[4.01; 23.09]	4.9%
leterogeneity: 7 = 100 %, t = 0.0130, p < 0							
Middle Africa							
Sao Tome and Principe, 2009	162	1544			10.49	[9.01; 12.07]	1.6%
Cameroon, 2011	328	5860	_ •		5.60	[5.02; 6.20]	1.6%
Equatorial Guinea, 2011	34	1094			3.11	[2.16; 4.23]	1.6%
Sabon, 2012 Republic of Congo 2012	271	4591	- T.		5.29	[2.75, 3.00]	1.6%
Democratic Republic of the Congo. 2014	713	9030			7.90	[7.35: 8.46]	1.6%
Chad, 2015	1411	10854			13.00	[12.37; 13.64]	1.6%
Random effects model		36829	-	-	6.68	[4.26; 9.59]	11.5%
Heterogeneity: I ² = 99%, τ ² = 0.0051, p < 0.0	01						
Northorn Africa							
Forunt 2014	1142	13601			8 40	[7 94·8 87]	1.6%
Random effects model		13601		•	8.40	[7.94: 8.87]	1.6%
Heterogeneity: not applicable							
Southern Africa	.	00.45	-		0.50	14.00 0.10	4.001
swatini, 2007	/4	2940			2.52	[1.98; 3.12]	1.6%
_esotho. 2014	52	2207 1869	_ # 1	T	2.78	[2.08: 3.58]	1.6%
South Africa, 2016	33	1384			2.38	[1.64; 3.26]	1.6%
Random effects model		8480	-		3.34	[1.87; 5.21]	6.5%
Heterogeneity: $I^2 = 95\%$, $\tau^2 = 0.0021$, $p < 0.0$	D1						
Surguz Republic 2012	447	1007			2 70	1224.2201	1 6%
Taiikistan 2017	368	4337	- × -		2.70	[4.97 6.07]	1.6%
Random effects model	500	11021	-	-	3.99	[1.70: 7.19]	3.3%
Heterogeneity: $I^2 = 98\%$, $\tau^2 = 0.0025$, $p < 0.0$	D1					,	
Polynesia	440	0400			1 40	1116: 1.003	1 69/
Samoa, 2008 Pandom effects model	118	8422 8422			1.40	[1.10; 1.66]	1.0%
Heterogeneity: not applicable		0422			1.40	[1.10, 1.00]	1.0 /0
, , , , , , , , , , , , , , , , , , ,							
Random effects model		625230		►	6.27	[4.61; 8.16]	100.0%
Heterogeneity: $I^2 = 100\%$, $\tau^2 = 0.0213$, $p = 0$			і — Т		_		
Residual heterogeneity: $I^2 = 99\%$, $p = 0$			U 5	10 15 20 25	>		
			Prev	arence or wasting (%)			

Figure 3. Forest plot of wasting prevalence by UN subregions of LMICs: event values represent the number of cases of wasting expressed as a percentage. Blue squares and their corresponding lines are the point estimates and 95% confidence intervals (95% CI). Maroon diamonds represent the pooled estimate of the prevalence for each subgroup (width denotes 95% CI). Weights are from the random-effects meta-analysis model described by DerSimonian and Laird²⁰ (*p* for interaction comparing the different subgroups < 0.0001).

Country, Year of survey	Underweight	Total		Events per 100 observations	Events	95%-CI	Weight
Southern Europe Albania, 2018 Random effects model Heterogeneity: not applicable	33	2367 2367	•		1.39 1.39	[0.96; 1.91] [0.96; 1.91]	1.6% 1.6%
Western Africa							
Burkina Faso, 2010	1797	6994			25.69	[24.68; 26.72]	1.6%
Guinea, 2012	636	3531		-	18.01	[16.76; 19.30]	1.6%
Niger, 2012	1995	5481			36.40	[35.13; 37.68]	1.6%
Gambia, 2013 Liberia, 2013	546 528	3372			16.19	[14.97; 17.46]	1.6%
Mali, 2013	1239	4857			25.51	[24.29; 26.75]	1.6%
Nigeria, 2013	7517	26190		•	28.70	[28.16; 29.25]	1.6%
Ghana, 2014	835 318	2895	-	•	10.98	[15.39; 17.42]	1.6%
Togo, 2014	525	3282	_	=	16.00	[14.76; 17.27]	1.6%
Angola, 2016 Sepagal 2017	1382	7468			18.51	[17.63; 19.39]	1.6%
Benin, 2018	2130	12832		0	16.60	[15.96; 17.25]	1.6%
Random effects model Heterogeneity: I^2 = 100%, τ^2 = 0.0078, p = 0		100007		-	19.15	[15.63; 22.93]	23.0%
Western Asia							
Azerbaijan, 2006	152	1979			7.68	[6.55; 8.90]	1.6%
Turkey, 2013	131	2519		_	5.20	[4.37; 6.10]	1.6%
Yemen, 2013 Armenia, 2016	5391	13823		•	39.00	[38.19; 39.81]	1.6%
Random effects model	00	19930			10.85	[0.18; 34.49]	6.5%
Heterogeneity: $I^2 = 100\%$, $\tau^2 = 0.0885$, $p = 0$							
Southern Asia Bangladesh, 2014	2386	7318			32.60	[31,54: 33 681	1.6%
India, 2016	78467	219796		0	35.70	[35.50; 35.90]	1.6%
Nepal, 2016	656	2428			27.02	[25.27; 28.80]	1.6%
Pakistan, 2018	342 808	2327 3622		-	14.70	[13.29; 16.17] [20.97; 23.68]	1.6%
Random effects model	000	235491			26.12	[19.54; 33.28]	8.2%
Heterogeneity: $I^2 = 100\%$, $\tau^2 = 0.0080$, $p < 0$.01						
Latin America and the Caribbean			_				
Bolivia, 2008 Guyana, 2009	362	8422	۹ _–		4.30	[3.88; 4.74]	1.6% 1.6%
Colombia, 2009	534	15702			3.40	[3.12; 3.69]	1.6%
Honduras, 2012	712	10167			7.00	[6.51; 7.51]	1.6%
Peru, 2012	312	9168			3.40	[3.04; 3.78]	1.6%
Guatemala, 2015	1583	12567	- T - E		12.60	[12.02; 13.18]	1.6%
Haiti, 2017	625	6646			9.40	[8.71; 10.12]	1.6%
Random effects model Heterogeneity: I^2 = 99%, τ^2 = 0.0057, $p < 0.0$)1	67813	-		6.41	[4.08; 9.21]	13.1%
Eastern Africa							
Madagascar, 2009		5436					0.0%
Mozambique, 2011	1537	10313		0	14.90	[14.22; 15.60]	1.6%
Kenva. 2014	2088	18986			11.00	[10.56: 11.45]	1.6%
Zambia, 2014	1825	12328			14.80	[14.18; 15.44]	1.6%
Rwanda, 2015 Zimbabwe, 2015	355	3813			9.31	[8.41; 10.25]	1.6%
Ethiopia, 2016	2459	10552		•	23.30	[22.50; 24.12]	1.6%
Malawi, 2016	660	5786			11.41	[10.60; 12.24]	1.6%
Tanzania, 2016	1335	9886 5136		•	13.50	[12.84; 14.18]	1.6%
Burundi, 2017	1887	6464	-		29.19	[28.09; 30.31]	1.6%
Random effects model Hotomagnetity $l^2 = 100\%$ $r^2 = 0.005\%$ $r = 0$		105906	-	-	14.23	[11.22; 17.53]	18.0%
Preterogeneity: $T = 100\%$, $\tau = 0.0038$, $p = 0$							
Cambodia, 2014	1169	4893			23.89	[22.71; 25.10]	1.6%
Myanmar, 2016	775	4100			18.90	[17.72; 20.12]	1.6%
Timor-Leste, 2016	2875	7206		•	39.90	[38.77; 41.03]	1.6%
Heterogeneity: $l^2 = 100\%$, $\tau^2 = 0.0161$, $p < 0$.01	10155			27.15	[13.47, 40.72]	4.3 /0
Middle Africa							
Sao Tome and Principe, 2009	202	1544	4	<u>t</u>	13.08	[11.45; 14.81]	1.6%
Equatorial Guinea, 2011	42	1094			3.84	[2.78: 5.06]	1.6%
Gabon, 2012	231	3856			5.99	[5.26; 6.76]	1.6%
Republic of Congo, 2012	533	4591			11.61	[10.70; 12.55]	1.6%
Chad, 2015	3126	10854		- o	28.80	[27.95; 29.66]	1.6%
Random effects model Heterogeneity: $l^2 = 100\%$, $\tau^2 = 0.0159$, $p = 0$		36829			13.36	[7.67; 20.34]	11.5%
Northern Africa	749	12601			E E0	1 5 12 5 901	1.6%
Random effects model	740	13601	- T		5.50	[5.12; 5.89]	1.6%
Heterogeneity: not applicable							
Southern Africa							
Eswatini, 2007	159	2940			5.41	[4.62; 6.26]	1.6%
Namibia, 2013	304	2287		•	13.29	[11.93; 14.72]	1.6%
Lesotho, 2014 South Africa, 2016	193	1869			10.33	[8.99; 11.75]	1.6%
Random effects model	02	8512	-		8.43	[5.02; 12.61]	6.5%
Heterogeneity: $l^2 = 98\%$, $\tau^2 = 0.0047$, $p < 0.0047$)1						
Central Asia							
Kyrgyz Republic, 2012	147	4337			3.39	[2.87; 3.95]	1.6%
lajikistan, 2017 Random effects model	504	6716			7.50	[6.89; 8.15]	1.6%
Heterogeneity: $I^2 = 99\%$, $\tau^2 = 0.0042$, $p < 0.0$)1	11033	_		J.20	[1.33, 10.00]	J.J /0
Polynesia							
Samoa, 2008	362	8422			4.30	[3.88; 4.74]	1.6%
Random effects model		8422	•		4.30	[3.88; 4.74]	1.6%
Heterogeneity: not applicable							
Random effects model		626130		-	13.71	[10.85; 16.85]	100.0%
Heterogeneity: $I^2 = 100\%$, $\tau^2 = 0.0302$, $p = 0$				20 20 40			
readual neterogeneity: r = 100%, p = 0			 Prevale 	∠u 3u 4U 50 nce of underweight (%)			

Figure 4. Forest plot of underweight prevalence by UN subregions of LMICs: event values represent the number of cases of underweight expressed as a percentage. Blue squares and their corresponding lines are the point estimates and 95% confidence intervals (95% CI). Maroon diamonds represent the pooled estimate of the prevalence for each subgroup (width denotes 95% CI). Weights are from the random-effects meta-analysis model described by DerSimonian and Laird²⁰ (*p* for interaction comparing the different subgroups < 0.0001).



Figure 5. Correlation of HDI with undernutrition: A moderate negative correlation exists between HDI and stunting (Spearman's rho; -0.65, *p* value < 0.0001, first column), wasting (Spearman's rho; -0.43, *p* value = 0.0006, second column) and underweight (Spearman's rho; -0.67, *p* value < 0.0001, third column).

	Stunting		Wasting		Underweight					
Variable	Odds ratio (95%CI)	<i>p</i> value	Odds ratio (95%CI)	<i>p</i> value	Odds ratio (95%CI)	<i>p</i> value				
UN subregion										
Africa										
Eastern Africa	1.88 (1.06, 3.36)	0.032	0.87 (0.42, 1.81)	0.718	1.79 (0.95, 3.36)	0.073				
Western Africa	1.36 (0.76, 2.43)	0.296	1.65 (0.80, 3.42)	0.176	2.35 (1.25, 4.40)	0.008				
Middle Africa	1.82 (1.01, 3.29)	0.047	1.57 (0.75, 3.30)	0.235	2.39 (1.25, 4.55)	0.008				
Southern Africa	1.84 (0.97, 3.48)	0.062	0.82 (0.36, 1.84)	0.629	1.65 (0.82, 3.31)	0.160				
Northern Africa	2.11 (0.84, 5.30)	0.110	3.60 (1.14, 11.36)	0.029	2.21 (0.82, 6.00)	0.119				
Asia										
Western Asia	1.44 (0.76, 2.75)	0.266	2.90 (1.29, 6.52)	0.010	2.53 (1.25, 5.12)	<.00001				
Southern Asia	2.41 (1.30, 4.46)	0.005	3.49 (1.61, 7.58)	0.002	7.44 (3.80, 14.57)	<.00001				
Southeastern Asia	2.57 (1.31, 5.04)	0.006	3.28 (1.41, 7.63)	0.006	6.82 (3.28, 14.15)	<.00001				
Central Asia	Reference		Reference		Reference					
Americas										
Latin America and the Carib- bean	1.41 (0.79, 2.54)	0.247	0.45 (0.22, 0.95)	0.036	1.56 (0.83, 2.96)	0.170				
Oceania										
Polynesia	2.89 (1.15, 7.23)	0.024	0.56 (0.17, 1.78)	0.323	1.71 (0.63, 4.64)	0.295				
Europe										
Southern Europe	0.99 (0.39, 2.49)	0.976	0.58 (0.18, 1.93)	0.376	0.53 (0.19, 1.52)	0.239				
HDI	0.60 (0.50, 0.72)	<.00001	0.63 (0.50, 0.79)	<.00001	0.49 (0.40, 0.60)	<.00001				

Table 1. Maximum likelihood estimates with associated 95% confidence intervals (CI) for the generalized linear mixed-effects model's regression coefficients. Odds ratios and their 95% confidence intervals for the association of United Nations regions and Human Development Index with undernutrition (stunting, wasting and underweight). The multivariable model included UN subregions and human development index. R² is the coefficient of determination indicating the amount of variation explained by the variables. The R² when HDI and UN subregions are included in the random-effects model is 54%, 56%, and 66% for stunting, wasting and underweight. Higher HDI was associated with lower odds of all three forms of undernutrition.

Lower HDI is associated with higher rates of infectious diseases²⁹⁻³². The interaction between undernutrition and infection creates a lethal cycle of worsening illness and deteriorating nutritional status. In regions that are profoundly affected by undernourishment, infection prevalence rates are also high. SSA and South and Southeastern Asia are disproportionately affected by malaria, human immunodeficiency virus (HIV), and tuberculosis (TB), all of which are associated with worsening the nutritional status of the child³³. Above and beyond the immediate outcomes such as death, infectious diseases expose children to a complex constellation of developmental risk factors embedded in contexts of biological stress and psychosocial disadvantage that perpetuates child growth failure. Infections induce a nutritional demand on biological systems through fevers and increased catabolic states by mediators of inflammation³⁴. This results in increased energy expenditure so that fewer calories are available to support growth. Furthermore, infections often decrease food intake and alter digestion and absorption, further worsening the nutritional status. Undernutrition increases susceptibility to infection, and on the other hand, infections propagate deterioration of nutritional status, resulting in a synergistic cycle of undernutrition-infection that ultimately leads to severe undernutrition³⁵.

Public health relevance and recommendation. Adopted by the UN General Assembly in the September of 2015, the Sustainable Development Goals represent a new coherent way of thinking about how issues as diverse as ending poverty (goal 1), ending hunger (goal 2), health promotion (goal 3), achieving quality education for all (goal 4), achieving gender equality (goal 5), and climate change, fit together to foster international development³⁶. Particularly, SDG 2.2 calls for an end to all forms of malnutrition by 2030, and is inseparable from many of the other SDG³⁷. The updated quantitative assessments of levels of undernutrition indicators presented here should inform strategies at the regional and national levels targeted at achieving these SDGs. Lastly, the substantial unexplained variance in our findings emphasizes the need for further characterization of the correlates of undernutrition in LIMCs including genomics, gut microbiota, ethnicity, diet composition, micronutrients, climate change and weather variability³⁸⁻⁴³. Such analysis should also focus on using more granular subnational data.

Strengths and limitations. The strengths of our study include that we only analyzed anthropometric measurement using the new modified 2006 WHO international standards of child growth measurements⁷. The nutrition analyses done using older growth references do not adequately represent early childhood growth^{44,45}. In addition, we analyzed all three indicators of undernutrition at the global, regional, and subregional scale. These three indicators are distinct on their effect on the physical and neurocognitive outcome of the child, and therefore should be presented separately⁴⁶.

Our study had some limitations. First, because we employed an ecological (aggregated data) study design, our findings may suffer bias and confounding from the ecological fallacy (drawing conclusions on individuals using aggregated data, even though the relevant individuals may not share such characteristics)⁴⁷. Second, WHO international child growth standards data used were a global median. Because height is heritable there are likely genetic variations in height at the country-level not accounted for^{48,49}. Nevertheless, at younger ages, differences in height and weight in preschool children that is explained by ethnic background are relatively small—3% for height and about 6% for weight, and the differences in these anthropometric measures at such ages are primarily driven by nutrition status of the child^{50,51}. The WHO attempted to overcome this limitation by representing the major global geographic regions in generating new growth curves¹¹. Thirdly, although we limited the analysis to the recent nationwide DHS data, the twelve-year window (2006–2018) is quite broad and the relative metrics of under nutrition continue to evolve with time. Finally, due to the bidirectional and complex relationship between major infectious diseases (including malaria, HIV, TB) and undernutrition⁵², we did not include these diseases in the meta-regression analysis since we cannot establish a causal link especially with the cross-sectional nature of the present study. Future longitudinal studies of individual-level data should estimate the contribution of major endemic infectious diseases to the risk of childhood undernutrition in multiple LMICs.

Conclusion

In summary, substantial regional, subregional and country-level disparities of stunting, wasting and underweight still exist. The updated quantitative assessments of levels of undernutrition indicators presented here should inform strategies at the global, regional and national levels targeted at further reducing the remaining substantially undernourished populations.

Data availability

The analyzed dataset is freely available from: https://dhsprogram.com/data/available-datasets.cfm. R code and data to reproduce the results in the present manuscript are archived at https://github.com/Schiff-Lab/Globa l-Malnutrition-Burden-Scientific-Reports-2021.

Received: 14 August 2020; Accepted: 15 February 2021 Published online: 04 March 2021

References

- 1. Gakidou, E. *et al.* Global, regional, and national comparative risk assessment of 84 behavioural, environmental and occupational, and metabolic risks or clusters of risks, 1990–2016: A systematic analysis for the Global Burden of Disease Study 2016. *The Lancet* **390**, 1345–1422. https://doi.org/10.1016/S0140-6736(17)32366-8 (2017).
- Lim, S. S. et al. A comparative risk assessment of burden of disease and injury attributable to 67 risk factors and risk factor clusters in 21 regions, 1990–2010: A systematic analysis for the Global Burden of Disease Study 2010. The Lancet 380, 2224–2260. https:// doi.org/10.1016/S0140-6736(12)61766-8 (2012).
- Akombi, B. J., Agho, K. E., Merom, D., Renzaho, A. M. & Hall, J. J. Child malnutrition in sub-Saharan Africa: A meta-analysis of demographic and health surveys (2006–2016). *PLoS ONE* 12, e0177338 (2017).
- 4. United Nations Children's Fund (UNICEF), W. H. O., International Bank for Reconstruction and Development/The World Bank. Levels and trends in child malnutrition: key findings of the 2019 Edition of the Joint Child Malnutrition Estimates. Geneva: World Health Organization. https://www.who.int/nutgrowthdb/jme-2019-key-findings.pdf?ua=1 (2019).
- McDonald, C. M. et al. Stunting and wasting are associated with poorer psychomotor and mental development in HIV-exposed tanzanian infants. J. Nutr. 143, 204–214. https://doi.org/10.3945/jn.112.168682 (2012).

- Victora, C. G. et al. Maternal and child undernutrition: Consequences for adult health and human capital. The Lancet 371, 340–357 (2008).
- Group, W. M. G. R. S. WHO Child Growth Standards based on length/height, weight and age. Acta paediatrica (Oslo, Norway: 1992). Supplement 450, 76 (2006).
- 8. Ties Boerma, J. & Sommerfelt, A. E. Demographic and health surveys (DHS: contributions and limitations (1993).
- 9. Demographic and Health Survey Program. Demographic and Health Survey Program. https://dhsprogram.com (2020).
- 10. Jain, A. K. & Hausman, R. E. Stratified multistage sampling. Wiley StatsRef: Statistics Reference Online (2014).
- World Health Organization. WHO Child Growth Standards: Length/Height for Age, Weight-for-Age, Weight-for-Length, Weight-for-Height and Body Mass Index-for-Age, Methods and Development (World Health Organization, 2006).
- de Onis, M., Onyango, A. W., Van den Broeck, J., Chumlea, W. C. & Martorell, R. Measurement and standardization protocols for anthropometry used in the construction of a new international growth reference. *Food Nutr. Bull.* 25, S27–S36 (2004).
- Dibley, M. J., Goldsby, J. B., Staehling, N. W. & Trowbridge, F. L. Development of normalized curves for the international growth reference: Historical and technical considerations. *Am. J. Clin. Nutr.* 46, 736–748 (1987).
- 14. Kuczmarski, R. J. 2000 CDC Growth Charts for the United States: Methods and development (Department of Health and Human Services, Centers for Disease Control and..., 2002).
- 15. Bhan, M. K. & Norum, K. R. The WHO multicentre growth reference study (MGRS): Rationale, planning, and implementation. *Food Nutr. Bull.* **25**, 66 (2004).
- Cole, T. J. & Green, P. J. Smoothing reference centile curves: The LMS method and penalized likelihood. Stat. Med. 11, 1305–1319 (1992).
- 17. United Nations Development Programme. Human development index (HDI). http://hdr.undp.org/en/data (2020).
- Boyle, M. H. et al. The influence of economic development level, household wealth and maternal education on child health in the developing world. Soc. Sci. Med. 63, 2242–2254 (2006).
- Fagbamigbe, A., Kandala, N. & Uthman, A. Demystifying the factors associated with rural-urban gaps in severe acute malnutrition among under-five children in low-and middle-income countries: A decomposition analysis. *Sci. Rep.* 10, 1–15 (2020).
- DerSimonian, R. & Laird, N. Meta-analysis in clinical trials. Control. Clin. Trials 7, 177–188. https://doi.org/10.1016/0197-2456(86)90046-2 (1986).
- 21. Viechtbauer, W. Conducting meta-analyses in R with the metafor package. J. Stat. Softw. 36, 1-48 (2010).
- 22. Schwarzer, G., Chemaitelly, H., Abu-Raddad, L. J. & Rücker, G. Seriously misleading results using inverse of Freeman-Tukey double arcsine transformation in meta-analysis of single proportions. *Res. Synth. Methods* (2019).
- 23. Higgins, J. P., Thompson, S. G., Deeks, J. J. & Altman, D. G. Measuring inconsistency in meta-analyses. BMJ 327, 557-560 (2003).
- 24. United Nations. Definition of Regions. https://population.un.org/wpp/DefinitionOfRegions/ (2019).
- 25. Schwarzer, G., Carpenter, J. R. & Rücker, G. Meta-Analysis with R (Springer, Berlin, 2015).
- Jensen, S. K., Berens, A. E. & Nelson, C. A. 3rd. Effects of poverty on interacting biological systems underlying child development. Lancet Child Adoles. Health 1, 225–239 (2017).
- 27. 27Summit, N. M. Millennium Development Goals. Human Development Report https://www.un.org/millenniumgoals/ (2000).
- Gil, J. D. B. et al. Sustainable development goal 2: Improved targets and indicators for agriculture and food security. Ambio 48, 685–698 (2019).
- 29. Goodarzi, E. et al. Geographical distribution of falciparum malaria in the world and its relationship with the human development index (HDI): countries based on the WHO report in 2017. Journal of Public Health **66**, 1–10 (2020).
- Gupta, S. et al. Part III. Analysis of data gaps pertaining to enterotoxigenic Escherichia coli infections in low and medium human development index countries, 1984–2005. Epidemiol. Infect. 136, 721–738 (2008).
- 31. Lou, L.-X., Chen, Y., Yu, C.-H., Li, Y.-M. & Ye, J. National HIV/AIDS mortality, prevalence, and incidence rates are associated with the human development index. *Am. J. Infect. Control* **42**, 1044–1048 (2014).
- Rodrigues-Júnior, A. L., Ruffino-Netto, A. & Castilho, E. A. d. Spatial distribution of the human development index, HIV infection and AIDS-Tuberculosis comorbidity: Brazil, 1982–2007. Revista Brasileira de Epidemiologia 17, 204–215 (2014).
- 33. Murray, C. J. *et al.* Global, regional, and national incidence and mortality for HIV, tuberculosis, and malaria during 1990–2013: A systematic analysis for the Global Burden of Disease Study 2013. *The Lancet* **384**, 1005–1070 (2014).
- 34. Hotamisligil, G. S. & Erbay, E. Nutrient sensing and inflammation in metabolic diseases. Nat. Rev. Immunol. 8, 923 (2008).
- Keusch, G. T. The history of nutrition: Malnutrition, infection and immunity. J. Nutr. 133, 336S-340S. https://doi.org/10.1093/ jn/133.1.336S (2003).
- Lee, B. X. et al. Transforming our world: Implementing the 2030 agenda through sustainable development goal indicators. J. Public Health Policy 37, 13–31 (2016).
- Fullman, N. et al. Measuring progress and projecting attainment on the basis of past trends of the health-related Sustainable Development Goals in 188 countries: An analysis from the Global Burden of Disease Study 2016. The Lancet 390, 1423–1459 (2017).
- Ssentongo, P. et al. Association of vitamin A deficiency with early childhood stunting in Uganda: A population-based cross-sectional study. PLoS ONE 15, e0233615 (2020).
- 39. Ssentongo, P., Ba, D. M., Fronterre, C. & Chinchilli, V. M. Village-level climate and weather variability, mediated by village-level crop yield, is associated with linear growth in children in Uganda. *BMJ Glob. Health* 5, e002696 (2020).
- Ssentongo, P. et al. Associations of malaria, HIV, and coinfection, with anemia in pregnancy in sub-Saharan Africa: A populationbased cross-sectional study. BMC Pregn. Childb. 20, 1–11 (2020).
- Ba, D. M., Ssentongo, P., Liao, D., Du, P. & Kjerulff, K. H. Non-iodized salt consumption among women of reproductive age in sub-Saharan Africa: A population-based study. *Public Health Nutr.* 6, 1–11 (2020).
- 42. Ba, D. M. et al. Adherence to Iron Supplementation in 22 sub-Saharan African countries and associated factors among pregnant women: A large population-based study. Curr. Dev. Nutr. 3, 20 (2019).
- Blanton, L. V., Barratt, M. J., Charbonneau, M. R., Ahmed, T. & Gordon, J. I. Childhood undernutrition, the gut microbiota, and microbiota-directed therapeutics. *Science* 352, 1533–1533 (2016).
- Black, R. E. et al. Maternal and child undernutrition: Global and regional exposures and health consequences. The Lancet 371, 243–260 (2008).
- 45. Black, R. E. *et al.* Maternal and child undernutrition and overweight in low-income and middle-income countries. *The Lancet* **382**, 427–451 (2013).
- 46. Amoah, B., Giorgi, E., Heyes, D. J., van Burren, S. & Diggle, P. J. Geostatistical modelling of the association between malaria and child growth in Africa. *Int. J. Health Geogr.* **17**, 7 (2018).
- 47. Sedgwick, P. Understanding the ecological fallacy. BMJ 351, h4773 (2015).
- Alves, A. C. *et al.* GWAS on longitudinal growth traits reveals different genetic factors influencing infant, child, and adult BMI. *Sci. Adv.* 5, eaaw3095 (2019).
- 49. Jorde, L. B. & Wooding, S. P. Genetic variation, classification and 'race'. Nat. Genet. 36, S28-S33 (2004).
- Habicht, J.-P., Yarbrough, C., Martorell, R., Malina, R. & Klein, R. Height and weight standards for preschool children: How relevant are ethnic differences in growth potential?. *The Lancet* 303, 611–615 (1974).
- 51. Dubois, L. et al. Genetic and environmental contributions to weight, height, and BMI from birth to 19 years of age: An international study of over 12,000 twin pairs. PLoS ONE 7, e30153 (2012).

 Oldenburg, C. E., Guerin, P. J., Berthé, F., Grais, R. F. & Isanaka, S. Malaria and nutritional status among children with severe acute malnutrition in Niger: A prospective cohort study. *Clin. Infect. Dis.* 67, 1027–1034 (2018).

Author contributions

P.S. has full access to all of the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis. Concept and design: P.S., V.M.C., S.J.S.. Data acquisition, analysis and interpretation: P.S., C.F., A.E.S., D.M.B., V.J.E.E., M.N., X.G., V.M.C., S.J.S.. Writing the manuscript: P.S., S.J.S.. Statistical analysis: P.S., C.F., V.M.C., X.G., S.J.S. Supervision: S.J.S., V.M.C.. All authors read and approved the final version of this manuscript. This study is part of the first author's doctoral dissertation with the College of Medicine, Pennsylvania State University, United States of America.

Funding

Support was provided by NIH Director's Pioneer Award 1DP1HD086071 (SJS, PS). The funder of the study had no role in study design, data collection, data analysis, data interpretation, or writing of the report. The corresponding author had full access to all the data in the study and had final responsibility for the decision to submit for publication.

Competing interests

The authors declare no competing interests.

Additional information

Supplementary Information The online version contains supplementary material available at https://doi.org/10.1038/s41598-021-84302-w.

Correspondence and requests for materials should be addressed to P.S.

Reprints and permissions information is available at www.nature.com/reprints.

Publisher's note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by/4.0/.

© The Author(s) 2021