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# Association of Cycling With All-Cause and Cardiovascular Disease Mortality Among Persons With Diabetes The European Prospective Investigation Into Cancer and Nutrition (EPIC) Study

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**IMPORTANCE** Premature death from all causes and cardiovascular disease (CVD) causes is higher among persons with diabetes.

**OBJECTIVE** To investigate the association between time spent cycling and all-cause and CVD mortality among persons with diabetes, as well as to evaluate the association between change in time spent cycling and risk of all-cause and CVD mortality.

**DESIGN, SETTING, AND PARTICIPANTS** This prospective cohort study included 7459 adults with diabetes from the European Prospective Investigation into Cancer and Nutrition study. Questionnaires regarding medical history, sociodemographic, and lifestyle information were administered in 10 Western European countries from 1992 through 2000 (baseline examination) and at a second examination 5 years after baseline. A total of 5423 participants with diabetes completed both examinations. The final updated primary analysis was conducted on November 13, 2020.

**EXPOSURES** The primary exposure was self-reported time spent cycling per week at the baseline examination. The secondary exposure was change in cycling status from baseline to the second examination.

**MAIN OUTCOMES AND MEASURES** The primary and secondary outcomes were all-cause and CVD mortality, respectively, adjusted for other physical activity modalities, diabetes duration, and sociodemographic and lifestyle factors.

**RESULTS** Of the 7459 adults with diabetes included in the analysis, the mean (SD) age was 55.9 (7.7) years, and 3924 (52.6%) were female. During 110 944 person-years of follow-up, 1673 deaths from all causes were registered. Compared with the reference group of people who reported no cycling at baseline (O min/wk), the multivariable-adjusted hazard ratios for all-cause mortality were 0.78 (95% CI, 0.61-0.99), 0.76 (95% CI, 0.65-0.88), 0.68 (95% CI, 0.57-0.82), and 0.76 (95% CI, 0.63-0.91) for cycling 1 to 59, 60 to 149, 150 to 299, and 300 or more min/wk, respectively. In an analysis of change in time spent cycling with 57 802 person-years of follow-up, a total of 975 deaths from all causes were recorded. Compared with people who reported no cycling at both examinations, the multivariable-adjusted hazard ratios for all-cause mortality were 0.90 (95% CI, 0.71-1.14) in those who cycled and then stopped, 0.65 (95% CI, 0.46-0.92) in initial noncyclists who started cycling, and 0.65 (95% CI, 0.53-0.80) for people who reported cycling at both examinations. Similar results were observed for CVD mortality.

**CONCLUSION AND RELEVANCE** In this cohort study, cycling was associated with lower all-cause and CVD mortality risk among people with diabetes independent of practicing other types of physical activity. Participants who took up cycling between the baseline and second examination had a considerably lower risk of both all-cause and CVD mortality compared with consistent noncyclists.

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Editor's Note

Supplemental content

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Corresponding Author: Mathias Ried-Larsen, PhD, Centre for Physical Activity Research, Rigshospitalet, Blegdamsvej 9, Copenhagen 2100, Denmark (mathias.ried-larsen@ regionh.dk). remature death from all causes and cardiovascular disease (CVD) is higher among people with diabetes. Regular physical activity is a critical behavioral target in the management of diabetes, but only structured exercise, in contrast with advice only, has been shown to improve CVD risk factors. Thus, it is necessary to investigate the influence of engagement in specific unstructured physical activities on mortality in this patient population.

Cohort studies in populations with diabetes have reported inverse associations between overall physical activity, leisure-time physical activity (LTPA), and walking with all-cause and CVD mortality. 7,8 However, associations with walking have been inconsistent, likely because only moderate-intensity walking appears to be associated with a reduced risk of all-cause and CVD mortality.9 Meeting the physical activity recommendations both in terms of total physical activity volume as well as intensity is a major challenge, especially in people with diabetes. 10-12 Because lack of time is often quoted as a barrier, incorporating activities into everyday life may be an effective strategy. Cycling is a potential activity to replace motorized transport for shortto-medium distance trips (eg, during commuting to work without a substantial effect on time use). As moderate-tohigh intensities are reached during cycling at self-selected paces in adults, 13-16 cycling could decrease the risk of premature mortality. It may also be a feasible strategy because cycling is one of the preferred activities in people with type 2 diabetes. 17,18 It is well established that there is a strong inverse association between cycling and the risk of all-cause and cause-specific mortality, such as CVD, in the general population. 19-21 However, to our knowledge, there are no studies that have examined the role of cycling in preventing premature mortality in people with diabetes.

The primary aim of this cohort study was to investigate the associations between cycling and all-cause and CVD mortality among persons with diabetes. A secondary aim was to study the associations between change in cycling over a 5-year period and all-cause and CVD mortality.

## Methods

## **Study Design and Setting**

This study was a prospective cohort study of people with diabetes at baseline in the European Prospective Investigation into Cancer and Nutrition (EPIC) study cohort. <sup>22</sup> In EPIC, 23 centers in 10 Western European countries (France, Italy, Spain, United Kingdom, the Netherlands, Greece, Germany, Sweden, Denmark, and Norway) collected information on nutrition, lifestyle, anthropometry, and medical history from more than 521 234 participating men and women. <sup>22</sup> Medical history, sociodemographic, and lifestyle information was assessed by questionnaires at baseline between 1992 and 1998 (baseline examination) and at a second examination between 1996 and 2011. Data were only available from 22 centers because data from Greece were not released for this study. The ethical review boards from the International Agency for Research on Cancer and all local participating centers approved this study. Informed consent was obtained from all participants.

## **Key Points**

**Question** Is cycling associated with risk of all-cause and cardiovascular disease mortality among persons with diabetes?

**Findings** In this cohort study of 7459 persons with diabetes, cycling was associated with at least a 24% lower all-cause mortality rate when compared with noncyclists, independent of other physical activity and putative confounders. Taking up cycling over a 5-year period was associated with at least a 35% lower risk of all-cause mortality when compared with consistent noncyclists.

**Meaning** Cycling could be encouraged as an activity for persons with diabetes to lower the risk of premature mortality.

#### **Study Population**

From the entire EPIC population, people with diabetes at the baseline assessment were included in the present study. Diabetes was based on self-report on a single occasion or by a second source (at least 1), including repeated self-report, by a general physician, linkage to register or medical record at a later point, prescription of use of glucose-lowering or diabetes-related medication, or baseline glycated hemoglobin at least 6.0% (42 mmol/mol [to convert to proportion of total hemoglobin, multiply by 0.01]).<sup>7</sup>

#### **Data Collection**

Study procedures have been described in detail elsewhere.<sup>22</sup> Briefly, height, weight, and waist circumference were measured using similar protocols across study centers.<sup>22</sup> Body mass index was calculated as weight in kilograms divided by height in meters squared. Central obesity was defined according the International Diabetes Federation criteria.<sup>23</sup> Diabetes duration was calculated as the time from self-reported age or calendar year of medical diagnosis to baseline.

Dietary intake, including alcohol consumption, was assessed by a questionnaire (quantitative, semiquantitative, or a combination) and 7-day or 14-day record, and individual energy and nutrient intake was based on the standardized EPIC nutrient database. <sup>22,24</sup> Because the Mediterranean diet is associated with improved metabolic control and decreased risk of diabetes, <sup>25-27</sup> it was included as a covariate expressed as the relative Mediterranean diet score. <sup>25,27,28</sup> Dietary data were only available for the baseline examination.

#### **Assessment of Physical Activity**

Information about physical activity habits was obtained from a lifestyle questionnaire and included information about duration and frequency of leisure-time and occupational physical activity. <sup>22,29</sup> Weekly time spent cycling to and/or from work and leisure time during winter and summer was averaged into a single variable of total annual cycling time and then categorized as 0, 1 to 59, 60 to 149, 150 to 299, and 300 or more min/wk. Change in total cycling from baseline to the second examination was categorized based on total time spent cycling at the 2 examinations: (1) noncycling: participants who reported zero minutes of cycling at both examinations; (2) stopped cycling: participants who reported cycling (any amount) at baseline but not at the second examination; (3)

started cycling: participants who did not report cycling at baseline but did report cycling (any amount) at the second examination; or (4) maintained cycling: those who were consistent cyclists at both examinations.

The LTPA energy expenditure (without cycling included) metabolic equivalent of tasks in hours per week (MET-h/wk<sup>15</sup>) was calculated at both examinations as the sum of energy expenditures from the following activities: gardening, do-it-yourself activities, stair climbing, housework activities, walking, and sports. Because information on stair climbing was only available from 4 study centers at the second examination, this activity was not included in the second examination LTPA variable. Occupational physical activity was reported in categories of sedentary occupation, standing occupation, manual or heavy manual work, or nonworker.<sup>29</sup>

## **Outcome Ascertainment**

The primary and secondary outcomes were all-cause and CVD mortality, respectively. Any deaths due to CVD were coded according to the *International Statistical Classification of Diseases and Related Health Problems, Tenth Revision*, using the codes IOO to I99. Vital status and cause of death were obtained through record linkage with registries, physicians or hospitals, or direct follow-up. <sup>30</sup> Follow-up end for the primary and secondary analyses was on March 26, 2015 (eTable 1 in Supplement 1).

#### **Statistical Analysis**

A statistical analysis plan was developed (Supplement 2) and preregistered (NCTO4171557) prior to commencing the analyses. The risks of all-cause and CVD mortality were computed as hazard ratios (HRs) with 95% CIs according to weekly time spent cycling at baseline estimated using stratified Cox proportional hazard regression models with age as the underlying time scale. Analyses were corrected for delayed entry. Participants were considered at risk from age at baseline examination in the primary analyses and from age at the second examination in the analyses of change in cycling. Participants leaving the study during follow-up owing to emigration or premature withdrawal were right censored at the age of emigration or withdrawal. A priori, participants with sufficient information to be included in the most extensive statistical models (models 3 and 2 for the primary and secondary analyses, respectively) were included in the primary models (models 2 and 1 for the primary and secondary analyses, respectively).

A crude model (model 1) was fitted with categories of cycling as exposure (O [reference], 1-59, 60-149, 150-299, and ≥300 min/wk), adjusted for sex and age (years), and further stratified by study center to adjust for confounding of this variable. The proportional hazards assumption for cycling was met within each stratum. Model 1 was further adjusted for attained educational level (no formal education, primary school, technical or professional school, secondary school, or university degree), smoking status (never smoker, former smoker, or current smoker), diabetes duration (years), adherence to the Mediterranean diet (relative Mediterranean diet score: low, 0-6 points; medium, 7-10 points; or high, 11-18 points), <sup>25,27</sup> total

energy intake (quartiles of kcal/d),<sup>24</sup> physical activity excluding cycling (quartiles of LTPA energy expenditure), and occupational physical activity (sedentary occupation, standing occupation, manual or heavy manual work, nonworker, or unknown status) (model 2 [main model]). Finally, history of prevalent stroke (yes or no), previous myocardial infarction (yes or no), prevalent cancer (yes or no), hypertension (yes or no), hyperlipidemia (yes or no), and central obesity (yes or no) at baseline were added as covariates (model 3). Effect modification by sex and diabetes duration (≥5 vs <5 years) was evaluated statistically using the likelihood-ratio test by comparing model 2, adding a multiplicative interaction term for sex or diabetes duration and cycling with a model including only main effects. Because several covariates did not meet the proportional hazard assumption in the multivariable models 2 and 3, we computed an extended Cox regression analysis in which we stratified by study center and energy intake in the analyses with all-cause mortality as the outcome. In the models with CVD mortality as the outcome, we stratified by study center, educational level, and LTPA (excluding cycling). We conducted a range of preplanned sensitivity analyses for the primary model (model 2) specified in the statistical analysis plan (Supplement 2) to investigate the association of residual confounding and reverse causality (excluding all deaths and CVD deaths within the first 2 years following the baseline examination).

In the preplanned secondary analysis, associations between all-cause and CVD mortality and change in cycling (from baseline to second examination) were investigated. The associations were initially adjusted for sex and age at the second examination and stratified by study center. A multivariable model was fitted additionally adjusting for educational level at baseline, smoking status at both examinations, diabetes duration at the time of second examination, LTPA (excluding cycling) at both examinations, and occupational physical activity at the second examination. The multivariable analyses were stratified by study center, baseline occupational physical activity, adherence to the relative Mediterranean diet, and total energy intake at baseline.

Because the associations between baseline and cycling and all-cause or CVD mortality were nonlinear, we computed (post hoc) restricted cubic splines of the respective associations with knot placements at the 10th, 50th, and 90th percentiles, as recommended by Harrell.31 These analyses were repeated using the average of baseline and second examinations. Because of large amounts of zero values in the cycling variable, the percentiles were computed excluding zero. Moreover, 10-year adjusted (standardized) cumulative mortality according to cycling at baseline or change in cycling status, consistent with the primary models, were estimated post hoc. Flexible parametric survival models<sup>32</sup> with additional postestimation of adjusted differences (95% CIs) in 10-year cumulative mortality comparing 0 min/wk of cycling at baseline and/or at the second examination to higher levels of cycling or stopping/ starting/maintaining cycling were used. Also, to assess the influence of unmeasured confounding, the E-value was calculated based on our estimated HR for all-cause and CVD mortality.<sup>33</sup> All analyses were conducted using STATA IC, version 16.1 (StataCorp), with 2-sided  $\alpha$  = .05 considered statistically significant.

#### Results

Of the 492 763 participants enrolled in the EPIC cohort, 10 995 had diabetes at the baseline examination. The analytic sample consisted of 7459 participants, of whom 4701 (63.0%) had confirmed diabetes and 4699 (63.0%) were noncyclists. Among the sample, the mean (SD) age was 55.9 (7.7) years, the mean (SD) diabetes duration was 7.7 (8.1) years, and 3924 (52.6%) were female (Table 1). The main reason for exclusion from the analytic sample was missing covariate or exposure information (eFigure 1 in Supplement 1). Reasons for missing data are reported in eTable 2 in Supplement 1.

The participants were followed for a mean (SD) of 14.9 (4.4) years (110 944 person-years) with 1673 deaths from all causes and 811 deaths attributable to CVD. A subset of participants (n = 5423) also completed the second examination and were included in the analysis of change in cycling. This analysis had a mean (SD) of 10.7 (4.3) years of follow-up, accumulating a total of 57 802 person-years with 975 deaths from all causes and 429 from CVD.

#### Baseline Cycling and All-Cause and CVD Mortality

The HR for all-cause mortality was lower for people reporting any cycling (>0 min/wk) when compared with noncyclists (model 1; Table  $2^{34}$ ). Cycling was also associated with a reduced risk of CVD mortality (Table 2). Adjusting for educational level, lifestyle risk factors, and diabetes duration did not materially change the association between cycling and all-cause mortality (model 2; Table 2). The 10-year cumulative mortality risk difference relative to 0 min/wk of cycling for ascending cycling categories (1-59, 60-149, 150-299, and ≥300 min/wk) were −1.9%, −2.0%, −2.7%, and −2.1% for all-cause mortality and −1.2%, −1.2%, −2.2%, and −1.0% for CVD mortality, respectively (eTable 3 in Supplement 1), with the lowest category of cycling consistently having the highest cumulative mortality (eFigure 2 in Supplement 1).

No statistically significant multiplicative interactions of sex or diabetes duration and cycling were observed for all-cause nor CVD mortality (eTable 4 in Supplement 1). Further adjustment for existing conditions and CVD risk factors only slightly attenuated the associations (model 3; Table 2). Sensitivity analyses investigating residual confounding by smoking, sports participation, self-reported diabetes, and reverse causality broadly confirmed the associations between cycling and both all-cause and CVD mortality (eTable 5 in Supplement 1), and E-values ranged from 1.66 to 2.91 in categories of baseline cycling (eTable 6 in Supplement 1).

The dose-response relationship with baseline cycling as a continuous variable for both all-cause and CVD mortality was modeled post hoc; for comparison, the association for LTPA (excluding cycling) is provided (**Figure 1**<sup>34</sup>). This revealed a reversed J-shaped association between both outcomes and cycling, and a linear association for LTPA (excluding cycling) (HRs per 10 MET-h increase per week: 0.97 [95% CI, 0.95-0.98] and

0.96 [95% CI, 0.94-0.98] for all-cause and CVD mortality, respectively) (Figure 1). The distribution of cycling and LTPA among observations with and without events underlying these analyses are found in eFigure 3 in Supplement 1. The associations between average cycling at baseline and second examination approximated a curve-linear dose-response relationship (eFigure 4 and eTables 7 and 8 in Supplement 1).

## Change in Cycling and All-Cause and CVD Mortality

For both all-cause and CVD mortality, HRs were 0.65 or less among participants who started or maintained cycling (Figure 2) relative to noncyclists (risk differences relative to noncyclists were –3.7% for all-cause mortality and –2.7% for CVD mortality; eTable 9 and eFigure 5 in Supplement 1). After excluding deaths within the first 2 years from the second examination, the associations were unchanged: HRs for all-cause mortality were 0.93 (95% CI, 0.72-1.19), 0.66 (95% CI, 0.45-0.95), and 0.64 (95% CI, 0.51-0.80) for people who stopped, started, or maintained cycling, respectively, compared with noncyclists), and corresponding HRs for CVD mortality were 1.08 (95% CI, 0.77-1.51), 0.56 (95% CI, 0.31-0.99), and 0.54 (95% CI, 0.39-0.75). E-values were more than 2 for the groups that started cycling and maintained cycling (eTable 6 in Supplement 1).

### Discussion

We observed that time spent cycling was associated with a lower risk of all-cause and CVD mortality in people with diabetes, independent of other physical activities and other putative confounders. However, the dose-response relationships were ambiguous.

The association between cycling and all-cause and CVD mortality in this study of person with diabetes was of the same magnitude and direction as observed in the healthy population. 13,20,21,35-41 This is moreover in line with previous observations by Andersen et al,8 who observed that the associations between physical activity and all-cause mortality seemed to be similar between the healthy population and persons living with chronic disease, including cardiovascular, lung, and musculoskeletal diseases. However, their investigation did not include persons with diabetes nor did it specifically investigate the interaction between chronic disease state and associations between cycling and all-cause mortality.8 The lower risk of all-cause and CVD mortality associated with overall physical activity as well as walking among persons with diabetes is well established. 7,9,42-48 The present investigation extends the level of evidence within this field by documenting that cycling and taking up cycling may offer specific health benefits in people with diabetes over and above other physical activities, including walking. Mixed-mode commuting (walking and/or cycling) has been associated with decreased mortality in persons with diabetes. 49 However, the association was weaker as compared with our observations. Because physical activity intensity is important in mediating the health benefits from walking among people with type 2 diabetes, 9,50 a lower intensity of physical activity, such as walking, when

Table 1. Baseline Characteristics of Sample Subgroups and the Total Cohort

	Participants, No. (%)						
	Time spent cyclin						
Characteristic	0	1-59	60-149	150-299	≥300	Total	
Participants	4648	422	999	736	654	7459	
Annual cycling, median (IQR), min/wk	0 (0-0)	30 (30-30)	90 (60-120)	180 (165-240)	420 (330-600)	0 (0-90)	
Sex							
Male	1972 (42.4)	251 (59.5)	561 (56.2)	401 (54.5)	350 (53.5)	3535 (47.4)	
Female	2676 (57.6)	171 (40.5)	438 (43.8)	335 (45.5)	304 (46.5)	3924 (52.6)	
Age, mean (SD), y	55.6 (7.8)	55.0 (7.4)	56.0 (7.9)	56.4 (7.1)	57.3 (6.9)	55.9 (7.7)	
Diabetes duration, mean (SD), y	7.7 (8.1)	7.3 (7.4)	7.7 (8.4)	7.7 (8.1)	7.5 (8.0)	7.7 (8.1)	
ducational level							
None	1086 (23.4)	6 (1.4)	42 (4.2)	27 (3.7)	18 (2.8)	1179 (15.8)	
Primary school	1642 (35.3)	179 (42.4)	354 (35.4)	295 (40.1)	287 (43.9)	2757 (37.0)	
Technical/professional school	793 (17.1)	98 (23.2)	277 (27.7)	210 (28.5)	193 (29.5)	1571 (21.1)	
Secondary school	390 (8.4)	43 (10.2)	107 (10.7)	66 (9.0)	58 (8.9)	664 (8.9)	
University degree	737 (15.9)	96 (22.7)	219 (21.9)	138 (18.8)	98 (15.0)	1288 (17.3)	
Smoking status							
Never	2363 (50.8)	161 (38.2)	400 (40.0)	304 (41.3)	251 (38.4)	2757 (46.6)	
Former	1243 (26.7)	164 (38.9)	373 (37.3)	272 (37.0)	268 (41.0)	1571 (31.1)	
Current	1042 (22.4)	97 (23.0)	226 (22.6)	160 (21.7)	135 (20.6)	664 (22.3)	
BMI, a mean (SD)	29.4 (5.1)	28.0 (4.5)	28.2 (4.8)	28.6 (4.6)	28.6 (4.7)	29.0 (5.0)	
Vaist circumference, mean (SD), cm <sup>b</sup>							
Male	101.4 (11.3)	98.8 (10.7)	98.9 (10.8)	99.1 (10.1)	99.3 (11.2)	100.4 (11.1)	
Female	92.2 (13.0)	86.7 (12.5)	89.5 (13.5)	92.0 (13.6)	91.6 (12.9)	91.6 (13.1)	
Central obesity						,	
Yes	3681 (79.2)	280 (66.4)	707 (70.8)	558 (75.8)	487 (74.5)	5713 (76.6)	
No	967 (20.8)	142 (33.6)	292 (29.2)	178 (24.2)	167 (25.5)	1746 (23.4)	
TPA (excluding cycling), median (IQR), //ET-h/wk	(====)	( ,		2. 2 (2.3.2)		2 (20)	
Gardening	0.0 (0.0-8.0)	4.0 (0.0-10.0)	4.0 (0.0-14.0)	4.0 (0.0-14.0)	4.0 (0.0-16.0)	0.0 (0.0-11.0)	
Do-it-yourself activities	0.0 (0.0-4.5)	4.5 (0.0-13.5)	4.5 (0.0-9.0)	2.3 (0.0-9.0)	0.0 (0.0-13.5)	0.0 (0.0-6.8)	
Stair climbing	0.9 (0.0-2.1)	1.0 (0.4-2.6)	1.3 (0.4-2.6)	1.3 (0.3-2.6)	1.3 (0.4-2.6)	1.0 (0.3-2.3)	
Housework	30.0 (3.0-84.0)	12.0 (0.0-42.0)	12.0 (6.0-42.0)	15.0 (6.0-45.0)	21.0 (6.0-45.0)	21.0 (3.0-63.0	
Walking	15.0 (6.0-27.0)	10.5 (4.5-21.0)	12.0 (6.0-21.0)	15.0 (9.0-27.0)	21.0 (12.0-36.0)	15.0 (6.0-27.0	
Sports	0.0 (0.0-0.0)	0.0 (0.0-6.0)	0.0 (0.0-9.0)	0.0 (0.0-11.3)	0.0 (0.0-12.0)	0.0 (0.0-6.0)	
Occupational physical activity							
Sedentary occupation	993 (21.4)	121 (28.7)	258 (25.8)	172 (23.4)	119 (18.2)	1663 (22.3)	
Standing occupation	964 (20.7)	80 (19.0)	179 (17.9)	137 (18.6)	110 (16.8)	1470 (19.7)	
Manual work	433 (9.3)	49 (11.6)	112 (11.2)	84 (11.4)	69 (10.6)	747 (10.0)	
Nonworker	2204 (47.4)	169 (40.0)	442 (44.2)	331 (45.0)	336 (51.4)	3482 (46.7)	
Unknown	54 (1.2)	3 (0.7)	8 (0.8)	12 (1.6)	20 (3.1)	97 (1.3)	
Energy intake, mean (SD), kcal/d	2025.1 (640.5)	2154.6 (641.9)	2103.0 (631.9)	2072.6 (640.9)	2120.9 (664.6)	2056.0 (642.9	
Adherence to relative Mediterranean diet	(0.0.3)		(001.0)	(0 .0.0)	(000)	(0 .2	
Low	772 (16.6)	89 (21.1)	301 (30.1)	212 (28.8)	203 (31.0)	1577 (21.1)	
Medium	2007 (43.2)	231 (54.7)	499 (49.9)	378 (51.4)	312 (47.7)	3427 (45.9)	
High	1869 (40.2)	102 (24.2)	199 (19.9)	146 (19.8)	139 (21.3)	2455 (32.9)	
Comorbidities	1005 (40.2)	102 (24.2)	199 (19.9)	140 (13.0)	133 (21.3)	2733 (32.3)	
Prevalent cancer	204 (4.4)	17 (4.0)	38 (3.8)	34 (4.6)	35 (5.4)	328 (4.4)	
Prevalent stroke	156 (3.4)	11 (2.6)	36 (3.6)	17 (2.3)	16 (2.4)	236 (3.2)	
Myocardial infarction	225 (4.8)	23 (5.5)	56 (5.6)	38 (5.2)	35 (5.4)	377 (5.1)	
Hyperlipidemia	1943 (41.8)	194 (46.0)	361 (36.1)	278 (37.8)	252 (38.5)	3028 (40.6)	
Hypertension	2200 (47.3)	203 (48.1)	485 (48.5)	359 (48.8)	348 (53.2)	3595 (48.2)	

Abbreviations: BMI, body mass index; IQR, interquartile range; LTPA, leisure-time physical activity; MET-h/wk, metabolic equivalent of tasks in hours per week.

<sup>&</sup>lt;sup>a</sup> Calculated as weight in kilograms divided by height in meters squared.

 $<sup>^{\</sup>rm b}$  To convert centimeters to inches, divide by 0.39.

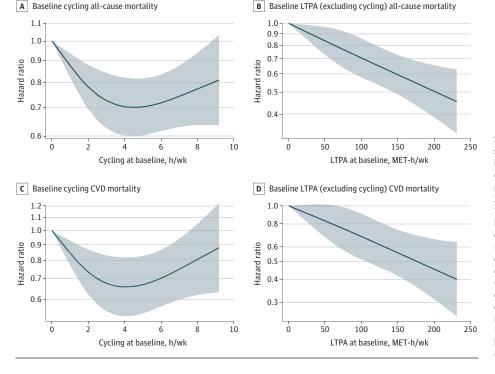
Table 2. Association Between Total Volume of Cycling at the Baseline Examination and All-Cause and CVD Mortality

	Cycling at first examination, HR (95% CI), min/wk							
Variable	0	1-59	60-149	150-299	≥300			
MET-h <sup>a</sup>	0	0.1-7.4	7.5-18.6	18.7-37.4	≥37.5			
Participants, No.	4648	422	999	736	654			
Total person-years	70 741	5994	14 395	10 521	9292			
Cases of all-cause mortality	1059	79	231	150	154			
Mortality rate/1000 person-years	14.97 (14.10-15.90)	13.18 (10.57-16.43)	16.04 (14.11-18.25)	14.26 (12.15-16.37)	16.57 (14.15-19.14)			
Model 1 <sup>b</sup>	1 [Reference]	0.76 (0.60-0.96)	0.72 (0.62-0.84)	0.65 (0.55-0.78)	0.71 (0.60-0.85)			
Model 2 <sup>c</sup>	1 [Reference]	0.78 (0.61-0.99)	0.76 (0.65-0.88)	0.68 (0.57-0.82)	0.76 (0.63-0.91)			
Model 3 <sup>d</sup>	1 [Reference]	0.81 (0.64-1.03)	0.77 (0.66-0.90)	0.70 (0.59-0.84)	0.81 (0.68-0.97)			
Cases of CVD mortality	499	41	119	66	86			
Mortality rate/1000 person-years	7.05 (6.46-7.70)	6.84 (5.04-9.29)	8.27 (6.91-9.89)	6.27 (4.93-7.89)	9.26 (7.49-11.43)			
Model 1 <sup>b</sup>	1 [Reference]	0.72 (0.52-1.00)	0.72 (0.59-0.89)	0.55 (0.42-0.71)	0.75 (0.59-0.96)			
Model 2 <sup>e</sup>	1 [Reference]	0.79 (0.56-1.11)	0.75 (0.60-0.93)	0.57 (0.44-0.76)	0.80 (0.62-1.03)			
Model 3 <sup>f</sup>	1 [Reference]	0.83 (0.59-1.18)	0.78 (0.63-0.98)	0.61 (0.46-0.81)	0.91 (0.70-1.17)			

Abbreviations: CVD, cardiovascular disease; HR, hazard ratio; LTPA, leisure-time physical activity; MET-h, metabolic equivalent of tasks in hours.

duration, adherence to the Mediterranean diet, LTPA (excluding cycling), occupational physical activity, prevalent stroke, myocardial infarction, prevalent cancer, hyperlipidemia, hypertension, and central obesity.

Figure 1. Post hoc Analyses of the Associations Between Cycling or Leisure-Time Physical Activity (LTPA) (Excluding Cycling) and All-Cause or Cardiovascular Disease (CVD) Mortality



Solid lines are hazard ratios, and shading is the upper and lower bounds of the 95% CIs. Restricted cubic splines were applied (knot placements in the analyses were 0.5, 2.0, and 7.5 h/wk, and 21.1, 74.3, and 150.8 metabolic equivalent of tasks in hours per week [MET-h/wk] for cycling and LTPA, respectively). The MET-h/wk for cycling can be calculated based on the Compendium of Physical Activities using the code 01015 "Bicycling, general" with an estimated intensity of 7.5 MET-h.  $^{34}$  To convert hours of cycling to MET-h, multiply by 7.5. The corresponding MET-h values for 2, 4, 6, 8, and 10 hours of weekly cycling are 15, 30, 45, 60, and 75, respectively.

<sup>&</sup>lt;sup>a</sup> The MET-h is calculated based on the Compendium of Physical Activities using the code O1015 "Bicycling, general" with an estimated intensity of 7.5 MET-h.<sup>34</sup> To convert minutes of cycling to MET-h, divide by 60 and multiply by 7.5. Exposure variables were obtained at the baseline examination.

<sup>&</sup>lt;sup>b</sup> Stratified according to study center and adjusted for sex and age.

<sup>&</sup>lt;sup>c</sup> Stratified according to study center and total energy intake (quartiles of kcal/wk). Adjusted for sex, age, educational level, smoking status, diabetes duration, adherence to the Mediterranean diet, LTPA (excluding cycling), and occupational physical activity.

<sup>&</sup>lt;sup>d</sup> Stratified according to study center and total energy intake (quartiles of kcal/wk). Adjusted for sex, age, educational level, smoking status, diabetes

e Stratified according to study center, educational level, and LTPA (excluding cycling). Adjusted for sex, age, smoking status, diabetes duration, adherence to the Mediterranean diet, total energy intake (quartiles of kcal/wk), and occupational physical activity.

f Stratified according to study center, educational level, and LTPA (excluding cycling). Adjusted for sex, age, smoking status, diabetes duration, adherence to the Mediterranean diet, total energy intake (quartiles of kcal/wk), occupational physical activity, prevalent stroke, myocardial infarction, prevalent cancer, hyperlipidemia, hypertension, and central obesity.

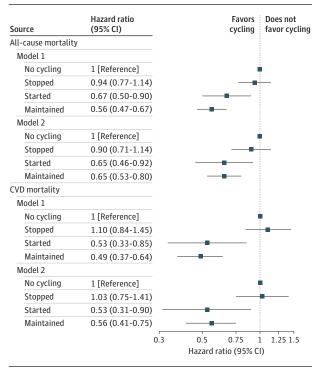
compared with cycling, may account for differences.<sup>15</sup> The lower risk of all-cause and CVD mortality observed in consistent cyclists or persons initiating cycling may be mediated by improvements in aerobic fitness, which is associated with all-cause and CVD mortality.<sup>48,51</sup>

The shapes of the dose-response curves were ambiguous. The uptick in the point estimate at high volumes of cycling can be associated with an increased risk of fatal injuries with increased cycling (eg, in urban settings or in those with increased risk of CVD or respiratory diseases owing to exposure to air pollutants during cycling in settings with dense motorized traffic). 52-54 In addition, the beneficial effects of physical activity on cardiovascular risk factors with increasing air pollutants may be attenuated. 55-58 However, previous cohort studies<sup>36,59,60</sup> have reported that levels of traffic-related air pollution did not modify the inverse association of outdoor physical activity with mortality or incidence of heart disease. Although air pollution during exercise may decrease lung function acutely, 55,57,58,61 it seems that the benefits of physical activity on the risk of asthma and chronic obstructive pulmonary disease are maintained when performed in moderately polluted settings such as urban environments. 62 Because cycling is associated with an increased risk of fatal injuries compared with passive transportation, 52 this may also explain the small uptick in the point estimate for all-cause mortality risk with increased cycling, but it cannot explain the corresponding shape of the curve for CVD mortality. Of note, commuter cycling may increase the risk of injuries and hospital admissions compared with nonactive commuting in the general population. 52 However, the health benefits of cycling may outweigh the increased risk of injuries owing to a decreased risk of morbidities in cyclists.  $^{63}\,\mathrm{While}$  data on traumatic deaths due to cycling would have enabled a benefit-harm analysis in this study, those data were unfortunately not collected. Another explanation for the uptick may be associated with potential adverse effects of very high volumes of physical activity. While a recent study does not support this in the healthy population, 64 it may constitute a concern in persons with diabetes. Participants who cycle many hours per week (eg, ≥5 h/wk) may do so at a higher intensity and thus accumulate a greater volume of physical activity compared with participants who cycle less. However, looking at the total volume of cycling (up to 75 MET-h/wk) in the context of the total volume of LTPA, it seems unlikely that this should explain the uptick because this is not observed for LTPA even at substantially higher volumes than observed with cycling. Finally, because the post hoc analyses using the average of the baseline and second examination as exposure to limit misclassification of cycling owing to change in cycling habits during follow-up to some extent attenuated the minor uptick observed when using the baseline exposure only, this suggests that the uptick might partly be a consequence of measurement error on the exposure variable. This is in line with previous observations.65-67

## Limitations

Limitations of the study include the observational design that limits causal inferences. We were unable to distinguish be-

Figure 2. Association Between All-Cause or Cardiovascular Disease (CVD) Mortality and Changes in Cycling From Baseline to the Second Examination



Among noncyclists, those who stopped cycling, those who started cycling, and those who maintained cycling, person-years of follow-up were 35 674, 5923, 3571, and 12 635, respectively; cases of all-cause mortality were 598, 138, 49, and 190, respectively; and cases of CVD mortality were 247, 78, 19, and 85, respectively. Median (interquartile range) minutes of weekly cycling at baseline were 0 (0-0), 90 (60-180), 0 (0-0), and 150 (90-300) minutes for noncyclists, those who stopped cycling, those who started cycling, or those who maintained cycling, respectively. Median (interquartile range) minutes of weekly cycling at the second survey were 0 (0-0), 0 (0-0), 90 (60-210), and 150 (90-300) minutes for noncyclists, those who stopped cycling, those who started cycling, or those who maintained cycling, respectively. All-cause mortality rates per 1000 person-years were 16.8 (95% CI, 15.5-18.2), 23.3 (95% CI, 19.7-27.5), 13.7 (95% CI, 10.4-18.2), and 15.0 (95% CI, 13.0-17.3) for noncyclists, those who stopped cycling, those who started cycling, or those who maintained cycling, respectively; the corresponding incidence rates per 1000 person-years for CVD mortality were 7.0 (95% CI, 6.1-7.8), 13.2 (95% CI, 10.6-16.4), 5.3 (95% CI, 3.4-8.3), and 6.7 (95% CI, 5.4-8.3), respectively. Model 1 was stratified by study center and adjusted for sex and age (second examination). Model 2 was stratified according to study center, baseline adherence to the Mediterranean diet, baseline occupational physical activity, and total energy intake, and adjusted for sex, age (second examination), baseline educational level, smoking status at both surveys, diabetes duration at the second survey, leisure-time physical activity (excluding cycling) at both examinations, and occupational physical activity at the second examination.

tween type 1 and type 2 diabetes, but generally type 2 diabetes accounts for 90% of all diabetes in adults. <sup>68</sup> To maximize the analytic sample, we chose to include both persons with self-reported diabetes and persons with diabetes confirmed through other sources, which increases the risk of misclassification. However, only few numerical differences were observed in the characteristics between persons with confirmed and self-reported diabetes (eTable 10 in Supplement 1), and the analysis restricted to those with confirmed diabetes only supported the overall findings. Although we adjusted the analyses

for a range of potential confounders, these were mostly selfreported and thus prone to misclassification. Although slightly attenuated, the associations observed for all-cause and CVD mortality were confirmed in sensitivity analyses, when those who had ever smoked and people reporting engaging in any sports were excluded. This suggests that residual confounding by smoking and sports-related physical activity may be minor, although the 95% CIs for the latter were wide for CVD mortality. A concern may be confounding by concomitant pharmacological intervention. However, as pharmacological intervention intensifies with increasing diabetes duration,<sup>69</sup> and as we consistently adjusted for diabetes duration, we may, to some extent, have addressed this issue. Because the prevalence of microvascular and macrovascular complications are highly prevalent among persons with diabetes, 70 persons with a history of CVD at baseline were included in the primary analyses to increase the generalizability of the findings. While this may increase the risk of reverse causality, the 2-year benchmark sensitivity analysis did not suggest reverse causation. Evalue calculations showed that to explain away the estimated associations of baseline and change in cycling with

mortality with adjustment for the measured covariates, an unmeasured confounder would need to have an HR of at least 1.66 and 1.94 with both cycling and mortality, respectively. We decided a priori only to include participants with complete data for all statistical models, which could have introduced selection bias and limited generalizability. While rerunning the analyses with missing data statistically imputed for cycling and confounders confirmed the present findings, this may not fully overcome the biases. Finally, the results may not be generalizable to people using electric cycles.

### Conclusions

Results of this cohort study showed that engaging in cycling was associated with a lower risk of all-cause and CVD mortality among people with diabetes after considering other physical activities as well as other risk factors. Studies with repeated measurement of cycling and documentation of cycling-related accidents are needed to elaborate on the doseresponse relationship between cycling and mortality.

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#### **REFERENCES**

1. Roglic G, Unwin N. Mortality attributable to diabetes: estimates for the year 2010. *Diabetes Res* 

## *Clin Pract*. 2010;87(1):15-19. doi:10.1016/j.diabres. 2009.10.006

- 2. Colberg SR, Sigal RJ, Yardley JE, et al. Physical activity/exercise and diabetes: a position statement of the American Diabetes Association. *Diabetes Care*. 2016;39(11):2065-2079. doi:10.2337/dc16-1728
- 3. Umpierre D, Ribeiro PA, Kramer CK, et al. Physical activity advice only or structured exercise training and association with HbA1c levels in type 2 diabetes: a systematic review and meta-analysis. *JAMA*. 2011;305(17):1790-1799. doi:10.1001/jama. 2011.576
- 4. Boulé NG, Haddad E, Kenny GP, Wells GA, Sigal RJ. Effects of exercise on glycemic control and body mass in type 2 diabetes mellitus: a meta-analysis of controlled clinical trials. *JAMA*. 2001;286(10):1218-1227. doi:10.1001/jama.286.10.1218
- **5**. Chudyk A, Petrella RJ. Effects of exercise on cardiovascular risk factors in type 2 diabetes: a meta-analysis. *Diabetes Care*. 2011;34(5):1228-1237. doi:10.2337/dc10-1881
- **6.** Hayashino Y, Jackson JL, Fukumori N, Nakamura F, Fukuhara S. Effects of supervised exercise on lipid profiles and blood pressure control in people with type 2 diabetes mellitus: a meta-analysis of randomized controlled trials. *Diabetes Res Clin Pract*. 2012;98(3):349-360. doi:10.1016/j.diabres.2012.10. 004
- 7. Sluik D, Buijsse B, Muckelbauer R, et al. Physical activity and mortality in individuals with diabetes mellitus: a prospective study and meta-analysis. Arch Intern Med. 2012;172(17):1285-1295. doi:10. 1001/archinternmed.2012.3130
- **8**. Andersen LB, Schnohr P, Schroll M, Hein HO. All-cause mortality associated with physical activity during leisure time, work, sports, and cycling to work. *Arch Intern Med.* 2000;160(11):1621-1628. doi:10.1001/archinte.160.11.1621
- 9. Gregg EW, Gerzoff RB, Caspersen CJ, Williamson DF, Narayan KM. Relationship of walking to mortality among US adults with diabetes. *Arch Intern Med*. 2003;163(12):1440-1447. doi:10.1001/archinte.163.12.1440
- **10.** Zhao G, Ford ES, Li C, Mokdad AH. Compliance with physical activity recommendations in US adults with diabetes. *Diabet Med.* 2008;25(2):221-227. doi:10.1111/j.1464-5491.2007.02332.x
- 11. Morrato EH, Hill JO, Wyatt HR, Ghushchyan V, Sullivan PW. Physical activity in U.S. adults with diabetes and at risk for developing diabetes, 2003. *Diabetes Care*. 2007;30(2):203-209. doi:10.2337/dc06-1128
- 12. van der Berg JD, Stehouwer CD, Bosma H, et al. Associations of total amount and patterns of sedentary behaviour with type 2 diabetes and the metabolic syndrome: The Maastricht Study. *Diabetologia*. 2016;59(4):709-718. doi:10.1007/s00125-015-3861-8
- **13.** Blond MB, Rosenkilde M, Gram AS, et al. How does 6 months of active bike commuting or leisure-time exercise affect insulin sensitivity, cardiorespiratory fitness and intra-abdominal fat? a randomised controlled trial in individuals with overweight and obesity. *Br J Sports Med.* 2019;53 (18):1183-1192. doi:10.1136/bjsports-2018-100036
- **14.** Berntsen S, Malnes L, Langåker A, Bere E. Physical activity when riding an electric assisted bicycle. *Int J Behav Nutr Phys Act.* 2017;14(1):55. doi: 10.1186/s12966-017-0513-z

- **15.** Ainsworth BE, Haskell WL, Whitt MC, et al. Compendium of physical activities: an update of activity codes and MET intensities. *Med Sci Sports Exerc.* 2000;32(9)(suppl):5498-S504. doi:10.1097/00005768-200009001-00009
- **16.** Oja P, Mänttäri A, Heinonen A, et al. Physiological effects of walking and cycling to work. *Scand J Med Sci Sports*. 1991;1:151-157. doi:10.1111/j. 1600-0838.1991.tb00288.x
- 17. Forbes CC, Plotnikoff RC, Courneya KS, Boulé NG. Physical activity preferences and type 2 diabetes: exploring demographic, cognitive, and behavioral differences. *Diabetes Educ.* 2010;36(5): 801-815. doi:10.1177/0145721710378538
- **18.** Wanko NS, Brazier CW, Young-Rogers D, et al. Exercise preferences and barriers in urban African Americans with type 2 diabetes. *Diabetes Educ.* 2004;30(3):502-513. doi:10.1177/014572170403000322
- **19**. Kelly P, Kahlmeier S, Götschi 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. doi:10.1186/s12966-014-0132-x
- 20. Nordengen S, Andersen LB, Solbraa AK, Riiser A. Cycling and cardiovascular disease risk factors including body composition, blood lipids and cardiorespiratory fitness analysed as continuous variables: part 2—systematic review with meta-analysis. *Br J Sports Med*. 2019;53(14):879-885. doi:10.1136/bjsports-2018-099778
- 21. Nordengen S, Andersen LB, Solbraa AK, Riiser A. Cycling is associated with a lower incidence of cardiovascular diseases and death: part 1—systematic review of cohort studies with meta-analysis. *Br J Sports Med*. 2019;53(14):870-878. doi:10.1136/bjsports-2018-099099
- **22.** Riboli E, Hunt KJ, Slimani N, et al. European Prospective Investigation into Cancer and Nutrition (EPIC): study populations and data collection. *Public Health Nutr*. 2002;5(6B):1113-1124. doi:10.1079/PHN2002394
- 23. Alberti KG, Zimmet P, Shaw J; IDF Epidemiology Task Force Consensus Group. The metabolic syndrome—a new worldwide definition. *Lancet*. 2005;366(9491):1059-1062. doi:10.1016/ S0140-6736(05)67402-8
- **24.** Slimani N, Deharveng G, Unwin I, et al. The EPIC nutrient database project (ENDB): a first attempt to standardize nutrient databases across the 10 European countries participating in the EPIC study. *Eur J Clin Nutr.* 2007;61(9):1037-1056. doi:10.1038/sj.ejcn.1602679
- 25. Romaguera D, Guevara M, Norat T, et al; InterAct Consortium. Mediterranean diet and type 2 diabetes risk in the European Prospective Investigation into Cancer and Nutrition (EPIC) study: the InterAct project. *Diabetes Care*. 2011;34 (9):1913-1918. doi:10.2337/dc11-0891
- **26.** Esposito K, Maiorino MI, Ciotola M, et al. Effects of a Mediterranean-style diet on the need for antihyperglycemic drug therapy in patients with newly diagnosed type 2 diabetes: a randomized trial. *Ann Intern Med.* 2009;151(5):306-314. doi:10. 7326/0003-4819-151-5-200909010-00004
- **27**. Buckland G, Agudo A, Luján L, et al. Adherence to a Mediterranean diet and risk of gastric adenocarcinoma within the European Prospective Investigation into Cancer and Nutrition (EPIC)

- cohort study. *Am J Clin Nutr*. 2010;91(2):381-390. doi:10.3945/ajcn.2009.28209
- 28. Trichopoulou A, Costacou T, Bamia C, Trichopoulos D. Adherence to a Mediterranean diet and survival in a Greek population. *N Engl J Med*. 2003;348(26):2599-2608. doi:10.1056/NEJMoa025039
- 29. Wareham NJ, Jakes RW, Rennie KL, et al. Validity and repeatability of a simple index derived from the short physical activity questionnaire used in the European Prospective Investigation into Cancer and Nutrition (EPIC) study. *Public Health Nutr.* 2003;6(4):407-413. doi:10.1079/PHN2002439
- **30**. Pischon T, Boeing H, Hoffmann K, et al. General and abdominal adiposity and risk of death in Europe. *N Engl J Med*. 2008;359(20):2105-2120. doi:10.1056/NEJMoa0801891
- **31.** Harrell FE Jr. Regression Modeling Strategies: With Applications to Linear Models, Logistic and Ordinal Regression, and Survival Analysis. Springer; 2001.
- **32**. Lambert PC, Royston P. Further development of flexible parametric models for survival analysis. *Stata J.* 2009;9(2):265-290. doi:10.1177/1536867X0900900206
- **33.** VanderWeele TJ, Ding P. Sensitivity analysis in observational research: introducing the E-value. *Ann Intern Med.* 2017;167(4):268-274. doi:10.7326/M16-2607
- **34.** Ainsworth BE, Haskell WL, Herrmann SD, et al. 2011 Compendium of Physical Activities: a second update of codes and MET values. *Med Sci Sports Exerc*. 2011;43(8):1575-1581. doi:10.1249/MSS. 0b013e31821ece12
- **35**. Grøntved A, Rasmussen MG, Blond K, Østergaard L, Andersen ZJ, Møller NC. Bicycling for transportation and recreation in cardiovascular disease prevention. *Curr Cardiovasc Risk Rep.* 2019; 13(26). doi:10.1007/s12170-019-0623-z
- **36.** Andersen ZJ, de Nazelle A, Mendez MA, et al. A study of the combined effects of physical activity and air pollution on mortality in elderly urban residents: the Danish Diet, Cancer, and Health Cohort. *Environ Health Perspect*. 2015;123(6):557-563. doi:10.1289/ehp.1408698
- **37**. Koolhaas CM, Dhana K, Schoufour JD, et al. Physical activity and cause-specific mortality: the Rotterdam Study. *Int J Epidemiol*. 2018;47(5):1705-1713. doi:10.1093/ije/dyy058
- **38.** Oja P, Kelly P, Pedisic Z, et al. Associations of specific types of sports and exercise with all-cause and cardiovascular-disease mortality: a cohort study of 80 306 British adults. *Br J Sports Med*. 2017;51(10):812-817. doi:10.1136/bjsports-2016-096822
- **39.** Sahlqvist S, Goodman A, Simmons RK, et al. The association of cycling with all-cause, cardiovascular and cancer mortality: findings from the population-based EPIC-Norfolk cohort. *BMJ Open.* 2013;3(11):e003797. doi:10.1136/bmjopen-2013-003797
- **40**. Østergaard L, Jensen MK, Overvad K, Tjønneland A, Grøntved A. Associations between changes in cycling and all-cause mortality risk. *Am J Prev Med*. 2018;55(5):615-623. doi:10.1016/j. amepre.2018.06.009
- **41**. Schnohr P, Marott JL, Jensen JS, Jensen GB. Intensity versus duration of cycling, impact on all-cause and coronary heart disease mortality: the

- Copenhagen City Heart Study. Eur J Prev Cardiol. 2012;19(1):73-80. doi:10.1177/1741826710393196
- **42**. Hu G, Jousilahti P, Barengo NC, Qiao Q, Lakka TA, Tuomilehto J. Physical activity, cardiovascular risk factors, and mortality among Finnish adults with diabetes. *Diabetes Care*. 2005;28(4):799-805. doi:10.2337/diacare.28.4.799
- **43**. Jonker JT, De Laet C, Franco OH, Peeters A, Mackenbach J, Nusselder WJ. Physical activity and life expectancy with and without diabetes: life table analysis of the Framingham Heart Study. *Diabetes Care*. 2006;29(1):38-43. doi:10.2337/diacare.29.01. 06.dc05-0985
- **44.** Nelson KM, Boyko EJ, Koepsell T. All-cause mortality risk among a national sample of individuals with diabetes. *Diabetes Care*. 2010;33 (11):2360-2364. doi:10.2337/dc10-0846
- **45**. Smith TC, Wingard DL, Smith B, Kritz-Silverstein D, Barrett-Connor E. Walking decreased risk of cardiovascular disease mortality in older adults with diabetes. *J Clin Epidemiol*. 2007; 60(3):309-317. doi:10.1016/j.jclinepi.2006.06.013
- **46**. Tanasescu M, Leitzmann MF, Rimm EB, Hu FB. Physical activity in relation to cardiovascular disease and total mortality among men with type 2 diabetes. *Circulation*. 2003;107(19):2435-2439. doi: 10.1161/01.CIR.0000066906.11109.1F
- **47**. Trichopoulou A, Psaltopoulou T, Orfanos P, Trichopoulos D. Diet and physical activity in relation to overall mortality amongst adult diabetics in a general population cohort. *J Intern Med*. 2006;259 (6):583-591. doi:10.1111/j.1365-2796.2006.01638.x
- **48**. Wei M, Schwertner HA, Blair SN. The association between physical activity, physical fitness, and type 2 diabetes mellitus. *Compr Ther.* 2000;26(3):176-182. doi:10.1007/s12019-000-0006-9
- **49**. Hu G, Eriksson J, Barengo NC, et al. Occupational, commuting, and leisure-time physical activity in relation to total and cardiovascular mortality among Finnish subjects with type 2 diabetes. *Circulation*. 2004;110(6):666-673. doi:10.1161/01.CIR.0000138102.23783.94
- **50**. Boulé NG, Kenny GP, Haddad E, Wells GA, Sigal RJ. Meta-analysis of the effect of structured exercise training on cardiorespiratory fitness in type 2 diabetes mellitus. *Diabetologia*. 2003;46(8): 1071-1081. doi:10.1007/s00125-003-1160-2
- **51.** Church TS, Cheng YJ, Earnest CP, et al. Exercise capacity and body composition as predictors of mortality among men with diabetes. *Diabetes Care*. 2004;27(1):83-88. doi:10.2337/diacare.271.83

- **52.** Gaither TW, Sanford TA, Awad MA, et al. Estimated total costs from non-fatal and fatal bicycle crashes in the USA: 1997-2013. *Inj Prev.* 2018;24(2):135-141. doi:10.1136/injuryprev-2016-042281
- **53.** Pope CA III, Ezzati M, Dockery DW. Fine-particulate air pollution and life expectancy in the United States. *N Engl J Med*. 2009;360(4):376-386. doi:10.1056/NEJMsa0805646
- **54.** Di Q, Wang Y, Zanobetti A, et al. Air pollution and mortality in the Medicare population. *N Engl J Med*. 2017;376(26):2513-2522. doi:10.1056/NEJMoa1702747
- **55.** Sinharay R, Gong J, Barratt B, et al. Respiratory and cardiovascular responses to walking down a traffic-polluted road compared with walking in a traffic-free area in participants aged 60 years and older with chronic lung or heart disease and age-matched healthy controls: a randomised, crossover study. *Lancet*. 2018;391(10118):339-349. doi:10.1016/S0140-6736(17)32643-0
- **56**. Kubesch NJ, de Nazelle A, Westerdahl D, et al. Respiratory and inflammatory responses to short-term exposure to traffic-related air pollution with and without moderate physical activity. *Occup Environ Med*. 2015;72(4):284-293. doi:10.1136/oemed-2014-102106
- **57.** Laeremans M, Dons E, Avila-Palencia I, et al. Short-term effects of physical activity, air pollution and their interaction on the cardiovascular and respiratory system. *Environ Int.* 2018;117:82-90. doi:10.1016/j.envint.2018.04.040
- **58**. Kubesch N, De Nazelle A, Guerra S, et al. Arterial blood pressure responses to short-term exposure to low and high traffic-related air pollution with and without moderate physical activity. *Eur J Prev Cardiol*. 2015;22(5):548-557. doi:10.1177/2047487314555602
- **59.** Kubesch NJ, Therming Jørgensen J, Hoffmann B, et al. Effects of leisure-time and transport-related physical activities on the risk of incident and recurrent myocardial infarction and interaction with traffic-related air pollution: a cohort study. *J Am Heart Assoc.* 2018;7(15):e009554. doi:10.1161/JAHA.118.009554
- **60**. Sun S, Cao W, Qiu H, et al. Benefits of physical activity not affected by air pollution: a prospective cohort study. *Int J Epidemiol*. 2020;49(1):142-152. doi:10.1093/ije/dyz184
- **61**. Laeremans M, Dons E, Avila-Palencia I, et al. Black carbon reduces the beneficial effect of

- physical activity on lung function. *Med Sci Sports Exerc*. 2018;50(9):1875-1881. doi:10.1249/MSS. 00000000000001632
- **62**. Fisher JE, Loft S, Ulrik CS, et al. Physical activity, air pollution, and the risk of asthma and chronic obstructive pulmonary disease. *Am J Respir Crit Care Med.* 2016;194(7):855-865. doi:10.1164/rccm.201510-2036OC
- **63.** Welsh C, Celis-Morales CA, Ho F, et al. Association of injury related hospital admissions with commuting by bicycle in the UK: prospective population based study. *BMJ*. 2020;368:m336. doi:10.1136/bmj.m336
- **64.** Blond K, Brinkløv CF, Ried-Larsen M, Crippa A, Grøntved A. Association of high amounts of physical activity with mortality risk: a systematic review and meta-analysis. *Br J Sports Med*. 2020;54 (20):1195-1201. doi:10.1136/bjsports-2018-100393
- **65.** Lee DH, Rezende LFM, Ferrari G, et al. Physical activity and all-cause and cause-specific mortality: assessing the impact of reverse causation and measurement error in two large prospective cohorts. *Eur J Epidemiol*. 2021;36(3):275-285. doi: 10.1007/s10654-020-00707-3
- **66**. Hutcheon JA, Chiolero A, Hanley JA. Random measurement error and regression dilution bias. *BMJ*. 2010;340:c2289. doi:10.1136/bmj.c2289
- **67**. Andersen LB. Relative risk of mortality in the physically inactive is underestimated because of real changes in exposure level during follow-up. *Am J Epidemiol*. 2004;160(2):189-195. doi:10.1093/aje/kwh195
- **68.** Saeedi P, Petersohn I, Salpea P, et al; IDF Diabetes Atlas Committee. Global and regional diabetes prevalence estimates for 2019 and projections for 2030 and 2045: results from the International Diabetes Federation Diabetes Atlas, 9th edition. *Diabetes Res Clin Pract.* 2019;157:107843. doi:10.1016/j.diabres.2019.107843
- **69**. Espeland MA, Glick HA, Bertoni A, et al; Look AHEAD Research Group. Impact of an intensive lifestyle intervention on use and cost of medical services among overweight and obese adults with type 2 diabetes: the action for health in diabetes. *Diabetes Care*. 2014;37(9):2548-2556. doi:10.2337/dc14-0093
- **70**. Zimmet P, Alberti KG, Magliano DJ, Bennett PH. Diabetes mellitus statistics on prevalence and mortality: facts and fallacies. *Nat Rev Endocrinol*. 2016;12(10):616-622. doi:10.1038/nrendo.2016.105