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Effect of rural-to-urban within-country migration on cardiovascular risk factors in low and middle income countries: A systematic review

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Abstract

Context—Limited information is available of effects of rural-to-urban within-country migration on cardiovascular (CV) risk factors in low and middle income countries (LMIC).

Objective—We performed a systematic review of studies evaluating these effects and having rural and/or urban control groups.

Study Selection—Two teams of investigators searched observational studies in MEDLINE, Web of Science and Scopus until December 2010. Studies evaluating international migration were excluded.

Data Extraction—Three investigators extracted the information stratified by gender. We captured information on 17 known CV risk factors.

Results—Eighteen studies (n=58,536) were included. Studies were highly heterogeneous with respect to study design, migrant sampling frame, migrant urban exposure, and reported CV risk factors. In migrants, commonly reported CV risk factors –systolic and diastolic blood pressure, body mass index, obesity, total cholesterol, and LDL– were usually higher or more frequent than the rural group, and usually lower or less frequent than the urban group. This gradient was usually present in both genders. Anthropometric (waist-to-hip ratio, hip/waist circumference, triceps

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skinfolds) and metabolic (fasting glucose/insulin, insulin resistance) risk factors usually followed the same gradient, but conclusions are weak due to information paucity. Hypertension, HDL, fibrinogen and C-reactive protein did not follow any pattern.

Conclusions—In LMIC, most but not all CV risk factors have a gradient of higher or more frequent in migrants than in the rural groups but lower or less frequent than the urban groups. Such gradients may or may not be associated to differential CV events and long-term evaluations remain necessary.

Keywords

emigration and immigration; cardiovascular system; risk factors; income; meta-analysis as topic

INTRODUCTION

Non-communicable diseases (NCDs) kill people mostly in low- and middle-income countries (LMIC).[1] The Grand Challenges report highlights the need to study the impact of poverty and urbanization on NCDs.[2] Urbanization is occurring at a fast rate in LMIC, which may be associated with less physical activity, poorer nutritional habits, and rapidly growing prevalence of cardiovascular (CV) risk factors, CV disease (CVD) and other NCDs. [3] Rural-to-urban migration is one of the key larger drivers of urbanization in LMIC. Thus, the evaluation of the impact of rural to urban migration on CV risk factors is relevant due to its huge public health consequences.

The effect of international migration to developed countries on CV risk factors and CVD has been systematically studied.[4] As a result of the unique pattern and rate at which migration is occurring within LMIC, international migration results cannot be inferred to these settings. In LMIC, differences in design and patient characteristics make it difficult to reach a definitive conclusion on the effects on CV risk factors [5-9]. Also, it is not known whether the effect of rural to urban migration is uniform across CV risk factors and across LMIC. Finally, the effect on emerging CV risk factors such as fibrinogen, HOMA insulin resistance, or C-reactive protein could also be evaluated in the most recent studies.

We conducted a systematic review of observational studies evaluating the effect of rural-tourban within-country migration on CV risk factors in LMIC. We hypothesized that migrants would have a worse CV risk profile than rural individuals and better CV risk profile than urban individuals.

METHODS

Study selection

A comprehensive literature search using PubMed-Medline, The Web of Science, and Scopus until December 31, 2010 was conducted. The following keywords were used: migration, emigration, immigration, residential mobility, transient and migrants, rural population, urban population, cardiovascular risk factors and cardiovascular disease. The search strategy of PubMed is available in the Appendix (Web only). We searched observational studies published in any language, performed in LMIC as defined by the World Bank.[10] We excluded studies of international migration or those from high income countries. A formal protocol was developed for this project.

Rural to urban migration was defined as the individual's self-report or objective information of birth in a rural setting and, at the time of the study, self-report or objective ascertainment of urban residence. Three types of control groups were possible: a) a rural comparison group, where migrants originated from the same area, b) an urban comparison group, where migrants share the same urban environment and c) both a rural and an urban comparison group. This strategy excluded studies focusing only on differences between rural and urban settings. Sampling frames for migrants were classified as: i) random sample of migrants from urban population, ii) cohort of people born in a rural area who moved to an urban area and were traced, iii) rural individuals selected and their family members followed up in urban area (sib-pair); and iv) population level survey where migration was based on a retrospective question ("were you born here?").

A list of retrieved articles was reviewed independently by 2 groups of investigators based in USA and Peru in order to choose potentially relevant articles, and disagreements were discussed and resolved. When multiple articles for a single study had been published, we used the most relevant publication and supplemented it, if necessary, with data from the other publications.

Data extraction

Data extraction was performed independently by 3 investigators (AVH, VP, AD). Disagreement was resolved by consensus. Using a standardized data extraction form, we collected information on lead author, year of study or publication year, study design, sample size, sampling frames for migrants, length of urban exposure, age of migration, average age, percentage of male participants, and 17 CV risk factors for migrants and control groups.

Outcomes

CV risk factors we collected were systolic blood pressure (SBP), diastolic blood pressure (DBP), hypertension; total cholesterol (TC), low density lipoprotein (LDL) cholesterol, high density lipoprotein (HDL) cholesterol; body mass index (BMI), obesity; triceps skinfolds thickness, waist circumference, hip circumference, waist-to-hip ratio (WHR); fibrinogen, C-reactive protein (CRP); fasting glucose, fasting insulin, and homeostasis model assessment (HOMA) scores, a validated measure of insulin resistance. Extracted information was stratified by gender. In the case of repeated measures, extracted information corresponded to the longest follow-up.

Study quality assessment

The order of quality of studies was considered as follows: 1) prospective cohort study, 2) retrospective cohort study, 3) case-control study, and 4) cross-sectional study. Also, we systematically assessed other key points of study quality proposed by the MOOSE collaboration.[11] These key points were: 1. clear identification of study population, 2. clear definition of outcome and outcome assessment, 3. independent assessment of outcome parameters (i.e. ascertainment of outcomes done by researchers other than the ones involved in the study), 4. selective loss during follow-up, and 5. important confounders and/or prognostic factors identified. Each point was rated as Yes/No. If the description was unclear, we considered that this as 'no'.

Statistical analysis

Our systematic review followed the Preferred Reporting Items for Systematic Reviews and Meta-analyses (PRISMA) statement.[12] Differences between migrants and either rural or urban control groups are shown as absolute differences (i.e. migrant minus control) and its 95% Confidence Intervals (CI) for both continuous and categorical CV risk factors. Non-normally distributed risk factors were described in some studies as geometric means, and we extracted them as published.

A high degree of heterogeneity among studies was expected and therefore we did not combine all studies in a formal meta-analysis. Taking into account the sources of heterogeneity, three subgroup meta-analyses were pre-specified: 1. studies with similar characteristics including cross sectional design, random sampling frame, and >5 years of urban exposure; 2. cross-sectional studies; and 3. studies published after 1990. Subgroup meta-analyses were stratified by gender, as differences were expected between genders.

DerSimonian and Laird random effects models were used for meta-analyses.[13] For dichotomous CV risk factors, we used the Mantel-Haenzel (M-H) method to calculate pooled odds ratios (OR) and 95% CIs. For continuous CV risk factors, we used the Inverse Variance method to calculate Mean Differences (MD) and 95% CIs. Statistical heterogeneity was evaluated with the Cochran Chi-square (χ^2) and the I² statistics. Funnel plots were used to evaluate publication bias for the meta-analyses. All analyses were stratified by gender. We used Review Manager (RevMan 5.0, Oxford, UK; The Cochrane Collaboration, 2008).

RESULTS

Study Characteristics

A total of 394 citations were identified and screened, of which 80 were retrieved for detailed assessment. Of these, 62 did not fit our criteria (Figure 1). Thus, 18 studies were chosen in the qualitative synthesis,[5-9,14-26] including 58,536 individuals (Table 1). Studies included mostly adults, with very few individuals <15 years-old.

Fourteen studies were cross-sectional, one was a retrospective cohort, [26] and three were prospective cohorts. [7,17,23] Studies were heterogeneous with respect to several characteristics. These studies were reported between 1964 and 2010. Fifteen studies compared the migrant group to the rural control group, and nine studies reported the years of urban exposure for the migrant group (urban exposure 6 months, and mostly >5 years). Sampling frames for migrants were also heterogeneous: random sampling in eight studies, a traced cohort in six studies, a population survey in three studies, and a sib-pair strategy in one study.

All studies clearly identified the study population and defined the outcome and outcome assessment (Table 1w, Web only). None of studies independently assessed the outcomes, and four studies had a selective loss of patients during follow-up.[7,17,23,26] Six studies identified important confounders or prognostic factors[6,8,9,18,25,26] and adjusted the association between migration and the CV risk factors.

The most commonly reported CV risk factors were SBP (13 studies), DBP (13 studies), hypertension (11 studies), BMI (11 studies), obesity (four studies), TC (six studies), LDL (five studies) and HDL (four studies). Hypertension was defined as BP 140/90 in only six studies, all of them published after 1990. Other CV risk factors including WHR, waist and hip circumference, triceps skinfolds, fasting glucose and insulin, HOMA scores, fibrinogen and CRP, were reported by fewer studies.

Effect of rural-to-urban migration on commonly reported CV risk factors

In most of the studies, values or rates in migrants were higher than the rural group, and lower than the urban group. Nine out of 12 studies showed higher SBP levels in migrants vs. rural (range 0.9 to 11.9 mmHg, significant differences in seven), and five out of six studies shower lower SBP in migrants vs. urban (range -0.2 to -8.3 mmHg, significant differences in two) (Table 2). Ten out of 12 studies reported higher DBP in migrants vs. rural (range 1.5 to 13.9 mmHg, significant differences in nine), and three out of five studies showed lower

DBP in migrants vs. urban (range -1.2 to -4.9 mmHg, significant differences in two) (Table 3). Eight out of nine studies showed higher hypertension rates for migrants vs. rural (range 2.3% to 25.3%, significant differences in five), and two out of four studies reported non-significant lower hypertension rates for migrants vs. urban (range -0.7% to -16.6%) (Table 2w, Web only).

Nine out of 10 studies reported higher BMI values in migrants vs. rural (range 0.2 to 3.8 kg/m², significant differences in six), and all of the four studies showed significant lower BMI values in migrants vs. urban (range -0.3 to -1.3 kg/m²) (Table 4). Two studies showed higher obesity rates in migrants vs. rural (range 0.2% to 18.1%, one with significant differences), and two studies showed significant lower obesity rates in migrant vs. urban (range -3.9% to -13.1%) (Table 3w, web only).

All of the five studies showed higher TC levels in migrants vs. rural (range 4.0 to 35.1 mg/ dL, significant differences in four); all of the three studies showed non-significant lower TC levels in migrants vs. urban (range -1.9 to -4.1 mg/dL) (Table 4w, Web only). Three out of four studies reported significantly higher LDL levels in migrants vs. rural (range 5.3 to 30.3 mg/dL); two out of three studies reported non-significant lower LDL levels in migrants vs. urban (range -1.0 to -3.9 mg/dL) (Table 5w, Web only). Finally, two out of three studies showed lower HDL levels in migrants vs. rural (range -0.1 to -1.8 mg/dL, one significant difference); one out of two studies showed lower HDL levels in migrants vs. urban (absolute difference 1.2 mg/dL) (Table 6w, Web only).

Effect of rural-to-urban migration on uncommonly reported CV risk factors

In two studies[8,9] WHR values in migrants were significantly higher than in the rural and urban (range 0.01 to 0.06, and -0.01, respectively). Three studies[7,9,20] showed larger waist circumference levels in migrants vs. rural (range 0.6 to 12.0 cm, significant differences in two); one study[9] showed shorter waist circumference levels in migrants vs. urban (difference -3.3 cm, 95% CI -1.4 to -5.2). In one study,[9] a significant larger hip circumference was shown in migrants vs. rural (difference 7.0 cm, 95% CI 6.1-8.0) and significant shorter hip circumference in migrants vs. urban (difference -4.5 cm, 95% CI -2.9 to -6.1). Two studies[9,16] showed larger triceps skinfolds thickness in migrants vs. rural (range 0.1 to 10.9 mm, one significant difference); one study reported significantly shorter triceps skinfolds thickness in migrants vs. urban (difference -8.9 mm, 95% CI -5.5 to -12.3).

Two out of three studies[7,9,20] reported higher mean or geometric mean glucose values in migrants vs. rural; two studies[9,24] reported lower mean or geometric mean glucose values in migrants vs. urban. Three studies[8,9,20] showed higher mean or geometric mean fasting insulin levels in migrants vs. rural; two of the studies[8,9] also showed lower fasting insulin levels in migrants vs. urban. The mean or geometric means of HOMA scores were significantly higher in migrants vs. rural in two studies;[9,20] in one study[9] the geometric mean of the HOMA score was significantly lower in migrants vs. urban. Geometric means of fibrinogen and CRP in migrants were significantly higher vs. rural, and similar to the values in urban.[9]

The observed gradient for most of commonly and uncommonly reported CV risk factors among migrants, rural and urban groups was found in both males and females separately.

Meta-analyses in subgroups of studies

Due to limited availability of CV risk factors, we only included SBP, DBP, and BMI in these meta-analyses. Substantial heterogeneity of effects among studies and no evidence of publication bias were seen in all three sets of meta-analyses.

Five studies[6,92,20-22] fulfilled our first pre-specified criteria of being cross-sectional studies, with a random sampling frame for migrants, and with at least 5 years of urban exposure for migrants. No differences were found between migrants and rural individuals in terms of SBP, DBP or BMI. A pattern of significantly lower SBP (MD -3.5 mmHg, 95% CI -5.6 to -1.4) and lower DBP (MD -3.0 mmHg, 95% CI -4.2 to -1.72) in migrants vs. urban was seen overall and for both males and females.

Fourteen studies were cross-sectional (Table 1) fulfilling our second pre-specified criteria, and thirteen studies were published after 1990 (Table 1) fulfilling our third criteria. Gradients were similar to the overall results for SBP, DBP and BMI on these subgroups.

DISCUSSION

Main findings

Studies conducted in LMIC evaluating the effect of within-country migration on CV risk factors showed substantial heterogeneity with respect to design, sample size, time of urban exposure, migrant sampling frame, and reported CV risk factors between migrants and rural or urban individuals. In general, when observing the absolute differences between migrants and comparison groups, a gradient for most of the commonly reported CV risk factors was noted: higher values or rates in migrants in comparison to rural individuals, and lower values or rates in migrants in comparison to urban individuals. This gradient was also seen in most of the studies when evaluating males and females separately. Nevertheless, against our hypothesis, some CV risk factors such as hypertension rates, HDL, fibrinogen and CRP levels did not follow any gradient.

What the current literature reports

The effect of international migration to developed countries on CV risk factors has been systematically studied by McKay et al.[4] These migrants are exposed not only to increased consumption of saturated fats and sugars and sedentary behavior but also to stressful life conditions. International migrants have poorer health and more disadvantaged CV risk factor profile than non-migrants, and this profile may worsen with increasing duration of stay in the urban environment.

Rural-to-urban within-country migration is a very common phenomenon in LMIC countries, [5] largely due to economic reasons. Its effects on CV risk factors have been poorly studied, [26] mostly for blood pressure/hypertension and BMI/obesity. Even less information is available for lipid profiles[27] or emerging and newer CV risk factors (e.g. CRP, fasting insulin, HOMA scores, fibrinogen).[9]

In the context of LMIC, urban-rural comparisons are of limited relevance in examining the effects of urban migration as the urbanization process in these countries is due to growth of existing urban populations, expansion of urban boundaries, and rural-to-urban migration. [28] Also particular genetic, cultural and life-style backgrounds of migrants and urban individuals further limit the value of urban-rural comparisons. However, several urban-rural comparisons of CV risk factors have been published in the literature. Comparisons of urban and rural areas in sub-Saharan Africa[29] and India[30] showed higher rates of hypertension, obesity and adverse lipid profiles for urban individuals; however, no differences between urban and rural areas for these risk factors have lately been found in China.[31]

What our study adds to current literature

We hypothesized that there would be a gradient with worse CV profile for the urban individuals than for migrants and worse for migrants than for rural individuals. This was the case for most of the CV risk factors; in most cases reported differences were significant. Some CV risk factors such as hypertension rates, HDL, fibrinogen and CRP levels did not follow any gradient. It seems implausible that these CV risk factors are not modified by migration given significant modifications of major risk factors, but recent reports, at least for blood pressure levels, suggest that the patterns of change following migration are very complex and do not necessarily follow the expected gradient.[23,32] An alternative explanation may include the scarcity of studies. Our chosen studies did not evaluate the effect of modified CV risk factors on CV events, and the gradients may or may not be associated to differential CV events across groups in the future. Thus, long-term longitudinal evaluations are necessary.

The gradient urban-migrant-rural for most of CV risk factors seems relevant in the context of studies with substantial heterogeneity. Recognizing the sources of heterogeneity, we secondarily analyzed three subgroups of studies. Cross-sectional and after 1990 subgroups of studies showed similar gradients between urban, migrant and rural as seen in all studies. The subgroup of cross-sectional studies with random sampling of migrants and at least 5 years of urban exposure for migrants showed lower SBP and DBP for migrants in comparison to urban individuals and non-significant differences between migrants and rural individuals. Although subgroup results should be taken with caution as heterogeneity of effects remained significant in all three subgroups of studies, they may highlight potential changes of gradient with longer urban exposure for migrants.

There was a notorious paucity of information on other important anthropometric risk factors (e.g. WHR, hip circumference, triceps skinfolds thickness) or metabolic and inflammation risk factors (e.g. glucose, fasting insulin, insulin resistance scores, fibrinogen, CRP). Our conclusions for these infrequently reported risk factors are weak at this moment and deserve further reevaluation in the future.

Only 6 studies[6,8,9,18,25,26] provided adjusted values of CV risk factors or adjusted estimates of the differences between migrants and controls. Adjusters included age, BMI, gender, socioeconomic status, education, occupation, marital status, physical activity, initial health status, and altitude. Most of these studies were published in 2009 and 2010, with the exception of the Yi Migrant[6] and Yi People[18] studies. Our main and secondary analyses were based on unadjusted values of CV risk factors, and therefore some bias may be present in our association measures between migration groups. Combination of adjusted metrics was not possible given the different sets of confounders adjusted for, the heterogeneity of studies, and the limited number of studies.

Limitations

First, we included studies that were heterogeneous with respect to several characteristics and therefore a meta-analysis of all studies was not possible. Pre-specified subgroups of more homogeneous studies also showed significant heterogeneity and subgroup results should be taken with caution. Second, we evaluated unadjusted differences between groups, as only unadjusted values were published by authors for most of the studies. Few recent studies provided adjusted values for a few of the CV risk factors we used in our analyses. Third, publication bias is always a concern in a systematic review; however we decreased it in our study by having no language restrictions, by using a comprehensive study search strategy in 4 literature engines, and by involving 2 groups of investigators with at least 2 researchers in each group. Fourth, we expect some differences on the effect of within-country migration on

CV risk factors across different countries and continents. We could not explore this hypothesis given the few numbers of studies available. Finally, the scarcity of reporting of several metabolic and inflammatory risk factors did not allow reaching stronger conclusions of the effect of migration on them.

Conclusions

Studies investigating the effect of rural-to-urban within-country migration on CV risk factors in LMIC are highly heterogeneous. Most of CV risk factors in migrants follow a gradient: higher or more frequent than in the rural groups, and lower or less frequent than the urban groups. Furthermore, some CV risk factors, such as hypertension rates, HDL, fibrinogen and CPR levels did not follow a pattern. Such gradients may or may not be associated to differential CV events across groups and long-term longitudinal evaluations of such associations remain necessary.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Table 1

Characteristics of studies included in the systematic review.

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Traced DBP cohort DBP, Random BP, cholesterol, weight, bematocrit, mental health BP, Random hypertension, obesity (ponderal index)	Traced DBP cohort DBP, cholesterol, Random weight, Sample hematocrit, mental health BP, Random hypertension, obesity (ponderal index) Traced BP, BMI, weight, cohort skinfolds	Traced DBP cohort DBP control and the second, weight, hematocrit, mental health BP, cholesterol, weight, hematocrit, mental health BP, weight, ponderal index) Traced BP, BMI, weight, cohort BP, weight, pulse rate, skinfolds skinfolds cohort and potassium, potassium, and creatinine	Traced DBP cohort DBP sample BP, cholesterol, weight, mental health BP, weight, mental health BP, weight, pubertension, obesity (ponderal index) Traced BP, BMI, weight, skinfolds stime odium, potassium, ad creatinine sample BP, weight, pulse rate, serum sodium, potassium, ad creatinine sample BP, hypertension, potassium, ad creatinine sample BP, hypertension,
Cross- Sectional Cross- Sectional Cross- Sectional	Cross- Sectional Cross- Sectional Sectional Cross- Sectional	Cross- Sectional Cross- Sectional Cross- Sectional Prospective Cohort	Cross- Sectional Cross- Sectional Cross- Sectional Prospective Cohort Cross- Sectional
Not reported Reported Not reported	Not reported Reported n reported Reported	Not reported Reported Not Reported Reported	Not reported Reported Not Reported Reported 2y 10y, 14.9y
N/A A cluster-random sample of non- migrants from East Azebaijan, living in West Teheran	N/A A cluster-random sample of non- migrants from East Azerbaijan, living in West Teheran N/A	N/A A cluster-random sample of non- migrants from East Azerbaijan, living in West Teheran N/A N/A	N/A A cluster-random sample of non- migrants from East Azerbaijan, living in West Teheran N/A N/A N/A N/A N/A A cluster-random sample of Han sample of Han people residing in Xichang city or Country seats (Butuo, Meigu, Zhaojue)
A cluster-random sample of native Serers living in one of 65 villages in Niakhar county A cluster-random sample of rural areas of East Azerbaijan	A cluster-random sample of native Serers living in one of 65 villages in Niakhar county A cluster-random sample of rural areas of East Azerbaijan A random sample of individuals living in Siaya District, Nyanza Province, Kenya	A cluster-random sample of native Serens living in one of 65 villages in Niakhar county A cluster-random sample of rural areas of East Azerbaijan fast Azerbaijan a random sample of individuals living in Siaya District, Nyanza Province, Kenya Province, Kenya esc, were selected from the same Luo tribe villages	A cluster-random sample of native 5 serets living in one of 65 villages in Niakhar county A cluster-random sample of rural areas of East Azerbaijan fast Azerbaijan fast Azerbaijan arandom sample of individuals living in Siaya District, Nyanza Province, Kenya Province, Fourter Province, Southwester Province, Southwester
Random sample drawn from a census of native Serers, from a particular tribe, living in certain parts of Dakar Cluster-random sample from East Azerbaijan living in West Teheran, Iran	Random sample drawn from a census of native Serens, from a particular tribe, living in certain parts of Dakar Cluster-random sample from East Azerbaijan living in West Teheran, Iran Random sample of Luo Tribe members originated from Siaya District, Nyanza Province, Kenya, and now living in poor urban slums of Nairobi, Kenya drawn from a census	Random sample drawn from a census of native Seres, from a particular tribe, living in certain parts of Dakar Cluster-random sample from East Azerbaijan living in West Teheran, Iran West Teheran, Iran Random sample of Luo Tribe members originated from Siaya District, Nyanza Province, Kenya, and now living in poor urban slums of Nairobi, Kenya drawn from a census All individuals who nuralWestern Kenya (Luo Tribe) to Nairobi (poor slum communites) on a permanent basis was identified and followed-up in Nairobi, Kenya	Random sample drawn from a census of native Seres, from a particular tribe, living in certain parts of Dakar Cluster-random sample from East Azerbaijan living in West Teheran, Iran Random sample of Luo Tribe members originated from Siaya District, Nyanza Province, Kenya, and now live, Nyanza Province, Kenya, and now liveng in poor urban slums of Nairobi, Kenya drawn from a census All individuals who nurban slums of Nairobi, Kenya drawn from a census All individuals who nuralWestem Kenya (Luo Tribe) to Nairobi (poor slum communites) on a permanent basis was identified and followed-up in Nairobi, Kenya All Yi people who had migrated to the county seats (Butuo, Meigu and Zhaojue) or years prior to the survey
40	40 40	60 41 40	61 60 41 40
2270 ~20 years-un 40-49y: 815 (57%); 50-59y: 613 (43%) 0% <20 years-old	22.70 < 20 years-010 40-49y: 815 (57%); 50-59y: 613 (43%) 0% <20 years-old 34y: 571 (49%); 355y: 215 (18%) 0% <20 years-old	 22.70 Curved Service 40-499; 815 (57%); 50-59y; 613 (43%) 0% <20 years-old 35y; 215 (18%) 0% <20 years-old 0% <20 years-old 30y; 260 (41%); 30y; 288 (45%); 30y; 89 (4%) 1% <15 years-old 	22.70 < 20 years-out 40-49y: 815 (57%); 50-59y: 613 (43%) 0% <20 years-old 0% <20 years-old 0% <20 years-old 1% <15 years-old 1% <15 years-old 0% <15 years-old 0% <15 years-old
1428	1428 1171	1428 1171 206	1428 1171 206 14505
1976, Iran	1976, Iran 1980, Kenya	1976, Iran 1980, Kenya 1981-1985, Kenya	1976, Iran 1980, Kenya Kenya 1981-1985, China
Nadim ⁵ , 1978	Nadim ⁵ , 1978 Poulter ¹⁶ , 1984	Nadim ⁵ , 1978 Poulter ¹⁶ , 1984 Poulter ¹⁷ ,	Nadim ⁵ , 1978 Poulter ¹⁶ , 1984 1990 He Migrant ⁶ ,
	Poulter ¹⁶ , 1980, 1171 34y: 571 (49%); 555: 54y: 385 (33%); 34y: 571 (49%); 0riginated from Siaya originated from Siaya ori	Poulae ¹⁶ , 1984, 1990, 1984, 1	$ Hent^{4}, 1980, 1171 35.54; 571 (49\%), 101 35.54; 571 (49\%), 101 35.54; 571 (49\%), 101 135.54; 571 (49\%), 101 135.54; 571 (49\%), 101 135.54; 571 (49\%), 101 135.54; 571 (49\%), 101 135.54; 571 (49\%), 101 135.54; 571 (49\%), 101 135.54; 571 (49\%), 101 135.54; 571 (49\%), 101 135.54; 571 (49\%), 101 135.54; 571 (49\%), 101 135.54; 571 (49\%), 101 135.54; 571 (49\%), 101 $

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Reported CV risk factors	electrolyte serum and urine concentrations	BMI, physical activity	 BP, BMI, waist circumference, % body fat, body fat, body fat, budy fat, body fatorse, insulin, HOMA index, HOMA index, 	BP, BMI, weight, mental health	BP, BMI, waist circumference, plasma glucose, total cholesterol, trigiycerides, LDI, HDL, skinfolds, wais/hip ratio, abdomen/hip ratio, body fat, physical activity	BP, BMI, weight, cholesterol, urinary electrolytes (urea, sodium, potassium, calcium, magnesium)
Sampling frame		Traced cohort	Random Sample	Random Sample	Traced cohort	Random Sample
Study design		Cross- Sectional	Cross- Sectional	Cross- Sectional	Prospective cohort	Cross- Sectional
Urban exposure		Average: 15y	89% for >20y	>5 y	Not Reported	Average: 20y
Urban population	power plant, a paper mill, a hospital, a middle school, and the government agency in Puge county seat.	Volunteers from three settlements in Port Moresby selected to match the age distribution of their village counterparts	A/A	N/A	A/A	N/A
Rural population	remote mountain districts drawn from a census	N/A	Convenience sample of women residing in small rural villages at high altitude in the Cuzco region	Kuna Amerinds residing in their indigenous home in the island of Ailigandi (Caribbean island off the coast of Panama)	People residing in or near their native villages and who were participants of a longitudinal growth and development study (1969 and 1977) and born in rural East Guatemala	Kuna Amerinds residing in their indigenous home in the island of Ailigandi (Caribbean island off the coast of Panama)
Migrant population	mill, hospital, middle school, and the government agency, in Puge county seat	Volunteers from two villages in Tari Basin, comprising of almost the entire adult population	Women of indigenous Andean ascent (Quechua) who had been residing in northern suburbs of Lima, Peru	Random sample drawn from a census of people of Kuna origin (O positive blood) and long term residents (5 years) of Abayala, Panama city	People living in Guatemala City and who were participants of a longitudinal growth and development study (1969 and 1977) and born in rural East Guatemala	A random sample of Kuna families residing in Vera Cruz, Panama City drawn from a census
% Male		52	0	43	45	39
Mean age or age groups / % of young individuals		20-30y: 18 (32%); 30-40y: 25 (45%); >40y: 13 (23%) 0% <20 years-old	35.5 % of young individuals not available	38.3 0% < 18 years-old	32.3 0% <18 years-old	37.4 0% <18 years-old
Total sample		56	210	458	1311	311
Years conducted, Country		1994-1995, Papua New Guinea	N/A, Peru	1999, Panama	1969-2004, Guatemala	1999, Panama
Study, Publication year		Y amauchi ¹⁹ , 2001	Lindgarde ²⁰ , 2004	Hollenberg ²¹ , 2005	Ramirez- Zea ⁷ , 2005	McCullough ²² , 2006

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Reported CV risk factors	BP, BMI, weight, waist, cholesterol, triglycerides, LDL, HDL, HbA1c	BP, BMI, waist/hip ratio, skinfolds, cholesterol, HDL, LDL, triglycerides, glucose	Overweight, Obesity	BMI, waist/hip ratio, % body fat, BP, cholesterol, triglycerides, blood glucose, fasting insulin, HOMA score, fat intake, metabolic equivalent tasks (METs)	Hypertension, mental health	BP, hypertension, BMI, skinfolds, waist-to-hip ratio, fasting glucose, diabetes, hyperotien protie, hypercholesterol emia C-reactive protein, fibrinogen
Sampling frame	Traced	Population Survey	Population Survey	Sib-Cohort	Population Survey	Random Sample
Study design	Prospective Cohort	Cross- Sectional	Cross- Sectional	Cross- Sectional	Retrospective Cohort	Cross- Sectional
Urban exposure	6 то	Not Reported	Not reported	86% for >10y	Not Reported	Average: 32y
Urban population	N/A	Convenience sample of people living in Wroclaw, Poland, those who were born in Wroclaw or migrated before 16 years of age	Women who were living in an urban area when<12-old, and did not move to urban city	A 25% random sample of nonmigrants invited to participate. Nonmigrants were also asked to invite a sib who resided in the same city but did not work in the factory	N/A	Stratified random sample of individuals born and permanently living in urban shanty town of Lima, Peru
Rural population	For each migrant, an age and sex matched non-migrant was identified in Morogoro, Tanzania	N/A	N/A	Each migrant worker and spouse invited one nonmigrant full sibling of the same sex and closest to them in age still residing in their rural place of origin	Identified from the 1997 and 2000 waves of the Indonesia Family Life Survey, representing 83% of the population	Stratified random sample of local residents of Sai Jose de Secce in Ayacucho was drawn from an updated census
Migrant population	Individuals living in the rural district of Morogoro, Tanzania, who migrated to Dar es Salaam for at least 6 months	Convenience sample of people living in Wroclaw, Poland who migrated after the age of 16y and recruited during a health screening program	Women who had moved from countryside or town to urban center, or who lived in country-side town when <12y-old	Factory workers for 4 cities and their coresident spouses were recurited. They were rural-urban migrants using employer records as a sampling frame	Identified from the 1997 and 2000 waves of the Indonesia Family Life Survey, representing 83% of the population	Stratified random sample of individuals born in Ayacucho but now residing in urban shanty town of Lima, Peru, drawn from updated census
% Male	48	43	0	58	46	47
Mean age or age groups / % of young individuals	29.2 0% <15 years-old	43 0% <16 years-old	30.7 12% <20 years-old and 0% <10 years-old	41.1 0% <17 years-old	29.7 0% <18 years-old	48 0% <30 years-old
Total sample	323	863	2807	6510	20122	686
Years conducted, Country	N/A, Tanzania	2006, Poland	2004, Bangladesh	2005-2007, India	1997-2000, Indonesia	2007, Реги
Study, Publication year	Unwin ²³ , 2006	Szklarska ²⁴ , 2008	Khan ²⁵ , 2009	Ebrahim ⁸ , 2010	Lu ²⁶ , 2010	Miranda ⁹ , 2010

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Table 2

Systolic Blood Pressure (SBP) in mmHg for the migrant, rural and urban groups

Study		SBP Migrants Mean (SD) [n]			SBP Rural Mean (SD) [n]			SBP Urban Mean (SD) [n]		Absolute difference M vs. R (95% CI)	Absolute difference M vs. U (95% CI)
	Male	Female	АП	Male	Female	ЯΙ	Male	Female	ЧЛ		
Nadim ⁵	134.9 (23.1)[278]	141.8 (24.9)[414]	138.1 (24.6)[692]	119.7 (20.2)[176]	127.9 (26.2)[203]	126.2 (23.7)[379]	137.3 (21.1)[117]	136.5 (23.1)[240]	138.4 (24.1)[357]	11.90 (8.89,14.91)	-0.30 (-3.40,2.80)
Poulter ¹⁶	128.2 (15.2)[220]	119.5 (13.7)[90]	124.7 (14.7)[310]	122.1 (18.3)[264]	120.0 (18.3)[597]	121.4 (18.7)[861]	I	I	I	3.30 (1.24,5.36)	ı
Poulter ¹⁷	125.6 (10.9)[43]	113.7 (9.9)[20]	121.0 (13.3)[63]	117.0 (11.4)[82]	106.2 (9.2)[61]	112.2 (12.1)[143]	ı	ı	I	8.80 (4.96,12.64)	ı
He Migrant ⁶	113 (14)[1656]	106 (12.8)[919]	111.0 (13.6)[2575]	110.8 (12.3)[5023]	111.2 (12.6)[3218]	110.8 (12.5)[8241]	114.5 (14.6)[2173]	108.1 (15.3)[1516]	112.2 (15.2)[3689]	0.20 (-0.39,0.79)	-1.20 (-1.92,-0.48)
He People	107.3 (11.6)[316]	101.6 (13.0)[201]	105.7 (12.6)[517]	101.2 (9.7)[2522]	101.1 (9.4)[2436]	101.1 (9.5)[4958]	108.3 (12.1)[638]	102.1 (11.2)[505]	105.0 (12.4)[1143]	4.60 (3.48,5.72)	0.70 (-0.60,2.00)
Lindgarde ²⁰	ï	96 (11)[105]	96 (11) [105]	ı	97 (10) [105]	97 (10)[105]	ı	ı	I	-1.00 (-3.84,1.84)	ı
Hollenberg ²¹	115[146]	100.5[179]	107.0 (17.7)[325]	104.4[51]	95.2[82]	98.7 (13.8)[133]	ı	ı	ı	8.30 (5.27,11.33)	ı
Ramirez- Zea ⁷	118.4 (11.6)[143]	113.0 (15.3)[202]	115.4 (13.7)[345]	116 (11)[424]	107 (12)[542]	111.3 (12.1)[966]	ı	ı	ı	4.10 (2.47,5.73)	ı
McCullough ²²	,		102 (14.7)[178]	ı	·	98.2 (13.8)[133]	ı	ı	ı	3.80 ($0.61, 6.99$)	ı
Unwin ²³	ı	·	116.2 (21.3)[153]	ı	ï	118.9 (18.9)[170]	I	ı	I	-2.70 (-7.11,1.71)	ı
Szklarska ²⁴	128.3 (15.6)[99]	115.2 (16.2)[178]	120.9 (17.8)[277]	I	ı	ı	128.6 (16.1)[268]	118.3 (16.8)[318]	123.1 (17.8)[586]	,	-2.20 (-4.74,0.34)
Ebrahim ^{8 *}	125.1 (17.1)[1127]	118.9 (16.0)[985]	122.7 (17.0)[2112]	122.9 (15.6)[1459]	118.9 (16.9)[652]	121.8 (16.2)[2111]	125.7 (17.7)[1201]	119.4 (16.8)[1086]	122.9 (17.8)[2287]	$0.90 \\ (-0.10, 1.90)$	-0.20 ($-1.23,0.83$)
Miranda ⁹	123.5 (14.9) [280]	116.6 (16.9)[309]	119.9 (16.4)[589]	125.2 (17.4)[95]	117.1 (19.1)[106]	120.9 (18.7)[201]	132.2 (23.2)[92]	124.7 (22.2)[107]	128.2 (22.9)[199]	-1.00 (-3.90,1.90)	-8.30 (-11.75,-4.85)
* adjusted for occuj	pation, age, age	group, and fact	ory including a 1	random effect of	f sibling pair.						

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Table 3

Diastolic Blood Pressure (DBP) in mmHg for the migrant, rural and urban groups

-4.90(-6.67, -3.13) 0.60 (-0.42,1.62) 1.30(-0.68,3.28) -1.20(-2.70,0.30) (-2.26, -1.14)Absolute difference M vs. U (95% CI) -1.7013.90 (12.08,15.72) (-4.38, -1.42)12.20(9.07,15.33) 2.60 (-1.15,6.35) 2.60 (2.08,3.12) (-5.32, -0.48)4.60 (3.71,5.49) 1.50(0.26,2.74) 5.60 (2.88,8.32) 7.00 (4.93,9.07) Absolute difference M vs. R (95% CI) (7.80, 11.60)(1.07, 4.33)-2.90 -2.90 9.70 2.70 , 84.7 (15.3)[357] 70.4 (10.5)[3689] 76.2 (11.5)[199] 80.0 (12.1)[586] (9.0)[1143]66.6 ЧI 1 DBP Urban Mean (SD) [n] 67.7 (10.3)[1516] 77.6 (10.6)[318] 73.7 (10.6)[107] 63.4 (7.6)[505] (16.0)[240]Female 84.7 72.6 (10.4)[2173] 84.8 (11.2)[268] 85.5 (15.6)[117] 69.2 (9.1)[638] 79.1 (11.8)[92] Male 72.1 (13.7)[379] 62.6 (7.1)[4958] 73.2 (10.0)[170] (12.1)[8241] 84.2 (9.4)[129] (11.7)[143] 74.2 (9.2)[201] (11.3)[861] (9) [105] (8.1)[133] (6.9)[133] (9.3)[966]70.9 53.5 ЧI 67.4 66.1 59 59.4 58.4 DBP Rural Mean (SD) [n] 65.8 (11.8)[3218] 62.5 (7.0)[2436] 73.6 (15.2)[203] 72.5 (9.2)[106] (11.2)[597] (9) [105] (9.6)[61]69 (9)[542] Female 57.9[82] 54.9 66.8 59 71.5 (14.2)[176] (11.4)[264]66.3 (12)[5023] (7.2)[2522] 53.1 (12.2)[82] 76.1 (8.9)[95] 72 (9)[424] 61.7[51] Male 63.0 68.1 68.7 (11.6)[2575] 67.2 (10.1)[517] 69.1 (13.7)[325] 72.4 (10.4)[345] 70.3 (12.0)[153] 86.0 (16.0)[692] (13.0)[310]64 (16) [178] 71.3 (9.3)[589] 86.8 (12.2)[50] 66 (6) [105] [10.0)[63]78.8 (9.6)[277] 65.7 ШV 70.1 DBP Migrants Mean (SD) [n] 65.2 (10.1)[919] 71.5 (11.0)[202] 63.2 (8.8)[201] (16.3)[414]75.5 (10)[178] 69 (8.7)[309] (11.4)[90](9.3)[20]66 (6) [105] 64.9[179] Female 61.7 67.2 87.1 70.9 (11.6)[1656] 73.7 (10.5)[143] 69.6 (10.1)[316] 83.5 (15.3)[278] 66.3 (14.0)[43] (14.5)[220] 74 (9.2)[280] 82.2 (9.5)[99] 74.2[146] Male 70.8 1 He People¹⁸ McCullough²⁷ Hollenberg²¹ Cruz-Coke¹⁴ Szklarska²⁴ He Migrant⁶ Lindgarde²⁰ Poulter¹⁶ Poulter¹⁷ Miranda⁹ Ramirez-Unwin²³ Nadim⁵ Study Zea⁷

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Table 4

Body mass index (BMI) in Kg/m^2 for the migrant, rural, and urban groups

Study		3MI Migrant: Mean (SD) [n	s []		BMI Rural Mean (SD) [n]			BMI Urban Mean (SD) [n]		Absolute difference M vs. R (95% CI)	Absolute difference M vs. U (95% CI)
	Male	Female	All	Male	Female	ЧI	Male	Female	ЧI		
Poulter ¹⁶	20.5 (2.6)[220]	21.8 (3.0)[90]	20.8 (2.9)[310]	20.0 (2.9)[264]	20.1 (3.0)[597]	20.2 (3.0)[861]			,	0.60 (0.22,0.98)	Ţ
He People ¹⁸	20.2 (2.3)[316]	20.9 (2.6)[201]	20.3 (2.5)[517]	18.9 (2.3)[2522]	18.9 (2.5)[2436]	18.9 (2.4)[4958]	20.5 (2.3)[638]	20.6 (2.3)[505]	20.6 (2.4)[1143]	1.40 (1.17,1.63)	-0.30 ($-0.56, -0.04$)
Yamauchi ¹⁹	26.4 (3.3)[14]	25.5 (3.1)[15]	25.9 (4.0)[29]	24.8 (1.6)[11]	23.2 (3)[9]	24.4 (2.2)[20]	ı	ı	ı	1.50 (-0.25,3.25)	ı
Lindgarde ²⁰		25.1 (4.7)[105]	25.1 (4.7)[105]	ı	24.6 (3.9)[105]	24.6 (3.9)[105]	ı	ı	ı	$0.50 \\ (-0.67, 1.67)$	ı
Hollenberg ²¹	22.6[146]	23.6[179]	23.1[325]	22.4[51]	22.5[82]	22.5[133]	ı	ı	ı	0.6^{**}	ï
Ramirez- Zea ⁷	26.0 (4.0)[143]	27.0 (3.8)[202]	26.5 (3.8)[345]	24.3 (3.4)[424]	26.8 (4.9)[542]	25.4 (4.6)[966]	ı	ı	ı	1.10 (0.61,1.59)	ı
McCullough ²²			23.4 (4)[178]	ı	ı	22.6 (3.5)[133]	ı	ı	ı	0.80 (-0.04,1.64)	ı
Unwin ²³		ï	24.0 (1.6)[153]	ı	ı	23.6 (1.0)[170]	ı	ı	ı	0.40 (0.11,0.69)	ı
Szklarska ²⁴	26.9 (3.1)[99]	24.4 (4.4)[178]	25.1 (4.1)[277]	ı	ı	ı	27.2 (4.1)[268]	24.5 (4.8)[318]	25.9 (4.5)[586]	ı	-0.80 (-1.40,-0.20)
Ebrahim ^{8 *}	24 (3.4)[1127]	25.2 (3.2)[985]	24.6 (3.4)[2112]	21.9 (3.9)[1459]	22.5 (3.9)[652]	22.0 (4.0)[2111]	24.3 (3.5)[1201]	25.9 (3.4)[1086]	25.0 (3.5)[2287]	2.60 (2.38,2.82)	-0.40 ($-0.60, -0.20$)
Miranda ⁹	25.9 (3.5)[280]	28 (4.7)[309]	27 (4.3)[589]	22.9 (2.1)[95]	23.5 (3.2)[106]	23.2 (2.7)[201]	26.8 (4)[92]	29.5 (6.1)[107]	28.3 (5.4)[199]	3.80 (3.29,4.31)	$^{-1.30}_{(-2.13,-0.47)}$
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adjusted for occupation, age, age group, and factory including a random effect of sibling pair.

** No standard errors available for groups.