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The effect of physical activity on mortality and cardiovascular disease in 130000 people from 17 high-income, middle-income, and low-income countries

Citation for published version:

Citation for published version: Lear, SA, Hu, W, Rangarajan, S, Gasevic, D, Leong, D, Iqbal, R, Casanova, A, Swaminathan, S, Anjana, RM, Kumar, R, Rosengren, A, Wei, L, Yang, W, Chuangshi, W, Huaxing, L, Nair, S, Diaz, R, Swidon, H, Gupta, R, Mohammadifard, N, Lopez-Jaramillo, P, Oguz, A, Zatonska, K, Seron, P, Avezum, A, Poirier, P, Teo, K & Yusuf, S 2017, 'The effect of physical activity on mortality and cardiovascular disease in 130000 people from 17 high-income, middle-income, and low-income countries: the PURE study', *The Lancet*, vol. 390, no. 10113, pp. 2643-2654. https://doi.org/10.1016/S0140-6736(17)31634-3

Digital Object Identifier (DOI):

10.1016/S0140-6736(17)31634-3

Link:

Link to publication record in Edinburgh Research Explorer

Document Version: Peer reviewed version

Published In: The Lancet

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2	The effect of physical activity on mortality and cardiovascular disease in 130,000 people from 17
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4	
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- 56

- 57 ABSTRACT
- 58

59 **Background:** It is not known whether the protective effects of physical activity (PA) on

60 cardiovascular disease (CVD) reported in high income countries (mainly recreational PA) is also

observed in lower income countries (mainly non-recreational PA). We examined whether

62 different levels and types of PA are associated with lower mortality and cardiovascular disease

63 (CVD) in countries at different economic levels.

64 Methods: In this prospective cohort study total PA was assessed using the International Physical

Activity Questionnaire in 130 843 participants from 17 countries. Mortality and CVD were

recorded during 6.9 years of follow-up. The effects of PA were adjusted for socio-demographic

67 factors and other risk factors taking into account household, community and country clustering.

Findings: Compared to low PA levels (<600 METS per week or <150 minutes/week of

69 moderate intensity PA), moderate (600 to 3000 or 150 to 750 minutes/week) and high PA levels

70 (>3000 or >750 minutes/week) were associated with a graded reductions in mortality (hazard

ratios =0.80 [0.74, 0.87] and 0.65 [0.60, 0.71], p<0.0001 for trend), and major CVD

72 (myocardial infarction, stroke and heart failure; (0.86 [0.78, 0.93], p<0.001 for trend).

73 Increasing PA was associated with lower risk of CVD and mortality in high, middle and low

⁷⁴ income countries. The adjusted population attributable fraction for not meeting the PA guidelines

75 was 8.0% and 4.6%, and not meeting high PA levels was 13.0% and 9.5% for mortality and

76 major CVD, respectively. Both recreational and non-recreational PA were associated with

77 benefits.

78 Interpretation: Increasing recreational and non-recreational PA are associated with a lower risk

of mortality and CVD events in low, middle and high income countries. Enhancing PA is a

simple, widely applicable, low cost global strategy which could avoid about 10% of deaths and

- 81 CVD in middle age.
- **Funding:** Funding sources listed at end of paper.
- **KEY WORDS:** physical activity, cardiovascular disease, global health, all-cause mortality

86	Cardiovascular disease (CVD) is the leading cause of death worldwide ¹ and a major economic
87	global burden. ² Despite reductions in CVD mortality in high income countries (HIC), global
88	CVD mortality increased by 41% between 1990 and 2013; largely driven by rises in low and
89	lower-middle-income countries. ³ Indeed, 70% of global CVD deaths come from low and middle
90	income countries where it is the largest cause of death. ^{4,5} It has been estimated that 23% of the
91	world's population was insufficiently active ⁶ and the WHO has recommended a decrease in
92	insufficient PA of 10% by 2020.7
93	
94	Numerous studies from HIC have reported significant inverse associations of PA with mortality
95	and CVD morbidity.8 but such data from low and middle income countries are sparse and limited
96	to a few small studies.9-11 In addition, studies of PA have focused primarily on recreational PA
97	(which is more common in HIC but less common in poorer countries), ^{8,11} with less evidence on
98	the benefits of other forms of PA such as during transportation, ¹² house work and occupational
99	PA. ¹³
100	
101	In the Prospective Urban Rural Epidemiologic (PURE) study being conducted in 17 high, middle
102	and low income countries, we examined whether PA is associated with lower risk of mortality
103	and CVD in countries at varying economic levels and whether these associations differ by type
104	of PA.
105	
106	METHODS
107	The PURE study includes participants from three HICs (Canada, Sweden, United Arab
108	Emirates); seven upper middle income countries ([UMIC] Argentina, Brazil, Chile, Poland,

109 Turkey, Malaysia, South Africa); three lower middle income countries ([LMIC] China,

110 Colombia, Iran); and four low income countries ([LIC] Bangladesh, India, Pakistan,

111 Zimbabwe).¹⁴ Country economic level was based on World Bank classifications in 2006. The

112 choice and number of countries selected in PURE reflects a balance between involving a large

113 number of communities in countries at different economic levels with substantial heterogeneity

in social and economic circumstances versus the feasibility of centres to successfully achievelong-term follow-up.

116

Within each country, urban and rural areas in and around selected cities and towns were 117 identified to reflect the geographical diversity of the countries. Communities were defined based 118 on the geographical clustering of common characteristics (sharing culture [as people of a similar 119 culture reside in close geographic proximity], socioeconomic status, and use of amenities, goods, 120 and services) such as through a set of contiguous postal code areas or group of streets or a small 121 122 village. The number of communities selected in each country varied. In some countries (eg, India, China, Canada, and Colombia), communities from several states/provinces were included 123 to capture diversity, socioeconomic status, culture, and environments. In other countries (eg, 124 125 Iran, Poland, Sweden, and Zimbabwe), fewer communities were selected. This strategy facilitated recruitment of individuals from different economic, cultural and geographical settings 126 127 (rural and urban) around the globe.

128

Within each defined community, households were approached and individuals between 35 and 70 years of age who intended to live at their current address for at least another four years were invited to participate in the study. The method of approaching households differed between

132 countries based on feasibility but was consistent across sites within each country. For example, in rural areas of India and China, a community announcement was made to the village through 133 contact of a community leader, followed by in-person door-to-door visits of all households. In 134 contrast in Canada, initial contact was by mail followed by telephone invitations to attend a 135 central clinic. For each household at least three attempts at contact were made. All eligible 136 137 residents of the household were invited and those who provided informed consent were recruited. Household participation rate was 86%. Recruitment started in a vanguard phase in Karnataka, 138 India in January 2003; however most communities recruited between January 2005 and 139 December 2010. See Supplementary Appendix provides more detail on sampling. The study was 140 approved by local institutional research ethics boards. 141

142

Socio-demographic factors, medical history, lifestyle behaviours and risk factors were recorded 143 using standardized measures and procedures.¹¹ One-week total PA was assessed using the long-144 form International Physical Activity Questionnaire (IPAQ)¹⁵ and calculated as a total of 145 occupation, transportation, housework and recreational activity reported in metabolic equivalents 146 (MET) * minutes/ week. Physical activity is also reported in minutes/week of moderate intensity 147 148 PA using the equation where minutes reported in each PA domain on the IPAQ by the participant are weighted relative to moderate intensity PA: 149 Minutes/week = 0.825*walking minutes + 1*moderate minutes + 1.375*garden minutes + 150 151 1.5*cycling minutes + 2*vigorous minutes Total PA was categorized as low (<600 MET*minute/week), moderate (600-3000 152

153 MET*minute/week) and high (\geq 3000 MET*minute/week) PA levels, which corresponds to <150

154 minutes/week, 150-750 minutes/week and \geq 750 minutes per week of moderate intensity PA.

155 Physical activity was also dichotomized as meeting or not meeting current PA guidelines (meeting guideline is PA >600MET*minute/week as per the IPAQ¹⁵ or PA >150 minutes/week 156 of moderate intensity PA as per the WHO¹⁶) with periods of less than 10 minutes of PA not 157 included as per the IPAQ guidelines.¹⁵ ¹⁶We also further categorized the high PA category into a 158 lower-high PA level and an upper-high PA level by the median value of the high PA category of 159 160 6453 MET*minutes/week to investigate whether the effect of very high PA levels was graded. PA was categorized into recreational PA and non-recreational PA (occupational, transportation 161 and housework). 162

163

The clinical outcomes of interest during follow-up were: mortality plus major CVD (CVD 164 mortality plus incident MI, stroke, and heart failure), either as a composite or separately 165 (Supplementary Appendix). In most low and middle income countries there was no central 166 system of death or event registration. We therefore obtained information on prior medical illness 167 and medically certified cause of death where available, and recorded best available information 168 from close family or friends in order to arrive at a probable diagnosis or cause of death. Death 169 certificates (available in 100% of deaths), medical records (MI: 49.4%, stroke 80.8% and heart 170 171 failure: 76.2%), household interviews and other sources of information was used. We also used Verbal Autopsies to ascertain cause of death in addition to medical records which were reviewed 172 by a health professional.¹⁷ To ensure a standard approach and accuracy for classification of 173 174 events across all countries and over time, a selection of cases from each country annually was adjudicated both locally and also by the adjudication chair, and if necessary further training was 175 176 provided.

177

178 Statistical analyses

The primary analyses were conducted excluding participants who reported having CVD at 179 baseline. Baseline characteristics were described for the entire cohort and stratified by low, 180 moderate, and high level of total PA. Total and domain specific PA values were not normally 181 distributed and are presented as median and inter-quartile range (IQR).¹⁸ Kruskal-Wallis test and 182 Jonckheere -Terpstra test^{19,20} were used to test the heterogeneity and trend across the four 183 country income levels (HIC, UMIC, LMIC, or LIC), respectively. For the two categorical PA 184 variables: level of total PA and whether meeting PA guideline, we calculated frequencies and 185 compared their difference and trend across the country income levels using Chi-square test and 186 Cochrane-Armitage test,^{21,22} as appropriate. 187

188

Age-sex-standardized incidence rates for all outcomes were calculated for levels of total PA and 189 whether meeting PA guidelines.²³ To examine the association between PA variables and 190 outcomes, we used the marginal Cox proportional hazard model.^{24,25} Models were adjusted for 191 age, sex, education, country income level, residency (urban vs. rural), family history of CVD and 192 smoking status (current and ever smoker vs. never smoker- cigarette, cigar and pipe smoking) 193 taking into account three levels of clustering. We conducted further analyses using wealth index 194 and household income (in separate models) in place of education but these did not change the 195 196 results. In addition, we further adjusted for body mass index (BMI).

197

Adjusted population attributable fractions related to not meeting PA guidelines and not achieving
high PA levels were calculated to quantify the benefit of PA, using the method developed by
Chen et al.²⁶ To minimize the potential for reverse causation, we conducted sensitivity analyses

by excluding participants who experienced CVD events in the first two years of follow-up.
Additional analyses were also conducted including PURE participants who had CVD at baseline
(n=141 945) and these yielded similar results. We estimated the effect of total PA on the
outcomes by country income level, sex, age (<50 or ≥50 years), BMI (<25 kg/m² or ≥25 kg/m²),
waist-hip-ratio (above 0.85 for female, 0.90 for male), and smoking, hypertension, and diabetes.

To assess and compare the effect of recreational PA vs. non-recreation PA, we fitted the adjusted 207 marginal Cox model with restricted cubic spline with four knots at the 5th, 35th, 65th and 95th 208 percentiles for overall and non-recreational PA.²⁷ Because 55% participants had no recreational 209 PA, we chose 50th, 65th, 80th and 95th percentile as the knots. We also examined whether the 210 association between PA and outcomes varied by country income and by type of PA (total, 211 212 recreational or non-recreational) using tests of interaction to compare the effects between HIC and UMIC vs LMIC and LIC. All analyses were conducted using SAS 9.4, for UNIX operating 213 214 system (SAS Institute, Cary, US) and R software, version 3.2.5, for Windows system.

215

216 **RESULTS**

A total of 168 916 participants were enrolled, of whom 141 945 had completed the IPAQ and the analyses were limited to the 130 843 participants without pre-existing CVD. Table 1 presents participant characteristics stratified by low, moderate, and high PA levels. Participant characteristics were not materially different in most features across the three groups with the exception of a lower proportion of males in the moderate PA group compared to the others and a greater proportion of family history of CVD in the high PA group. The prevalence of

hypertension and diabetes were lower with higher PA. There was no association between dietscore or body mass index with PA levels.

225

Table 2 presents PA by country income levels in both MET*minutes/week and minutes/week of moderate intensity PA. There was a trend towards lower total PA and recreational PA from HIC to LIC (p<0.0001 for both), but not for non-recreational PA. A large majority of participants met the PA guidelines, but fewer than half of the participants reached high levels of PA. During the mean follow-up of 6.9 ± 3.0 years there were 5334 deaths,1294 CVD deaths and

4040 non-CVD deaths, 1987 individuals with incident MI, 2086 with incident stroke, and 386 with new heart failure (Supplemental Table 1). When stratified by PA level, there was a graded reduction in age and sex adjusted event rates for all outcomes from low to moderate to high PA (p<0.0001 for trend for all events except for stroke (p=0.0010)) except heart failure (Table 3).

Those meeting the guidelines for minimal PA had lower age and sex adjusted rates of all

237 outcomes (Table 3).

238

Participating in PA at or above the PA guidelines was associated with significant lower rates of
outcomes compared to those participants not meeting the PA guidelines ≥600

241 MET*minute/week as per the IPAQ¹⁵ or PA \geq 150 minutes/week of moderate intensity PA as per

the WHO¹⁶). In fully adjusted models, meeting PA guidelines was associated with hazard ratios

243 (HR) [95% confidence interval] of 0.78 [0.74, 0.83] for mortality plus major CVD, 0.72 [0.67,

244 0.77] for mortality and 0.80 [0.74, 0.86] for major CVD (p<0.0001 for all).

246 In fully adjusted models increasing levels of PA (moderate and high) were associated with lower HR for mortality plus major CVD, mortality and major CVD compared to those with low levels 247 of total PA (p<0.0001 p<0.0001 and p=0.0005 for trend, respectively; Figure 1). When adjusted 248 249 for either wealth index or household income in place of education, HR did not change (Supplemental Tables 2 and 3). In addition, high PA was associated with a lower HR compared 250 251 to moderate PA for all outcomes. Dichotomizing high PA levels above or below the median value in this group off 6453 MET*minutes/week did not show further reductions in risk 252 (Supplemental Table 4). Increasing PA was also associated with lower HR in CVD mortality, 253 254 non-CVD mortality and MI (Supplemental Table 5). With further adjustment for BMI, the HR were slightly attenuated but remained significant (Supplemental Table 6). Excluding participants 255 who had a CVD event within the first two years of follow-up (to account for potential reverse 256 257 causality if sick individuals were less active), the results were consistent for all outcomes (Supplemental Table 7). 258

259

Survival curves (Supplemental Figure 1) for low, moderate, and high PA levels for our three
primary outcomes of mortality plus major CVD, mortality and major CVD indicated a lower risk
as PA increased (p<0.0001).

263

The five-year adjusted population attributable fraction of not meeting the PA guidelines was 5.3%, 8.0% and 4.6% for mortality plus major CVD, mortality and major CVD, respectively (Supplemental Figure 2). These values were higher (10.3%, 13.0% and 9.5%, respectively), for not achieving high PA.

269	Increasing PA was associated with lower risk mortality in a range of subgroups (Figure 2).
270	Compared to low levels of PA, moderate and high levels of PA were associated with a lower
271	graded risk for mortality regardless of sex, age, and in the presence of risk factors.
272	
273	Increasing PA was associated with significantly lower risk up to approximately 3000
274	MET*minutes/week (or 750 minutes/week of moderate intensity PA) with more modest benefits
275	above that PA level (p<0.0001). (Figure 3). For recreational PA, increasing PA was associated
276	with significantly lower risk up to approximately 600 MET*minutes/week (or 150 minutes/week
277	of moderate intensity PA) ($p=0.01$) (as few had levels of PA higher than this), while for non-
278	recreational PA, increasing PA was associated with significantly lower risk up to approximately
279	5000 MET*minutes/week (or 1250 minutes/week of moderate intensity PA) with more modest
280	benefits above that PA level ($p < 0.0001$).
281	
282	Increasing PA was associated with significantly lower risk for mortality plus major CVD in

283 UMIC and LIC, mortality in UMIC, LMIC and LIC and major CVD in UMIC and LIC (Table 284 4). When stratified by country income level (HIC + UMIC versus LMIC + LIC) there was a 285 significant interaction between country income level and PA levels for total (p=0.0012) and 286 recreational PA (p=0.0063) such that the HIC + UMIC had a lower risk with increasing PA level 287 (Figure 4). This was less clear for non-recreational PA (p=0.063).

288

289 DISCUSSION

290 In this study involving 3 HIC, 7 UMIC, 3 LMIC, and 4 LIC countries, increasing PA was

associated with a lower risk for mortality and incidence of major CVD. This lower risk was

292 present even at moderate levels of PA compared to low levels of PA, and was more marked at 293 higher PA levels . The benefit of PA was present independent of the type of PA (recreational or 294 non-recreational), a range of socio-economic and CVD risk factors, and was similar in various 295 countries with differing income levels.

296

297 In our study population of predominantly non-HIC residing participants, meeting the PA guidelines (150 minutes/week of moderate intensity of PA) was associated with a 22%, 28%, and 298 20% lower risk for all-cause mortality plus major CVD, mortality and major CVD, respectively, 299 300 resulting in adjusted population attributable fractions of $5 \cdot 3\%$, $8 \cdot 0\%$ and $4 \cdot 6\%$, respectively. These attributable fractions are similar to those reported by Lee et al.; 9.4% for mortality and 301 5.8% for CVD in HIC.⁹ In addition, we observed a graded effect such that participants at higher 302 levels of PA had a lower risk than those participants engaging in moderate levels of PA. For 303 example, compared to people at moderate levels of PA, participating in high levels of PA 304 conferred an additional reduction in risk of 15%, 19% and 12% for mortality plus major CVD, 305 mortality and major CVD, respectively. This benefit plateaued only at very high levels of PA 306 (approximately 1250 minutes/week of moderate intensity of PA). Similar to previous studies of 307 recreational PA in HIC,^{28,29} we did not observe any adverse efforts of PA on our outcomes even 308 in the approximately 9000 participants who reported over 2500 minutes/week of moderate 309 310 intensity of PA (equivalent to 17 times that of the PA guidelines). Therefore while participating 311 at even low levels of PA confers benefit (30 minutes per day for five days a week), the benefit continues to increase up to high levels of total PA. Given these findings and that the affordability 312 of other CVD interventions such as consuming fruits and vegetables,³⁰ and taking generic CVD 313

drugs is beyond the reach of many people in low and middle income countries,³¹ participating in
PA represents a low cost approach to CVD prevention .

316

317 When stratified by country income level there was a consistent reduction of risk with increasing PA. For HIC, meeting the PA guidelines was associated with a 30% lower risk for mortality, 318 319 which is lower than the 11% reported in a meta-analysis of walking from a study conducted predominantly in HIC.¹² However, this earlier study did not include participants in the high PA 320 level of >3000 MET*minutes/week as ours did in which we found a continued benefit with 321 322 increasing PA levels. Studies in Iran and China also reported PA to be significantly associated with lower mortality in a dose-dependent manner.^{32,33} Notably, Matthew et al. reported a HR of 323 0.61 [0.51-0.73] at the highest levels of PA of the Chinese women, which is similar to our HR of 324 325 0.65 [0.60, 0.71]. These findings are consistent with what we found in the LMIC category of countries of which Iran and China accounted for the overwhelming majority of participants in 326 our LMIC group. 327

328

Increasing PA levels, was associated with a lower risk of mortality plus major CVD in higher and UMIC compared to lower middle and LIC for both total and recreational PA; there were less clear differences with respect to non-recreational PA. It is unclear why recreational PA may be less effective in the lower middle and LIC, however, very few participants from the countries participated in any recreational PA and so these findings may be swayed by a very small number of participants who are atypical of the general populations in poorer countries.

336 Few studies have assessed the association of non-recreational PA with outcomes. The available studies are relatively small and report inconsistent results.^{12,34-38} We found that increasing levels 337 of both recreational PA and non-recreational PA were independently associated with lower risk 338 with our composite of all-cause mortality plus major CVD, indicating that PA of any type is 339 beneficial. Of note, high levels of PA was only possible in those individuals participating in non-340 341 recreational PA. Indeed, only 2.9% of our study population participated in high level of PA (≥3000 MET*minutes per week or ≥750 minutes/week of moderate intensity of PA) derived 342 exclusively from recreational PA compared to 37.9% of participants who attained this level 343 through non-recreational PA. This reflects the challenges inherent with participating in high 344 levels of recreational PA in that it is, by definition, conducted during discretionary hours of the 345 day outside of occupational and domestic duties. In contrast, incorporating PA into one's daily 346 lifestyle whether through active transportation, occupation and/or domestic duties has the 347 potential to achieve higher levels of PA that are associated with even lower risk for mortality and 348 CVD events. 349

350

To address concerns related to "reverse causality", we excluded those with known CVD and then conducted a sensitivity analysis further excluding those who had events within the first two years of follow-up. Our results were unchanged for our main study outcomes. In addition, we also conducted subgroup analyses stratified by sex, age, body mass index, smoking, presence of hypertension and presence of diabetes and observed consistent results. We also observed that increasing PA was associated with reduced CVD and non-CVD mortality. Regular PA has been reported to be associated with lower mortality from certain cancers^{28,34,36,39} and respiratory

358 conditions.⁴⁰ With continued follow-up, we anticipate accruing enough events to reliably

investigate the effects of PA on specific categories of non-CVD mortality.

360

361 Limitations

While PA determined from the self-reported IPAQ has been found to modestly overestimate PA, 362 363 it demonstrates good reliability and moderate validity compared to accelerometers such that higher IPAQ values correspond to higher levels of PA measured by accelerometers, thus 364 providing good internal validity.^{15,41,42} If PA is overestimated by the IPAQ, then the potential 365 366 benefits of PA may be more marked and may occur at lower PA levels than reported here. In addition, the IPAQ has been tested across a range of countries similar to the PURE study¹⁵ and 367 the use of self-reported measures for assessing PA in large studies is considered acceptable in 368 low resource settings.⁴³ 369

370

While it was not feasible to collect a proportionate sampling of the globe's population, our 371 selection of countries and communities ensured that our population was typical of the regions 372 from which participants were recruited with only modest differences compared to national data (373 Supplementary Appendix for comparative data).⁴⁴Although we did not recruit a random sample 374 of individuals, our approach minimized biases in selection of individuals once the communities 375 were identified. Given the range of countries across five continents at different economic levels 376 377 the large number of communities, and the large size of our study, our results are globally applicable. Given our method of event ascertainment, it is possible that some events may have 378 379 been misclassified. However, we believe this to be of very limited numbers as the majority of 380 events were ascertained using supporting documents, standardized definitions and adjudication

using standardized definitions providing a high level of confidence in the validity. Lastly, in such
a large study, it is not uncommon to report low p values that may not be clinically relevant,
therefore p values should be interpreted with caution unless they are extreme (p<0.001). Given
the magnitude and consistency of the effects observed across the different analyses, we can be
confident in our main findings.

386

387 Conclusions

Our findings demonstrate that PA (both recreational and non-recreational) is associated with a 388 389 lower risk for mortality and major CVD events., which was independent of the type of PA, other risk factors and seen in all major regions of the world and various country economic levels. In 390 particular we demonstrate that increasing PA is associated with lower risk in LMIC and LIC for 391 which limited data existed previously. . Even meeting the minimal PA guidelines such as 392 walking for as little as 30 minutes on most days of the week had a substantial benefit, while 393 higher levels of PA (up to and beyond 17 times the recommended PA guidelines) were 394 associated with even lower risks. . As participating in PA (especially in daily life) is 395 inexpensive, PA is a low cost approach to reducing deaths and CVD that is applicable globally 396 397 with potential large impact. The results of our study provide robust evidence to support public health interventions to increase all forms of PA levels in countries of different socioeconomic 398 circumstances.45 399

	Overall	Low Physical Activity	Moderate Physical Activity	High Physical Activity
	(n = 130 843)	$(n = 23 \ 631)$	$(n = 49\ 348)$	(n = 57 864)
Age (years)	$50{\cdot}2\pm9{\cdot}7$	$51{\cdot}0\pm10{\cdot}1$	50.5 ± 9.7	$49{\cdot}7\pm9{\cdot}5$
Male	54 621 (41.7%)	11 080 (46.9%)	18 224 (36.9%)	25 317 (43.8%)
Urban resident	69 993 (53·5%)	12 983 (54.9%)	28 525 (57.8%)	28 485 (49·2%)
Country Income Level				
High	13 546 (10.4%)	1435 (6.1%)	4991 (10.1%)	7120 (12·3%)
Upper Middle	34 625 (26.5%)	7479 (31.6%)	11 922 (24·2%)	15 224 (26·3%)
Lower Middle	53 841 (41.1%)	8620 (36.5%)	22 648 (45.9%)	22 573 (39.0%)
Low	28 831 (22.0%)	6097 (25.8%)	9787 (19.8%)	12 898 (22.4%)
Education				
None, Primary or Unknown	54 635 (41.9%)	10 642 (45·2%)	19 085 (38.8%)	24 908 (43.1%)
Secondary	50 500 (38.7%)	9035 (38·3%)	19 746 (40.1%)	21 719 (37.6%)
Trade, College or University	25 396 (19.5%)	3885 (16.5%)	10 412 (21.1%)	11 099 (19·2%)
Family History of Heart Disease or Stroke	36 812 (31.3%)	4911 (23.5%)	13 605 (30.5%)	18 296 (35.0%)
Hypertension	47 752 (39.0%)	9053 (42.6%)	18 364 (39.7%)	20 335 (36.9%)
Diabetes	12 740 (9.7%)	2898 (12.3%)	5102 (10.3%)	4740 (8.2%)
Smoker (current and former)	40 955 (31.5%)	7093 (30·3%)	13 695 (28.0%)	20 167 (35.0%)
Alternate Healthy Eating Index score	35.1 ±8.0	34.9 ±7.6	35.5 ±7.9	34.8 ±8.3
Body Mass Index (kg/m ²)	25·7 ±5·1	$25 \cdot 9 \pm 5 \cdot 4$	25.9 ± 5.0	$25 \cdot 4 \pm 5 \cdot 1$

Table 1: Participant characteristics stratified by levels of total physical activity (data presented as means \pm SD or counts and percentage).

Low physical activity <600 MET*minute/week and <150 minutes/week of moderate intensity physical activity Moderate physical activity 600-3000 MET*minute/week and 150-750 minutes/week of moderate intensity physical activity High physical activity \geq 3000 MET*minute/week and \geq 750 minutes/week of moderate intensity physical activity Column percentage presented for categorical variables.

	High Income	Upper Middle	Lower Middle	Low Income	P value (for	P value
	Countries	Income Countries	Income Countries	Countries	heterogeneity)	(for trend)
	(n=13 546)	(n=34 625)	(n=53 841)	(n= 28 831)		
Total Physical Activity*					<0.0001	<0.0001
MET*minutes/week	3227 [1485-6426]	2436 [750-5979]	2340 [960-5177]	2520 [721-6442]		
Minutes/week	(807 [371-1607])	(609 [188-1495])	(585 [240-1294])	(630 [180-1611])		
Recreational Physical					<0.0001	<0.0001
Activity*						
MET*minutes/week	518 [50-1386]	0 [0-320]	99 [0-693]	0 [0-0]		
Minutes/week	(130 [12-347])	(0 [0-80])	(25 [0-173])	(0 [0-0])		
Non-Recreational					<0.0001	0.7762
Physical Activity*						
MET*minutes/week	2115 [806-4980]	1983 [578-5400]	1748 [693-4186]	2297 [594-6222]		
Minutes/week	(529 [202-1245])	(496 [144-1350])	(437 [173-1047])	(574 [149-1556])		
Low Physical Activity	1435 (10.6%)	7479 (21.6%)	8620 (16.0%)	6097 (21.1%)	<0.0001	
Moderate Physical	4991 (36.8%)	11 922 (34·4%)	22 648 (42.1%)	9787 (33.9%)		
Activity						
High Physical Activity	7120 (52.6%)	15 224 (44.0%)	22 573 (41.9%)	12 947 (44.9%)	•	
Meeting Physical	12 111 (89.4%)	27 146 (78.4%)	45 221 (84.0%)	22 734 (78.9%)	<0.0001	<0.0001
Activity Guidelines †						

Table 2: Physical activity by country income level).

* presented as median [inter-quartile range (IQR)] in MET*minutes/week and in minutes/week of moderate intensity physical activity Low physical activity <600 MET*minute/week and <150 minutes/week of moderate intensity physical activity Moderate physical activity 600-3000 MET*minute/week and 150-750 minutes/week of moderate intensity physical activity High physical activity ≥3000 MET*minute/week and ≥750 minutes/week of moderate intensity physical activity
† meeting physical activity guidelines ≥600 MET*minute/week and ≥150 minutes/week of moderate intensity physical activity
P value for heterogeneity was calculated by Chisq test for categorical variable and Kruskal-Wallis for continuous variable. P value for trend was calculated by Cochrane-Armitage test for categorical variable and Jonckheere-Terpstra test for continuous variable.

	Mortal	ity plus	Mor	tality	Major	CVD^	CVD M	lortality	Non-	CVD	Myoc	ardial	Str	oke	Heart	Failure
	Major	CVD^		-				-	Mor	tality	Infar	ction				-
	Events	Rate	Events	Rate	Events	Rate	Events	Rate	Events	Rate	Events	Rate	Events	Rate	Events	Rate
Low	1941	9.46	1396	6.37	1000	5.13	377	1.75	1019	4.63	496	2.64	427	2.09	84	0.42
physical		[8·99,		[5.99,		[4·78,		[1.55,		[4·30,		[2·39,		[1.87,		[0.32,
activity (n =		9.94]		6.76]		5.48]		1.94]		4.96]		2.89]		2.31]		0.52]
23 549)																
Moderate	3002	7.14	1881	4.25	1682	4.13	480	1.12	1401	3.13	730	1.86	820	1.93	144	0.34
physical		[6·86,		[4.04,		[3.91,		[1.01,		[2.95,		[1.71,		[1.79,		[0.27,
activity ($n =$		7.43]		4·47]		4.34]		1.23]		3.32]		2.00]		2.08]		0.40]
49 245)																
High	3233	6.60	2057	4.11	1718	3.53	437	0.87	1620	3.24	761	1.58	839	1.68	158	0.30
physical		[6·36,		[3.92,		[3.35,		[0.78,		[3.07,		[1.47,		[1.56,		[0.25,
activity ($n =$		6.84]		4.30]		3.70]		0.96]		3.40]		1.70]		1.80]		0.36]
57 725)																
P value for		p<0.00		p<0.00		p<0.00		p<0.00		p<0.00		p<0.00		p=0.00		p=0.09
trend		01		01		01		01		01		01		10		97
Not meeting	1941	9.46	1396	6.37	1000	5.13	377	1.75	1019	4.63	496	2.64	427	2.09	84	0.42
guidelines		[8.99,		[5.99,		[4.78,		[1.55,		[4·30,		[2·39,		[1.87,		[0.32,
(n=23 549)		9.94]		6.76]		5.48]		1.94]		4.96]		2.89]		2.31]		0.52]
Meeting	6235	6.86	3938	4.19	3400	3.80	917	0.98	3021	3.21	1491	1.71	1659	1.79	302	0.32
guidelines		[6.68,		[4.05,		[3.66,		[0.92,		[3.08,		[1.62,		[1.70,		[0.28,
(n=106 970)		7.05]		4.33]		3.94]		1.05]		3.33]		1.81]		1.88]		0.35]
P value for		p<0.00		p<0.00		p<0.00		p<0.00		p<0.00		p<0.00		p=0.00		p=0.03
trend		01		01		01		01		01		01		88		76

Table 3: Summary of fatal and non-fatal events rates (per 1000 person years) and 95% CI stratified by different levels of physical activity, and those meeting, or not meeting, the recommended levels of physical activity.

Event rates are standardized for age and sex.

Low physical activity <600 MET*minutes/week; moderate physical activity 600-3000 MET*minutes/week; high physical activity \geq 3000 MET*minutes/week

CVD = cardiovascular disease

[^]Major CVD = CVD mortality plus incident myocardial infarction, stroke, and heart failure

A maximum of one event per participant is tabulated for each outcome.



Hazard Ratios and 95% CI

Figure 1: Hazard ratios and 95% CI for the pairwise comparison between levels of total physical activity, adjusted for age, sex, education, country income level, urban/rural residency, family history of CVD and smoking status taking into account household, community and country clustering. There were 3155, 2041 and 1723 events for all-cause mortality and major CVD, all-cause mortality, and major CVD, respectively.

Low physical activity level <600 MET*minute/week; moderate physical activity 600-3000 MET*minute/week; high physical activity ≥3000 MET*minute/week

CVD = cardiovascular disease

Major CVD = CVD mortality plus incident MI, stroke, and heart failure. The p values of the first column show the significance of each comparison. P-values of the second column show the significance of the overall effect of PA level.



Figure 2: Hazard ratios and 95% CI of level of total physical activity for mortality (adjusted for age, sex, education, country income level, urban/rural residency, family history of cardiovascular disease and smoking status taking into account household, community and country clustering). Based on data for 115 436 participants with complete data.

Note: Low physical activity level (<600 MET*minute/week) is the reference group.

Moderate physical activity 600-3000 MET*minute/week; high physical activity ≥3000 MET*minute/week

BMI = body mass index; WHR = waist to hip ratio (high WHR was defined as above 0.85 for female and above 0.9 for male)



Relationship between PA Type with Mortality and Major CVD expressed MET*min/week

Relationship between PA Type with Mortality and Major CVD expressed min/week of Moderate Intensity PA



Figure 3: Relationship between increasing total physical activity (PA; left panels), recreational PA (middle panels) and nonrecreational PA (occupational, transportation and housework PA; right panels) with mortality and major cardiovascular disease (CVD) presented in MET*minutes/week (top panels) and minutes/week of moderate intensity PA (bottom panels). Models were adjusted for age, sex, education, country income level, urban/rural residency, family history of CVD and smoking status, taking into account household, community and country clustering. Recreational PA was truncated at 3000 MET*minutes/week and 750 minutes/week of moderate intensity PA due to sparse observation above that level.

 Major CVD = CVD mortality plus incident MI, stroke, and heart failure

HR = hazard ratio

		Mortality plus Major CVD^	Mortality	Major CVD [^]
High Income Countries	Events	548	259	335
	Moderate PA	0.70 [0.54, 0.91]	0.69 [0.48, 1.00]	0.62 [0.44, 0.86]
	High PA	0.58 [0.45, 0.76]	0.54 [0.38, 0.78]	0.53 [0.38, 0.72]
	P value	0.0550	0.0818	0.1891
Upper Middle Income Countries	Events	1665	1150	836
	Moderate PA	0.82 [0.72, 0.93]	0.77 [0.66, 0.89]	0.86 [0.72, 1.03]
	High PA	0.65 [0.57, 0.74]	0.63 [0.54, 0.73]	0.64 [0.54, 0.77]
	P value	<0.0001	0.0056	0.0004
Lower Middle Income Countries	Events	2811	1343	1852
	Moderate PA	0.99 [0.89, 1.10]	0.94 [0.81, 1.08]	0.94 [0.82, 1.07]
	High PA	0.92 [0.82, 1.02]	0.79 [0.68, 0.92]	0.94 [0.83, 1.08]
	P value	0.0741	0.0043	0.8913
Low Income Countries	Events	1579	1203	804
	Moderate PA	0.76 [0.67, 0.87]	0.73 [0.63, 0.85]	0.83 [0.69, 1.00]
	High PA	0.61 [0.53, 0.69]	0.58 [0.50, 0.66]	0.63 [0.53, 0.75]
	P value	0.0002	0.0010	0.0013

Table 4. Summary of risk (hazards ratio) of mortality and major CVD events stratified by country income level and physical activity levels.

Hazard ratios (and 95% CI) were adjusted for age, sex, education, country income level, urban/rural residency, family history of CVD and smoking status taking into account household, community and country clustering.

Note: Low physical activity level (<600 MET*minute/week) is the reference group. Moderate physical activity 600-3000 MET*minute/week; high physical activity \geq 3000

MET*minute/week

CVD = cardiovascular disease





HIC+UMIC = high income countries plus upper middle income countries

LMIC+LIC = lower middle income countries plus low income countries

	Mortal	ity plus	Mor	tality	Major	CVD^	CVD N	lortality	Non-	CVD	Myoc	ardial	Str	oke	Heart	Failure
	Major			1		1		1	Mor	lanty	Infar	ction		1		1
	Events	Rate	Events	Rate	Events	Rate	Events	Rate	Events	Rate	Events	Rate	Events	Rate	Events	Rate
High	548	4.06	259	1.79	335	2.57	30	0.22	229	1.56	155	1.24	142	1.03	50	0.34
Income		[3.67,		[1.53,		[2·25,		[0.13,		[1.33,		[1.02,		[0.84,		[0.22,
Countries		4.45]		2.04]		2.88]		0.32]		1.80]		1.46]		1.23]		0.45]
Upper	1690	7.08	1171	4.65	843	3.66	270	1.07	901	3.57	407	1.79	311	1.33	139	0.57
Middle		[6.71,		[4·36,		[3.40,		[0.94,		[3.32,		[1.61,		[1.17,		[0.47,
Income		7.44]		4.94]		3.92]		1.21]		3.83]		1.98]		1.48]		0.68]
Countries																
Lower	2863	6.23	1371	2.92	1884	4.11	293	0.64	1078	2.28	583	1.31	1206	2.59	127	0.27
Middle		[5·98,		[2.75,		[3.91,		[0.56,		[2.13,		[1.19,		[2.43,		[0.21,
Income		6.48]		3.09]		4·31]		0.72]		2.43]		1.42]		2.74]		0.32]
Countries				_		_		_		_		_		_		_
Low Income	3075	10.75	2533	8.49	1338	4.87	701	2.37	1832	6.12	842	3.12	427	1.48	70	0.24
Countries		[10.34,		[8·13,		[4.59,		[2.18,		[5.81,		[2.89,		[1.33,		[0.18,
		11.16]		8.86]		5.15]		2.57]		6.43]		3.34]		1.63]		0.30]
Total	8176	7.32	5334	4.57	4400	4.03	1294	1.12	4040	3.46	1987	1.87	2086	1.84	386	0.33
		[7.15,		[4·44,		[3.90,		[1.05,		[3·34,		[1.79,		[1.76,		[0.30,
		7.49]		4·71]		4.16]		1.19]		3.57]		1.96]		1.93]		0.37]

Supplemental Table 1: Summary of fatal and non-fatal events rates (per 1000 person years) and 95% CI stratified by country income level.

Event rates are standardized for age and sex.

CVD = cardiovascular disease

[^]Major CVD = CVD mortality plus incident myocardial infarction, stroke, and heart failure

A maximum of one event per participant is tabulated for each outcome.

Supplemental Table 2: Summary of risk (hazard ratios) for mortality and major CVD events compared by physical activity (PA) levels (adjusted for wealth index in place of education).

	Mortality plus	plus Major CVD^ Mortality		tality	Major CVD [^]		
	HR	P value	HR	P value	HR	P value	
Moderate PA vs. Low PA	0.85 [0.80,0.91]	<0.0001	0.80 [0.74,0.87]	<0.0001	0.86 [0.79,0.94]	0.0008	
High PA vs. Low PA	0.73 [0.68,0.78]	<0.0001	0.66 [0.60,0.71]	<0.0001	0.76 [0.69,0.83]	<0.0001	
High PA vs. Moderate PA	0.85 [0.81,0.90]	<0.0001	0.82 [0.76,0.88]	<0.0001	0.88 [0.82,0.95]	0.0007	

Hazard ratios (and 95% CI) were adjusted for age, sex, wealth index, country income level, urban/rural residency, family history of CVD and smoking status taking into account household, community and country clustering.

Low PA level <600 MET*minute/week; Moderate PA 600-3000 MET*minute/week; High physical activity ≥3000 MET*minute/week;

MET*minute/week

CVD = cardiovascular disease

[^]Major CVD = CVD mortality plus incident MI, stroke, and heart failure

Supplemental Table 3: Summary of risk (hazard ratios) for mortality and major CVD events compared by physical activity (PA) levels (adjusted for household income in place of education).

	Mortality plus	Major CVD^	Mortality		Major CVD [^]		
	HR	P value	HR	P value	HR	P value	
Moderate PA vs. Low PA	0.85 [0.80,0.91]	<0.0001	0.80 [0.74,0.88]	<0.0001	0.87 [0.79,0.95]	0.002	
High PA vs. Low PA	0.72 [0.68,0.78]	<0.0001	0.64 [0.59,0.70]	<0.0001	0.77 [0.70,0.84]	<0.0001	
High PA vs. Moderate PA	0.85 [0.80,0.90]	<0.0001	0.80 [0.74,0.87]	<0.0001	0.88 [0.82,0.95]	0.0013	

Hazard ratios (and 95% CI) were adjusted for age, sex, household income, country income level, urban/rural residency, family history of CVD and smoking status taking into account household, community and country clustering.

Low PA level <600 MET*minute/week; Moderate PA 600-3000 MET*minute/week; High physical activity ≥3000 MET*minute/week

CVD = cardiovascular disease

Supplemental Table 4: Summary of risk (hazard ratios) for mortality and major CVD events compared by physical activity (PA) levels.

	Mortality plus	Major CVD^	Mor	tality	Major	CVD^
	HR	P value	HR	P value	HR	P value
Moderate PA vs.	0.85 [0.80, 0.91]	<0.0001	0.80 [0.74 ,0.87]	<0.0001	0.86 [0.79, 0.93]	0.0004
Low PA						
Lower high PA	0.88 [0.82 ,0.94]	0.0002	0.83 [0.76 ,0.91]	<0.0001	0.92 [0.84, 1.00]	0.0528
vs. Moderate PA						
Upper high PA	0.93 [0.86, 1.01]	0.0807	0.95 [0.85, 1.05]	0.2896	0.91 [0.82, 1.01]	0.0817
vs. Lower high						
PA						

Hazard ratios (and 95% CI) were adjusted for age, sex, education, country income level, urban/rural residency, family history of CVD and smoking status taking into account household, community and country clustering.

Low PA level <600 MET*minute/week; Moderate PA 600-3000 MET*minute/week; Lower high PA 3000-6453 MET*minute/week; Higher high PA ≥6453 MET*minute/week

CVD = cardiovascular disease

11	2				
	CVD Mortality	Non-CVD Mortality	Myocardial Infarction	Stroke	Heart Failure
Moderate PA	0.75 [0.64, 0.88]	0.83 [0.75, 0.91]	0.78 [0.68, 0.89]	0.93 [0.82, 1.06]	0.83 [0.63, 1.11]
High PA	0.58 [0.49, 0.68]	0.68 [0.62, 0.75]	0.67 [0.59, 0.76]	0.85 [0.75, 0.96]	0.76 [0.58, 1.01]
P value for trend	p=0.0010	p<·0001	p=0.0100	p=0.0716	p=0·4794

Supplemental Table 5: Summary of risk (hazard ratios) for major CVD events stratified by physical activity (PA) level.

Hazard ratios (and 95% CI) were adjusted for age, sex, education, country income level, urban/rural residency, family history of CVD and smoking status taking into account household, community and country clustering.

Note: Low physical activity level (<600 MET*minute/week) is the reference group.

Moderate physical activity 600-3000 MET*minute/week; high physical activity $\geq 3000 \text{ MET*minute/week}$ CVD = cardiovascular disease

Supplemental Table 6: Summary of risk (hazard ratios) for mortality and major CVD events stratified by physical activity (PA) level with the additional adjustment for body mass index.

	Mortality plus Major CVD^	Mortality	Major CVD ^	CVD Mortality	Non-CVD Mortality	Myocardial Infarction	Stroke	Heart Failure
Moderate PA	$\begin{array}{c} 0.88 \ [0.82, \\ 0.94] \end{array}$	0·84 [0·77, 0·92]	0·88 [0·80, 0·96]	0·80 [0·67, 0·96]	0.85 [0.77, 0.94]	0·81 [0·71, 0·93]	0·96 [0·84, 1·09]	0·83 [0·62, 1·12]
High PA	0·76 [0·71, 0·81]	0·69 [0·63, 0·75]	0·79 [0·72, 0·86]	0.64 [0.54, 0.76]	0·71 [0·64, 0·78]	0·71 [0·62, 0·82]	0·87 [0·77, 0·99]	0·81 [0·61, 1·08]
P value for trend	p<0.0001	p<0.0001	p=0.0056	p=0.0059	p<0.0001	p=0.0358	p=0.0926	p=0.8712

Hazard ratios (and 95% CI) were adjusted for age, sex, education, country income level, urban/rural residency, family history of CVD, body mass index and smoking status taking into account household, community and country clustering.

Note: Low physical activity level (<600 MET*minute/week) is the reference group.

Moderate physical activity 600-3000 MET*minute/week; high physical activity ≥3000 MET*minute/week

CVD = cardiovascular disease

Supplemental Table 7: Summary of risk (hazard ratios) for mortality and major CVD events stratified by physical activity (PA) level (*excluding* participants with CVD events in first two years of follow-up).

	Mortality plus	Mortality	Major CVD ^	CVD	Non-CVD	Myocardial	Stroke	Heart Failure
	Major CVD^			Mortality	Mortality	Infarction		
Moderate PA	0·89 [0·83, 0·96]	0·85 [0·78, 0·93]	0·87 [0·79, 0·96]	0·76 [0·63, 0·93]	0·88 [0·79, 0·98]	0·80 [0·69, 0·93]	0·95 [0·83, 1·09]	0·77 [0·56, 1·06]
High PA	0·77 [0·71, 0·83]	0·69 [0·63, 0·76]	0·80 [0·73, 0·89]	0·64 [0·53, 0·77]	0·71 [0·64, 0·79]	0·75 [0·65, 0·87]	0·88 [0·77, 1·02]	0·78 [0·57, 1·06]
P value for trend	p<0.0001	p<0.0001	p=0.0492	p=0.0533	p<0.0001	p=0·3588	p=0·1996	p=0.9531

Hazard ratios (and 95% CI) were adjusted for age, sex, education, country income level, urban/rural residency, family history of CVD and smoking status taking into account household, community and country clustering.

Note: Low physical activity level (<600 MET*minute/week) is the reference group.

Moderate physical activity 600-3000 MET*minute/week; high physical activity $\geq 3000 \text{ MET*minute/week}$ CVD = cardiovascular disease



Supplemental Figure 1: Adjusted survival curves for mortality and major cardiovascular disease (CVD) (left panel), mortality (middle panel) and major CVD (CVD mortality plus incident MI, stroke, and heart failure; right panel) stratified by level of physical activity (PA). All models adjusted for age, sex, education, country income level, urban/rural residency, family history of CVD and smoking status, taking into account household, community and country clustering.

Low physical activity <600 MET*minute/week; moderate physical activity 600-3000 MET*minute/week; high physical activity ≥3000 MET*minute/week

P values corresponding to testing heterogeneity of the three curves in each panel



Supplemental Figure 2. Adjusted 5-year population attributable fraction and 95% CI of not meeting physical activity (PA) guideline (gray bars) and of not participating high PA (black bars). Meaning a proportion reduction in the outcomes by 5 years if the entire population met PA guidelines (gray bars) and if entire population achieved high physical activity (\geq 3000 MET*minute/week; black bars). (Adjusted for age, sex, education, country income level, urban/rural residency, family history of CVD and smoking status, taking into account household, community and country clustering).

CVD = cardiovascular disease

ACKNOWLEDGEMENTS

SA Lear holds the Pfizer/Heart and Stroke Foundation Chair in Cardiovascular Prevention at St. Paul's Hospital. S Yusuf is supported by the Mary W Burke endowed chair of the Heart and Stroke Foundation of Ontario.

AUTHOR CONTRIBUTIONS

S Lear wrote the analysis plans and had the primary responsibility for writing this paper S Yusuf designed and supervised the study, data analysis, interpreted the data & reviewed and commented on drafts K Teo reviewed and commented on the data analysis and drafts S Rangarajan coordinated the worldwide study & reviewed and commented on drafts W Hu conducted the analysis & reviewed and commented on drafts A Casanova reviewed and commented on the data analysis and drafts D Leong reviewed and commented on the data analysis and drafts D Gasevic reviewed and commented on the data analysis and drafts All other authors coordinated the study in their respective countries and provided comments on drafts of the manuscript.

CONFLICT OF INTEREST

The authors declare no conflicts.

FUNDING SOURCES

The PURE Study is an investigator-initiated study that is funded by the Population Health Research Institute, the Canadian Institutes of Health Research, Heart and Stroke Foundation of Ontario, Support from CIHR's Strategy for Patient Oriented Research, through the Ontario SPOR Support Unit, as well as the Ontario Ministry of Health and Long-Term Care and through unrestricted grants from several pharmaceutical companies [with major contributions from Astra Zeneca (Canada), Sanofi-Aventis (France and Canada), Boehringer Ingelheim (Germany & Canada), Servier, and GSK], and additional contributions from Novartis and King Pharma and from various national or local organizations in participating countries.

These include: **Argentina:** Fundacion ECLA; **Bangladesh:** Independent University, Bangladesh and Mitra and Associates; **Brazil:** Unilever Health Institute, Brazil; **Canada:** Public Health Agency of Canada and Champlain Cardiovascular Disease Prevention Network; **Chile:** Universidad de la Frontera; China: National Center for Cardiovascular Diseases; Colombia: Colciencias, Grant number:6566-04-18062; **India:** Indian Council of Medical Research; **Malaysia:** Ministry of Science, Technology and Innovation of Malaysia Grant Nbr 100 - IRDC / BIOTEK 16/6/21 (13/2007), Grant Number 07-05-IFN-BPH 010, Ministry of Higher Education of Malaysia Grant Nbr 600 - RMI/LRGS/5/3 (2/2011), Universiti Teknologi MARA, Universiti Kebangsaan Malaysia (UKM-Hejim-Komuniti-15-2010); **occupied Palestinian territory:** the United Nations Relief and Works Agency for Palestine Refugees in the Near East (UNRWA), occupied Palestinian territory; International Development Research Centre (IDRC), Canada; **Philippines:** Philippine Council for Health Research & Development (PCHRD); **Poland:** Polish Ministry of Science and Higher Education grant Nr 290/W-PURE/2008/0, Wroclaw Medical University; **Saudi Arabia:** The Deanship of Scientific Research at King Saud University, Riyadh, Saudi Arabia (Research group number: RG -1436-013); **South Africa:** The North-West

University, SANPAD (SA and Netherlands Programme for Alternative Development), National Research Foundation, Medical Research Council of SA, The SA Sugar Association (SASA), Faculty of Community and Health Sciences (UWC); **Sweden**: Grants from the Swedish state under the Agreement concerning research and education of doctors; the Swedish Heart and Lung Foundation; the Swedish Research Council; the Swedish Council for Health, Working Life and Welfare, King Gustaf V:s and Queen Victoria Freemasons Foundation, AFA Insurance, Swedish Council for Working Life and Social Research, Swedish Research Council for Environment, Agricultural Sciences and Spatial Planning, Grant from the Swedish State under (LäkarUtbildningsAvtalet) Agreement, Grant from the Västra Götaland Region (FOUU); **TURKEY**: Metabolic Syndrome Society, Astra Zeneca, Turkey, Sanofi Aventis, Turkey; **UAE**: Sheikh Hamdan Bin Rashid Al Maktoum Award For Medical Sciences and Dubai Health Authority, Dubai UAE.

Role of Sponsor: The funders and sponsors had no role in the design and conduct of the study; in the collection, analysis, and interpretation of the data; in the preparation, review, or approval of the manuscript; or in the decision to submit the manuscript for publication.

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REFERENCES

1. Global, regional, and national age-sex specific all-cause and cause-specific mortality for 240 causes of death, 1990-2013: a systematic analysis for the Global Burden of Disease Study 2013. *Lancet* 2015; **385**(9963): 117-71.

2. Bloom DE, Cafiero ET, Jané-Llopis E, et al. The Global Economic Burden of Noncommunicable Diseases. Geneva: World Economic Forum; 2011.

3. Roth GA, Forouzanfar MH, Moran AE, et al. Demographic and epidemiologic drivers of global cardiovascular mortality. *N Engl J Med* 2015; **372**(14): 1333-41.

4. Benziger CP, Roth GA, Moran AE. The Global Burden of Disease Study and the Preventable Burden of NCD. *Global heart* 2016; **11**(4): 393-7.

5. Beaglehole R, Bonita R. Global public health: a scorecard. *Lancet* 2008; **372**(9654): 1988-96.

6. Sallis JF, Bull F, Guthold R, et al. Progress in physical activity over the Olympic quadrennium. *Lancet* 2016; **388**(10051): 1325-36.

7. Sixty-sixth World Health Assembly. Follow-up to the Political Declaration of the Highlevel Meeting of the General Assembly on the Prevention and Control of Non-communicable Diseases: United Nations, 2013.

8. Nocon M, Hiemann T, Muller-Riemenschneider F, Thalau F, Roll S, Willich SN. Association of physical activity with all-cause and cardiovascular mortality: a systematic review and meta-analysis. *Eur J Cardiovasc Prev Rehabil* 2008; **15**(3): 239-46.

9. Lee IM, Shiroma EJ, Lobelo F, Puska P, Blair SN, Katzmarzyk PT. Effect of physical inactivity on major non-communicable diseases worldwide: an analysis of burden of disease and life expectancy. *Lancet* 2012; **380**(9838): 219-29.

10. Milton K, Macniven R, Bauman A. Review of the epidemiological evidence for physical activity and health from low- and middle-income countries. *Global public health* 2014; **9**(4): 369-81.

11. Yusuf S, Hawken S, Ounpuu S, et al. Effect of potentially modifiable risk factors associated with myocardial infarction in 52 countries (the INTERHEART study): case-control study. *Lancet* 2004; **364**(9438): 937-52.

12. Kelly P, Kahlmeier S, Gotschi T, et al. Systematic review and meta-analysis of reduction in all-cause mortality from walking and cycling and shape of dose response relationship. *Int J Behav Nutr Phys Act* 2014; **11**: 132.

13. Johnsen AM, Alfredsson L, Knutsson A, Westerholm PJ, Fransson EI. Association between occupational physical activity and myocardial infarction: a prospective cohort study. *BMJ Open* 2016; 6(10): e012692.

14. Teo K, Chow CK, Vaz M, Rangarajan S, Yusuf S. 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): 1-7 e1.

15. Craig CL, Marshall AL, Sjostrom M, et al. International physical activity questionnaire: 12-country reliability and validity. *Med Sci Sports Exerc* 2003; **35**(8): 1381-95.

16. World Health Organization. Global recommendations on physical activity for health. Geneva: World Health Organization, 2010.

17. Gajalakshmi V, Peto R, Kanaka S, Balasubramanian S. Verbal autopsy of 48 000 adult deaths attributable to medical causes in Chennai (formerly Madras), India. *BMC Public Health* 2002; **2**: 7.

18. Guidelines for data processing and analysis of the International Physical Activity Questionnaire (IPAQ)-Short and Long Forms. 2005.

http://www.ipaq.ki.se/dloads/IPAQ%20LS%20Scoring%20Protocols_Nov05.pdf

19. Jonckheere AR. A distribution-free k-sample test against ordered alternatives. *Biometrika* 1954; **41**: 133-45.

20. Terpstra TJ. The asymptotic normality and consistency of Kendall's test against trend, when ties are present in one ranking. *Indagationes Mathematicae* 1952; **14**: 327-33.

21. Armitage P. Tests for linear trends in proportions and frequencies. *Biometrics* 1955; **11**: 375-86.

22. Cochran WG. Some methods of strengthening the common 2 x tests. *Biometrics* 1954; **10**: 417-51.

23. Schenker N, Parsons VL, Lochner KA, Wheatcroft G, Pamuk ER. Estimating standard errors for life expectancies based on complex survey data with mortality follow-up: A case study using the National Health Interview Survey Linked Mortality Files. *Stat Med* 2011; **30**(11): 1302-11.

24. Lee EW, Wei LJ, Amato DA. Cox-Type Regression Analysis for Large Numbers of Small Groups of Correlated Failure Time Observations. In: Goel JPKaPK, ed. Survival Analysis: State of the Art. Dordrecht, Netherlands: Kluwer Academic Publishers; 1992: 237-47.

25. Lin DY. Cox Regression Analysis of Multivariate Failure Time Data: The Marginal Approach. *Statistics in Medicine* 1994; **13**: 2233-47.

26. Chen L, Lin DY, Zeng D. Attributable fraction functions for censored event times. *Biometrika* 2010; **97**(3): 713-26.

27. Harrell FE. Regression Modeling Strategies: with applications to linear models, logistic regression, and survival analysis. New York, USA: Springer-Verlag New York, Inc.; 2001.

28. Arem H, Moore SC, Patel A, et al. Leisure time physical activity and mortality: a detailed pooled analysis of the dose-response relationship. *JAMA Intern Med* 2015; **175**(6): 959-67.

29. Moore SC, Patel AV, Matthews CE, et al. Leisure time physical activity of moderate to vigorous intensity and mortality: a large pooled cohort analysis. *PLoS medicine* 2012; **9**(11): e1001335.

30. Miller V, Yusuf S, Chow CK, et al. Availability, affordability, and consumption of fruits and vegetables in 18 countries across income levels: findings from the Prospective Urban Rural Epidemiology (PURE) study. *The Lancet Global health* 2016; **4**(10): e695-703.

31. Khatib R, McKee M, Shannon H, et al. Availability and affordability of cardiovascular disease medicines and their effect on use in high-income, middle-income, and low-income countries: an analysis of the PURE study data. *Lancet* 2016; **387**(10013): 61-9.

32. Etemadi A, Abnet CC, Kamangar F, et al. Impact of body size and physical activity during adolescence and adult life on overall and cause-specific mortality in a large cohort study from Iran. *Eur J Epidemiol* 2014; **29**(2): 95-109.

33. Matthews CE, Jurj AL, Shu XO, et al. Influence of exercise, walking, cycling, and overall nonexercise physical activity on mortality in Chinese women. *Am J Epidemiol* 2007; **165**(12): 1343-50.

34. Yu R, Leung J, Woo J. Housework reduces all-cause and cancer mortality in Chinese men. *PLoS One* 2013; **8**(5): e61529.

35. Besson H, Ekelund U, Brage S, et al. Relationship between subdomains of total physical activity and mortality. *Med Sci Sports Exerc* 2008; **40**(11): 1909-15.

36. Autenrieth CS, Baumert J, Baumeister SE, et al. Association between domains of physical activity and all-cause, cardiovascular and cancer mortality. *Eur J Epidemiol* 2011; **26**(2): 91-9.

37. Sabia S, Dugravot A, Kivimaki M, Brunner E, Shipley MJ, Singh-Manoux A. Effect of intensity and type of physical activity on mortality: results from the Whitehall II cohort study. *Am J Public Health* 2012; **102**(4): 698-704.

38. Hu G, Jousilahti P, Antikainen R, Tuomilehto J. Occupational, commuting, and leisuretime physical activity in relation to cardiovascular mortality among finnish subjects with hypertension. *Am J Hypertens* 2007; **20**(12): 1242-50.

39. Wu CY, Hu HY, Chou YC, Huang N, Chou YJ, Li CP. The association of physical activity with all-cause, cardiovascular, and cancer mortalities among older adults. *Prev Med* 2015; **72**: 23-9.

40. Vaes AW, Garcia-Aymerich J, Marott JL, et al. Changes in physical activity and allcause mortality in COPD. *The European respiratory journal* 2014; **44**(5): 1199-209.

41. Bauman A, Bull F, Chey T, et al. The International Prevalence Study on Physical Activity: results from 20 countries. *Int J Behav Nutr Phys Act* 2009; **6**: 21.

42. Wanner M, Probst-Hensch N, Kriemler S, Meier F, Autenrieth C, Martin BW. Validation of the long international physical activity questionnaire: Influence of age and language region. *Preventive medicine reports* 2016; **3**: 250-6.

43. Strath SJ, Kaminsky LA, Ainsworth BE, et al. Guide to the assessment of physical activity: Clinical and research applications: a scientific statement from the American Heart Association. *Circulation* 2013; **128**(20): 2259-79.

44. Corsi DJ, Subramanian SV, Chow CK, et al. Prospective Urban Rural Epidemiology (PURE) study: Baseline characteristics of the household sample and comparative analyses with national data in 17 countries. *Am Heart J* 2013; **166**(4): 636-46.e4.

45. Reis RS, Salvo D, Ogilvie D, Lambert EV, Goenka S, Brownson RC. Scaling up physical activity interventions worldwide: stepping up to larger and smarter approaches to get people moving. *Lancet* 2016; **388**(10051): 1337-48.