BMJ Open Cohort profile: the BangladEsh Longitudinal Investigation of Emerging Vascular and nonvascular Events (BELIEVE) cohort study

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ABSTRACT

Purpose Bangladesh has experienced a rapid epidemiological transition from communicable to noncommunicable diseases (NCDs) in recent decades. There is, however, limited evidence about multidimensional determinants of NCDs in this population. The BangladEsh Longitudinal Investigation of Emerging Vascular and nonvascular Events (BELIEVE) study is a householdbased prospective cohort study established to investigate biological, behavioural, environmental and broader determinants of NCDs.

Participants Between January 2016 and March 2020, 73883 participants (aged 11 years or older) were recruited from 30817 households across urban, urban-poor ('slum') and rural settings in Bangladesh. A structured questionnaire was administered by trained personnel recording participants' demographic, socioeconomic, behavioural, medical, environmental and other factors. Anthropometric measurements and blood pressure were recorded for each participant. Biological specimens were collected and aliquoted for long-term storage and analysis. **Findings to date** Of the 73 883 study participants (mean [SD] baseline age: 39 [15] years), 43 470 (59%) were females, and 38 848 (52%) had no or only primary-level education. Focusing only on the 65 822 adult participants aged 20-79 years at baseline, 15 411 (23%) reported being diagnosed with hypertension; 10 578 (16%) with type 2 diabetes and 7624 (12%) with hypercholesterolaemia. Age and sex-

STRENGTHS AND LIMITATIONS OF THIS STUDY

- ⇒ Largest bioresource for non-communicable diseases (NCDs) and related traits in Bangladesh, configured to investigate multidimensional determinants of NCDs across urban, slum and rural settings.
- ⇒ Comprehensive recording of questionnaire-based data, including demographic, socioeconomic, behavioural, medical, environmental and other factors, as well as physical measurements.
- ⇒ Collection of a range of biological samples—including serum, plasma, whole blood and nail samples stored in long-term repositories.
- ⇒ Use of stored biological samples to enable the study of many genomic and molecular factors together with other information, laying the foundations to advance scientific discovery and public health action.
- ⇒ The study is not necessarily representative of the general Bangladesh population and could be limited by misclassification of disease outcomes and by loss of contact with some participants during follow-up.

standardised prevalences of these conditions were much higher in urban than slum and rural settings. Overall, the mean (SD) body mass index (BMI) was 25 (5) kg/m², with 10 442 (16%) participants aged 20–79, classified as obese (ie, BMI \geq 30 kg/m²). Mean BMI was also higher in urban than slum and rural areas. **Future plans** The collection of information during the baseline visit was completed in 2020. Regular longitudinal follow-up is ongoing for ascertainment and adjudication of a range of fatal and non-fatal health outcomes among participants. This cohort will provide a powerful resource to investigate multidimensional determinants of incident NCDs across diverse settings in Bangladesh, helping to advance scientific discovery and public health action in an archetypal low-middle-income country with pressing public health needs.

INTRODUCTION

Non-communicable diseases (NCDs)-including cardiovascular diseases, diabetes, cancer and chronic respiratory diseases-are the leading causes of premature death in low-income and middle-income countries (LMICs).¹² In particular, South Asia has recorded a higher number of life-years lost due to premature NCDs than any other global region, a situation which reflects both the region's large population and the relatively young age at which NCD deaths tend to occur in this region.¹³ Furthermore, while NCD mortality rates have decreased during recent decades in most countries, they remain at a high level in the South East Asia region.⁴⁵ Better understanding of the multidimensional determinants of NCDs in South Asia should, therefore, inform the development of appropriately tailored strategies for disease prevention and control.

There is, however, limited evidence available on NCD determinants for many South Asian populations.² ⁶ For example, Bangladesh, a country with 170 million people, is one of the least studied most densely-populated countries with regard to NCDs, despite steep and sustained increases in NCDs risk factors and incidence during recent decades.⁷⁸ In 2021, 74% of all adult deaths in Bangladesh were attributed to NCDs.⁷⁹¹⁰ The high burden of NCDs in Bangladesh is not just of local public health concern, as mortality due to NCDs has been reported to be more than two times higher among Bangladeshis living in western regions compared with host populations.¹¹ An important challenge is, therefore, to establish informative epidemiological resources in a rigorous manner to evaluate risk factors among Bangladeshis.

The present report provides a description of the methods used in the establishment of the BangladEsh Longitudinal Investigation of Emerging Vascular and nonvascular Events (BELIEVE) study. It also describes the baseline characteristics of the study population recruited so far and outlines the rationale for the study's further development.

Cohort description

Study design and participants

Between January 2016 and March 2020, the BELIEVE prospective cohort study recruited 73883 participants from 30817 households across three different settings in Bangladesh: urban (Mirpur-Dhaka), rural (Matlab-Chandpur) and urban-poor ('slum'; Bauniabadh-Dhaka) (figure 1). Households were initially identified through

complete surveys that had previously mapped households' information across the three study sites. Individuals living in the identified households were eligible for enrolment into the study if they (1) were aged 11 years or older; (2) had lived in their current household for at least 3 years; (3) were intending to reside in the study site for at least a further 5 years and (4) were willing to provide written informed consent and prospective follow-up information. For all participants, the recruitment procedure first involved trained study personnel visiting all identified households to provide written and verbal information about the study. On completion of the household visit, all eligible household members were invited to attend a local study clinic to complete an individual-level baseline assessment (figure 2).

Questionnaire administration and physical measurements at baseline assessment

The study questionnaire was adapted to the Bangladesh context based on validated questionnaires used in previous large-scale studies.¹²⁻¹⁴ The initial version of the structured questionnaire underwent preliminary testing with a small group of individuals from the target population to uses rela evaluate clarity, relevance and cultural appropriateness. Following this, a larger group of individuals was involved for further validation and refinements, and adjustments were made to enhance readability and ensure logical flow, before finalising the questionnaire for the main study. To đ ensure both linguistic accuracy and cultural relevance, e the questionnaire was translated into Bengali using a rigorous translation and back-translation process. During the household visit, trained personnel administered the ð structured questionnaire to collect household-level information and family structure characteristics (table 1) using a bespoke Android interface operated through handheld touchscreen tablet devices. During the clinic visit, trained personnel used a computer-based prepiloted epidemiological questionnaire to collect self-reported information on >300 items related to socioeconomic and demographic characteristics, consanguinity, behaviours (eg, tobacco and alcohol consumption, dietary intake, physical activity), female health, self-reported personal and family medical history and medication use (table 1; a copy of the study questionnaire is available in online supplemental appendix). Direct computer entry by participants, technolog rather than interviews, was employed to enhance privacy when answering sensitive questions, with the option to skip these questions.

Self-reported information on medical history and medication use was supplemented by medical records, drug prescriptions and by asking participants to bring their medications to the clinic visit. To assess local dietary patterns, benchmark food-frequency questionnaires previously developed and validated in Bangladesh were adapted for BELIEVE.¹⁵ These questionnaires estimated the standard portion size assigned to each food item and drinking water source. Participants' phone numbers (or, for participants younger than 18 years, a contact number



Figure 1 Location of the BELIEVE study sites. BELIEVE, BangladEsh Longitudinal Investigation of Emerging Vascular and nonvascular Events.

of a legal parent, guardian or caretaker) were routinely collected at the baseline visit for future contact and to collect follow-up information.

Standardised procedures and equipment were used to assess height, weight, waist and hip circumference, body composition, systolic and diastolic blood pressure, heart rate and upper body strength (online supplemental table 1). Briefly, blood pressure was measured using an automated device (Omron HEM 7130) on the right arm twice with a 3 min interval between each measurement, least 5 minutes before the first evaluation. Height was measured using the ShorrBoard ICA Measuring Board to within 1 cm and weight was measured using the Tanita HD-661 scale to within 0.1 kg. Waist circumference was assessed using a non-stretchable soft standard measuring tape over the abdomen at the widest diameter between the costal margin and the iliac crest, and hip circumference at the level of the greater trochanters (ie, the widest diameter around the buttocks). Anthropometric



Figure 2 Study recruitment and follow-up procedures.

Table 1	Questionnaire-based information,	, physical measurements and biological samples collected at baseline in the)
BELIEVE	study		

Characteristics	Description		
House and household			
House information	Location, type of accommodation, house construction materials, number of bedrooms, cooking source, sanitation, water source, indoor and outside environment		
Household information	Number of occupants, contact details, family size		
Individual			
Demographics	Age, sex, ethnicity, religion, marital status, consanguinity		
Behavioural factors	Tobacco consumption (cigarette and non-cigarette), passive tobacco exposure, cooking habits, food frequency assessment, alcohol consumption, physical activity, sleeping habits, mobile phone usage		
Socioeconomic factors	Education, occupation, income, remittances, loans, assets owned (including mobile phone, television, refrigerator, digital versatile disc player, air conditioning, bicycle, motorcycle, car, livestock, land, bank account)		
Psychosocial factors	Stress at work and at home, social support, Centre for Epidemiological Studies Depression symptoms score, Generalised Anxiety Disorder score, sleeping habits, life events		
Female health	Age at menarche, hormonal contraceptive usage, menstrual and pregnancy history		
Personal and family medical history	Myocardial infarction, angina, hypertension, other vascular diseases, type 2 diabetes mellitus, atrial fibrillation, cancers, hypercholesterolaemia, chronic liver disease, chronic kidney disease, chronic obstructive respiratory disease, mental disorders, neurological diseases, infectious diseases, childhood disorders, major surgery, chest and limb pain, current medication usage		
Physical measurements	Blood pressure, heart rate, respiratory rate, height, weight, body composition, upper body strength, waist and hip circumferences.		
Biological samples	Serum, plasma and whole blood samples, finger and toe-nail clippings		
BELIEVE, BangladEsh Longitudinal Investig	ation of Emerging Vascular and nonvascular Events.		

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measurements were performed in a standing position. Body composition was assessed by bioimpedance using the Tanita MC-780MA analyser and handgrip strength using a Jamar Plus Digital Hand Dynamometer.

To enhance consistency in the collection of data, we trained staff extensively and adopted standardised approaches, validated instruments and electronic data collection methods with built-in validity checks and queries. For example, the paper-free digital data collection platform involved extensive computerised checks to restrict missing values, duplications, inconsistencies and outliers. Information was transferred daily in a secure manner to the study's central database at Cambridge University, with a copy also kept at the local recruitment centres for additional checks and queries. Reports were generated by data managers and reviewed by study Principal Investigators (PIs) and coordinators on a regular basis.

Collection, storage and initial use of biological samples

Participants provided non-fasting blood samples for research with the time of the last meal recorded. Up to 23 mL of non-fasted whole blood samples were collected from each participant aged 18 years and above in two tubes (11 mL EDTA and 12 mL serum). For participants aged 11-17 years, 12mL of non-fasted whole blood samples were collected in two tubes (6mL EDTA and 6 mL serum). Samples were centrifuged within 45 min of venipuncture (at 6000 rpm for 15 min). Isolated serum, EDTA plasma and whole blood samples were stored at -20°C before being transferred to a -86°C freezer at the end of each working day (online supplemental figure 1). To enable the measurement of additional potential risk factors (eg, metal contaminants), finger-nail and toe-nail clippings were collected in a subset of participants, kept separately in plastic bottles and stored at room temperature. Biological samples have been stored in long-term dual repositories located in Bangladesh and the UK to enable further assays. A range of biochemical and genomic analyses on the stored samples is currently ongoing.

Follow-up procedure and outcome ascertainment

Study participants are being actively followed up indefinitely for cause-specific mortality and incidence of selected non-fatal health outcomes (figure 2). Follow-up involves contact with participants by the study team at regular intervals (eg, every 18–24 months), during which trained personnel use an electronic questionnaire to collect information on medical events reported since the last contact. Based on the responses to the questionnaire (or if phone contact is unsuccessful), a household visit is scheduled to collect additional information and medical documents. During the household visit, trained personnel collect further information for selected major categories of non-fatal events (including myocardial infarction, diabetes, stroke and common cancers) using a structured questionnaire and a bespoke Android electronic interface operated through handheld touchscreen tablet devices.

Open access

As the majority of individuals in the study settings who have received medical attention tend to have paper copies of medical documents in their homes, study personnel visiting households collect digital photographs of any relevant medical documents available (including discharge summaries, diagnostic test reports, medications and death certificates) using tablet devices. If medical documents are unavailable, written informed consent is obtained to retrieve these from the relevant hospitals. Nearly all deaths in the study areas will have involved some ŝ form of medical attention, with their underlying causes certified by a medical doctor. Verbal autopsy is conducted 8 in all deaths by trained personnel to help determine the most likely cause from symptoms or signs described by family members.¹⁶ To help adjudicate and classify health outcomes (eg, into definite, probable and possible categories), the information collected described above is periodically reviewed by medically trained personnel.

The quality and completeness of both mortality and morbidity follow-up data in each study site are regularly checked during the study period by the study coordinating centres. This involves monitoring the number of people who have died or are lost to follow-up, assessment of overall mortality patterns in the study cohort, levels of diagnostic criteria for individual diseases and the proportion of deaths with unknown causes.

Patient and public involvement

In keeping with this study's principles of cocreation and ð Ĩťa community engagement, we have implemented a participatory strategy, collaborating closely with local residents and key stakeholders (some of whom were study participants). Community consultations were conducted in communal spaces to communicate the study's objectives and actively seek feedback from community members. Invitations were extended to local leaders for these sessions, ß facilitating a deeper understanding of the community's needs and concerns and allowing us to tailor the study accordingly. In addition, we employed local community members as field team personnel, who were trained to collect data and assist study participants. This approach not only provided employment opportunities for local people but also enhanced the cultural appropriateness of the study approaches we used. As well as cocreating the **o** study design, study participants were provided information about the study's purpose, with the assurance that **g** their information would be exclusively used for research purposes.

Sample size considerations and statistical analysis

Sample size considerations have been guided by a combination of pragmatic constraints (eg, the availability of resources) and statistical power calculations. Assuming 5% of the population develops a disease condition during follow-up, a study of 70000 individuals should provide 80% power to detect an HR of about 1.1 per 1 SD higher value at 5% significance level. In this report, continuous variables were summarised as mean (SD) or median and IQR, and categorical variables were summarised as frequencies and percentages. A complete case analysis was used to handle missing data. Variables were summarised and compared across relevant population subgroups (eg, study site, sex and age group) using the t-test for continuous variables and χ^2 test for categorical variables. The prevalences of chronic conditions at baseline were summarised as overall crude prevalences, and by site, sex and 20-year age group. To make comparisons between sites analysis was conducted restricted to data from participants aged between 20 and 79 years to compare prevalences of conditions standardised according to the Bangladesh 2022 population structure.¹⁷ All statistical tests were two sided, and the significance level was set at p<0.05. Robust SEs were estimated to allow for the clustering of participants by household. Analyses were performed using STATA (V.16, StataCorp) and R (V.4.0.1, R Foundation for Statistical Computing). Future statistical analyses will be developed following relevant guidelines for cohort data with household clustering (eg, Strengthening the Reporting of Observational Studies) and will be presented elsewhere.

Findings to date

Demographic, behavioural and physical characteristics in the overall study population

Overall, 59846 (81%) participants were recruited from Mirpur-Dhaka (urban site), 5332 (7%) from Bauniabadh-Dhaka (slum site) and 8705 (12%) from Matlab-Chandpur (rural site). 43470 (59%) participants were female; the mean (SD) age at recruitment was 39 (15) years; 4122 participants (5.6%) were aged between 11 and 17 years inclusive (table 2). At the time of the baseline survey, 38848 (52%) participants had either no formal education or primary level education only; 17% were current smokers; 23% were current users of chewable tobacco products and only 1.3% of the participants reported consuming alcohol regularly.

Several characteristics varied by study site. For example, 55% of participants in the urban site had a secondary or higher level of education compared with only 16% and 18% in slum and rural sites, respectively (online supplemental table 2). Whereas the percentages of participants self-reporting as current smokers were similar across the sites (17%, 17% and 13% in urban, slum and rural sites, respectively), reported use of chewable tobacco and alcohol was higher in the slum site (62% and 12%, respectively, vs 18% and 0.4% in the urban site, and 34%and 0.4% in the rural site). Additional conventional risk factor values differed by study site, with a tendency towards more adverse risk factor profiles in urban participants (online supplemental table 2). For example, mean (SD) systolic blood pressure was 125 (21) mm Hg in participants from the urban site vs 112 (20) and 120 (20) mm Hg in the slum and rural sites, respectively. Mean (SD) body

ble 2 Baseline characteristics of the	e 73833 BELIEVE
	Mean (SD) or
ey characteristics	number (%)
mographic and behavioural characte	eristics
e (years)	39 (15)
dividuals aged <18 years	4122 (5.6)
male	43 470 (59)
udy site of residence	
Urban site	59846 (81)
Urban slum site	5332 (7.2)
Rural site	8705 (12)
arried	54386 (74)
ucation level	
None/preprimary	16594 (22)
Primary	22254 (30)
Secondary	21 367 (29)
Bachelors or higher degree	13 105 (18)
ırrent smoker	12450 (17)
irrent chewable tobacco user	16982 (23)
irrent alcohol consumer	927 (1.3)
ysical measurements	
stolic blood pressure (mm Hg)	123 (21)
astolic blood pressure (mm Hg)	77 (12)
ody mass index (BMI, kg/m²)	25 (5)
aist circumference (cm)	86 (12)
aist to hip ratio	0.92 (0.08)
dy fat percentage (%)	27 (9)
ssing data by variable: Married (n=3), edu ioking/chewable tobacco use (n=4), alcoh stolic blood pressure (n=23), diastolic bloo II (n=45), waist circumference (n=26), wai dy fat percentage was not measured in th sruitment and therefore was not available LIEVE, BangladEsh Longitudinal Investig- scular and nonvascular Events.	acation (n=563), nol consumption (n=42), od pressure (n=22), ist to hip ratio (n=30). he early stages of in 15049 participants. ation of Emerging
ss index (BMI) was 26 (5) kg/m ² (5) kg/m ² in participants from t al sites, respectively, with other m cluding, waist-to-hip ratio and be owing a similar trend. As the tren n accelerates in Bangladesh, such tor profiles portend implications f NCDs. ^{19 20} This contrast in risk fa- ings also suggests the need for c as to help better prevent and contra- tribute towards this goal through ce of risk factors and NCD incide nort, as well as the repurposing co- ele to support applied backs	² , 23 (5) kg/m ² and he urban, slum and heasures of adiposity ody fat percentage, nd towards urbanisa h differences in risk for striking increases actor profiles across context-specific solu rol NCDs. We plan to longitudinal surveil nce in the BELIEVF of the cohort frame

Certain baseline characteristics also differed between males and females (online supplemental table 3). Females were, on average, younger (38 vs 41 years), had spent fewer years in education (45% vs 50% had a secondary or higher level of education), were less likely to report any tobacco smoking (<1% vs 41%) and consuming alcohol (<1% vs 3%). Systolic blood pressure was higher among males (127 vs 121 mm Hg), whereas females had a higher BMI (26 vs 24 kg/m²).

Overweight, obesity and chronic health conditions in adults aged 20–79

Among adult participants aged between 20 and 79 years (online supplemental table 4-6) and using WHO-defined BMI categories, 39% of individuals were considered overweight $(BMI 25-29 \text{ kg/m}^2)$ and 16% obese (ie, $BMI \ge 30 \text{ kg/m}^2$) m² online supplemental table 4). After standardisation of the sex and age distribution of the Bangladeshi population for the same age range, the corresponding proportions were 37% and 14%, respectively (online supplemental figure 2). Proportions of overweight and obese participants were higher in the urban site, compared with the slum and rural sites, and also higher in females compared with males (online supplemental figure 2-3). There are likely to be multiple factors contributing to excess adiposity, including dietary patterns, sedentary lifestyles and urbanisation.¹²¹ Additionally, socioeconomic disparities and lack of awareness regarding the importance of diet and physical activity may play roles. As overweight and obesity are associated with multiple chronic health conditions, including cardiovascular diseases, diabetes and certain cancers,^{22–24} addressing this challenge is necessary to improve the health of the Bangladeshi population and alleviate healthcare system burdens.^{25 26}

The most prevalent self-reported chronic health conditions recorded at baseline among adults aged 20-79 were hypertension, type 2 diabetes and hypercholesterolaemia (online supplemental table 6), with standardised prevalences of 21%, 15% and 11%, respectively (figure 3). The prevalence of these conditions was higher among participants recruited in the urban site compared with those in slum and rural sites. For example, the standardised prevalence of hypertension was 23%, 16% and 14% in urban, slum and rural sites, respectively. Corresponding prevalences were 16%, 10% and 5.7% for type 2 diabetes, and 12%, 8% and 1.5% for hypercholesterolaemia. The overall prevalence of type 2 diabetes in this study was higher than those previously reported for Bangladesh¹⁹ and varied by age and sex, with the highest prevalences Bu among older participants and females (online supplemental figure 4–6). These findings may suggest potential $\vec{\mathbf{Q}}$ uses age-specific and sex-specific differences in susceptibility to type 2 diabetes or disparities in healthcare access and re management or both. We plan to contribute towards understanding such potential age-specific and sex-specific differences in determinants of chronic diseases, as it should inform the development of targeted preventive



Figure 3 Standardised prevalences of selected chronic conditions at recruitment in BELIEVE for participants aged 20–79. Prevalence was standardised using the sex and age distribution of the Bangladesh 2022 population aged 20–79. BELIEVE, BangladEsh Longitudinal Investigation of Emerging Vascular and nonvascular Events.

measures and tailored interventions to address the specific risk factors affecting different populations.

Strengths and limitations

Although Bangladesh is experiencing substantial increases in NCDs,⁷ there is limited evidence about the multidimensional risk factors and drivers of NCDs in the country, thereby preventing a deeper understanding of the web of causation for NCDs and limiting development of optimal prevention and control approaches. The BELIEVE study has been established as a long-term epidemiological bioresource to help address this gap, including an investigation of Bangladesh's contrasting urban and rural settings. Configured to investigate biological, behavioural, environmental and broader determinants of NCDs, the BELIEVE study is a household-based prospective cohort study that spans urban, slum and rural settings. To our knowledge, it represents the largest bioresource for NCDs and related traits in Bangladesh, involving active longitudinal follow-up of participants for new-onset health outcomes.

The BELIEVE study has demonstrated the feasibility of employing modern epidemiological methods at scale in a population-based study located across multiple different settings in Bangladesh. The study has used electronic data collection methods, using study forms implemented through bespoke software applications ('apps'), allowing comprehensive recording of questionnaire-based data, as well as physical measurements. The study has also collected a range of biological samples, including serum, plasma, whole blood and nail samples stored in long-term repositories. The assay of these samples (which already includes genotyping, sequencing, multiple molecular 'omics, and soluble clinical biomarkers) enables the study of many molecular factors, laying the foundations for an advanced understanding of disease pathways and potential therapeutic targets for the treatment and prevention of NCDs in Bangladesh and beyond.

The potential limitations of the BELIEVE study merit consideration. Within each of the three study settings, every household was invited to participate in the cohort. As the household participation rates have been high (>80%), they suggest that study participants should be broadly representative of the source population in the participating sites. However, the participants, households and sites included in the study were not necessarily representative of the general Bangladesh population. Nevertheless, it should be reasonable to infer that findings from this cohort can be generalised to other similar settings in Bangladesh because the study involves participants with a wide range of diverse characteristics (eg, in relation to age, sex, socioeconomic status, education level and occupation), recruited from a variety of different settings (urban, slum and rural areas) characteristic of contemporary Bangladesh.

A further potential limitation relates to the scope for misclassification of risk factors and health outcomes recorded at baseline and during study follow-up because of inaccuracies in participant self-reports. To help limit the

effects of such potential misclassification, the BELIEVE study is supplementing self-reported data with the use of objective measurements (eg, assay of low-density lipoprotein -cholesterol (LDL-C), glycated haemoglobin (HbA1c) and arsenic metabolites), inspection of health records kept by participants during household follow-up visits, use of previously validated 'verbal autopsy' methods, and exploration of emerging potential linkages of study participants with digital health records kept at community healthcare and hospital levels.

Finally, there is potential in this prospective study **u** for underestimation of the strength of the associations observed between risk factors and incident health outcomes due to fluctuations in risk factor levels within individuals over time (ie, 'regression dilution' bias). To help limit such bias, the BELIEVE study plans to conduct operiodic resurveys of risk factor levels in randomly right, including selected subsets of the study participants.

researchers. Data are available on application to the study's Steering and Data Access Committee.

CONCLUSION

By providing a powerful resource to investigate multidimensional determinants of NCDs across diverse settings in Bangladesh, the BELIEVE study should help advance scientific understanding and inform public health action in an archetypal LMIC with pressing public health needs.

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Patient consent for publication Consent obtained directly from patient(s).

Ethics approval The BELIEVE study has received approvals from the relevant institutional review boards of the Bangladesh Medical Research Council, the National Heart Foundation Hospital and Research Institute, icddr,b and Bangabandhu Sheikh Mujib Medical University (BMRC/NREC/2013-2016/390; BMRC/NREC/2016-2019/243; BSMMU/2019/1184; BSMMU/2019/1185; PR-18051; HBREC.2019.09). Written informed consent has been obtained from each participant (or by a parent or guardian for participants under the age of 18), including for future use of data and stored samples and invitation to further research studies. The data collected in this research are subject to the core data protection principles and requirements of the UK Data Protection Act 1998. The investigators and institutional review boards are committed to ensure that research is conducted according to the latest version of the Declaration of Helsinki, Universal Declaration on the Human Genome and Human Rights adopted by the United Nations Educational, Scientific and Cultural Organization (UNESCO) and other legislation.

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REFERENCES

- 1 GBD 2019 Diseases and Injuries Collaborators. Global burden of 369 diseases and injuries in 204 countries and territories, 1990-2019: a systematic analysis for the Global Burden of Disease Study 2019. *Lancet* 2019;396:1204–22.
- 2 Bennett JE, Stevens GA, Mathers CD, et al. NCD Countdown 2030: worldwide trends in non-communicable disease mortality and progress towards Sustainable Development Goal target 3.4. The Lancet 2018;392:1072–88.
- 3 Siegel KR, Patel SA, Ali MK. Non-communicable diseases in South Asia: contemporary perspectives. *Br Med Bull* 2014;111:31–44.
- 4 Martinez R, Lloyd-Sherlock P, Soliz P, *et al.* Trends in premature avertable mortality from non-communicable diseases for 195 countries and territories, 1990-2017: a population-based study. *Lancet Glob Health* 2020;8:e511–23.
- 5 Shu J, Jin W. Prioritizing non-communicable diseases in the postpandemic era based on a comprehensive analysis of the GBD 2019 from 1990 to 2019. *Sci Rep* 2023;13:13325.
- 6 The Academy of Medical Sciences. Science to tacklenoncommunicable diseases in South Asia and beyond in the SDG era, 2020. Available: https://acmedsci.ac.uk/file-download/8141929 [Accessed 1 Nov 2023].
- 7 GBD 2019 Bangladesh Burden of Disease Collaborators. The burden of diseases and risk factors in Bangladesh, 1990-2019: a systematic analysis for the Global Burden of Disease Study 2019. *Lancet Glob Health* 2019;11:e1931–42.

- 8 Riaz BK, Islam MZ, Islam ANMS, et al. Risk factors for noncommunicable diseases in Bangladesh: findings of the populationbased cross-sectional national survey 2018. BMJ Open 2020;10:e041334.
- 9 Ahmed A, Nahian MA, Rahman MM, et al. Adult mortality trends in Matlab, Bangladesh: an analysis of cause-specific risks. *BMJ Open* 2023;13:e065146.
- 10 Chowdhury SR, Islam MN, Sheekha TA, *et al.* Prevalence and determinants of non-communicable diseases risk factors among reproductive-aged women: Findings from a nationwide survey in Bangladesh. *PLoS One* 2023;18:e0273128.
- 11 Harding S, Rosato M, Teyhan A. Trends for coronary heart disease and stroke mortality among migrants in England and Wales, 1979-2003: slow declines notable for some groups. *Heart* 2008;94:463–70.
- 12 Teo K, Chow CK, Vaz M, *et al.* The Prospective Urban Rural Epidemiology (PURE) study: examining the impact of societal influences on chronic noncommunicable diseases in low-, middle-, and high-income countries. *Am Heart J* 2009;158:1–7.
- 13 Sudlow C, Gallacher J, Allen N, et al. UK biobank: an open access resource for identifying the causes of a wide range of complex diseases of middle and old age. *PLoS Med* 2015;12:e1001779.
- 14 Chowdhury R, Alam DS, Fakir II. The Bangladesh Risk of Acute 526 Vascular Events (BRAVE) Study: objectives and design. *Eur J Epidemiol* 2015;30:577–87.
- 15 Chen Y, Ahsan H, Parvez F, et al. Validity of a food-frequency questionnaire for a large prospective cohort study in Bangladesh. Br J Nutr 2004;92:851–9.
- 16 Murray CJ, Lopez AD, Black R, et al. Population Health Metrics Research Consortium gold standard verbal autopsy validation study: design, implementation, and development of analysis datasets. *Popul Health Metrics* 2011;9:27.
- 17 Population pyramids of the world from 1950 to 2100, bangladesh 2022 population: 539. 2022. Available: https://www. populationpyramid.net/bangladesh/2022 [Accessed 10 Nov 2023].
- 18 Elm E von, Altman DG, Egger M, et al. Strengthening the reporting of observational studies in epidemiology (STROBE) statement: guidelines for reporting observational studies. *BMJ* 2007;335:806–8.
- 19 STEPS WHO. Bangladesh ncd risk factor survey 2018. Available: https://cdn.who.int/media/docs/default-source/searo/bangladesh/ bangladesh-ncd-risk-factor-survey-2018.pdf?sfvrsn=266ad1da_1 [Accessed 1 Nov 2023].
- 20 Rawal LB, Biswas T, Khandker NN, et al. Non-communicable disease (NCD) risk factors and diabetes among adults living in slum areas of Dhaka, Bangladesh. PLoS One 2017;12:e0184967.
- 21 Bishwajit G. Nutrition transition in South Asia: the emergence of noncommunicable chronic diseases. *F1000Res* 2015;4:8.
- 22 Adams J. Addressing socioeconomic inequalities in obesity: Democratising access to resources for achieving and maintaining a healthy weight. *PLoS Med* 2020;17:e1003243.
- 23 Chowdhury MZI, Rahman M, Akter T, et al. Hypertension prevalence and its trend in Bangladesh: evidence from a systematic review and meta-analysis. *Clin Hypertens* 2020;26:10.
- 24 Dinsa GD, Goryakin Y, Fumagalli E, et al. Obesity and socioeconomic status in developing countries: a systematic review. Obes Rev 2012;13:1067–79.
- 25 Biswas T, Pervin S, Tanim MIA, *et al.* Bangladesh policy on prevention and control of non-communicable diseases: a policy analysis. *BMC Public Health* 2017;17:582.
- 26 Islam K, Huque R, Saif-Ur-Rahman KM, et al. Implementation status of non-communicable disease control program at primary health care level in Bangladesh: Findings from a qualitative research. Pub Health Pract (Oxf) 2022;3:100271.

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