## Optimal number of steps per day to prevent all-cause mortality in people with prediabetes and diabetes

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## SUPPLEMENTAL MATERIALS

**Table S1.** Baseline characteristics of the participants in the study (n=1194 participants with pre-diabetes; n=493 participants with diabetes)

**Figure S1.** Dose–response association (Adjusted<sup>a</sup> hazard ratios—solid lines and associated 95% confidence interval band—dashed lines) between accelerometer-derived steps per day and all-cause mortality in participants with pre-diabetes or diabetes (n=1687; events=338).

**Figure S2.** Dose–response association (Adjusted<sup>a</sup> hazard ratios—solid lines and associated 95% confidence interval band—dashed lines) between accelerometer-derived steps per day and all-cause mortality in participants with pre-diabetes (n=1102; events=183) and diabetes (n=454; events=122) and with 3 or more days of valid accelerometry.

**Figure S3.** Dose–response association (Adjusted<sup>a</sup> hazard ratios—solid lines and associated 95% confidence interval band—dashed lines) between accelerometer-derived steps per day and CVD mortality in participants with pre-diabetes (n=1194; events=49) and diabetes (n=493; events=36).

pre-diabetes; n=493 participants with diabetes)		
	Pre-diabetes (n=1194)	Diabetes (n=493)
Age (years)	54.7 (17.9)	61.6 (13.8)
Sex, female (%)	528 (44.2)	243 (49.3)
Ethnicity (%)		
Mexican American	253 (21.2)	120 (24.3)
Other Hispanic	33 ( 2.8)	12 ( 2.4)
Non-hispanic, white	549 (46.0)	183 (37.1)
Non-hispanic, black	303 (25.4)	164 (33.3)
Other	56 ( 4.7)	14 ( 2.8)
Education (%)		
Less Than 9th Grade	159 (13.9)	102 (20.8)
9-11th Grade	179 (15.6)	86 (17.5)
High School Grad/GED or Equivalent	291 (25.3)	138 (28.1)
Some College or Associated Arts Degree	334 (29.1)	112 (22.8)
College Graduate or above	185 (16.1)	53 (10.8)
Body Mass Index (Kg/m <sup>2</sup> )	30.2 (7.3)	32.3 (7.7)
Alcohol (grams)	9.59 (24.69)	5.62 (18.58)
Cotinine (ng/mL)	5.71 (0.52)	7.36 (1.81)
Diet quality, HEI score (1-100)	52.0 (13.6)	55.0 (13.1)
Glycohemoglobin (%)	5.71 (0.52)	7.36 (1.81)
Diabetes medication (%)	0.08 (0.27)	0.57 (0.50)
Major chronic conditions, no (%)	566 (49.0)	166 (33.8)
Accelerometer varaibles		
Steps per day (n)	8949.6 (4171.8)	7151.2 (3930.6)
Valid daily wear time (min)	873.7 (147.5)	850.8 (142.1)
Valid wear days (n)	5.5 (1.7)	5.6 (1.7)
Values are shown as mean (SD) unless otherwise specified.		
HEI, Healthy Eating Index. Lower scores represent lower diet quality		

**Table S1.** Baseline characteristics of the participants in the study (n=1194 participants with pre-diabetes; n=493 participants with diabetes)

**Figure S1.** Dose–response association (Adjusted<sup>a</sup> hazard ratios—solid lines and associated 95% confidence interval band—dashed lines) between accelerometer-derived steps per day and all-cause mortality in participants with pre-diabetes or diabetes (n=1687; events=338).



Participants with pre-diabetes and diabetes

<sup>a</sup>Adjusted for age, sex, ethnicity, education, smoking, alcohol, diet, diabetes medication, and valid daily wear time. To prevent over-influential outliers from affecting the analyses, values in all step-based metrics that were greater than the 99<sup>th</sup> percentile of the variable distribution were top-coded (i.e., they were substituted with the value of the 99<sup>th</sup> percentile) and values that were below the 1<sup>st</sup> percentile of the variable distribution were bottom-coded (i.e., they were replaced with the value of the 1<sup>st</sup> percentile). Darker colors in the lower bars represent a higher sample clustering. Dose-response associations were assessed with restricted cubic splines with knots at 10<sup>th</sup> (reference), 50<sup>th</sup>, and 90<sup>th</sup> centiles of the distribution of the exposure of interest. Hazard ratios are in logarithmic scale.

**Figure S2.** Dose–response association (Adjusted<sup>a</sup> hazard ratios—solid lines and associated 95% confidence interval band—dashed lines) between accelerometer-derived steps per day and all-cause mortality in participants with pre-diabetes (n=1102; events=183) and diabetes (n=454; events=122) and with 3 or more days of valid accelerometry.



<sup>a</sup>Adjusted for age, sex, ethnicity, education, smoking, alcohol, diet, diabetes medication, and valid daily wear time. To prevent over-influential outliers from affecting the analyses, values in all step-based metrics that were greater than the 99<sup>th</sup> percentile of the variable distribution were top-coded (i.e., they were substituted with the value of the 99<sup>th</sup> percentile) and values that were below the 1<sup>st</sup> percentile of the variable distribution were bottom-coded (i.e., they were replaced with the value of the 1<sup>st</sup> percentile). Darker colors in the lower bars represent a higher sample clustering. Dose-response associations were assessed with restricted cubic splines with knots at 10<sup>th</sup> (reference), 50<sup>th</sup>, and 90<sup>th</sup> centiles of the distribution of the exposure of interest. Hazard ratios are in logarithmic scale.

**Figure S3.** Exploratory dose–response association (Adjusted<sup>a</sup> hazard ratios—solid lines and associated 95% confidence interval band—dashed lines) between accelerometer-derived steps per day and CVD mortality in participants with pre-diabetes (n=1194; events=49) and diabetes (n=493; events=36).



The associations between between accelerometer-derived steps per day and CVD mortality was conducted as an exploratory analysis. Fine and Gray models were used to account for competing risks with other causes of mortality. <sup>a</sup>Adjusted for age, sex, ethnicity, education, smoking, alcohol, diet, diabetes medication, and valid daily wear time. To prevent over-influential outliers from affecting the analyses, values in all step-based metrics that were greater than the 99<sup>th</sup> percentile of the variable distribution were top-coded (i.e., they were substituted with the value of the 99<sup>th</sup> percentile) and values that were below the 1<sup>st</sup> percentile of the variable distribution were replaced with the value of the 1<sup>st</sup> percentile). Darker colors in the lower bars represent a higher sample clustering. Doseresponse associations were assessed with restricted cubic splines with knots at 10<sup>th</sup> (reference), 50<sup>th</sup>, and 90<sup>th</sup> centiles of the distribution of the exposure of interest. Hazard ratios are in logarithmic scale.